# Microsatellite scanning of the Immunogenome for Associations with Graft-versus-Host Disease following Haematopoietic Stem Cell Transplantation 

A thesis submitted to the University of Newcastle in accordance with the requirements for the degree of Doctor of Philosophy (PhD)

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## Candidate's Declaration

I, Christian Harkensee, hereby certify that this thesis has been written by me, that it is the record of work carried out by me (unless stated otherwise) and that it has not been submitted in any previous application for a higher degree.

Dedicated to the memory of

Akira Sasaki

Without whom this work would never have been undertaken.

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The work described in this thesis is exclusively my own, unless stated otherwise.

Finally, this work could not have been done without the support of my family, my wife Ena and my children Aila and Kai. The time should have been yours...

## Publications

The work described in this thesis has been published in part.

## Peer-reviewed journal publications

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#### Abstract

Non-HLA gene polymorphisms contribute to the immune response, leading to complications of haematopoietic stem cell transplantation (HSCT). A systematic approach using 4,321 microsatellite (MS) markers typing for 2,909 immune response genes ('immunogenome') on pooled DNA of 922 Japanese donors and recipients of HSCT was used to identify recipient and donor risk loci for graft-versus-host disease (GVHD).

Splitting the population into discovery and confirmation cohorts (460/462 pairs), DNA pools were created for a 2-step pooled DNA screening. Fisher's exact test for $2 \times 2$ (each MS allele) and $2 \times m$ Chi Square tests were performed, comparing allele frequencies of recipient/donor pools with GVHD grade 0-1 with those of GVHD grade 2-4.

The independent, 2-step pooled DNA screening process has effectively reduced false-positive associations. In the final pooled DNA analysis, 17 (recipient) and 31 (donor) MS loci remained associated with risk or protection from GVHD and were further investigated by individual genotyping in the combined cohorts. Ten of these loci were confirmed to have consistent associations with GVHD; of these, two associations remained when applying multiple testing correction and multivariate statistics: D6S0035i (MAPK14, $p=0.00035, O R=0.68)$ and D1S0818i (ELTD1, $p=0.000078, \mathrm{OR}=1.52$ ).

These findings implicate important new immunoregulatory genes with the process of moderate to severe acute GVHD. These data show that genetic susceptibility to GVHD following HSCT is complex and depends on multiple recipient and donor risk loci. Large-scale genomic screening with microsatellites on pooled DNA, here described for the first time in a HSCT population, is a useful method for the systematic evaluation of multigeneic traits.


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## 1. INTRODUCTION

1.1. Haematopoietic Stem Cell Transplantation and Graft versus Host Disease
1.2. Pathophysiology and pathobiology of GVHD
1.3. The genetics of HSCT
1.4. Summary and conclusion; aim of this study
1.5. Outline of study plan/brief history of the project

### 1.1. Haematopoietic Stem Cell Transplantation and Graft versus Host Disease

Graft versus Host Disease (GVHD) and its consequences remain the single most important contributor to morbidity and mortality in patients following haematopoietic stem cell transplantation (HSCT). Despite progress made in Human Leukocyte Antigen (HLA) matching, and the use of pharmacologic immunosuppression as GVHD prophylaxis after myeloablative transplantation, moderate to severe acute GVHD (grades IIIV) occured in $25 \%$ to $60 \%$ of matched related donor transplant recipients, and up to $45 \%$ to $70 \%$ in unrelated donor recipients (Horowitz, 2004, Andre-Schmutz et al., 2002, Grewal et al., 2003, Laughlin et al., 2001, Morishima et al., 2002). The unpredictability of occurrence and severity of GVHD is the main obstacle today that prevents the wider application of HSCT.

The first successful human bone marrow transplant (BMT) was performed 1959 by Thomas in the US on a patient with leukaemia, using syngeneic bone marrow from his identical twin (Thomas et al., 1959) .

The history of BMT began with the work of Jacobson and Lorenz in the early 1950's, demonstrating that infusion of marrow or spleen cells could 'rescue' mice after a lethal radiation dose (Jacobson et al., 1949, Lorenz et al., 1951).

Later in the 1950's, van Bekkum and De Vries established murine models that provided fundamental knowledge of the biology of marrow transplantation (van Bekkum and De Vries, 1967). Billingham was the first to describe a condition that was initially termed 'Secondary Disease’ (because it occurred after successful engraftment), 'runt disease' or 'wasting disease', and later re-named 'Graft-versus-Host Disease', after it became clear that this was an immunological reaction of donor T-cells against host tissues (Billingham and Brent, 1959). In 1957, Uphoff (Uphoff,
1957) was the first to suggest a genetic cause for the graft-versus-host reaction in allogeneic transplants.

In 1958, a first classification of HLA groups as the most important marker of an individuals biological identity was established (van Rood et al., 1958, Dausset, 1958). The capability of the immune system to generate antibodies against antigens on the leukocyte surface was already described in 1954 (Miescher and Fauconnet, 1954). Dausset demonstrated that MHC (Major Histocompatibility Complex) genes (which encode for HLA) were required for the presentation of peptide antigens to T-cell receptors, playing an important role in transplant immunology. In humans, the MHC cluster on chromosome 6 was identified as the coding region for HLA. Further research lead to the detection of the different gene loci (HLA A,B,C,DR, DP, DQ) and a large number of HLA alleles (Shiina et al., 2004). Two main pathways of antigen recognition were detected: A direct response, in which CD8+ cytotoxic T-cells interact with HLA, and an indirect response in which CD4+ T-cells are activated to induce a delayedtype hypersensitivity reaction, cell-mediated toxicity and alloantibody production. Major events were the discovery of the role of lymphocytes and the thymus in the ontogeny of the immune system (1961), the delineation of the human MHC (1963), distinction of B/T-cell subsets (1968) and demonstration of the MHC restricted nature of the adaptive immune response (reviewed in: (Thomas, 1994).

In the 1960's, studies in canine models laid the foundations for conditioning regimens, GVHD prophylaxis and genetic matching. Observations included that total body irradiation (TBI) did not cure leukaemia nor did it prevent GVHD or bone marrow recovery. HLA mismatch was likely to result in rejection or GVHD, and methotrexate was introduced as an immunosuppressive agent (Thomas et al., 1962). By the end of the 1960's, the supportive care had also seen significant improvement.

The increasing knowledge of histocompatibility lead to a renewed interest in allogeneic transplantation, using fully or partially matched siblings as donors. While some success was achieved in the transplantation of
patients with immunodeficiency (Gatti et al., 1968), the initial results of BMT for leukaemic patients were not encouraging (Thomas et al., 1975a, Thomas et al., 1975b). The reason for this was thought to be patient selection: BMT was seen as a 'last resort' for patients with advanced disease, or after multiple chemotherapy failures. A small percentage of patients, however, achieved long-term cure, indicating that BMT had the potential to cure even very advanced disease. Outcome for leukaemic patients improved greatly from the late 1970's onwards after the introduction of transplantation after first remission, or in early first relapse (Thomas et al., 1979a, Beutler et al., 1979). The increased use of allogeneic bone marrow lead to the first bone marrow donor registries (UK 1975, US 1984).

Transplant conditioning regimens, given as an immunosuppression and for tumour eradication, had come a long way from the early, simple TBI regimen to the reduced intensity conditioning (RIC) regimens of today. Observations with TBI, as well as cyclophosphamide (CY) as single therapies in the 1950's and 1960's were such that leukaemia relapsed rapidly after transplantation. The combination of TBI and CY lead to the first observed long term remissions in the mid 1970's (Thomas et al., 1975a, Thomas et al., 1975b). Introduction of newer chemotherapeutic agents such as busulphan, and fractioned irradiation have reduced toxicity and improved survival. Nevertheless, these myeloablative regimens proved still far too toxic for elderly patients or those with co-morbidities, in which transplant-related mortality (TRM) would reach unacceptable high levels. The last two decades have opened up transplant opportunities for this age and risk group by the development of RIC regimens (Koh and Chao, 2008).

Whilst immunosuppressive regimens achieved far-reaching control of transplant rejection, GVHD and its consequences remained the single most important complication of stem cell transplantation. Studies in animal models and observations in human patients clarified important checkpoints in the pathophysiology of GVHD (Korngold and Sprent, 1978, Shlomchik, 2007). In the 1980's, even with HLA matched sibling donors up to $50 \%$ of
recipients developed GVHD. In the 1960's and 1970's, methotrexate and steroids were first used for treatment, later for prevention with only limited effect (Thomas et al., 1979b). The introduction of cyclosporin A, an inhibitor of T-cell activation and proliferation, in 1980, used in combination with a short course of methotrexate, resulted in improved prevention of GVHD (Deeg et al., 1982). T-cell depletion, first described in the early 1980's (Martin et al., 1985), was found to prevent GVHD, however, at the expense of the loss of the GVL (Graft versus Leukaemia) effect, risk of graft failure and delayed immunologic recovery. Newer agents used in prophylaxis include Tacrolimus and Sirolimus, believed to be more potent in blocking T-cell proliferation, and Mycophenolate Mofetil, active against both T-cells and B-cells.

Over the last two decades, improved techniques of molecular genetics replaced the HLA serum typing with genotyping, allowing for matching at individual allele level, further reducing the incidence of GVHD (Little, 2007). Advanced molecular techniques also permitted the identification of minor histocompatibility antigens (mHags), which play an important role in otherwise fully HLA-matched transplants (Goulmy, 2006). The process of GVHD also seemed to be modified by polymorphisms in immunoregulatory genes (Mullighan and Bardy, 2007). Presence of active infection represented a risk factor for GVHD, while in turn GVHD (and its treatment) produced profound immune suppression, increasing the risk of infection.

The concept of the GVL effect was first pointed out in the 1950's (Barnes et al., 1956), and observed in mouse experiments during the 1960's. It was first described in human patients in the mid-1970's when it became apparent that patients with GVHD had a reduced risk of leukaemia relapse (Weiden et al., 1979). Around the same time, animal experiments demonstrated that the GVL effect could be separated from GVHD (Bortin et al., 1979). The introduction of post-transplant donor lymphocyte infusions (DLI) in the second half of the 1980's was the first established method with the aim to enhance the GVL in patients with haematological and other malignancies. The potential of DLI to induce long-lasting
remission was demonstrated, however, at the expense of increased GVHD and prolonged bone marrow suppression (Kolb et al., 1990).

Current directions in progress in BMT can be described as four big areas. Firstly, there is continuing improvement in supportive care, anti-infectious therapy, and reduction of toxicity of induction regimens. Secondly, the study of clinical and genetic (HLA and non-HLA) risk factors may lead to a more reliable prediction of transplant complications. Thirdly, separation of GVL from GVHD and fourthly, enhancement of immune reconstitution, could be summarized under the term 'Adoptive Immunotherapy'. Some methods applied, amongst others, include in vivo and in vitro tumorspecific alloimmunization, the use of immunomodulatory cells (T-regulatory cells, dendritic cells, Mesenchymal stem cells), cytokines, growth hormones, non-specific immune regulators and monoclonal antibodies to suppress GVHD, enhance GVL and stimulate the reconstitution of a healthy immune function. With GVHD being the major hurdle of the application of HSCT for other indications, advances in its control are likely to expand the future role of HSCT.

## SUMMARY

- The history of HSCT spans five decades, during which the procedure has become more effective and safe.
- Progress in timing, genetic matching, conditioning regimen, GVHD prophylaxis and clinical care has widened the applicability and indications of HSCT


### 1.2 Pathophysiology and pathobiology of GVHD

### 1.2.1. Overview

Pathophysiologically, GVHD involves the recognition of target tissues as being foreign by immunocompetent donor cells, resulting in the induction of an inflammatory response. Acute GVHD has been described in three different phases: (I) presence of a profoundly damaged host milieu prior to contact with donor tissue, characterized by excessive endo- and epithelial damage due to preparing chemotherapy regimes, irradiation, damaged and dying malignant cells and infections, (II) donor T cell activation by host antigen-presenting cells (APC), and (III) the inflammatory response consisting of activation of donor cytotoxic T cells, natural killer cells (NKC) and monocytes. Cytokine toxicity and viral and bacterial infections also contribute to the inflammatory response (Ferrara and Levine, 2006, Ferrara et al., 1999).

### 1.2.2. GVHD phase 1: Preceding tissue damage, pre-transplant conditioning, and activation of antigen-presenting cells

## Preceding local tissue damage

The damage to recipient tissues caused by conditioning regimens and previous chemotherapy or radiation, the underlying malignancy, and concurrent infections had a major impact on the incidence and severity of acute GVHD (Xun et al., 1994, Perez-Simon et al., 2005, Couriel et al., 2004, Gratama et al., 1987). The local micro-environment of inflammatory mediators determined the degree and specificity of APC activation, and subsequently, induction of naïve T-cells. An important mechanism appeared to be the increased cytokine secretion in the local environment, leading to increased expression of adhesion molecules and MHC, increased antigen presentation by APC, increased antigen recognition by $T$ cells, and subsequently increased T cell activation and proliferation. TNF, IL1 and IL6 were the most important cytokines involved in stimulating local
tissue inflammation in GVHD (Nestel et al., 1992, Imamura et al., 1994, Remberger et al., 1995, Cooke et al., 1998)
Tissue insults by toxic effects of chemotherapeutics, irradiation, dying diseased cells, or infection with bacterial, viral, fungal or protozoal pathogens, were causing a mixed pattern of cells either weakened by stress, or cell apoptosis and cell necrosis with disruption of tissue architecture. Those tissues which were environmentally exposed, or rapidly dividing and renewing, such as the epithelia of the liver, intestinal tract and skin, were the most vulnerable.

There is evidence that the initial immune response to the tissue damage were triggered by endogenous stress or danger signals, powerful enough to induce activation of local APC in a paracrine fashion (Lotze et al., 2007, Skoberne et al., 2004, Gallucci et al., 1999, Gallucci and Matzinger, 2001). In contrast to pathogen-associated proteins, which induce a specific immune response, endogenous proteins from injured tissue represented a potentially unlimited source of self-antigen. While apoptotic cells were engulfed in quiescence and their antigens presented to T-cells without costimulation, hence inducing tolerance; proteins from necrotic cells presented as antigens lead to T-cell activation (Shlomchik, 2007). Nevertheless, a mouse model (Teshima et al., 2002) demonstrated that alloantigen presentation on host epithelium was not a requirement for acute GVHD, but that GVHD could be abrogated by TNFa blockade, supporting the notion of the importance of the cytokine micro-environment.

## Activation of host APC

The first circle of augmentation of the pro-inflammatory response occurred still in the injured tissue itself. When local APC became activated, they secreted further pro-inflammatory cytokines like tumour necrosis factor alpha (TNF $)$, interleukin 1 (IL-1), granulocyte-macrophage stimulating factor (GM-CSF) and Interferon gamma (IFNY) (Ferrara et al., 1999), which in turn recruited more epithelial cell and APC for cytokine secretion. Increased cytokine concentrations eventually reached systemic circulation, providing a powerful chemoattractant stimulus. Increased cytokine expression stimulates the excretion of chemokines and their receptors,
which had an important role in dendrictic cell (DC) activation and maturation, as well as CD8+ T-cell homing to GVHD target tissues (New et al., 2002, Mapara et al., 2006). This micro-climate, in turn, attracted and activated DC and other antigen-presenting cells and broke the ground for mature donor T-cells, which after transplantation recognised increasingly expressed MHC molecules and/or mHags (Matzinger, 2002).

The consequences of tissue damage and subsequent immune system activation were even more imminent in the intestinal mucosal surface. Here, local TNF secretion negatively affected the surface integrity (Laster et al., 1988), enhancing the potential injuries caused by intramucosal displacement of bacterial endotoxins and lipopolysaccharides (LPS), which could potentiate even further the pro-inflammatory cytokine production by gut-associated macrophages and lymphocytes (Nestel et al., 1992). The intestinal Peyer's Patches (PP) had been identified as a key lymphoid compartment for the development of acute GVHD (Murai et al., 2003) by demonstrating that acute GVHD was abrogated in mice which were PPdeficient or whose T-cell homing had been interrupted by C-chemokine receptor 5 (CCR5) -blockade; suggesting that GVHD induction was determined by the local inflammatory environment, rather than tissuespecific mHags. On the other hand, PP were redundant, hence T-cells primed elsewhere in the body were capable of inducing intestinal GVHD (Welniak et al., 2006).
APC were either tissue-specific and residing (e.g. Langerhans' cell in skin, Kupfer cell in liver) or circulated through the body's blood and lymphoid system in their naïve and immature form. Their key function for the adaptive immune system was their capability of inducing a specific, adaptive immune response by presenting a specific antigen to naïve or memory T-cells through their MHC receptor. DC, the most important APC in GVHD induction, could be activated in a specific (antigen, foreignrecognition) or non-specific (micro-environmental stimuli, danger recognition) manner. The antigen was internalised and processed in endolysosomes to peptide which was loaded onto MHC molecules. Physiologically, immature DC relentlessly sampled endogenous antigen, which was presented to T-cells rendering those tolerant to self-antigen.

Antigen recognised as 'non-self', however, lead to DC maturation and antigen presentation to T-cells accompanied by appropriate co-stimulatory signals. DC were able to suppress the response of T-regulatory cells, and control the blood flow to the lymph node into which they home. In the lymph node, the matured DC interacted with naïve or memory CD4 T-cells through the MHC/T-cell receptor (TCR) 'immunological synapse', resulting in effective activation and proliferation of an antigen-specific T-cell clone (Lee and Iwasaki, 2007).

It was thought that in early GVHD, tissue antigen was presented to the infused donor T cells by host DC which were critical, and sufficient, to induce a GVHD response (Shlomchik et al., 1999, Duffner et al., 2004). Host Langerhans cells in the skin had been shown to be critical mediators of skin GVHD (Merad et al., 2004). Antigen presentation by host APC to donor T-cells, also called direct recognition, was the predominant mode of allorecognition in the MHC mismatched transplant setting (Ruggeri et al., 2002).

Removal of APC from their specific organ could potentially abrogate GVHD in the same organ (Zhang et al., 2002b). The same authors suggested that host APC localised in specific target organs recruited mature donor T-cells, while DC and macrophages homed to lymph nodes and other secondary lymphoid organs were critical for the activation of a cytotoxic CD8+T-cell response.

Of the different subsets of APC, DC probably was the most critical one in the development of GVHD, supported by its role as the most important 'professional' APC, and also by observation in experimental GVHD settings (Duffner et al., 2004, Zhang et al., 2002a).
APC regulation in the context of GVHD is not fully understood. TBI as part of an HSCT conditioning regimen, in combination with G-CSF exacerbated APC activation (Morris et al., 2009), while үס T-cells, host Natural Killer Tcells (NKT), natural killer (NK) cells and B cells reduced activation (Paczesny et al., 2009a).

Activated host APC relocated into secondary lymphoid organs, where following HSCT they met donor T cells. Donor T-cells recognized antigen presented by host APC (foreign antigen or the host MHC receptor itself direct presentation) or donor APC (recognition of the foreign antigen indirect presentation) (Shlomchik, 2003, Sayegh and Carpenter, 1996). Donor T-cells required co-stimulation of their T-cell receptor in the binding to the MHC receptor of the APC in order to become activated (Appleman and Boussiotis, 2003), with CD28, ICOS, CD40, OX40 (activation) and CTLA4 (inhibition) being the most important co-stimulatory molecules (Paczesny et al., 2009a).

Different subsets of T-cells had varying roles in induction of GVHD. The two main subsets, CD4+ and CD8+ T cells were both capable of inducing GVHD. In the absence of HLA mismatch, this potency was determined by host mHags which could lead to specific T-cell clones in either CD4+ or CD8+ subset (Goulmy, 2006, Wu and Ritz, 2006). Selective elimination of either subset from grafts did not lead to a reduction in GVHD, and research on selecting specific antigeneic clones for elimination is ongoing (Bondanza et al., 2006).
Naïve donor T-cells (CD62L+ CD44+) also had the potential of inducing GVHD (Anderson et al., 2003). Non-alloreactive donor memory T-cells (CD62L-) did not induce GVHD, but were able to mediate GVL effects through memory transfer (Zheng et al., 2008). Alloreactive donor T-cells, however, were a main cause of GVHD (Zhang et al., 2005).
Regulatory T-cells (Treg) had been the focus of more recent research. Several studies had demonstrated that Treg are capable of suppressing the expansion of activated donor T-cells, and therefore reduce the risk of GVHD, whilst preserving GVL reactions (Yan and Da, 2006, Salomon et al., 2006, Ruggeri et al., 2002). Host and donor NKT also had GVHD-reducing properties (Pillai et al., 2007), by shifting cytokine responses to a T-helper type 2 (Th2) profile (Lowsky et al., 2005), or eliminating host APC (Morris et al., 2005).

In the late 1980's it was discovered that following activation subsets of mainly CD4+ T-cells had antagonistic cytokine excretion profiles, described as Th1 (T-helper type 1) and Th2 (T-helper type 2) cells (Mosmann et al., 1986). Th1 cells secreted pro-inflammatory cytokines like IL2 (interleukin 2), TNF, IFNY (interferon gamma), while the Th2 (T-helper 2 cell) had anti-inflammatory properties, secreting cytokines like IL4 (interleukin 4) and IL10 (interleukin 10). These phenotypes followed each other over time (in the initial phase of an inflammatory reaction, the response was predominantly pro-inflammatory, followed by an antiinflammatory response to counteract and prevent excessive inflammation). There is good evidence that genetic polymorphisms resulting in over- or under-expression of certain cytokines could tilt this balance to either a more pro-inflammatory or an anti-inflammatory response (see below). In Phase 2, donor T cells stimulated by APC secrete IL2 and IFNy (Mosmann et al., 1986), which was central in the control and amplification of the immune reaction against the foreign antigen. IFNy primed macrophages to produce and secrete IL1 (interleukin 1) and TNF (Nestel et al., 1992), induced the skin and gut pathology of GVHD (Dickinson et al., 1991), and impaired T-lymphocyte function (Huchet et al., 1993). The levels of IFNY were predictive of GVHD severity (Tanaka et al., 1994). This could be described as a Th1-type response, promoting GVHD. On the other side, Th2-type cytokines like IL4 and IL10 had antagonistic effects on IL2 and IFNy secretion (Seder and Le Gros, 1995), dampening down reactions leading to acute GVHD and 'shifting' the immune response towards chronic GVHD (Krenger and Ferrara, 1996). Nevertheless, the biology of cytokines in GVHD is likely to be more complex, as opposing effects of the same cytokine (e.g. IL2 or IFNy) had been observed, depending on factors like timing, concentration and tissue location (Wang et al., 1995, Krenger et al., 1996, Baker et al., 1995, Yang et al., 1998, Brok et al., 1997). Sun et. al. (Sun et al., 2007) postulated that "early Th1 polarization of donor T-cells and Th1 cytokines are critical for GVHD induction, whereas inadequate production could modulate acute GVHD through a breakdown of negative feedback mechanisms for activated T-cells".

IL17 (interleukin 17) Th-cells were a more recently described subset of CD4+ T-cells, characterized by the production of IL17. Their role in GVHD remained controversial (Sun et al., 2007, Paczesny et al., 2009a), as studies had shown that IL17 deficient T- cells enhanced Th1 skewing augmenting acute GVHD, whereas Th17 cells caused severe GVHD in vitro.

### 1.2.4. GVHD Phase 3: Effector phase

Once activated donor T-cells migrate from secondary lymphoid tissue into target organs, where they cause tissue damage. Potentially any organ is capable of expressing alloantigen and therefore to become a target organ for GVHD, however, skin, gut, liver and thymus are the most commonly affected target tissues.

Recent studies have shed some light onto mechanisms that control the 'homing' of alloreactive T-cells into their target tissues. Chemokines appeared to be one of the key players. Inflammatory chemokines were expressed by a wide variety of tissues and cells (endothelial cells, fibroblasts, DC, monocytes, NK cells and T-lymphocytes) and regulated trafficking of donor T-cells towards the lymphoid organs where they interacted with APC, as well as target tissues (reviewed in: (Wysocki et al., 2005). Activated T-lymphocytes expressed chemokine receptors which, at least in part, determined their destination by homing to tissues where the according ligands were expressed (e.g.: liver: receptors CXCR3, CCR2, CCR5, ligands: CXCL9/10/11, CCL2/3; gut: receptors CXCR3, CCR9, ligands CXCL9/10/11, CCL25; skin: receptors CXCR3, ligands CXCL9/10/11, CCL2/5/17).

Other mechanisms of T-cell trafficking included selectins and integrins (also reviewed in (Wysocki et al., 2005). E, P and L-selectin were expressed in various tissues including on cells of the myeloid and lymphoid system, and found their ligands, the peripheral node addressins (PNAds) expressed in chronically inflamed tissue. Integrins were transmembrane proteins expressed on immune cells which interacted with molecular
structures in tissues that lay exposed due to tissue damage (e.g. collagen, laminin, fibronectin) or specific expressed ligands like adhesion molecules (MADCAM1, VCAM1, ICAM). Expression of MADCAM1 and ICAM1, for example, was critical for induction of acute GVHD in the PP of the intestine and in the liver.

Donor T-cells exerted their deleterious effect on target tissues by direct cytotoxicity (van den Brink and Burakoff, 2002) using different pathways of apoptosis. CD4+ cytotoxic T-lymphocytes (CTL) applied mainly the FasFasL (TNF-receptor superfamily member 6) pathway (Via et al., 1996). Fas is widely expressed, its expression is inducible by TNF and IFNY and therefore enhanced in inflamed tissues. The same cytokines enhanced FasL expression on CTL, hence this mechanism appeared to be a selfaugmenting cycle of tissue damage, inflammation, CTL recruitment and apoptosis. CD8+ CTL worked preferentially through the perforinegranzyme pathway. Perforin is secreted by the CTL causing pores in the target cell membrane. This allows granzyme to enter and induce a celldeath sequence. In experimental GVHD, survival was better in peforin/granzyme deficient mice, but the cytotoxic effect was less pronounced than for Fas/FasL (Graubert et al., 1997). Experimental studies for both pathways, by inactivating important components, had shown that GVHD can be abrogated or delayed in onset (reviewed in: (Sun et al., 2007)).

Other apoptosis mechanisms include TRAIL (TNF-related apoptosisinducing ligand) (Pan et al., 1997), TWEAK (TNF-like weak inducers of apoptosis) and LT $\beta /$ LIGHT pathways (Brown et al., 2005).

Much of the tissue damage of the effector phase of GVHD could be attributed to an excessive release of pro-inflammatory cytokines, the socalled 'cytokine storm'. Local tissue damage, due to conditioning or infection, induced APC activation through TLR and non-TLR pathways, and caused chemoattraction to macrophages, which in response secreted TNF and IL1 (Antin and Ferrara, 1992). These cytokines could significantly augment T-cell activation, which in turn stimulated TNF, IL1 and IFNY secretion from an array of immune cells. In that sense, the degree of
priming of immune cells determined the severity of GVHD. In contrast, tissue damage in itself without the interaction of CTL was capable of inducing GVHD damage (Teshima et al., 2002).

TNF was the most important cytokine in the effector phase (Reddy and Ferrara, 2003). It had synergistic and pleiotrophic effects, causing cachexia, induce APC maturation, recruited T-effector cells, neutrophils and monocytes, and primed homing by chemokine induction. TNF could also cause direct tissue damage by inducing apoptosis and necrosis, and could activate T-cells directly through the receptors TNFR1 (TNF receptor 1) and TNFR2 (TNF receptor 2).

Other effector molecules that had been studied include IL1 and nitric oxide (NO). IL1 had effects very similar to TNF, but might be more organ specific to spleen and skin (Abhyankar et al., 1993). NO is a product of activated macrophages that caused direct tissue insult, and inhibited repair mechanisms. Development of GVHD correlated with levels of oxidation products of NO (Nestel et al., 2000, Weiss et al., 1995).

More recently, larger studies of protein components, genetic expression and genetic polymorphism were aiming to dissect the pathobiology of GVHD further and identifying molecules (biomarkers) that could predict GVHD (Hansen, 2008, Kaiser et al., 2004, Mohty et al., 2007, Paczesny et al., 2008, Srinivasan et al., 2006, Weissinger and Dickinson, 2009).

## SUMMARY

- GVHD is the most important cause of adverse outcome of HSCT and remains largely unpredictable.
- Host antigen recognition by donor T-cells is the key step in the induction of GVHD.
- However, modulation of this process is very complex and involves pathophysiological events before, during and after Tcell activation.


### 1.3. The Genetics of HSCT

### 1.3.1. Self/non-self genetics

## Human Leukocyte Antigens (HLA)

The HLA complex is the strongest known determinant of self/non-self recognition. Six HLA loci are now commonly used for donor/recipient matching: HLA-A, B, C (HLA class I) and HLA-DRB1, DQB1 and DPB1 (HLA class II). Disparity between donor and recipient HLA antigens results in either rejection of the graft (host-versus-graft reaction), or cellular toxicity of the graft against the host (GVHD and GVL).

Well into the 1980's, HLA matching was based on serologic typing at antigen level. Observations from this time (Kernan et al., 1993) showed that unrelated HSCT had a higher prevalence of GVHD and worse survival, compared to related HSCT.

The introduction of DNA-based high resolution typing since the 1990's did contribute a great deal to the understanding of HLA matching. More than 2,000 different alleles had been identified within HLA class I and II (Shiina et al., 2009), and large scale registry studies, primarily in the US and Japan, had analysed the effects of different mismatch combinations in unrelated HSCT. For the US registry (Petersdorf et al., 1998), the first high-resolution data showed that in fact $47 \%$ of serologically matched HSCT (HLA-A, B, C, DRB1, DQB1) had one allele level mismatch, and 25\% had 2 mismatches. Combined mismatches in HLA class I and class II significantly increased the risk of severe GVHD and death. Single class II, but not class I mismatches increased the risk of GVHD. HLA-DRB1 mismatch was the strongest predictor of GVHD. The first data of highresolution typing from Japan were published in the same year (Sasazuki et al., 1998), coinciding with the US data that combined HLA class I and II mismatches carry the highest risk for GVHD and death. However, single class II mismatches did not increase the GVHD risk, whereas HLA-A mismatch had the strongest association with GVHD and death. HLA-C mismatch was also associated strongly with GVHD, but not with survival risk. These data were expanded in 2002 (Morishima et al., 2002), showing
that single mismatches in any of HLA-A, B, C, DRB1 implicated a higher risk of GVHD, with multiple class I mismatches, in particular involving HLAC , resulting in the highest risk. HLA-A and/or B mismatches increased the risk of death, with combined mismatches of HLA-A or B + HLA-C + HLADRB1 or DQB1 showing the poorest survival.
Updates on the US registry confirmed a high GVHD risk for a single or combined HLA-A mismatch, and worse survival with single or combined HLA-A, B, C and DRB1 mismatches (Flomenberg et al., 2004). Conversely, allele-level HLA-A, B, C and DRB1 matching had the best survival, while even a single mismatch of any of those had a measurable effect on survival (Lee et al., 2007). As in previous studies, no effect of HLA-DQB1 and DPB1 was found.

The Japanese registry has recently advanced into identifying individual GVHD high-risk allele mismatch pairs (Kawase et al., 2007), assuming that not all mismatches would actually induce alloreactivity. This study found 29 high-risk allele mismatch combinations in HLA-A, B, C, DRB1 and DPB1. Following on from this work, high risk and low risk mismatch combinations for relapse of haematological malignancies were identified and correlated with high-risk GVHD allele mismatches. Eight mismatch combinations in HLA-DPB1 and HLA-Cw were found that have a very low relapse risk and no increased GVHD risk, elucidating the HLA-basis of the GVL effect (Kawase et al., 2009). The Japanese registry was also the first to describe highly conserved HLA haplotypes and their association with risk or protection from GVHD (Morishima et al., 2010)

Other self/non-self genetics: KIR, LILR, mHags

Killer immunoglobulin-like receptors (KIR) are cell surface receptors on NK cells. Their function is to recognize normal MHC class I receptor expression on cells, hence normal MHC expression leads to inhibition of NK cell activity whereas an abnormal expression ('missing self') releases the inhibition and results in killing of the target cell by an apoptotic signal. KIR are highly polymorphic. In mismatched related HSCT for leukaemia, this effect (termed the KIR ligand mismatch) could be exploited for a graft-
versus-tumour effect without a higher risk of GVHD (Ruggeri et al., 2007, Leung et al., 2004). However, for the unrelated HSCT setting the data were more controversial. Yabe et. al. (Yabe et al., 2008) and Morishima et.al. (Morishima et al., 2007) had described the effects of HLA and KIR matching for the Japanese registry. These data show that KIR2DL ligand mismatch in the GVHD direction increased the risk for GVHD and mortality, but dependent on HLA matching, underlying malignancy and administration of ATG.

Leukocyte immunoglobulin-like receptors (LILR) have a wider distribution as compared to KIR, but also recognize MHC class I molecules (Sloane et al., 2004). As KIR, LILR are predominantly inhibitory and also highly polymorphic. They had a role in controlling the maturation of DC (Young et al., 2008), and so far unpublished data from the Japanese registry showed associations of LILR with GVHD and survival (verbal communication Toshio Yabe, Kouyuki Hirayasu).

Non-HLA recipient proteins that resulted from gene polymorphisms that were disparate between donor and recipient represented Minor Histocompatibility Antigens (mHags) (Spencer et al., 2010). Donor T-cells recognized such antigens and responded with clonal expansion. If a mHag happened to derive from a malignant protein or cell, the donor T-cell response could exert a strong and specific graft-versus tumour effect, which could be beneficial (Goulmy, 2006). However, if the mHag derived from otherwise healthy tissue, severe GVHD could be the result. In theory, the potential number of mHags could be as vast as the polymorphic disparity between donor and recipient (Brickner, 2006), however, only few mHags induced donor T-cell responses (immunodominance) for reasons that are not fully understood. Experimental matching for known mHags did not result in reduction of GVHD (Warren, 2009). Gene polymorphisms on the Y-chromosome of male recipients of female grafts were potentially a source of many mHags, and several have been identified; female into male HSCT had been recognized as having a higher risk of GVHD and mortality (Randolph et al., 2004).

More recent efforts had attempted to capture mHags with genome-wide approaches (Hansen et al., 2010, Ogawa et al., 2008, Kawase et al., 2008).

### 1.3.2. Non-HLA genetics

Non-HLA genetics in HSCT is defined as the effect of functional gene polymorphisms that impact on outcome by modulating existing immune or metabolic responses, rather than having direct involvement in self/non-self recognition. HSCT outcomes like acute and chronic GVHD, relapse and survival are not rare events and vary between individuals who may otherwise be genetically similar; possibly comparable to how phenotypes of inflammation, infection and immunity vary in a normal population. It was therefore assumed that the non-HLA genetic effects on HSCT outcome were determined by common genetic variants.

Studies that were aiming at understanding the pathophysiology of GVHD (see above) had identified several immunoregulatory key players, like e.g. cytokines, adhesion molecules, regulators of innate immunity and chemokines. The first such study was published in 1998, implicating TNF and IL10 with GVHD (Middleton et al., 1998).

For this study, a systematic literature search was undertaken that identified 248 gene association studies with outcome of HSCT (list: supplementary file 1.1). At least 105 genes, including cytokines, regulators of innate and adaptive immunity, drug metabolism genes, DNA repair and metabolic genes were found to associate with any HSCT outcome (which included acute and chronic GVHD, relapse, rejection, survival, VOD, infection, drug toxicity). These findings are summarized in table 1.1. Of these, markers for 49 genes associated with acute or chronic GVHD, described in 141 studies.

Analyzing the methodology of these studies, the vast majority ( $\mathrm{n}=238$, $96 \%)$ were single cohort candidate gene association studies. Only seven studies sought independent confirmation of findings within the same study's setup (Xiao et al., 2010, Chien et al., 2006, Lin et al., 2003, Bochud et al., 2008, Elmaagacli et al., 2009, Mullighan et al., 2004, Espinoza et al., 2011), and only three studies used larger scale approaches. Mullally et. al.
reported a study of 1143 SNP for 220 candidate genes, identifying several chemokines associating with HSCT outcome (Mullally et al., 2008). JMDP had conducted a genome-wide association study with SNP markers, which failed to identify any non-HLA gene association but detected a possible mHag locus (Ogawa et al., 2008). Finally, the NMDP carried out a genome-wide association study (Hansen et al., 2010) the results of which have not been finally reported. Imputation of SNP previously associated, however, confirmed association of IL10 and IL6 with GVHD (Chien et al., 2012).

Given the rapid evolution of HSCT, it had been difficult in the past to build large-scale HSCT study cohorts. Limited availability of study subjects made consideration of demographic or clinical risk factors in study cohort selection difficult, despite the existence of these risks being well established in the literature (e.g. patient and donor age (Kollman et al., 2001, Loren et al., 2006, Wojnar et al., 2006), female donor to male recipient (Randolph et al., 2004, Gahrton, 2007), diagnosis and staging (Chaidos et al., 2007, Wojnar et al., 2006), prior chemotherapy (Hahn et al., 2008), conditioning regimen (Perez-Simon et al., 2005, Hahn et al., 2008), concurrent infections (Hahn et al., 2008, Ljungman, 2007, Young, 2008)). Previous studies often relied on study populations displaying different underlying ethnicities, underlying diagnosis, stem cell sources (related/unrelated), conditioning regimens and GVHD prophylaxis, weakening study power and leading to disparate results. HLA matching and HSCT from sibling donors were the most common measures applied in the study of non-HLA gene polymorphisms, presuming that reducing the 'noise' from genetic mismatching would make small effect-size non-HLA association more readily identifiable. Very few studies deliberately chose unrelated or HLA mismatched HSCT, therefore there is a paucity of data on these settings, although these represent the majority of HSCT. Also, earlier serotypical HLA matches may have actually represented mismatches at allele level (Weisdorf et al., 2008), hampering the comparison of results from different studies. Many of the early studies in particular lacked statistical power for the allele/genotype frequency reported; sample sizes <100 were not uncommon.

More recently, the collection of large HSCT cohorts has become reality. Both NMDP and JMDP have now conducted $>10,000$ unrelated donor HSCT, allowing for future studies with better stratification of genetic, demographic and clinical risk factors. Eventually, the availability of such study populations would allow for study designs that comply with recommendations for the design of genetic association studies (Colhoun et al., 2003, Gambaro et al., 2000, Hirschhorn et al., 2002, McCarthy et al., 2008, Lander and Schork, 1994, Schork, 1997).

| Gene | function | gene | function | gene | function | gene | function |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| ABO | haematopoietic system | ESR | innate immunity | IL15RA | cytokine | NOS2A | innate immunity |
| ACE | other effector | Factor V | haematopoietic system | IL17A | cytokine | OGGI | DNA repair |
| BAFF | haematopoietic system | Fas | adaptive immunity | IL18 | cytokine | P2X7 | Drug metabolism |
| BP1 | other effector | FCGR2A | adaptive immunity | IL1RA | cytokine | PAI1 | haematopoietic system |
| CASP8 | adaptive immunity | FCGR3B | adaptive immunity | IL2 | cytokine | PARP1 | DNA repair |
| CCL27 | chemokines | FCRL3 | adaptive immunity | IL23R | cytokine | PCAM1 | adhesion |
| CCL3 | chemokines | FOXP3 | adaptive immunity | IL4 | cytokine | PGP | Drug metabolism |
| CCL4 | chemokines | GRZB | adaptive immunity | IL4R | cytokine | PIR | innate immunity |
| CCL5 | chemokines | GSTA1 | Drug metabolism | IL6 | cytokine | Prothrombin | haematopoietic system |
| CCR5 | chemokines | GSTM1 | Drug metabolism | IL7R | cytokine | PTPN22 | haematopoietic system |
| CCR6 | chemokines | GSTP1 | Drug metabolism | IMPDH1 | Drug metabolism | PTPRC | haematopoietic system |
| CCR9 | chemokines | GSTT1 | Drug metabolism | LCT | metabolic | RFC1 | DNA repair |
| CD14 | adaptive immunity | HFE | haematopoietic system | LIG3 | DNA repair | TGFB1 | cytokine |
| CD3EAP | DNA repair | HLA-E | adaptive immunity | MADCAM1 | adhesion | TGFB1R | cytokine |
| CD86 | adaptive immunity | HLA-G | adaptive immunity | Mal | adaptive immunity | TLR1 | innate immunity |
| CPS1 | metabolic | HMGB1 | adaptive immunity | MASP2 | innate immunity | TLR4 | innate immunity |
| CTLA4 | adaptive immunity | HO1 | haematopoietic system | MBL | innate immunity | TLR9 | innate immunity |
| CXCL12 | chemokines | HP | haematopoietic system | MBL2 | innate immunity | TNF | cytokine |
| CYP2B6 | Drug metabolism | HPA5 | haematopoietic system | MCP1 | innate immunity | TNFRSF1 | cytokine |
| CYP2C19 | Drug metabolism | HPSE | haematopoietic system | MDR1 | Drug metabolism | TNFRSF2 | cytokine |
| CYP3A4 | Drug metabolism | HSP70 | innate immunity | MIF | innate immunity | VDR | innate immunity |
| CYP3A5 | Drug metabolism | ICOS | adaptive immunity | MPO | innate immunity | VEGF | adhesion |
| DAAM2 | metabolic | IFNg | cytokine | MTHFR | Drug metabolism | VLA4 | adhesion |
| DARC | chemokines | IL1 | cytokine | MUTYH | DNA repair | XRCC3 | DNA repair |
| DECTIN1 | innate immunity | IL10 | cytokine | NKG2D | adaptive immunity |  |  |
| DNAM1 | adaptive immunity | IL12 | cytokine | NLRP2 | innate immunity |  |  |
| ERC2 | DNA repair | IL13 | cytokine | NOD2 | innate immunity |  |  |

Table 1.1 (previous page): Genes associated with HSCT outcomes from 248 previous gene association studies. Forty-nine genes (in bold) have been implicated with acute or chronic GVHD.

## SUMMARY

- The genetics of HSCT outcome, including GVHD, involves multiple genetic mechanisms.
- Self/non-self recognition strongly predicts GVHD.
- Matching of the Human Leukocyte Antigen (HLA) complex reduces the risk of GVHD and is widely applied in clinical practice.
- Other self/non-self recognition mechanisms (e.g. KIR, LILR, mHag ) also influence the risk of GVHD but are less well understood and not commonly used in clinical practice.
- Non-HLA gene polymorphisms modulate innate and adaptive immune responses, >100 genes have been reported to associate with HSCT outcomes.
- Results for non-HLA gene polymorphisms are often inconclusive due to limited study quality, therefore findings have been applied little in clinical practice.


### 1.4. Summary and conclusion; aim of this study

HSCT has become an ever more important treatment option for an ever wider variety of indications, now moving well beyond malignant and nonmalignant conditions of the haematopoietic system. The procedure itself and surrounding medical and nursing care has evolved dramatically, improving cure of the underlying disease and survival, whilst reducing transplant-related mortality and morbidity.

GVHD remains the single major hurdle in wider application of HSCT. Even full HLA matching from sibling donors cannot guarantee its prevention, indicating that other genetic and non-genetic factors are at play which we are just beginning to understand. The research into the KIR and LILR systems are likely to provide a better understanding of self/non-self recognition in the future. A small number of immunodominant minor histocompatibility antigens will probably give explanation for some strong GVHD and GVI effects. NonHLA gene polymorphisms determine the 'milieu' in which self/non-self recognition occurs and may therefore be of important influence on the strength of the immune responses leading to GVHD or GVT.

Despite an abundance of data from >200 previous studies, effects of many non-HLA gene polymorphisms remain inconclusive, which is founded in the generally small effect size of associations of common alleles and genotypes, and issues with study design (heterogeneous study populations, lack of statistical power, lack of validation by confirmatory study on a similar population).

The aim of this study is to elucidate the role of non-HLA gene polymorphisms for the risk of GVHD in a more robust fashion, by applying recommendations for high-quality gene association study design. Key elements of the study include:

- Study population: genetically homogeneous background, control of clinical confounders, clinically relevant population, adequate sample size
- Gene targets: More indiscriminate approach (genome-wide/targeted), rather than a candidate gene approach
- More stringent study design: screening and independent confirmation cohorts


### 1.5 Outline of study plan/brief history of the project

The idea of this study was conceived in 2004. The author's plan was to conduct a high-quality association study in the field of immunogenetics; the search for an adequate study population led to co-operation with institutions in Japan.

In 2005, the author spent six months at the Division of Molecular Life Sciences at Tokai University, Kanagawa, Japan, at the invitation of Professor Hidetoshi Inoko, a renowned expert in the field of HLA genetics. Tokai University hosts the DNA sample collection of JMDP and is involved in many of the registry research projects. During these six months, the author finalised a proposal for a genome-wide association study into the non-HLA genetics of GVHD using microsatellite markers, a methodology also pioneered at Tokai University. Dr Peter Middleton and Dr Andrew Gennery would act as supervisors for a PhD thesis with Newcastle University, whilst Professor Inoko would provide local supervision.

After funding for this work was obtained (JSPS post-doctoral fellowship, KayKendall Leukaemia Fund international fellowship), the work started in Japan in March 2007. An initial assessment of the availability of DNA samples (March-June 2007), however, showed that a genome-wide association study would be unfeasible due to the lack of samples with a sufficient amounts of DNA. The study plan was adjusted to a targeted genomic screening focusing on the immune system.

The selection of an appropriate study population received particular attention in order to control confounding variables (see chapter 2), and a small-scale pilot study (October-December 2007) using 41 candidate SNP markers was conducted to ensure that a study based on the selected study population would be capable of demonstrating small effect-size non-HLA gene polymorphisms. After this was achieved, the large-scale approach was carried out. Main steps included the identification of the genes of the immunogenome and allocation of microsatellite markers (July-December 2007), pooling of DNA samples (January-March 2008 and DecemberFebruary 2009), and genotyping of pooled DNA in a screening (April-

December 2008) and confirmatory cohort (March-August 2009), followed by data analysis and evaluation (until February 2010).

## 2. METHODOLOGY

### 2.1 Aim and purpose

2.2 Objectives
2.3 Study question and hypothesis
2.4 Overview of Study design
2.5 Selection of the study population
2.6 Selection of genes and markers
2.7 Preparation of DNA
2.8 Construction of DNA pools
2.9 Procedure of individual sample DNA PCR
2.10 Procedure of pooled DNA PCR
2.11 DNA genotyping
2.12 Data retrieval and processing
2.13 Data analysis

### 2.1 Aim and purpose of the study

The aim of this study was to improve health, survival and quality of life of recipients of matched unrelated donor HSCT for acute leukaemia by identifying genetic risk factors that increase or decrease risk of GVHD. This study aimed to identify 'risk' and 'protection' recipient and donor non-HLA genetic polymorphisms that contribute to the severity of acute GVHD, enabling clinicians to stratify the risk of GVHD prospectively.

Findings were expected to be applicable to future patient care:

- identified "risk" alleles in donors/recipients that predict the occurrence and severity of acute GVHD
- identification of 'risk' donors or recipients as a guide for planning conditioning and GVHD prophylaxis regimens
- identified "risk" genes in donors aid donor selection for reducing acute GVHD

Potential broader future scope of results:

- contribution to the understanding of genetic pathology of acute GVHD
- facilitating the development of new, specific prophylaxis and treatment options for GVHD (monoclonal antibodies, gene therapy)
- impact on research and management of similar or related inflammatory disorders


### 2.2 Objectives

### 2.2.1. Objective

The objective of this study was to investigate allele frequency differences of microsatellite markers between cases of absent or mild acute GVHD (grade 0-1, controls) and moderate to severe acute GVHD (grade 2-4, cases). As cases we considered HSCT recipients (,intrinsic' risk of GVHD) as well as donors (,graft' risk of GVHD).

### 2.2.2 Key objective elements

## Variables

Genetic polymorphisms presenting in the form of microsatellite allele frequency differences between the study groups of different severity of acute GVHD (grade 0-1 versus 2-4) separately for donors and recipients.

## Outcome parameters

Pooled DNA genotyping outcomes:
Moderate to severe acute GVHD (grade 2-4) was the single outcome of the two-step pooled DNA screening (discovery and confirmatory cohorts).
Standard definitions and classifications were used to clinically define acute GVHD - the modified Glucksberg criteria according to international consensus (Glucksberg et al., 1974, Rowlings et al., 1997, Przepiorka et al., 1995).

From a clinician's perspective, acute GVHD grade 2 was seen as a cutoff point for starting active and aggressive intervention to stop progress of acute GVHD (Hara et al., 2007). Acute GVHD grade 2 or more severe
was associated with increased morbidity and mortality, and reduced quality of life (Pasquini, 2008, Kanda et al., 2012, Kodera et al., 1999, Morishima et al., 2007, Sasazuki et al., 1998, Yano et al., 2000).

Individual genotyping: outcomes

- Degree of severity of acute GVHD (grade 0 versus 1-4, grade 0-1 versus $2-4$, grade $0-2$ versus grade $3-4$, grade $0-3$ versus 4 )
- Degree of severity of chronic GVHD (absent chronic GVHD versus limited and extensive disease, absent and limited disease versus extensive disease)
- 100 day/one year/three years/five years survival rate, log rank test for survival
- Relapse rate


### 2.3 Study Question and Hypothesis

### 2.3.1. Study Question

"Are non-HLA microsatellite polymorphisms in unrelated HSCT donor and/or recipient immune system genes associated with graft versus host disease?"

### 2.3.2. Hypothesis

The hypothesis was based on the concept that genetic susceptibility to GVHD is the result of a complex genetic trait, involving multiple loci:

- GVHD has in part a complex genetic trait, and that common allele polymorphism of non-HLA genes in the patient and donor genomes contribute to the development of GVHD; these risk alleles may be detectable with a systematic genome scanning approach.
- Such non-HLA risk alleles can have an effect size that reaches that of certain HLA mismatches, hence can be consistent despite variation in clinical and genetic risk factors over time. Such risk alleles can be useful for prediction of acute GVHD risk in clinical practice.

Hypothesis:
"Allele frequency differences of microsatellite markers are associated with moderate-severe acute GVHD."

Null hypothesis:
"Allele frequency differences of microsatellite markers are not associated with moderate-severe acute GVHD."

### 2.4. Overview of study design

### 2.4.1. Key features of a robust genetic association study design

Considering the above methodological issues about genetic association studies, as discussed in the introduction part of this thesis, a more robust design should entail:

- Study and control cohorts should stem from a genetically homogenous population
- Confounding variables needed to be well controlled
- Cases and controls needed to be well defined, phenotypes well established and graded
- Environmental factors that influence gene function well established
- Genes and markers selected that are biologically meaningful
- Outcomes well defined, consistently reported
- Study design addressing error by chance, multiple testing issues: Design with at least a discovery and independent confirmation cohort, appropriate rigorous statistics
- Sufficiently powered cohorts - adequate sample size
- Systematic rather than random/candidate marker approach


### 2.4.2. Measures to achieve a more robust study design

This study has taken measures to address the above issues by:

- Identifying of a study population from a more homogeneous background
- Control of confounding parameters: All known genetic, demographic and clinical risk factors were carefully analyzed.
- 'Modelling' of hypothetic cohorts were used to design a study cohort with ideally minimal confounders and maximum statistical power.
- Definition of phenotypes: Ensuring that GVHD in all subjects of this study was defined using the modified Glucksberg criteria (Przepiorka et al., 1995)
- Estimating impact of environmental factors
- Defining the scope of approach: Weighing the advantages and disadvantages of different options: candidate gene approach, targeted screening, whole genome screening.
- Addressing type I (incorrectly rejecting the null hypothesis) and type II (false acceptance of the null hypothesis) errors by considering issues of statistical power, reproducibility, multiple testing.


### 2.4.3. Outline of the study design

- Modification of a whole-genome scanning approach with microsatellite markers (Tamiya et al., 2005).
- Descriptive, retrospective case-control study with two nested cohorts (discovery/confirmatory) of pooled DNA screening, followed by individual genotyping of the combined cohorts for confirmation.
- Selection of a genetically and clinically homogenous cohort of approximately $\mathrm{n}=1000$ donor-recipient pairs (see power calculation) from the JMDP registry
- Microsatellite markers as screening tool - indiscriminate approach (i.e. large scale rather than candidate gene approach)
- Focus on immune system genes would be expected to yield targets with a higher positive predictive value than a more indiscriminate approach. A review of genome-wide association studies on immune system disorders had shown that genetic associations are more likely to be located in immunoregulatory genes (Zhernakova et al., 2009)
- Estimating impact of environmental factors by dividing the cohort into two subsequent time frames (1993-2000, 2001-2005). HSCT practice was likely to have changed during these time periods, not all of these changes may have been recorded in the dataset.
- Introduction of a two-step independent screening, estimate of statistical power, use of statistical correction for multiple testing.
- A phased, 2-step pooled DNA screening: Splitting of cohort of $n=1000$ pairs into two groups of approximately $\mathrm{n}=500$ pairs each. Within each group, construction of four DNA pools (Donors GVHD 0-1, Donors GVHD 2-4, Recipients GVHD 0-1 and Recipients GVHD 2-4) using an established, highly accurate DNA pooling method. Pooled typing of the full MS marker set in the $1^{\text {st }}$ screening step, followed by retyping of positively associated MS markers only in the confirmatory step, to eliminate pseudo-positive markers.
- Individual genotyping of remaining associated MS markers on the combined cohort of approximately $\mathrm{n}=1000$ pairs, to eliminate artefacts introduced by DNA pooling.
- Analyses planned for pooled genotyping: two directions of analysis:
o Donors GVHD 0-1 with Donors GVHD 2-4 ('intrinsic risk of donor to induce severe GVHD in recipient')
o Recipients GVHD 0-1 with Recipients GVHD 2-4 ('intrinsic risk of recipient to develop severe GVHD')
- Significance of allele frequency differences would be determined by Fisher's Exact Test for each individual marker allele ( $2 \times 2$ test) and for alleles of a marker ( 2 xm test).


Figure 2.1: Overview of the design of this study. In a first screening step (Discovery Cohort), microsatellite markers representing the entire immunogenome are typed on pooled DNA of 460 HSCT recipients and their donors; aiming to detect allele frequency differences between those recipients and donors of no or low grade GVHD (grade 0-1) and those of moderate to severe grade GVHD (grade 2-4). Positive markers only will be taken over to a second screening step (Confirmation Cohort) with an identical setup. Markers still remaining associated with GVHD will then be typed on all individuals of the combined discovery and confirmation cohorts.

### 2.4.4. Ethical approval

Ethical approval to this study was granted by the Ethics Committee of the School of Medicine of Tokai University, approval number No 02-4-1, 9 June 2006.

This study was also approved by a meeting of the Research Committee of the Japan Marrow Donor Programme (JMDP). Use of DNA samples and clinical data was explicitly granted. Donors and recipients have given written informed consent at the time of graft harvesting or transplantation for DNA samples and data to be used for research purposes, according to the declaration of Helsinki.

Summary - study design

- Study cohort from a genetically homogenous population with little/no admixture, controlled confounding parameters, well defined phenotypes and outcomes
- Sample size with adequate statistical power
- Confirmatory testing of identified associations in two independent cohorts
- Targeted genome scanning approach, focusing on biologically meaningful genes
- Using microsatellite markers in a pooled DNA typing approach


### 2.5. Selection of the study population

### 2.5.1. Analysis of JMDP registry population and cohort selection

The JMDP registry confirmed that all selected HSCT pairs were of Japanese origin - hence there was no genetic admixture other than that of the overall Japanese population.

For the purpose of selection of study cohorts, an opportunity was given to analyze data from the JMDP registry of unrelated donor HSCT performed between 1993 and 2000 ( $n=2469$ HSCT pairs) for risk factors of acute GVHD grade $0-1$ versus grade $2-4$. This was followed by 'modelling' of potential study cohorts according to the identified risk factors, aiming for a clinically meaningful study population with an optimal control of confounding variables. Given here is a brief summary; the details of the analysis are available in supplementary file 2.1.

Univariate and backward multivariate logistic regression analyses were performed on the registry population. Univariate analysis showed that Tcell depletion, antithymoglobulin (ATG), HLA matching, GVHD prophylaxis with a tacrolimus-based regimen, standard dose cyclophosphamide/total body irradiation (Cy/TBI) conditioning regimen, donor age $\leq 30$ years, and underlying diagnosis other than acute lymphoblastic leukaemia (ALL) or chronic myeloid leukaemia (CML) are all protective of acute GVHD grade 2-4. Multivariate analysis was performed by stepwise logistic regression, including all the variables showing associations in univariate analysis. ATG administration, HLA matching, GVHD prophylaxis with tacrolimus, donor age $\leq 30$ years and Cy/TBI standard regimen were upheld in the final step of multivariate analysis.

Based on these findings, four model cohorts were devised, by selecting out for the established risk factors in a step-wise fashion. Decision criteria included the results of multivariate analyses of the models, clinical meaningfulness of the model, and available sample size. Eventually, the most suitable model for the discovery cohort was a
selection by diagnosis (acute leukaemia: ALL and acute nonlymphoblastic leukaemia (ANLL)), recipient age ( $\geq 4$ years, $\leq 40$ years) and no T-cell depletion; resulting in a cohort of approximately 1000 sample pairs. This selection would include all degrees of HLA matching and mismatching, and therefore somehow reflect a population as typically seen in HSCT practice.

### 2.5.2. Power estimation

A statistical power estimation (conducted by Dr Hirofumi Nakaoka, details in supplementary file 2.2) showed that a sample size of approximately 500 pairs per cohort would be required to provide sufficient statistical power (0.8) to demonstrate effect sizes of an Odd's ratio of 1.5 at allele frequencies of 0.3-0.6. Larger effect sizes would detect lower frequency allele associations (approximately allele frequency of 0.1 for $\mathrm{OR}=2.0$; allele frequency of 0.05 for $\mathrm{OR}=2.5$ ). Associations with an OR between 1.3-1.5 represented the lower limit of detection for the screening and confirmatory cohorts.

### 2.5.3. Construction of the actual study cohorts

The next step in constructing the actual study cohorts was an assessment of DNA sample availability (supplementary file 2.3). This assessment showed that of the initial $n=1000$ sample pairs, $n=112$ were depleted of DNA, and a further $\mathrm{n}=345$ sample pairs were transferred for use for another study. The DNA content of each available sample was determined by DNA quantification and multiplication by volume. This revealed that a genome-wide study would have been unfeasible because the vast majority of samples would not provide sufficient DNA. However, for a targeted genome scanning or a candidate gene approach enough samples would have been available. The options were discussed amongst all contributors and decided to opt for a targeted
genomic scanning (of immune system genes) in a two-step pooled DNA screening design. For the first step, samples of the 1993-2000 registry cohort would be included, for a second step samples from a later cohort (2001-2005) would be made available by JMDP.

Based on the estimates made of DNA amount required (see section 2.7.2. below), sample pairs were selected on the basis of DNA amount availability for both samples per pair (supplementary file 2.3). $\mathrm{N}=460$ pairs were chosen from the 1993-2000 registry cohort for the discovery cohort (first screening step), following the criteria: Diagnosis (acute leukaemia: ALL and acute non-lymphoblastic leukaemia (ANLL)), recipient age ( $\geq 4$ years, $\leq 40$ years), full bone marrow as stem cell source and no T-cell depletion.

The same criteria were used for selection of the confirmatory cohort, this time incorporating HSCT pairs from the time period 2001-2005. The HLA matching of the confirmatory cohort were to be 'adjusted' to that of the discovery cohort by pairing of each sample pair for HLA matches and mismatches at the same HLA locus, or combination of loci.

The characteristics of this population are shown in supplementary file 2.3.

## SUMMARY - cohort construction

- Careful univariate and multivariate analysis of a large stem population (here: JMDP HSCT registry 1993-2000, n=2469 HSCT) can provide a good understanding of demographic, clinical and genetic risk factors for the intended outcome (here: GVHD grade 2-4).
- Designing of hypothetical cohorts with repeat univariate and multivariate analysis can optimize control of confounding variables whilst maintaining adequate sample size for good statistical power.
- The heterogeneous nature of HSCT, lack of large numbers of study subjects, and variability of HSCT management over time makes control of confounding variables difficult.
- The cohort construction of this study faced challenges over the availability of DNA samples, demanding a flexible approach to cohort and study design.
- The process resulted in devising of two study cohorts with well established confounders (i.e. diagnosis, HLA mismatch as the most consistent), and a strategy for a study design (confirmation by independent cohort).


### 2.6. Selection of genes and markers

### 2.6.1 Systematic identification of genes

As outlined above, it was decided to adopt an approach of a targeted genome scanning, focusing on genes that are biologically meaningful in the context of GVHD. The pathobiology has been studied in some detail (Ferrara et al., 2003, Duran-Struuck and Reddy, 2008, Sun et al., 2007). Key pathophysiological pathways are located within the immune system, or are driven by immune responses:

- Development and maturation of immune cells
- Innate immunity
- Adaptive immunity
- Lymphocyte receptor repertoires, MHC, tyrosine kinases, protein kinases
- Pattern recognition - Toll/like Receptors
- Effector pathways of lymphocytes - apoptosis
- Intracellular mechanisms
- Extracellular mechanisms (cytokines, complement, chemokines)
- Modifiers of immune responses

The approach was inclusive of genes for which some involvement with the immune system was described, rather than exclusive of genes without described immune system involvement; because such a role may not yet have been investigated for many genes. Exclusion on the basis of absent evidence would be more arbitrary then inclusion of genes for which such evidence exists; recognizing, however, that such an approach had its limitations as potentially relevant genes may not have been included. The minimum inclusion criteria for admission into the immunogenome panel were a functional role of the gene in immunoregulation (i.e. function of the transcribed gene product or a genetic variation of the gene) and belonging to the same gene family of
such a gene. Genes that would not fulfil these criteria would not be included in this study. As the search approach was inclusive and aimed at identifying those genes that were eligible rather than those that were not, there was no active process of exclusion. A literature search was carried out using defined search terms with the objective to compile a complete 'Immunogenome'. This literature search included general textbooks (Janeway et al., 2005, Mak and Saunders, 2006) and used a wide variety of databases to broaden and deepen the search, as well as to include the very latest information from recently published journal papers. Overall information on candidate genes were extracted from >2000 journal papers.

In addition, genes specifically linked to GVHD and other HSCT outcomes were traced and categorised:

- Genes associated with GVHD and other HSCT outcomes in previous studies
- Genes whose expression has been associated with GVHD and other HSCT outcomes
- Genes that have been associated, by identification of polymorphism or gene expression, with immune processes that are highly relevant also in the GVHD pathophysiology
- Genetic susceptibility loci of acute leukaemia (as potential confounders as these have an impact on survival)
- Genes that may have implications for GVHD or transplant outcomes in a broader sense, e.g. enzymes and other metabolic genes influencing immune responses, drug metabolism genes, DNA repair genes, etc.

Gene names were initially compiled in a list, which was standardised to current nomenclature by identifying the official gene symbol from the GeneCard and NCBI databases. Additional information, such as the exact genomic location, was retrieved. Literature searches were ongoing throughout the duration of the project until completion of genotyping work (February 2010) to include cutting-edge research and new
associations. The compiled 'HSCT-specific Immunogenome' included eventually $\mathrm{n}=3093$ genes (see supplementary file 2.4). Finally, this 'immunogenome' was compared with a similar collation reported in the previous literature (Ortutay and Vihinen, 2006), finding that genes from this previous study were included, but that this study's gene inclusion was far more comprehensive than that of the previous one.

### 2.6.2. Selection of MS markers

With regards to larger scale and genome wide studies Lander et.al. pointed out that genetically younger, isolated populations have larger haplotype blocks with wider linkage desequilibrium, requiring fewer markers, and predicatively less disease alleles (Lander and Schork, 1994).

MS markers for this study were selected from the existing panel of approximately 30,000 markers routinely used for whole genome association studies.

This marker panel was collated over several years by a research team at Tokai University devoloping the above mentioned genome wide scanning methodology (Tamiya et al., 2005) with MS markers. Almost 70,000 markers were typed on different East-Asian populations (Japanese, Mongolian, Korean), and eventually 30,000 markers selected that were highly polymorphic, had a limited average number of alleles $(6.4+/-3.1)$ and an average heterozygosity of $0.67+0.16$. According to haplotype block structure of East-Asian populations, which was well preserved in Japanese in particular, an average marker linkage disequilibrium (LD) of 100 kB was estimated. Markers were also selected by position, aiming to chart the entire euchromatic genome at regular 100 kB intervals, thus providing overlapping/double coverage for each LD region. For $95 \%$ of the genome marker coverage with intermarker distance of <200kB was achieved.

For this study, markers from this panel were selected that would flank the candidate gene to provide overlapping cover within the range of LD,
estimated to be approximately 100 kb . Hence, if the two flanking markers would be no more than 100 kb apart, full overlapping coverage for the locus would have been provided.

In this study, of the $\mathrm{n}=3,093$ target genes, $\mathrm{n}=184$ (6\%) had to be excluded because these were located in regions lacking appropriate microsatellites or their exact genomic location was unclear. For n=34 further genes not represented by the marker panel we identified suitable microsatellites and designed primers accordingly (appendix 2.2).
The final selection included $n=2,909$ genes. Because many of these genes were located in clusters at close proximity, $n=2,297$ target genes were selected as representative for the $\mathrm{n}=2,909$ genes (table 2.1). These gene loci were tagged with $n=4,321$ microsatellite markers (supplementary file 2.5). When measuring the distance between the centre of the gene and the marker start point, for $88 \%$ of target genes full overlapping coverage within a 100 kb range was identified. A further $8 \%$ of genes had partial coverage within estimated LD, with one or two flanking markers at >100 kb but <200 kb range. For 3\% of target genes the range of one or two microsatellite markers was >200 kb (table 2.2). Each target gene locus was tagged with an average of 1.8 microsatellite markers. There was a broad variation in the number of markers covering a gene locus, between a single flanking marker pair covering several candidate genes in regions of high gene density, and up to ten markers covering a single, very large gene. It was estimated that the total LD range of the selected markers taken together may cover up to $15 \%$ of the genome (table 2.3), hence cover substantially more genes than the selected target genes. From the number of genes within the LD range of 65 microsatellites (associated at an interim step in pooled screening) we extrapolated that our selection of microsatellite markers may have covered up to a third of all human genes.

|  | $\mathrm{N}=$ | $\%$ |
| :--- | ---: | ---: |
| Selected immunogenome genes | $\mathbf{3 , 0 9 3}$ | 100 |
| Unknown gene location/no marker | 184 | 6 |
| Immunogenome genes included in this <br> study | 2,909 | $\mathbf{2 , 4 8 1}$ |
| Target genes selected to represent <br> immunogenome of 3,093 genes | $\mathbf{1 0 0}$ |  |
| Included: genes covered with markers <br> from MS panel | $\mathbf{2 , 2 6 3}$ | $\mathbf{1 0 0}$ |
| Included: genes for which markers were <br> designed | -35 | 1.4 |
| excluded: unknown gene location | -149 | -1.4 |
| excluded: no MS marker available | $\mathbf{- 6 . 0}$ |  |
| Total target genes included in study, <br> representing 2,909 immunogenome <br> genes | $\mathbf{2 , 2 9 7}$ |  |

Table 2.1: Gene and marker selection. As many MS markers had several genes within their LD range, one 'target gene' was selected for each such MS marker. For 7.6\% of gene loci appropriate marker cover could not be established.

| Markers | markers | $\%$ |
| :--- | ---: | :---: |
| Total markers selected: | 4,321 |  |
| Total genes selected: | , 2297 |  |
| Markers per gene average: | 1.88 |  |
| Markers intronic: | 3,320 | 30.55 |
| Markers outside genes: | 3,801 | 69.45 |
| markers/genes covered with 2 flanking <br> markers within 100 kb range | 516 | 87.97 |
| markers/genes not covered with 2 <br> flanking markers within 100 kb range | 4 | 11.94 |
| markers exact location unknown | 4,321 | 0.09 |

Table 2.2: Specification of the degree of LD coverage of genes provided by the selected MS markers

|  | $\mathrm{n}=$ | $\%$ |
| :--- | ---: | :---: |
| Base pairs human genome | $3,164,700,000$ | 100 |
| LD cover 4321 MS markers | $432,100,000$ | 13.7 |
| Total number of genes human <br> genome | 30,000 | 100 |
| Estimated genes covered by <br> 4321 MS markers | 10,301 | 34.3 |

Table 2.3: Estimation of total gene coverage of the selection of MS markers for this study. Data on the number of base pairs and genes of the genome are from the Human Genome Project website. LD of MS markers assumes 100 kb , and the estimated number of genes covered by this selection is an extrapolation from the number of genes within the 100 kb LD range of 65 MS markers from this study.

Summary - genes and markers

- As GVHD is a disorder of immunoregulation, associated polymorphic genes are more likely to be located in the immune system. Targeting functional and structural genes of the immune system would be expected to yield a higher positive predictive value for such associations than a more indiscriminate approach.
- A genomic screening of the immunogenome is feasible at high density with gene-flanking microsatellite markers.
- Due to their long range linkage disequilibrium, microsatellite markers cover large genomic areas around the target genes.


### 2.7. Preparation of DNA

### 2.7.1 Provision of DNA samples

All DNA samples for this study were provided by JMDP, DNA preparation was not part of this study. In brief, samples were obtained from patients and donors at the time of HLA matching confirmation and stored for research purposes with appropriate consent. Tokai University is hosting the sample collection for JMDP.
Fresh samples were centrifuged and the buffy coat removed, from which the DNA of nucleated cells was extracted using commercial DNA extraction sets (QIAmp DNA blood extraction kit®, QIAGEN).

### 2.7.2 Estimation of DNA requirements for this study

The requirement of DNA amount was determined mainly by two factors: The number of planned reactions (i.e. the number of microsatellite markers, approximately $\mathrm{n}=4,000$ ), and the DNA concentration of the DNA pool aimed for. The latter one depended on the DNA samples with the lowest DNA concentration to be included, and had its limitation in the composition of the PCR mixture. The standard PCR mixture for this experiment had a total volume of $20 \mu \mathrm{l}$, of which $8 \mu \mathrm{l}$ was dedicated for the DNA. Dilution of the DNA sample was limited by the need for a restricted use of TE buffer (as >10 $\mu$ l buffer per well is known to inhibit the PCR reaction). In addition, if the total amount of DNA in the reaction was $<40 \mathrm{ng}$, PCR might become more unstable and the capability to detect small allele frequency differences of MS markers decreases (observations by Dr Akira Oka of his own experiments).

As the lowest sample concentrations was approximately $10 \mathrm{ng} / \mu \mathrm{l}$, the estimated final DNA pool concentration would be approximately 6-8 $\mathrm{ng} / \mu \mathrm{l}$, therefore total amount of DNA per reaction well is 48-64 ng. Thus, the amount of DNA required from each sample varied with the total
number of individual samples in the pool - the higher the number of individual samples, the lower the amount of DNA required from each individual as the total amount of DNA per marker plate well was constant. Based on the measurements of DNA concentrations, estimates of total amount per sample, and a preliminary estimate of total DNA requirements for the study, only samples with a total DNA amount of >4 $\mu \mathrm{g}$ were 'shortlisted' for the first pooled screening, and samples with an amount of $>2.5 \mu \mathrm{~g}$ for the second pooled screening (details of the estimation of DNA amounts required: appendix 2.3).

### 2.7.3 Measurement of DNA concentration

Amount of DNA available was likely to be limited, given the age of DNA samples, the large number of previous studies performed on this collection, and the large amount of DNA required for this study.
In order to preserve the collection for future work, it was agreed with JMDP that the total DNA amount of each selected sample would be assessed by measurements of concentration and volume. DNA concentration of each selected sample would be determined by an established standardized method for measurement of DNA concentration (PICO Green®), and volume would be estimated by visual comparison with a standard volume set (identical sample tubes with volumes in steps of $50 \mu \mathrm{l}$, ranging from $50-1000 \mu \mathrm{l}$ ).

For the DNA concentration measurement with PICO green® dsDNA quantification kit (Molecular Probes, P-7589) a DNA dilution of 1:200 was used according to the maker's instructions. In preparation, $5 \mu \mathrm{l}$ of original DNA was diluted in $995 \mu \mathrm{l}$ of $1 / 10$ TE buffer pH 7.5 . Of this solution, $100 \mu$ l was pipetted onto the measurement plate and incubated for 3 min with a $100 \mu \mathrm{l}$ of a $1 / 200$ solution of PICO green fluorescent reactant. For quality control, a set of different concentrations of a standard DNA of a known concentration was divised: 1/1000, 1/300, $1 / 100,1 / 30,1 / 10$ dilutions as well as a control well containing buffer with
no DNA. Three independent measurements were then obtained using a Flouroskan Ascent CF (Thermo Labsystems) photometer, with settings: Integration time 20 ms , filter pair: Excitation 485nm, Emission 527nm, normal beam, single measurement. A customized excel worksheet was used for quality control and calculation of concentrations.

### 2.7.4. Estimation of total DNA amount of sample, allocation of pairs to screening steps

Availability of DNA concentrations and sample volumes were allowing for a more exact estimation of total available DNA amount per sample. The total content of a DNA per sample in ng/ $\mu$ l was estimated by a simple formula:

Total amount of DNA/sample in $n g=D N A$ concentration in $n g / \mu l \boldsymbol{x}$ sample volume in $\mu \mathrm{l}$

Sample pairs were then ranked according to DNA amount in both partners in descending order, grouping them in three distinct groups:

- Total amount $>4,000 \mathrm{ng}$ : For inclusion into $1^{\text {st }}$ pooled screening
- Total amount $>2,500 \mathrm{ng}$ : For inclusion into $2^{\text {nd }}$ pooled screening
- Total amount >500 ng: For inclusion into individual typing
- Total amount <500 ng: exclusion from the study

Decision on inclusion and exclusion from the study, and allocation to pools were based on the estimated requirements, the total amount of DNA available per sample, and the original sample concentration. Agreement with JMDP on sample handling, in- and exclusion criteria:

- At least a minimum of 500 ng of DNA should remain in each sample, hence samples with an amount of DNA of 500 ng or less were excluded from the study.
- Samples with a total amount of $4,000 \mathrm{ng}$ or more were included into the first screening, pipetting $3,500 \mathrm{ng}$
- Samples with a total amount between 2,500 and $4,000 \mathrm{ng}$ were included into the second screening, pipetting 2000 ng
- From samples with a total amount between 1000 and $2500 \mathrm{ng}, 500 \mathrm{ng}$ were pipetted to store for inclusion into individual MS or SNP typing.


### 2.8 Construction of DNA pools

### 2.8.1. Considerations for definition of DNA pools

At the outset, decisions had to be made on defining the pools ('which samples to pool together?') and how large the intended pool was going to be. Key aspects here are:

- Sufficient statistical power of individual pool: The power calculation (see above) suggested that a minimum pool size should be in the range $n=200-250$ samples (at a cohort size of 400-500).
- The definition of pools should be clinically meaningful

Although it would have been desirable to pool different degrees of GVHD (e.g. grades 0, 1-2 combined, 3-4 combined) separately, resulting pool sizes would not have provided a sufficient statistical power. From the sample numbers available, separating degrees of GVHD into two groups seemed the only feasible option. Donors and Recipients were pooled separately but accordingly.
Three scenarios were considered:

- Grade 0 GVHD versus Grade 1-4 GVHD. This approach would have separated samples at a $\sim 50: 50$ proportion, in view of statistical power the strongest option. Drawback: Would not have distinguished between degrees of GVHD, but only presence of GVHD yes/no.
- Grade 0-1 GVHD versus Grade 2-4 GVHD: Would have separated samples at a $\sim 2 / 3$ : $1 / 3$ proportion. Resulting in acceptable power, and would have distinguished groups with a survival advantage from groups with survival disadvantage, also marking the stage of clinical intervention. Drawback: May not have distinguished risk genes for severe GVHD very clearly.
- Grade 0-2 GVHD versus Grade 3-4 GVHD: Would have separated samples at a $\sim 3 / 4: 1 / 4$ proportion. Advantage: Clearly would have
distinguished severe GVHD. Disadvantages: Reduced statistical power for severe GVHD group, would have ignored distinction by survival and point of clinical intervention.

The decision to choose the Grade 0-1 GVHD versus Grade 2-4 GVHD was based on:

- The aim of the study to provide a risk predicting tool for clinical decision making - determining which genetic risks separated recipients with favourable and poor survival perspective, and which genetic risks separated recipients requiring treatment intervention from those who did not.
- The acceptable statistical power for this option.


### 2.8.2. Existing methods of DNA pooling

Methods of DNA pooling and their accuracy compared to individual typing and family typing for the study of complex genetic diseases had been described previously (Shaw et al., 1998, Barcellos et al., 1997, Craig et al., 2005, Hoffjan et al., 2006).

Here a high-accuracy pooling method was applied that was a standard procedure for genome wide association studies (GWAS) in the same department (Tamiya et al., 2005, Oka et al., 2003, Collins et al., 2000, Daniels et al., 1998), which had been modified further to increase accuracy (unpublished, internal validation data available).

In some aspects the application of the pooling method differed from the application to WGA studies:

- This study cohort consisted of paired samples - therefore inclusion of both partners was essential to reflect allele frequency differences.
- Genome wide association studies (GWAS) with microsatellites rarely used pool sizes >200 samples - there were no data on pooling accuracy for such a pool size.
- DNA amount, concentration and quality was very variable - compared to previous studies, samples of this study had rather low DNA concentrations.

The initial preparation of samples consisted of measuring sample DNA concentration measured by the PICO green® method (described above).

Key features to ensure a high accuracy of DNA pooling were:

- The use of calibrated pipettes for all pipetting
- Repeat measurements with PICO green®, acceptance of a narrow variation margin ( $<5 \%$ ) only for DNA pooling


### 2.8.3. Practical procedure of DNA pooling

Sequential steps were involved in the pooling process. The pooling process was divided into four phases.

Phase 1 - Individual sample measurement

- Pipette testing (procedure described in appendix 2.4)
- Dividing samples into groups of the intended pools (see above)
- Ranking of samples by DNA concentration in decreasing order
- Dividing of the sample group into sub-groups of 96 -well plate format size
- Choosing a target concentration for each group. To pool equal amounts of DNA at equal concentration, by definition the final pool concentration was determined by the lowest sample concentration, towards which the pooling process aimed. The limitation of this
approach lay within a minimum concentration of $6 \mathrm{ng} / \mu \mathrm{l}$ required for pooled DNA PCR. Thus, any samples with a lower concentration had to be excluded; and caution had to be applied not to dilute low concentration samples too much to render them unsuitable. As higher concentrations ranged from $\sim 30-200 \mathrm{ng} / \mu \mathrm{l}$, a step-wise approach in dilution was applied, diluting the first group of samples to a target concentration of $\sim 25 \mathrm{ng} / \mu \mathrm{l}$.
- Dilution to this initial target concentration (using non-calibrated pipettes) and measurement of sample concentration by PICO green (the PICO green assay set up by using calibrated pipettes) using 1/10 TE Buffer pH 8.0. Three measurements were performed on the same plate, and an average concentration calculated.
- Ranking of tested samples in decreasing order of concentration. Exporting result file into excel to identify subgroups of samples that lie within a +/- $2.5 \%$ range of concentration - separation of these samples for small pool construction.
- The remaining samples of all subgroups combined were again ranked in decreasing order of concentration. A new subgroup of plate format size is identified, a new target concentration chosen, dilution and concentration measurements performed. Again this subgroup was ranked by sample concentration, groups for intermediate pool construction identified and separated.
- This procedure was repeated until all samples were allocated to intermediate pools. This procedure required 3-5 rounds until all or most samples were resolved. Concentration of the last small pools often approached required minimum concentration of the final DNA pool. Occasionally, a very few samples with borderline-low concentration were allocated into the final DNA pool, rather than one of the intermediate pools.


## Phase 2 - Construction of intermediate pools

- Using the calibrated pipettes, between 5 and 13 intermediate pools were created from each group of samples representing a DNA pool. The average concentration of samples considered for each intermediate was calculated as a fixed volume to be pipetted into the intermediate pool. The volume depended on the intended DNA amount for the final DNA pool.


## Phase 3 - Construction of final DNA pool

- The concentration of intermediate pools was assessed by PICO green measurements (each pool in three independent wells, three measurements per well, calculation of average)
- Careful, stepwise dilution of intermediate pools was applied to adjust these pools to an equal concentration within a range of $+/-2.5 \%$
- Intermediate pools were then pooled together using calibrated pipettes to achieve a final DNA pool


## Phase 4 - Adjusting final DNA pool concentrations

- In order to ensure comparable conditions among all pools (i.e. measurement of true allele frequencies), final DNA pools in each screening step were also adjusted to an equal concentration, amount and volume.
- DNA pools were assessed in the same way as intermediate pools by multiple, repeat concentration measurement.
- Concentration was adjusted by a cautious stepwise dilution (2-3 steps) towards the pool with the lowest concentration, accepting a range of no more than $+/-2.5 \%$.

DNA pooling represented the most technically difficult step of this project.
To keep a consistently high standard and avoid human error, these considerations were applied:

- Use of calibrated pipettes for all pipetting actions (except the initial sample dilution, see appendix 2.4). All calculations of DNA amount and sample volumes were based on the actual volumes measured in pipette testing, rather than the nominal volume on the pipette.
- Careful tracking of all pipetted volumes, correcting at each step for volumes abstracted for concentration measurements.
- DNA protection by aseptic working conditions, light protection
- Adequate mixing at spinning at each handling step
- DNA-saving approach: Dilution to low concentration, early pooling into intermediate pools to avoid DNA-consuming repeat measurements (the Tokai standard protocol dilutes all samples to a relatively low concentration within a narrow margin of variation +/- 5\%) before constructing intermediate pools of equal sample numbers). To compensate for the possibly induced increased variation, we decreased the margin of variation to $+/-2.5 \%$.
- The pooling strategy resulted in a number of intermediate pools with a wider range of concentrations - again, we compensated by applying a more narrow margin for concentration variation (+/-2.5\% for intermediate and large pools)
- Requirement for DNA top-up of low concentration samples for inclusion - for a small number of samples, top-up with highly concentrated original DNA was necessary to increase concentration to a level suitable for inclusion.

Results of the pooling process

Due to the requirement of highest attainable accuracy, pooling of DNA was a time consuming process. Construction of each DNA pool took between 3 and 6 weeks.

Table 2.4 summarizes the eight DNA pools constructed, and Figure 2.2 illustrates a typical process of pool building.

1st Screen

| Pool | average concentration $\mathrm{ng} / \mu \mathrm{l}$ | No. individuals | Pool volume (ml) | DNA amount pool (ng) | DNA amount per individual ( ng ) |
| :---: | :---: | :---: | :---: | :---: | :---: |
| D01 | 6.78688 | 276 | 82382.57 | 559120.6 | 2025.799 |
| D24 | 6.6247 | 184 | 90477.92 | 599389.1 | 3257.549 |
| P01 | 6.426204 | 276 | 83366.22 | 535728.4 | 1941.045 |
| P24 | 6.487866 | 184 | 83667.12 | 542821.1 | 2950.115 |

Average all
6.581412

2nd screen

|  | Average <br> concentration <br> $\mathrm{ng} / \mu \mathrm{l}$ | No. <br> individuals | Pool volume <br> $(\mathrm{ml})$ | DNA amount <br> pool (ng) | DNA <br> per <br> (ng) |
| :--- | :--- | :--- | :--- | :--- | :--- |
| Pool |  |  |  |  |  |

Average all 20.19441

Table 2.4: Constructed DNA pools for $1^{\text {st }}$ and $2^{\text {nd }}$ screening. Concentration variation for each screening step is within a $\mathbf{2 . 5 \%}$ margin of the average concentration. Note that samples of the first screening step were diluted previously and had therefore a lower initial concentration than samples of the $2^{\text {nd }}$ screening, explaining the difference in concentration.


Fig 2.2: Example of a 'pooling tree', summarising the pooling process (here for the Donor GVHD grade 0-1 pool of the $2^{\text {nd }}$ screening). Individual samples are concentration-adjusted and then pooled into small pools (here: 12 small pools). These small pools are then concentration-adjusted again and pooled together into intermediate pools. In a stepwise process one large pool results. Some individual samples of low initial concentration have to run alongside to be added to intermediate pools or even the final large pool.

### 2.8.4. Quality control of pooled DNA

Test marker for pooled/individual PCR and genotyping

Prior to embarking on pooled genotyping, individual DNA samples and pooled DNA were tested for accuracy with the applied methods by typing of a microsatellite test marker.

Objectives of test marker typing:

- To ensure appropriate PCR conditions
- To investigate if the different amounts of DNA lead to appropriate PCR results
- To define the optimal typing dilution
- To investigate whether the results of pooled PCR typing reflect results of individual typing (= quality check of DNA pooling process)

Microsatellite marker 066B03 was a standard test marker that had been used in most previous microsatellite genome-wide association studies from this laboratory to study the quality of DNA pools. It amplified well under standard pooled PCR conditions, and typed well under standard typing conditions. It had six major alleles, and reflected therefore an 'average' microsatellite.

Important variables of the PCR and genotyping process (details see below) were the amount of DNA used in the PCR mixture, and the dilution of resulting DNA product used for the genotyping process. Both of these factors could influence the fluorescent signal in genotyping, which determined the allele frequency in the pool. For this experiment, a high, medium and low level DNA setup was chosen (appendix 2.5) for the PCR procedure, and the PCR product diluted for each of these x 10 , x20 and x 40 . This was performed separately for each of the DNA
pools, and then compared with results from typing all 922 HSCT pairs individually.

The results of the test marker typing were summarized in the supplementary file 2.6.

Comparison of allele frequencies between pooled and individual genotyping showed a very high concordance between pooled DNA and individual sample typing. Using a high amount of DNA gave results most accurately reflecting allele frequencies seen in the individual typing. Dilution of the PCR product had relatively little influence on results, although we saw increasing inaccuracies with using the higher dilution.

Using high amount DNA and a PCR product dilution x20, the mean difference in allele frequency for the test marker in the discovery cohort was $0.45 \%$ (standard deviation (SD): -2.61 - 0.942\%, range -5.42$6.3 \%$ ); for the confirmatory cohort $-0.08 \%$ (SD: - $0.62-0.26 \%$, range -$1.16-0.82 \%)$. The pooled genotyping was able to pick up allele frequencies as low as 0.02 . There were no significant differences in allele numbers for frequencies of 0.05 and above, which we therefore used as a lower cut off value for reporting allele frequency in this study.

For the process of pooled DNA PCR and genotyping, a high amount of DNA (93 ng/reaction) for PCR and a PCR product dilution x20 was chosen.

### 2.9. Procedure of individual sample PCR

As a first step, two sets of individual typing master plates of individual samples were created. Using concentration data from DNA pooling, the volume required was calculated to pipet an amount of 100 ng of DNA (an amount estimated to cover all individual genotyping needs of this study). Matching volumes of $\mathrm{dH}_{2} \mathrm{O}$ were added manually to achieve a concentration of $1 \mathrm{ng} / \mu \mathrm{l}$.

Using a Beckman MultiMek pipetting robot (Beckman Coulter Inc), 50 $\mu \mathrm{l} / \mathrm{well}$ of this solution was separated onto yet another set of plates. As a final step, again using MultiMek, both sets of plate were diluted by pipetting $50 \mu \mathrm{l} /$ well of $\mathrm{dH}_{2} \mathrm{O}$ to a final concentration of $0.5 \mathrm{ng} / \mu \mathrm{l}$.

Sequence of Individual PCR and genotyping:

- A set of reaction plates was prepared in accordance with the individual typing DNA masterplates
- Using Beckman MultiMek, $2 \mu \mathrm{l}$ of DNA were transferred from the master plate onto the reaction plate
- A PCR reaction mixture for individual typing was prepared in a tube at $-30^{\circ} \mathrm{C}$ :
dH2O
ABI 10x Buffer
ABI 2.0 M dNTP
Roche AmpliTaq Gold 5U/ $\mu \mathrm{l}$
Primer Mix $10 \mu \mathrm{M}$ each
$5.45 \mu \mathrm{l} / \mathrm{well}$
$1.00 \mu \mathrm{l} / \mathrm{well}$
$1.00 \mu \mathrm{l} / \mathrm{well}$
$0.05 \mu \mathrm{l} / \mathrm{well}$
$0.5 \mu \mathrm{l} / \mathrm{well}$
- Using a reservoir, $8 \mu / / w e l l$ of this solution was pipetted into each well using MultiMek.
- PCR conditions on an ABI gene amplification system (Applied Biosystems) were used as described in the protocol (figure 5); here $56^{\circ} \mathrm{C}$ were applied as annealing temperature for both markers.
- Individual genotyping was identical to the protocol for pooled genotyping (see above), a dilution of x20 was used for individual plates.


## Individual DNA Typing <br> (PCR)

Marker:

Basic Mixture

| Ingredient | per well | Total (x1030) | Lot |
| :--- | ---: | ---: | ---: |
| $\mathrm{dH2O}$ | 5.45 | 5613.5 |  |
| $10 \times$ Buffer | 1 | 1030 |  |
| 2.0 mM dNTP | 1 | 1030 |  |
| Ampli Taq Gold $5 \mathrm{U} / \mathrm{ul}$ | 0.05 | 51.5 |  |
| Primer mix 10 uM each | 0.5 |  |  |
| DNA (0.5ng/ul) | 2 |  |  |
| sum | 10 | 7725 |  |

Main Marker Mixture

| Ingredient | x1000 wells |
| :--- | ---: |
| Basic mixture | 7500 |
| dH2O | 400 |
| primer F-100uM | 50 |
| primer R-100uM | 50 |
| sum | 8000 |

$-8 u l / w e l l$

Control Marker Mixture

Marker 1:
Marker 2:

| Ingredient | x12 wells |
| :--- | ---: |
| Basic mixture | 90 |
| dH2O | 3 |
| Primer mix 20M each) | 3 |
|  | 96 |

x2

PCR settings

| 95 C | 9 min |
| :--- | :---: |
| $*$ | 1 min |
| 72 C | 1 min |
| 96 C | 45 sec |
| $*$ | 45 sec |
| 72 C | 1 min |
| 72 C | 5 min |
| 4 C | $\infty$ |$\quad 40$ cycles

Figure 2.3: Protocol and worksheet for individual MS PCR

### 2.10. Procedure of pooled DNA PCR

### 2.10.1. Primer preparation

Primers for microsatellite markers (Sigma-Genosys, Japan) were extracted manually from master plates of the MS marker collection for genome-wide association studies onto plates specific for this study. 15 $\mu \mathrm{l}$ of $10 \mu \mathrm{Mol}$ primer mix were pipetted into each well. A set of 44 marker plates was created for the first screening. On the original master plates, markers were located in order of chromosomes and known typing requirements (PCR product dilution $x 10, \times 20, x 40$ ) to reduce typing error. This order was disrupted when creating plates for this study, resulting in a higher rate of typing error as compared to the previous studies.

### 2.10.2. Steps of PCR procedure

- For each marker plate, a set of four reaction plates representing the four pools was created.
- Onto each reaction plate, $2 \mu \mathrm{l}$ of primer mix was pipetted into each well using a multi-channel pipette.
- A PCR mix of sufficient volume for all four plates was prepared in a bottle kept at $-20^{\circ} \mathrm{C}$. Volumes per well:

AB 10x PCR Buffer $15 \mathrm{mM} \quad 2.0 \mu \mathrm{l}$
AB Gene Amp dNTP mix $2 \mathrm{mM} \quad 2.5 \mu \mathrm{l}$
Roche AmpliTaq Gold 5U/ul $0.1 \mu \mathrm{l}$

- Prepared reaction plates were set onto frozen metal block trays.
- PCR mix and pooled DNA ( $13.4 \mu / /$ well) were mixed manually in a sample tray after appropriate mixing and spinning down of the ingredients.
- $18 \mu \mathrm{l} /$ well of the PCR mix/DNA mixture was then pipetted onto the reaction plate under intense manual mixing with the primer in the bottom of the well.
- Plates were sealed, spun down and immediately amplified on a ABI DNA amplification system using a standard amplification protocol.
- Following amplification, PCR products were sealed and stored at $-30^{\circ} \mathrm{C}$ for further processing, usually the following day.


## PCR Procedure for Pooled DNA

1. Preparations

Get reagents (dNTP, buffer) and marker plates out of freezer
Switch on PCR Thermocyclers (AB GeneAmp PCR System 9700)
PCR PE plates: Label/ clearly distinguish by colour
Get 2 large ice boxes
25 ml tube for PCR mix - into ice box
X1 pipetting tray on frozen block - into ice box
Pipettes and pipette tips for corresponding volumes
2. Preparation of mixture

Mix and spin down reagents.
Prepare PCR mixture according to number of plates:

AB Gene Amp dNTP mix 2mM lot:
AB 10x PCR Buffer 15mM lot:
Roche AmpliTaq Gold 5U/ul lot:

| Marker plates |  | 1 plate | 2 plates | 3 plates | 4 plates |  |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| Total pool plates | 1 well | 4 plates | 8 plates | 12 plates | 16 plates | 1 plate |
| Ingredient | vol ul/well $\times 410$ wells $\times 820$ wells $\times 1230$ wells $\times 1640$ wells $\times 100$ wells |  |  |  |  |  |
| $10 \times$ buffer | 2 | 820 | 1640 | 2460 | 3280 | 200 |
| 2.0 mM dNTP | 2.5 | 1025 | 2050 | 3075 | 4100 | 250 |
| AmpliTaq Gold | 0.1 | 41 | 82 | 123 | 164 | 10 |
| Total PCR mix all pools 4.6 | 1886 | 3772 | 5658 | 7544 | 460 |  |
|  |  |  |  |  |  |  |
| DNA/each pool | 13.4 | 1340 | 2680 | 4020 | 5360 | 1340 |
| Primer Mix | 2 | NA | NA | NA | NA | NA |
| PCR mix/each pool | NA | 460 | 920 | 1380 | 1840 | 460 |
| Total each pool mix | 20 | 1800 | 3600 | 5400 | 7200 | 1800 |

## 3. Preparation of plates

Get ice trays for plates from freezer - put plates on trays.
First step: divide marker plate into the four plates of each set of pools. Pipet marker into the bottom of each well. Use same pipette tips for each row for all four plates. Cover with strong sticky sheet.
Second step: Add PCR mix by rows, mix well using one set of pipette tips per row. Cover with clear rubber sheets. Spin down.

| vol mix/DNA <br> per well: |  | 18 ul |
| :--- | :--- | :--- |
| Vol primer per <br> well: |  | 2 ul |
| Total per well: |  | 20 ul |

PCR Run (Applied Biosystems GeneAmp PCR System 9700)

| $96^{\circ} \mathrm{C}$ | 9 min |
| :---: | :--- |
| $57^{\circ} \mathrm{C}$ | 1 min |
| $72^{\circ} \mathrm{C}$ | 1 min |
| $96^{\circ} \mathrm{C}$ | 45 sec |
| $57^{\circ} \mathrm{C}$ | 45 sec |
| $72^{\circ} \mathrm{C}$ | 1 min |
| $72^{\circ} \mathrm{C}$ | 5 min |
| $4^{\circ} \mathrm{C}$ | $\infty$ |

hold

30
cycles
hold

Figure 2.4: Pooled DNA PCR protocol and worksheet

### 2.11. DNA Genotyping

### 2.11.1. Protocol for individual and pooled DNA genotyping

Pooled and individual DNA genotyping followed the same protocol. A stored PCR product was diluted, dried up and denatured before a run on a DNA analyzer.

Sequence of pooled DNA genotyping:

- Set up of ABI 3730 DNA analyzer (Applied Biosystems)- fresh reagents (AB 3730 Buffer (10x), Polymer, $\mathrm{dH}_{2} \mathrm{O}$ ) for each typing lot
- Preparation of a typing plate for each PCR product plate
- Dilution of PCR product $\times 20 / \times 40$ (reasoning see below) with $\mathrm{dH}_{2} \mathrm{O}$ using the Beckman MultiMek pipetting robot. Briefly, the robot prepared an intermediate mixing solution of a variable amount of $\mathrm{dH}_{2} \mathrm{O}$ and $2 \mu \mathrm{l}$ of PCR product; and pipetted an aliquot of $2 \mu \mathrm{l}$ (representing the desired target dilution) of that solution onto the prepared typing plate
- Dry up of diluted DNA in a SpeedVac vacuum centrifuge for 10 min at $45^{\circ} \mathrm{C}$
- Preparation of a mixture of 1 ml of standard Formamide with $5 \mu \mathrm{l} A$ GeneScan 500LIZ Size Standard per plate
- Pipetting $10 \mu \mathrm{l}$ of this solution by Eppendorf Multi-pipette into each well, sealing of plate
- Denature of plate for 3 min at $95^{\circ} \mathrm{C}$ on a ABI DNA amplification system, followed by 5 min cooling on an ice tray
- Running of plates on an ABI 3730 DNA analyzer in gene mapper modus (protocol: GM_in2kV10sec_RV7_RT3500). Usually, two DNA analyzers were run in parallel, one running donor plates and one running recipient plates, swapping over donors and recipients on a daily base to randomise technical artefacts introduced by the individual machine.

At laboratory peak times, part of the pooled DNA genotyping work (overall $<10 \%$ ) was carried out by Ms Yamaguchi and Miss Matsushita, two laboratory technicians at Tokai University, following the protocols established by the author and under his supervision, using PCR products created by the author. Approximately one third of the individual microsatellite marker genotyping was undertaken by Ms Higuchi, a further laboratory technician at Tokai University, mostly in summer 2010 (following the return of the author to the UK), using the established protocols, under guidance from the author and direct supervision by Dr Akira Oka.

## Pooled Genotyping - Run on ABI 3730 DNA analyser <br> Preparations

- Get PCR products, Formamide, 500 LIZ out of freezer/fridge
- Switch on RTV400
- Switch on L,R pumps
- Switch on Speed vac, open valve, check rubber ring, set temp $45^{\circ} \mathrm{C}, \mathrm{t}=10 \mathrm{~min}$
- Switch on multimek, set up: water, trays
- Switch on $x 4$ PCR Thermocyclers, set for heat $95^{\circ} \mathrm{C}$
- Prepare ABI 3730 DNA analyser:

1. remove and clean water/wast/buffer plates
2. Mix new buffer: 15 ml of 3730 buffer plus 135 ml aqua dest (total 150)
3. Replace water/waste/buffer trays - ensure buffer tray cable secured
4. Fill glass bottle with buffer
5. Check polymer
6. Close, buffer tray back in to place
7. Insert sample sheets: Run 3730 data Collection - Tree: Plate Manager - connect USB Import - Mark All - open - ok. Tree: Run Scheduler - advanced search (put in plate date) search - add all - done

PCR Product Dilution Dilution Factor :x10 x20 x40

- Spin down PCR product plates
- Prepare a set of PCR PE plates - label
- Prepare a set of NUNC plates - no label
- Prepare plate covers - strong for PCR prod, tissue for new plates
- Make dilution on multimek according to plan
- Cover new plates/PCR product plates, discard NUNC plates

Dry up

- Speed Vac temp $45^{\circ} \mathrm{C}$, $\mathrm{t}=10 \mathrm{~min}$

Add Size Standard, Formamide

- Preparation: Get 1 bottle of Formamide $(1 \mathrm{ml})$ for each plate
- To each 1 ml of Formamide add 5 ul of 500LIZ
- Mix well and spin down
- Use multipipette to pipette 10 ul of Formamide/500LIZ into each well
- Cover with strong cover
- Spin down


## Denature

- Get 2 Ice boxes
- Switch on PCR thermocylers
- Denature for 3 min at $95^{\circ} \mathrm{C}$, Ice cooling for 5 min

Prepare 3730 run:

- Spin down
- Remove strong plate cover
- Put into 3730 plate tray, grey rubber cover
- Slot in according to order
- Start run - green arrow

Figure 2.5: Protocol/worksheet for Pooled PCR genotyping

### 2.12. Data input, retrieval and processing

### 2.12.1. Data input to 3730 DNA analyzer (Applied Biosystems):

- A Java-based application (SampleSheetMaker v1.0, Applied Biosystems) was used for data input for the genotyping process. This application produced a text file (.txt) that contained, besides the technical information for the run, details such as the marker name, amplicon size and allele size to facilitate further processing with the GeneMapper and MultiTyper softwares (see below).


### 2.12.2. Raw typing data retrieval and processing

- Peak signal quality was initially assessed using the 'capillary viewer' function in the Run 3730 Data Collection v2.0 software (Applied Biosystems®).
- Electrophoretic runs were analysed using the GeneMapper v3.5 software (Applied Biosystems®). In particular, peak signal and size standard quality were assessed. Settings for the analysis methods were such that the optimum analysis quality was achieved for peak height between 2000 and 11,000 fu (fluorescent units), with lowest recognition level at 200 fu and highest at 30,000 fu.
- Raw data were retrieved from GeneMapper in the form of 'fsa' (per well) and 'ser' file (containing the data analysis per plate)
- These data were combined using a Java-based application from ABI, called 'Fsa2Fsb'.


### 2.12.3. Assessing peak heights

- Fsa2Fsb files were imported into the MultiPeaks software (Applied Biosystems®), a further Java-based application. This
application could visually display allele size and peak heights of MS markers, based on the data input information, and allowed selection and logging of marker peaks.
- Within Multipeaks, peaks were selected manually in a simultaneous display of all four pools for each screening step. Information provided included the peak height (fu), the allele size (in base pairs), and the allele frequency (in \%). As markers names were coded by their plate location in the institutes primer stock (e.g. '136A04'), there was a blinding to the analyzer as to which candidate gene he was assessing.
- All markers were assessed by the same analyzer in at least triplicate at different time intervals, in order to reduce intraobserver variability.
- As a general rule, alleles with a frequency $<5 \%$ in all pools were excluded as such frequencies represent the limits of technical resolution and statistical power of this study. Allele frequencies $<5 \%$ were occasionally selected if the other three pools had a consistent frequency of this allele $>5 \%$, and if the quality of the allele signal was appropriate (>200fu).
- Microsatellite allele selection involved an element of judgment, consisting in recognition of a particular microsatellite pattern (size of repeat units) and certain known microsatellite artifacts. The repeated microsatellite analysis lead to a 'training effect' in the observer, with the result that almost all microsatellite patterns were recognized and alleles determined. The strategy for resolving unclear typing results involved:
o re-analysis by the same observer at a different time
o retrieval of repeat size information of the marker and attempt to identify the microsatellite pattern
o re-analysis by a different observer experienced in microsatellite analysis
o re-typing/re-PCR of the same marker and re-analysis
o exclusion of marker if no meaningful analysis can be obtained
o Using a source reference of MS graphs from previous studies, as archived in the laboratory.
- In almost all cases it was possible to identify the MS by its typical peak pattern and by the base pair distance between peaks. If the MS identification was unclear, one or more independent opinions from other experienced researchers in the team were sought.
- For individual genotyping, the software Multityper (Applied Biosystems ${ }^{\circledR}$ ) presented peak height data in a similar way for single datasets only.
- Results of chosen peaks were stored in a text file. Text files had to be manually edited (elimination of null and error well) before statistical analysis.


### 2.12.4. Genotyping artefacts

Microsatellite genotyping has a number of inherent artefacts which could affect the number of alleles, or allele allocation measured (Olejniczak and Krzyzosiak, 2006, Matsumoto et al., 2004, Miller and Yuan, 1997).

- Stutter alleles: these are artificial peaks that derive from product amplification one to two repeat units shorter than the correct sized PCR product or allele. The reason for this effect is slippage of Taq polymerase on the repeated sequence. In pooled typing, these stutter peaks are included with the correct sized alleles one or two units shorter, and therefore not identifiable. Stutter is more prevalent in dinucleotide repeats. Identification of stutter alleles requires at least some individual
typing to identify and quantify the stutter effect. A number of mathematical methods have been devised to analyze stutter alleles. The stutter effect is usually marker specific, very consistent and reproducible between pools, therefore it is not thought to influence consistency of pooled DNA typing by e.g. mimicking allele frequency differences. Nevertheless, it can lead to a wrong estimate of allele frequencies in pooled as well as individual genotyping.
- "+A peaks": An artificial fragment created by DNA polymerase adding a non-templated nucleotide at the 3 ' end of the DNA fragment. This results in artefacts one base pair longer than the true allele for each peak, true or stutter allele. These artefacts are often recognisable as a parallel pattern 'shifted' from the true microsatellite pattern by one base pair, and would become apparent on visual inspection of the peak graph.
- Differential amplification: Preferential amplification of a shorter allele or PCR product. Reason: larger alleles reanneal at a faster rate because of more repeat units), resulting in reduced PCR efficiency. Short sized PCR products, artefacts ("starter peaks") as well as short-sized alleles, can be grossly overestimated and lead to false-positive results. This is not a consistent effect and tends to vary with each genotyping hence the repeated independent typing (eight pools, if a marker passed through both screenings) is likely to have reduced a large proportion of such artefacts.
- Compound/interrupted microsatellite repeats, areas of gene copy number variation: As our marker panel is highly selected for informative markers, such microsatellites have largely been excluded previously.

Individual genotyping of markers that would remain associated with GVHD outcome would reveal the majority of such artefacts.

### 2.13. Data Analysis

### 3.13.1. Preparation of data

Data from text files, specifying the marker name, the allele positions and peak heights, were fused together at the level of the individual pools.

For the analysis a custom-made data analysis tool was used that had been applied in genome-wide microsatellite studies previously (Tamiya et al., 2005). The functions of this tool were:

- Conversion of peak height signals into allele frequencies by a mathematical algorithm on the basis of numbers of pool size and numbers of cases and controls
- Calculation of allele frequency differences
- Calculation of p-values using two types of Fisher's exact test for the $2 \times 2$ contingency tables for each individual allele and the $2 x m$ contingency tables for each locus, where $m$ referred to the number of marker alleles observed in a population. The Markov chain/Monte Carlo simulation method was employed to execute the Fisher's exact test for the $2 x m$ contingency table (Tamiya et al., 2005).

After the first screening a large number of false positive markers and alleles was expected (each marker has 2-20 alleles - statistically there is a 1:20 by-chance association of each allele). There was a deliberate non-application of multiple testing statistics at this point to retain a high sensitivity for small effect-size association. Measures for identifying those markers and alleles that have a consistent association with GVHD included:

- Direct comparison of associations between the first and second screening by $p$-value and direction of Odds Ratio (protective or risk in both screens consistent)
- Careful inspection of peak graphs of the remaining consistently associated markers to identify typing errors and artifacts
- Individual genotyping of those markers showing the highest technical quality and statistical consistency. Test-typing on a small number of samples in the first instance may identify errors relating to the pooled genotyping process and artifacts.

Analysis of individual genotyping would be conducted in SPSS for Windows v 17.0 (IBM®), including:

- Ensuring Hardy-Weinberg Equilibrium (http://genepop.curtin.edu.au/help input.html)
- Significance tests: 2-sided Fisher's Exact test, Kaplan-Meyer Analysis for alleles and genotypes
- Application of Bonferroni's correction for multiple testing
- Multivariate analysis (multiple logistic regression, Cox regression) in SPSS


## 3. Exploration study

3.1. Introduction
3.2. Aims, hypotheses, objectives and study design
3.3. Materials and methods
3.4. Results
3.5. Discussion

### 3.1. Introduction

Evidence from a large number of previous studies showed that non-HLA gene polymorphisms had an impact on the risk of HSCT outcomes, such as acute and chronic GVHD, relapse and survival. However, the review of the literature also showed that very few of these associations were of larger effect size or consistent amongst studies in different ethnic populations or clinical settings. Examining quality criteria of genetic association studies it emerged that a more stringent design, involving a discovery or screening cohort and an independent confirmation cohort, was necessary.

Having analyzed a large cohort of HSCT donor and recipient pairs, which encompasses almost all unrelated donor HSCT through JMDP in Japan between 1993 and 2000, an understanding of demographic, clinical and genetic risk factors within this population permitted the construction of a study cohort with improved control of confounding variables (supplementary file 2.2). Before embarking on a larger scale scanning of the extended genomic areas, which committed large resources, it was useful to test the study cohort with known determinants of HSCT outcome, such as SNP and MS markers that showed strong results in previous studies.

### 3.2. Aims, hypotheses, objectives and study design

### 3.2.1. Aims

This exploratory study had the aim to confirm or refute previously identified SNP associations with HSCT outcomes, which include acute GVHD, chronic GVHD, relapse and survival.

### 3.2.2. Hypotheses

A study population exists that allowed the identification of non-HLA genetic associations in a consistent fashion across two independent cohorts, even if the effect size of the association was low.

Polymorphisms in non-HLA genes are associated with HSCT outcomes like acute GVHD, chronic GVHD, relapse and survival.

### 3.2.3. Objectives

The objectives of this study were:

- Testing of a panel of SNP and MS markers previously associated with HSCT outcome, as a confirm/refute approach in a Japanese population
- Variables: Genotypes of polymorphic SNP and MS markers
- Outcomes: acute GVHD (grade 0 versus grade 1-4, grade 0-1 versus grade $2-4$, grade $0-2$ versus grade $3-4$, grade $0-3$ versus grade 4), chronic GVHD (no cGVHD versus limited and extensive disease, no cGVHD and limited disease versus extensive disease), relapse (yes versus no), survival (Kaplan-Meyer analysis)


### 3.2.4. Study design

This was a case-control study with a two-step screening/confirmation approach. A population was defined by modelling hypothetical cohorts with different risk factors and assessing them by multivariate analysis, and the
model with the least clinical confounding chosen (supplementary file 2.2). Selected markers were identified from the previous literature. Standardised laboratory methods were applied to PCR and genotyping. Statistical methods include Bonferroni's correction for the number of included markers, and an additional measure of effect size (previous studies showed that associations with an OR<0.5 or >2.0 have a higher likelihood of being consistent, regardless of p-value).

### 3.3 Materials and Methods

### 3.3.1. Population

Donor and recipient HSCT pairs were selected from the JMDP registry of unrelated HSCT. We chose pairs with a diagnosis of acute leukaemia. These form the largest subgroup within HSCT. Cohorts represented 2 samplings of the same national pool, taken from two distinct timeframes (1993-2000, 20012005). Inclusion criteria were diagnosis (acute lymphoblastic leukaemia, ALL; acute non-lymphoblastic leukaemia, ANLL), age (4-40 years), conditioning (myeloablative), and stem cell source (bone marrow). All transplants were Tcell replete and received GVHD prophylaxis with either cyclosporin A or tacrolimus with methotrexate and corticosteroids. Analysis of the source as well as the selected HSCT population showed that HLA mismatching, donor age and GVHD prophylaxis regimen (cyclosporin A versus tacrolimus) were the only confounders remaining significant in multivariate analysis (data not shown here).
All donor-recipient pairs were HLA-typed retrospectively to allele level at six loci (HLA-A, HLA-B, HLA-C, HLA-DRB1, HLA-DQB1, and HLA-DPB1). The distribution of HLA matching of the confirmatory cohort was adjusted to that of the screening cohort by matching each sample of the screening cohort with a confirmatory cohort sample of the same HLA class or HLA class combination according to the previous literature (Sasazuki et al., 1998, Morishima et al., 2002) and our own analyses of risk matches/mismatches within this study population (data not shown).
Table 4 in supplementary file 2.3 shows the demographic and clinical characteristics of the selected cohorts. There was no statistically significant difference between the cohorts in the baseline demographic criteria. Tables 5 and 6 in supplementary file 2.3 specify the degree of HLA matching and mismatching. For reasons of comparison, we have used the NMDP/CIBMTR classification of HLA matching (Weisdorf et al., 2008). According to this classification, 357 HSCT pairs have a 8/8 (HLA A, B, C, DRB1) highresolution allele match, 331 (35.9\%) are partially matched (1 mismatch within
these HLA loci), and 234 (25.4\%) are mismatched (two or more mismatches within these HLA loci). Considering the HLA DQ and DP loci also, only 78 HSCT pairs ( $8.5 \%$ ) had a $12 / 12$ allele match. In Japanese, HLA A, B, and C mismatches are associated with risk of acute GVHD. HLA C mismatches, however, have a protective effect on relapse (whilst HLA A, C and B mismatches associate with a risk of death) (Sasazuki et al., 1998, Morishima et al., 2002, Morishima et al., 2007). More recent research has focused on specific allele mismatches, rather than mismatches in loci, aiming to identify non-permissive mismatches for acute GVHD (Kawase et al., 2007) or protective mismatches against relapse (Kawase et al., 2009), as well as risk HLA haplotypes for GVHD (Morishima et al., 2010).

### 3.3.2. Gene and SNP marker selection

Selection of candidate markers was based on a search of the published literature on genetic associations with HSCT outcomes. As the TaqMan® SNP genotyping platform was used, selection was limited to markers for which standard assays were available for this system.
For some genetic loci the same markers that were associated in other populations were non-polymorphic in Japanese (NOD2, TGFB1). The HapMap data base (www.hapmap.org) was used to identify haploTag SNP for these loci. The SNP markers included in this study are detailed in table 3.1.

### 3.3.3. Genotyping

Assay information of the used genotyping assays below is supplied in appendix 3.1.

TaqMan genotyping

TaqMan® SNP genotyping assays (Applied Biosystems, Branchburg, USA) were applied for 38 selected SNP according to the maker's instructions.

Individual genomic DNA ( $2.1 \mathrm{ng} /$ well) was arranged onto 384 -well plates (EDR-384 SII) and samples dried up at $45^{\circ} \mathrm{C}$ in vacuum over 90 minutes. A 40x reaction mixture, consisting of $2 \mu \mathrm{l} /$ well of TaqMan Universal Master Mix, No Amp Erase ${ }^{\circledR}$ UNG (Applied Biosystems, Branchburg, USA), $0.05 \mu \mathrm{l} /$ well of 40x TaqMan® SNP genotyping assay mix, and $1.95 \mu / /$ well of purified water, was pipetted robotically directly onto the dried-up 384 well plates. Plates were covered with Micro Amp Optical Adhesive Film 4311971 (Applied Biosystems, Branchburg, USA) before running a polymerase chain reaction (PCR) on a Gene Amp PCR System 9700. PCR conditions consisted of an initial cycle of $95^{\circ} \mathrm{C}(10 \mathrm{~min}), 40$ cycles of $92^{\circ} \mathrm{C}(15 \mathrm{sec})$ and $60^{\circ} \mathrm{C}(1 \mathrm{~min})$, and a final cooling to $4^{\circ} \mathrm{C}$.

Runs were analysed on a 7900HT Sequence Detection System (Applied Biosystems) using the SDS 2.1 (Applied Biosystems) software. Results were exported as txt files into Microsoft office excel for compilation and further processing. Genotypes were allocated considering the forward/reverse direction of the primer, and checked against the genotype distribution of each SNP for a Japanese population in HapMap (www.hapmap.org). HardyWeinberg Tests were carried out using the genepop tool (http://genepop.curtin.edu.au/genepop op1.html).

## Luminex genotyping of IL10 SNP

The IL-10 promoter SNPs rs1800872 (-592A/C), rs1800871(-819T/C), and rs1800896 (-1082A/G) were genotyped by PCR-SSO using Luminex MultiAnalyte Profiling system (xMAP) (Luminex Corp., Austin, TX).A fragment of IL10 promoter region, containing SNPs, $-592,-819$, and -1082 , was amplified by PCR using 5' biotinylated primers (supplementary table 1). The PCR product was denatured and hybridized with the mixture of the six oligonucleotide probes, specific for each base of the corresponding biallelic SNP, immobilized on fluorescent coded microsphere beads. The hybridization mixture containing the PCR product, hybridization buffer (Wakunaga Pharmaceuticals, Hiroshima, Japan), beads mixture and SAPE (Wakunaga Pharmaceuticals, Hiroshima,

Japan) were incubated at $55^{\circ} \mathrm{C}$ for 30 minutes. After washing, the hybridized product was analyzed on the Luminex 100.

### 3.3.4. Statistical analysis

Genotype results were imported into SPSS Statistics v 17.0 (SPSS Inc). Because little is known about effects of non-HLA polymorphisms in HLAmismatched populations, we used three analytic approaches in order to identify significant associations: 2-sided Fisher's Exact Test (95\% confidence intervals) with Bonferroni's correction for significance testing, Odd's Ratio ( $95 \%$ confidence intervals) as a measure of effect size, and independent testing in a confirmatory cohort (without application of multiple testing correction). Variables were the three individual genotypes, and mismatch between donor and recipient genotypes. Outcomes were acute GVHD (0-4), acute GVHD grades 2-4, acute GVHD grades 3-4, acute GVHD grade 4, chronic GVHD, extensive chronic GVHD, relapse, death (overall, at 100 days/1 year/3 years) and survival (as log rank test in Kaplan Meier analysis). For the screening cohort we considered as significant a $p$-value of $p=0.05$ with Bonferroni's correction for the number of SNP markers tested. As the $p$-value is not a good surrogate marker for effect size, and often small in HSCToutcome association studies, associations showing Odd's Ratios of $\leq 0.5$ and $\geq 2.0$ (this follows observations of OR's of significant markers in previous studies) were included separately.
Screening and confirmatory cohort data were analysed on the overall cohort in the first instance. In order to reduce confounding by HLA mismatching, we conducted identical analyses on a subgroup with a higher degree of HLA matching ( $8 / 8$ allele matching at the HLA A, B, C, DRB1 loci, with additional exclusion of combined HLA-DQB1 and DPB1 mismatches; allowing for either a HLA-DQB1 or a HLA-DPB1 mismatch only), similar to previous reports from JMDP(Ogawa et al., 2008), resulting in cohorts of 160 (discovery) and 166 (confirmatory) pairs.
For the screening cohort, we would genotype all 41 chosen SNP markers (table 1) on both donor and recipient cohorts, and conduct overall and
subgroup analyses. Markers only that show a corrected $p$-value of $<0.05$ and/or an OR of $\leq 0.5$ and $\geq 2.0$ in either the overall or the subgroup analyses would be selected for confirmatory typing. If a marker that showed an association that was persisting applying when Bonferroni's correction, we tested other associations of the same marker in the confirmatory cohort even if these would not reach the multiple testing thresholds, in order to capture borderline significance or effect size of genotypes, building on the strength of testing in an independent confirmatory cohort.
Given the high degree of linkage between the CTLA4 as well as the IL10 SNPs in the study, unambiguous haplotypes could be determined directly without recourse to computational methods.

As the distribution of acute GVHD degrees of severity was significantly different between the screening and confirmation cohort, all associations with acute GVHD as outcome were re-analysed after randomising the study population into two different cohorts (using an online based tool for random assignment:
http://www1.assumption.edu/users/avadum/applets/RandAssign/GroupGen.ht ml).

Multivariate analysis was performed on the combined cohorts using STATA v 11.0. Odds ratio (OR) of acute GVHD for the selected SNP in multivariate analysis was estimated by a multivariate logistic regression analysis with the adjustment for recipient and donor ages, underlying diagnosis, the use of TBI, ATG, female donor into male transplant, GVHD prophylaxis (tacrolimus versus cyclosporin A), relapse and HLA mismatch to address possible confounding.

| Target gene | SNP | Target gene | SNP |
| :---: | :---: | :---: | :---: |
| CCL4 | rs2634508 | NOD2 | rs1077861 |
| CD86 | rs1129055 |  | rs1861757 |
| CTLA4 | rs231777 |  | rs1861759 |
|  | $\begin{array}{\|l} \hline \text { rs231775 (CTLA4- } \\ 49) \\ \hline \end{array}$ |  | rs6500328 |
|  | $\begin{aligned} & \text { rs3087243 (CTLA- } \\ & \text { CT60) } \end{aligned}$ |  | rs2111234 |
| FAS | $\begin{aligned} & \text { rs1800682 (FAS- } \\ & 670 \text { ) } \end{aligned}$ |  | rs2111235 |
| FCGR2A | rs1801274 |  | rs7203344 |
| HLA-E | $\begin{array}{\|l} \hline \text { rs1264457 (HLA-E } \\ \text { R128G) } \\ \hline \end{array}$ |  | rs17313265 |
|  | rs1800795 | TGFB1 | $\begin{aligned} & \text { rs1800469 (TGFB1- } \\ & 509) \\ & \hline \end{aligned}$ |
| HSP70/hom | rs2075800 |  | rs2241715 |
| IFNg | rs2069705 |  | rs2241716 |
| IL1A | $\begin{aligned} & \text { rs1800587 (IL1A- } \\ & \text { 889) } \\ & \hline \end{aligned}$ |  | rs4803455 |
| IL1B | rs16944 (IL1B-511) | TLR4 | rs12377632 |
| IL2 | $\begin{aligned} & \text { rs2069762 (IL2- } \\ & 330) \end{aligned}$ |  | rs1927907 |
| IL10 | $\begin{array}{\|l\|} \hline \text { rs1800896 (IL10- } \\ \text { 1082) } \\ \hline \end{array}$ | TNF | rs361525 (TNF-238) |
|  | $\begin{aligned} & \text { rs1800871 (IL10- } \\ & \text { 819) } \\ & \hline \end{aligned}$ |  | rs1799964 (TNF-1031) |
|  | $\begin{aligned} & \text { rs1800872 (IL10- } \\ & 592) \end{aligned}$ |  | rs1800629 (TNF-308) |
| IL15RA | $\begin{aligned} & \text { rs2228059 (IL15RA } \\ & \text { N182T) } \end{aligned}$ |  | rs1799724 (TNF-857) |
| IL23R | rs6687620 | TNFRSF1B | $\begin{aligned} & \text { rs1061622 (TNFR2 } \\ & \text { codon 196) } \\ & \hline \end{aligned}$ |
| MIF | rs755622 | $V D R$ | rs731236 |
| MTHFR | $\begin{aligned} & \text { rs1801133 (MTHFR } \\ & \text { C677T) } \end{aligned}$ |  |  |

Table 3.1: Selected candidate SNP markers of this study.

### 3.4. Results

### 3.4.1. Screening cohort

## All transplants ( $n=460$ pairs)

In the screening cohort, involving 460 bone marrow transplants performed between 1993 and 2000, 41 single nucleotide SNP markers were typed in both patient and donor cohorts. Of these, six markers were excluded from analysis, for technical (multiple clusters: rs1927907, rs4803455) and statistical reasons (minor allele frequency $<5 \%$ : rs1800795, rs6687620, rs361525, rs1800629). All 35 markers included in the analysis were in Hardy-Weinberg equilibrium (defined as $p$ value $>0.05$, with statistical correction for the number of tested markers).
Thirteen markers, plus the IL10 and CTLA4 haplotypes, showed an association with a HSCT outcome in the donor screening cohort (table 3.2). By significance testing applying Bonferroni's correction, only the marker IL101082 and the CTLA4 haplotype showed significant association, while three further markers were selected for confirmatory typing by their effect size (marker CTLA4 rs231775 also showed relevant effect size individually; marker CTLA4 rs231777, which showed no individual association, was included in the confirmatory cohort as part of the CTLA4 haplotype (not listed in table 3.2)). The recipient cohort (table 3.3) revealed 15 markers, plus the CTLA4 haplotype, that were associated with a HSCT outcome. The IL2-330 SNP and the CTLA4 haplotype revealed significant associations above the multiple testing thresholds, while five SNP markers had ORs $\leq 0.5$ and $\geq 2.0$.

## HLA-matched subgroup (n=160 pairs)

When analyzing the HLA-matched subgroups of these cohorts, 7 markers and the CTLA4 and IL10 haplotypes in the donor cohort (table 3.4) showed outcome associations, of which 5 markers and the CTLA4 haplotype were included for confirmatory typing. Only the CTLA4 haplotype had a p-value
significant when multiple testing correction was applied. In the HLA matched recipient subgroup, three markers showed an association with HSCT outcome, of which one was selected for the confirmation cohort by strength of OR (table 3.5).

### 3.4.2. Confirmatory cohort

## All transplants (n=462 pairs)

Seven markers for the donor cohort (CTLA4: rs231775, rs231777, rs3087243(included for forming the CTLA4 haplotype, only rs231775 and rs3087243 showed an association in the screening cohort); FAS: rs1800682; IL10: rs1800896; NOD2: rs2111235, rs6500328) and ten markers for the recipient cohort (CTLA4: rs231775, rs231777, rs3087243(part of CTLA4 haplotype, only rs231775 and rs231777 showed were associated in the screening cohort) ; FAS: rs1800682; IL2: rs2069762 ; NOD2: 17313265; TGFB1: rs2241716; TNF: rs1799964; TNFRSF1B: rs1061622) were selected for typing in the confirmatory cohort. Firstly; the aim was to confirm associations from the screening cohorts that had significant p-values after multiple testing correction (high significance), then associations that had ORs $\leq 0.5$ or $\geq 2.0$ (large effect size), and thirdly associations within these selected markers that were consistent in both screening and confirmatory cohort (independent cohort confirmation), regardless of multiple testing correction or effect size.

There were no consistent findings in the overall donor confirmatory cohort (table 3.2). In the overall recipient confirmatory cohort (table 3.3), the donorrecipient genotype mismatch of the TNF-1031 SNP (rs1799964) was consistently associated in both screening and confirmatory cohorts with a higher risk of severe acute GVHD (grade 4). The CC genotype of the same marker was associated with acute GVHD grade 4 in the screening cohort, and just escaped significance level in the confirmatory cohort ( $p=0.06$ ).

## HLA-matched subgroups (166 pairs)

In the donor HLA-matched subgroup (table 3.4), none of the markers typed in the confirmatory cohort showed any association. The HLA matched recipient cohort (table 5) revealed a consistent association between risk of chronic GVHD and the GT genotype of rs2069762 (IL2-330).

Table 3.6 summarises the consistent associations of this study, comprising the IL2-330 and TNF-1031 SNP.

| Gene | Marker | Discovery cohort - genotype \& association | Confirmatory cohort - genotype \& association |
| :---: | :---: | :---: | :---: |
| CTLA4 | rs231775 | AA aGVHD ( $p=0.0043$, OR: $=\mathbf{0 . 0 4 9}, \mathrm{Cl}: 0.028-0.083$ GG aGVHD ( $p=0.0071, \mathrm{OR}=1.90, \mathrm{CI}: 1.19-3.03$ | $\mathrm{n} / \mathrm{s}$ |
| CTLA4 | rs3087243 | GG aGVHD ( $p=0.0086, \mathrm{OR}=1.81, \mathrm{Cl} 1.18-2.78)$ | $\mathrm{n} / \mathrm{s}$ |
| CTLA4 | haplotype | CAA aGVHD ( $p=0.0025, \mathrm{OR}=0.59, \mathrm{Cl}: 0.42-0.82$ ) CGG aGVHD ( $\boldsymbol{p}=\mathbf{0 . 0 0 0 5 7 , \mathrm { OR } = 1 . 7 2 , \mathrm { Cl } : 1 . 2 7 - 2 . 3 4 \text { ) } ) ~ ( 1 )}$ | $\mathrm{n} / \mathrm{s}$ |
| FAS | rs1800682 |  | $\mathrm{n} / \mathrm{s}$ |
| IFNg | rs2069705 | $\begin{aligned} & \text { CC ext cGVHD }(p=0.035, \mathrm{OR}=0.57, \mathrm{Cl}: 0.33-0.96) \\ & \text { CC relapse }(p=0.04, \mathrm{OR}=0.60, \mathrm{Cl}: 0.37-0.96) \\ & \hline \end{aligned}$ | n/t |
| IL10 | rs1800896 | AA survival ( $p=0.001$ ) protective | $\mathrm{n} / \mathrm{s}$ |
| IL10 | haplotype | CCA survival ( $p=0.032$ ) protective | n/t |
| MTHFR | rs1801133 | CT cGVHD ( $p=0.03, \mathrm{OR}=0.63, \mathrm{Cl}=0.42-0.96$ ) | n/t |
| NOD2 | rs17313265 | CT survival ( $p=0.012$ ) risk CC survival ( $p=0.008$ ) protective | $\begin{aligned} & \mathrm{n} / \mathrm{t} \\ & \mathrm{n} / \mathrm{t} \end{aligned}$ |
| NOD2 | rs2111235 | TT aGVHD4 ( $p=0.016, \mathrm{OR}=\mathbf{0 . 3 3}, \mathrm{Cl}: 0.14-0.80$ ) | $\mathrm{n} / \mathrm{s}$ |
| NOD2 | rs6500328 | GG ext cGVHD ( $p=0.011, \mathrm{OR}=\mathbf{0 . 1 7 , \mathrm { Cl } : 0 . 0 2 3 - 0 . 7 8 ) ~}$ | $\mathrm{n} / \mathrm{s}$ |
| TGFB1 | rs1800469 | CC aGVHD2-4 ( $p=0.035, \mathrm{OR}=1.69, \mathrm{CI}: 1.09-2.61$ ) CT aGVHD2-4 ( $p=0.036$, OR=0.66, CI: 0.45-0.96) | $\begin{aligned} & \mathrm{n} / \mathrm{t} \\ & \mathrm{n} / \mathrm{t} \end{aligned}$ |
| TGFB1 | rs2241715 | GG aGVHD2-4 ( $p=0.047$, OR=1.64, CI: 1.06-2.53 GT survival ( $p=0.03$ ) protective <br> GT ext cGVHD ( $p=0.032, \mathrm{OR}=0.57, \mathrm{Cl}: 0.34-0.94$ ) <br> GT aGVHD2-4 ( $p=0.037, \mathrm{OR}=0.67, \mathrm{Cl}: 0.46-0.98$ ) | $\begin{aligned} & \text { n/t } \\ & \mathrm{n} / \mathrm{t} \\ & \mathrm{n} / \mathrm{t} \\ & \mathrm{n} / \mathrm{t} \end{aligned}$ |
| TNF | rs1799964 | TT relapse ( $p=0.041, \mathrm{OR}=1.71, \mathrm{Cl}: 1.04-2.82$ ) | $\mathrm{n} / \mathrm{t}$ |
| TNF | rs1799724 | CC survival ( $p=0.014$ ) protective | n/t |

Table 3.2 (previous page): Results of SNP genotyping on all donor samples. Explanation of abbreviations (apply also to the the donor HLA matched and recipient results tables): aGVHD= acute GVHD, aGVHD4= acute GVHD grade 4, aGVHD 2-4= acute GVHD grade 2-4, cGVHD= chronic GVHD, ext cGVHD= extensive chronic GVHD, survival= p-value for log rank test as explored by Kaplan-Meyer analysis, mismatch= genotype mismatch between donor and recipient, $p=p$-value by 2 -sided Fisher's Exact Test, OR= Odds Ratio, Cl= 95\% confidence intervals for OR, n/s= non-significant, n/t= not tested. Bold: Withstanding Bonferroni's multiple testing corrections or have OR $\leq 0.5$ or $\geq 2$, italic: consistent associations. Marker rs231777 had no individual association and is therefore not included in this table, but was included into the confirmatory cohort as part of the CTLA4 haplotype.

## Recipient - All

| Gene | Marker | Discovery cohort - genotype \& association | Confirmatory cohort - genotype \& association |
| :---: | :---: | :---: | :---: |
| CTLA4 | rs231775 | AA cGVHD ( $p=0.046, \mathrm{OR}=1.83, \mathrm{Cl}$ : 1.02-3.28) | $\mathrm{n} / \mathrm{s}$ |
| CTLA4 | rs231777 | mismatch aGVHD ( $p=0.004, \mathrm{OR}=1.91, \mathrm{Cl}: 1.24-2.96)$ | $\mathrm{n} / \mathrm{s}$ |
| CTLA4 | haplotype | CAA cGVHD ( $p=0.011$, OR=1.5, CI=1.11-2.03); CGG cGVHD ( $\boldsymbol{p}=\mathbf{0} .0013$, $\mathrm{OR}=0.62, \mathrm{Cl}: 0.47-0.83$ ) CGG aGVHD2-4 ( $p=0.019, \mathrm{OR}=0.70, \mathrm{Cl}: 0.52-0.94$ ) TAG aGVHD4 ( $p=0.0071, \mathrm{OR}=3.71, \mathrm{Cl}: 1.56-8.86$ ) | $\begin{aligned} & \mathrm{n} / \mathrm{s} \\ & \mathrm{n} / \mathrm{s} \\ & \mathrm{n} / \mathrm{s} \\ & \mathrm{n} / \mathrm{s} \end{aligned}$ |
| FAS | rs1800682 | CC relapse ( $p=0.017, \mathrm{OR}=1.68, \mathrm{Cl}: 1.03-2.74$ ) <br> CT relapse ( $p=0.0025, \mathrm{OR}=\mathbf{0 . 5 0}, \mathrm{Cl}: 0.33-0.78$ ), <br> CT aGVHD ( $p=0.009$, OR=1.79, CI: 1.15-2.77) <br> TT cGVHD ( $p=0.024, \mathrm{OR}=1.75, \mathrm{Cl}: 1.03-2.82$ ) <br> TT ext cGVHD ( $p=0.014$. OR=1.74, CI: 1.03-2.94) | $\begin{aligned} & \mathrm{n} / \mathrm{s} \\ & \mathrm{n} / \mathrm{s} \\ & \mathrm{n} / \mathrm{s} \\ & \mathrm{n} / \mathrm{s} \\ & \mathrm{n} / \mathrm{s} \end{aligned}$ |
| HLA-E | rs1264457 | mismatch survival ( $p=0.023$ ) risk | n/t |
| IL1A | rs1800578 | mismatch aGVHD2-4 ( $p=0.026, \mathrm{OR}=1.69, \mathrm{CI}$ : 1.11-2.56) | $\mathrm{n} / \mathrm{t}$ |
| IL1B | rs16944 | AA aGVHD ( $p=0.048, \mathrm{OR}=0.63, \mathrm{Cl}: 0.39-0.99$ ) GG aGVHD ( $p=0.032, \mathrm{OR}=1.75, \mathrm{Cl}: 1.08-2.82$ ) | $\begin{aligned} & \mathrm{n} / \mathrm{t} \\ & \mathrm{n} / \mathrm{t} \end{aligned}$ |
| IL15RA | rs2228059 | AC survival ( $p=0.024$ ) risk | n/t |
| IL2 | rs2069762 | GG aGVHD4 ( $\boldsymbol{p}=\mathbf{0 . 0 0 1 4 , ~ O R = 4 . 5 1 , ~ C l}: 1.91-10.6$ ) GT survival ( $p=0.0021$ ) protective, TT survival ( $p=0.0061$ ) risk | $\begin{aligned} & \mathrm{n} / \mathrm{s} \\ & \mathrm{n} / \mathrm{s} \\ & \mathrm{n} / \mathrm{s} \end{aligned}$ |
| NOD2 | rs17313265 | CC aGVHD2-4 ( $p=0.036, \mathrm{OR}=2.15, \mathrm{Cl}$ : 1.06-4.37) | $\mathrm{n} / \mathrm{s}$ |
| TGFB1 | rs1800469 | mismatch aGVHD2-4 ( $p=0.02$, OR=1.63, $\mathrm{Cl}: 1.1-6.4$ ) | n/t |
| TGFB1 | rs2241715 | mismatch aGVHD2-4 ( $p=0.015$, OR=1.61, CI: 1.09-2.39) mismatch cGVHD ( $p=0.035, \mathrm{OR}=1.58, \mathrm{Cl}: 1.04-2.41$ ) | $\begin{aligned} & \mathrm{n} / \mathrm{t} \\ & \mathrm{n} / \mathrm{t} \end{aligned}$ |
| TGFB1 | rs2241716 | AA ext cGVHD ( $p=0.0041, \mathrm{OR}=2.58, \mathrm{Cl}: 1.36-4.87$ ) | n/s |
| TNF | rs1799964 | mismatch aGVHD4 ( $p=0.022$, OR=2.53, Cl:1.16-5.53) CC aGVHD4 ( $p=0.041$, OR=4.92, CI:1.27-19.02) | mismatch aGVHD4 ( $p=0.0053$, OR=3.40, Cl:1.48-7.81) CC aGVHD4 trend ( $p=0.06$ ) |
| TNF | rs1799724 | CC survival ( $p=0.02$ ) protective, CT survival ( $p=0.02$ ) risk | $\begin{aligned} & \mathrm{n} / \mathrm{t} \\ & \mathrm{n} / \mathrm{t} \\ & \hline \end{aligned}$ |
| TNFRSF1B | rs1061622 | TT aGVHD4 ( $p=0.023, \mathrm{OR=4.69}$,Cl 1.1-20.11) | $\mathrm{n} / \mathrm{s}$ |

Table 3.3 (previous page): Significant Results of SNP genotyping on all recipient samples. Explanations of abbreviations please see table 2. The marker rs3087243 was not associated individually with chronic or acute GVHD and is not listed here, but was included in the confirmatory cohort forming part of the CTLA4 haplotype.

| Donor HLA |  |  |  |
| :---: | :---: | :---: | :---: |
| Gene | Marker | Discovery cohort - genotype \& association | Confirmatory cohort - genotype \& association |
| CTLA4 | rs231775 | GG aGVHD ( $p=0.026, \mathbf{O R = 2 . 0 2 , ~ C I : ~ 1 . 0 9 - 3 . 7 5 ) ~}$ | $\mathrm{n} / \mathrm{s}$ |
| CTLA4 | rs3087243 | GG aGVHD ( $p=0.021, \mathrm{OR}=1.97, \mathrm{Cl}$ : 1.11-3.50) | $\mathrm{n} / \mathrm{s}$ |
| CTLA4 | haplotype | CAA aGVHD ( $p=0.012, \mathrm{OR}=0.55, \mathrm{Cl}: 0.35-0.87$ ) CGG aGVHD ( $\boldsymbol{p}=\mathbf{0} .00097, \mathrm{OR}=2.06, \mathrm{CI}: 1.22-5.94$ ) | $\begin{aligned} & \hline \mathrm{n} / \mathrm{s} \\ & \mathrm{n} / \mathrm{s} \end{aligned}$ |
| IFNg | rs2069705 | CC ext cGVHD ( $p=0.036$, OR=0.42, CI:0.20-0.93) CT ext cGVHD ( $p=0.017$, OR=2.69, Cl: 1.22-5.94) | $\begin{aligned} & \mathrm{n} / \mathrm{s} \\ & \mathrm{n} / \mathrm{s} \end{aligned}$ |
| IL10 | rs1800896 | AA aGVHD ( $p=0.038, \mathrm{OR}=\mathbf{0 . 2 1 , ~ \mathrm { Cl } : 0 . 0 4 - 0 . 9 6 \text { ) }}$ | $\mathrm{n} / \mathrm{s}$ |
| IL10 | haplotype | CCG aGVHD ( $p=0.027$, OR=4.70, Cl:1.08-20.54) | $\mathrm{n} / \mathrm{s}$ |
| MTHFR | rs1801133 |  | n/t |
| NOD2 | rs17313265 | CT relapse ( $p=0.013, \mathrm{OR}=\mathbf{2 . 6 8}, \mathrm{Cl}: 1.02-7.09$ ) | $\mathrm{n} / \mathrm{s}$ |
| TNF | rs1799724 | CC survival ( $p=0.006$ ) protective | n/t |

Table 3.4: Results of SNP genotyping on HLA-matched donor samples. Explanations of abbreviations please see table 2.

Recipient - HLA

| Gene | Marker | Discovery cohort - genotype \& association | Confirmatory cohort - genotype \& association |
| :---: | :---: | :---: | :---: |
| FAS | rs1800682 | CT aGVHD ( $p=0.0024, \mathrm{OR}=0.39, \mathrm{Cl}=0.22-0.71$ ) | $\mathrm{n} / \mathrm{s}$ |
| IL1B | rs16944 | AA aGVHD ( $p=0.043, \mathrm{OR=0.51}, \mathrm{Cl:0.27-0.97} \mathrm{)} \mathrm{{ }}^{\text {a }}$ ( ${ }^{\text {a }}$ ( | n/t |
| IL2 | rs2069762 | GT survival ( $p=0.037$ ) protective <br> GT cGVHD ( $p=0.039, O R=1.97, C /=1.05-3.71$ ) <br> TT survival ( $p=0.039$ ) risk | $\mathrm{n} / \mathrm{s}$ GT $\boldsymbol{c G V H D}(p=0.00041, O R=3.24, C I: 1.69-6.20)$ $\mathrm{n} / \mathrm{s}$ |

Table 3.5: Results of SNP genotyping on HLA-matched recipient samples. Explanations of abbreviations please see table 2.

| marker | $\begin{array}{\|c} \hline \text { genotyp } \\ \text { e } \\ \hline \end{array}$ | cohort | outcome | $p=$ | total | cases all | control s all | cases <br> pos | cases <br> neg | control <br> s pos | control s neg | Odds ratio | OR CI (95\%) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| TNF-1031 | mismatch | Screening | aGVHD 4 | 0.022 | 448 | 28 | 420 | 12 | 16 | 96 | 324 | 2.53 | 1.16-5.53 |
| $\begin{array}{\|l} \text { rs1799964 } \\ \text { recipients (all) } \\ \hline \end{array}$ | mismatch | Confirmation | aGVHD 4 | 0.0053 | 460 | 24 | 436 | 12 | 12 | 99 | 337 | 3.40 | 1.48-7.81 |
| IL2-330 | GT | Screening | cGVHD | 0.039 | 160 | 72 | 88 | 39 | 33 | 33 | 55 | 1.97 | 1.05-3.71 |
| rs2069762 <br> recipients <br> (HLA matched) | GT | Confirmation | cGVHD | 0.00041 | 166 | 75 | 92 | 40 | 35 | 23 | 68 | 3.24 | 1.70-6.20 |
|  | GG | random 1 | aGVHD | 0.022 | 159 | 58 | 101 | 20 | 38 | 54 | 47 | 0.46 | 0.27-0.78 |
| rs3087243 <br> donors (HLA matched) | GG | random 2 | aGVHD | 0.045 | 166 | 53 | 11 | 22 | 31 | 67 | 46 | 0.49 | 0.29-0.83 |

Table 3.6: SNP markers showing significant association in recipient screening and cohorts.

### 3.4.3. Further analyses

To understand the mechanism of the associated genotype, the analysis was extended to all IL2-330 genotypes and chronic GVHD outcomes in the confirmatory cohort, and it was found that GT also associated with extensive chronic GVHD ( $p=0.00022$, OR: $5.18, \mathrm{Cl}: 2.37-11.39$ ). The TT genotype exerts a protective effect against extensive chronic GVHD ( $p=0.0029$, OR: 0.3, $\mathrm{Cl}: 0.13-0.67)$. This finding was replicated when combining screening and confirmatory cohorts (GT and extensive chronic GVHD: p=0.00055, OR: 2.90, $\mathrm{Cl}: 1.74-5.08$; TT and extensive chronic GVHD: $\mathrm{p}=0.001$, OR: 0.40 , $\mathrm{Cl}: 0.23-$ 0.71 ), suggesting that the GG genotype is likely to be the higher risk genotype. No significant association was found with the GG genotype, which was likely due to limited statistical power of this low frequency genotype. Mirroring the analysis by MacMillan et. al. (MacMillan et al., 2003) in the combined cohorts, the $G$ allele showed a trend with risk of extensive chronic GVHD ( $p=0.07$ ), but not with acute GVHD.

The extended analysis of the TNF-1031 CC genotype in the confirmatory cohort showed that it was also associated with acute GVHD grade 2-4 ( $\mathrm{p}=0.029$, OR=3.41, $\mathrm{Cl}: 1.99-5.82$ ). The TNF-1031 donor-recipient genotype mismatch was found to be a risk factor for acute GVHD grade 2-4 $(p=0.003$, $\mathrm{OR}=1.93, \mathrm{Cl}: 1.13-3.30$ ) and grade $3-4(\mathrm{p}=0.002$, $\mathrm{OR}=2.21, \mathrm{Cl}: 1.13-3.80$ ) in the confirmatory cohort.

The stratification applied in 'matching' the degree of HLA mismatch of the confirmatory cohort to that of the screening cohort may have introduced a bias (significantly different distribution of acute GVHD grades, see supplementary table 1). In order to address this, samples were randomly assigned to two cohorts, resolving any significant difference between time frames, and acute GVHD as an outcome measure. Re-analysis of the data for acute GVHD outcomes showed that the genotype mismatch of the TNF-1031 SNP as a risk factor for acute GVHD grade 4 would still hold up as significant ( $p=0.005$, $\mathrm{OR}=3.26, \mathrm{Cl}: 1.91-5.58 ; \mathrm{p}=0.021$, $\mathrm{OR}=2.60, \mathrm{Cl}: 1.52-4.45)$. The CTLA4-CT60 (rs3087243) SNP showed a consistent association of the GG genotype as protective against acute GVHD ( $p=0.022$, $\mathrm{OR}=0.46$, $\mathrm{CI}: 0.27-0.78$; $\mathrm{p}=0.045$,
$\mathrm{OR}=0.49, \mathrm{Cl}: 0.29-0.83$ ) in the random cohort analysis of the HLA-matched subgroup.

### 3.4.4. Multivariate analyses

Multivariate analyses (tables 3.7-3.9) were performed on the combined (screening and confirmatory) cohorts and showed that the TNF-1031 donorrecipient genotype mismatch (acute GVHD grade 4), the CC genotype (acute GVHD grade 4), and the IL2-330 GT genotype (chronic GVHD) are independent risk factors, while the CTLA4-CT60 GG genotype is independently protective against acute GVHD.

| IL2-330: chronic GVHD | Univariate |  |  |  |
| :--- | :--- | ---: | :--- | ---: |
| Variable | OR $(95 \% \mathrm{CI})$ | -valtivariate |  |  |
| Recipient age | $1.008(0.99-1.03)$ | OR $(95 \% \mathrm{CI})$ | $P$-value |  |
| Donor age | $1.024(0.99-1.05)$ | 0.481 | $1.008(0.98-1.03)$ | 0.528 |
| Female to male transplant | $0.900(0.52-1.57)$ | 0.106 | $1.020(0.99-1.05)$ | 0.195 |
| Diagnosis ANLL vs ALL | $1.087(0.70-1.69)$ | 0.71 | $0.876(0.48-1.60)$ | 0.664 |
| Total body irradiation (TBI) | $1.419(0.72-2.80)$ | 0.711 | $1.022(0.63-1.67)$ | 0.929 |
| Cyclosporine vs tacrolimus | $1.024(0.66-1.59)$ | 0.313 | $1.284(0.62-2.67)$ | 0.502 |
| Relapse | $0.526(0.32-0.86)$ | 0.916 | $0.996(0.61-1.62)$ | 0.987 |
| Genotype GT | $\mathbf{2 . 5 0 7}(1.60-3.93)$ | 0.011 | $0.573(0.34-0.96)$ | 0.033 |

Table 3.7: Multivariate analysis of the IL2-330 GT genotype as risk factor for chronic GVHD in the HLA-matched subgroup. The genotype is an independent risk factor.

| CTLA4-CT60: acute GVHD | Univariate |  | Multivariate |  |
| :--- | :--- | ---: | :--- | ---: |
| Variable | OR $(95 \% \mathrm{CI})$ | P-value | OR $(95 \% \mathrm{Cl})$ | $P$-value |
| Recipient age | $1.017(0.99-1.04)$ | 0.146 | $1.020(0.99-1.05)$ | 0.121 |
| Donor age | $0.995(0.97-1.03)$ | 0.763 | $0.997(0.97-1.03)$ | 0.854 |
| Female to male transplant | $1.644(0.93-2.89)$ | 0.085 | $1.630(0.89-2.97)$ | 0.111 |
| Diagnosis ANLL vs ALL | $1.280(0.81-2.03)$ | 0.296 | $1.129(0.69-1.85)$ | 0.631 |
| Total body irradiation (TBI) | $0.847(0.43-1.68)$ | 0.634 | $0.916(0.45-1.86)$ | 0.809 |
| Relapse | $1.255(0.77-2.06)$ | 0.369 | $1.330(0.80-2.24)$ | 0.273 |
| Genotype GG | $\mathbf{0 . 4 6 8 ( 0 . 2 9 - 0 . 7 5 )}$ | $\mathbf{0 . 0 0 2}$ | $\mathbf{0 . 4 9 7}(0.31-0.80)$ | $\mathbf{0 . 0 0 4}$ |

Table 3.8: Multivariate analysis of the CTLA4-CT60 GG genotype for acute GVHD (grade 1-4 versus no GVHD) in the HLAmatched subgroup, confirming this genotype as an independent risk factor.

| TNF-1031: acute grade 4 GVHD Variable | Univariate OR (95\% CI) | $P$-value | Multivariate OR (95\% CI) | $P$-value |
| :---: | :---: | :---: | :---: | :---: |
| Recipient age | 0.978 (0.95-1.01) | 0.109 | 0.975 (0.94-1.01) | 0.112 |
| Donor age | 1.038 (1.00-1.08) | 0.044 | 1.033 (0.99-1.07) | 0.105 |
| Female to male transplant | 0.610 (0.27-1.38) | 0.235 | 0.582 (0.24-1.42) | 0.236 |
| Diagnosis ANLL vs ALL | 1.001 (0.57-1.76) | 0.996 | 1.148 (0.60-2.18) | 0.673 |
| Total body irradiation (TBI) | 0.909 (0.40-2.07) | 0.819 | 0.992 (0.39-2.51) | 0.987 |
| Anti-thymoglobulin (ATG) | 3.562 (0.99-12.73) | 0.051 | 2.246 (0.45-11.15) | 0.322 |
| Cyclosporine vs tacrolimus | 1.336 (0.75-2.37) | 0.321 | 1.516 (0.80-2.86) | 0.198 |
| Relapse | 0.115 (0.03-0.48) | 0.003 | 0.154 (0.04-0.65) | 0.011 |
| HLA match | 0.465 (0.24-0.92) | 0.027 | 0.765 (0.35-1.67) | 0.502 |
| Genotype CC | 4.336 (1.7-11.1) | 0.002 | 3.888 (1.39-10.90) | 0.010 |
| Genotype mismatch | 2.905 (1.65-5.1) | 0.00023 | 2.307 (1.18-4.52) | 0.015 |

Table 3.9: Multivariate analysis of TNF-1031 genotype mismatch and CC genotype as a risk factors for acute GVHD grade 4 in the overall (HLA matched and mismatched) cohort. Both are independent risk factors, with competing effects from HLA matching and relapse.

### 3.5. Discussion

The exploration study has identified three non-HLA SNP associations with HSCT outcome: The TNF-1031 donor-recipient genotype mismatch with severe GVHD (grade 4, in the overall cohort), the recipient IL2-330 GT genotype with risk of chronic GVHD, and the CTLA4-CT60 GG genotype protective against acute GVHD (grade 1-4; the latter two associations were found in the HLA-matched subgroup only).

TNF $\alpha$ is a cytokine that has been associated with severity of acute GVHD in several previous genetic, gene expression and animal model studies. Teshima et. al. (Teshima et al., 2002) have demonstrated in an animal model that TNF is essential in the development of acute GVHD. Previous data from a Japanese population have shown that the TNF haplotype including TNF-1031 was associated with severe GVHD(Ishikawa et al., 2002), and the TNF-1031C allele was associated with higher TNF expression(Higuchi et al., 1998). A more recent study (Goyal et al., 2010) describes the C allele as a risk factor for grade 3-4 acute GVHD. Therefore an association of the TNF-1031 CC genotype with severe acute GVHD, as seen in this study, albeit showing only a trend in the confirmation cohort, would be biologically meaningful and replicate previous findings. However, the TNF-1031 CC genotype displays strong linkage disequilibrium with HLA, in particular with HLA-B61(Higuchi et al., 1998). This may explain our finding of the strong association between donor-recipient genotype mismatch and acute GVHD grade 4 in the overall cohort only, but not in the HLA matched subgroup. Our study did not have the power to elucidate if any particular TNF-1031 genotype mismatch combinations carry a higher risk. As the group affected with acute GVHD grade 4 is small (just above $5 \%$ ), further studies should confirm this result independently. The finding that genotype mismatch was also associated with grade 2-4 as well as grade 3-4 acute GVHD (which are larger groups) in the confirmatory cohort gives further indication that the genotype mismatch is likely to be a risk factor for acute GVHD. Nevertheless, the strength and consistency of this association mean that it is potentially a strong discriminator
for prediction of the most severe form of acute GVHD (grade 4), which could be exploited in clinical practice.

The IL2-330 (rs2069762) SNP has an almost identical genotype distribution between Caucasian and Japanese populations (Caucasian: TT: 0.536, GT: 0.464, GG: 0; Japanese (this study): TT: 0.450, GT: 0.440, GG: 0.110). The G allele is the known high-expressing allele, and high levels of IL2 have been described to correlate with severity of acute GVHD (Das et al., 2001, MacMillan et al., 2003). A previous study from North America on a cohort of similar time frame to our screening cohort (MacMillan et al., 2003) reported an association between the recipient IL2-330 G allele and acute GVHD; and a trend towards risk of chronic GVHD. In our study, we found an association of the GT genotype with risk of chronic GVHD. More detailed analysis showed that the low frequency GG genotype is likely to be the highest risk genotype for chronic GVHD, whilst GT associated with risk, and TT with protection. Our findings therefore confirm those of the previous study even across different ethnic populations, qualifying this marker as a predictor of chronic GVHD risk. The effect of the CTLA4-CT60 polymorphism on HSCT outcomes was studied previously, in settings of HLA matched sibling donors(Perez-Garcia et al., 2007, Murase et al., 2011) and matched unrelated donors (Vannucchi et al., 2007) in Caucasian populations. In HLA matched sibling transplants, the donor G allele was associated with increase of relapse and worse survival, while the AA genotype was linked to risk of acute GVHD. The findings in matched unrelated donor HSCT were similar, with the donor AA genotype associating with severe acute GVHD (grade 3-4), but risk of G allele or GG genotype with relapse or survival was not observed. Our findings are in accordance with these results, identifying the GG genotype as protective against acute GVHD (remarkably, the screening cohort result indicated a risk of the GG genotype with acute GVHD (see table 4) - a finding completely reversed by the randomisation). We could not establish any risk of the GG genotype with relapse or survival, or the AA genotype with acute GVHD. This may be explained by the fact that in the Japanese population, the GG genotype is more prominent than in Caucasians, while the AA genotype is more rare (HapMap data of genotypes: Caucasians: AA: 0.208, AG: 0.513, GG: 0.283; Japanese: AA: 0.047, AG: 0.389, GG: 0.542). The risk of acute

GVHD, relapse or survival associated with this marker may therefore be lower in the Japanese population, compared to Caucasians.

The results raise also some methodological questions which are beyond the scope of this study:

- By incorporating a measure of effect size into the statistical analysis, this study extends beyond previous approaches focussing on significance and correction for multiple testing. Our results suggest that this approach may be more sensitive, but because of limited power and small number of identified associations no conclusions could be made about the impact on sensitivity and specificity, and statistical multiple testing burden.
- Despite the effort to control variability of study population characteristics, reproducibility of associations remains low and appeared to be dependent on distribution of these characteristics amongst the cohorts. This may be due to the overall small effect size of the associations, confounders in the study cohort, or both. A more comprehensive typing (full typing of all markers on both screening and confirmation cohort) and analysis would be required.

Clinical and population characteristics of study cohorts may explain some of the contradictory results observed in previous studies, therefore careful design of study cohorts and control of confounders should receive more attention. The growing number of HSCT may facilitate in the future the availability of larger, genetically and clinically more homogeneous study cohorts; however, the changing and expanding indications of HSCT are likely to prove a challenge.

In conclusion, this study demonstrates that non-HLA genetic association with HSCT outcomes do exist and can be detected even in the HLA-mismatched setting. Such associations could be useful for application in future clinical practice in this clinically highly relevant population. These findings should be verified by larger studies also on populations of different ethnicities.

## 4. RESULTS

4.1. Pooled DNA PCR and genotyping - $1^{\text {st }}$ and $2^{\text {nd }}$ screening steps
4.2. Individual Genotyping
4.3. Further exploration of a susceptibility region by SNP typing
4.4. Genetic susceptibility regions for moderate-severe acute GVHD

### 4.1. Pooled DNA PCR and genotyping - $1^{\text {st }}$ and $2^{\text {nd }}$ screening steps

### 4.1.1. Technical quality aspects

In the first instance, the full set of marker plates for the first screening, involving $4,321 \mathrm{MS}$ markers, was typed in all four pools. The quality of peak signals was assessed within the Run 3730 Data Collection version 2.0 software (Applied Biosystems). The Capillary Viewer would indicate peak signals that were excessively high, adequate or absent.

Following import and analysis of typing data in GeneMapper version 3.5 (Applied Biosystems), peak sizes and quality of size standardisation were analysed. In particular peak sizes that were off-scale, and samples with inadequate size standard became apparent.

Finally, peak signal quality was evaluated in MultiPeaks version 0.21.1 (a Java- application also supplied by Applied Biosystems). For pooled DNA genotyping, consistency of peak sizes and quality amongst pools was of particular importance. Peak sizes $>30,000$ and <200 flourescence units (fu) were classified as typing error. For the purpose of consistency, however, stricter quality criteria were applied: A minimum peak height of 1000 fu for higher frequency alleles ( $>15 \%$ ) and 500 fu for lower frequency alleles ( $<15 \%$ ), absence of noise at the baseline, and no more than $50 \%$ peak height variation between the four pools. The overall peak pattern would be consistent amongst pools. Figure 4.1 shows a typical four-pool graph of a marker with a significant association.

The initial error rate was $11.36 \%$, the majority of these were high or low signal errors. With re-typing applying different DNA dilutions, and re-PCR, the error rate was reduced to $0.8 \%$ (details see appendix 4.1).

First Screen


## Second Screen



Figure 4.1: Example of a peak height graph of marker D6S0035i as displayed by the MultiTyper (Applied Biosystems ${ }^{\circledR}$ ) software. The images show the results of the four pools (top image: first screening, bottom image: second screening). Allele 2 has a higher peak height in the donor GVHD 0-1 group, suggesting a protective effect. This is replicated in the $2^{\text {nd }}$ screen.

### 4.1.2. Results of pooled DNA screening

## First pooled DNA screening (Discovery Cohort)

In the $1^{\text {st }}$ pooled DNA screening, 4,321 microsatellite markers were typed in four DNA pools (donors of recipients with GVHD grade 0 and 1, donors of recipients with GVHD grade 2-4, recipients with GVHD grade 0 and 1, and recipients with GVHD grade 2-4).
Allele frequency differences were analysed in two directions, separately for each individual allele (Fisher's exact test for $2 \times 2$ Chi Square test) and for each marker as a whole (Fisher's exact test for 2xm Chi Square test):

- Between donors of recipients with GVHD grade 0 and 1 and donors of recipients with GVHD grade 2-4
- Between recipients with GVHD grade 0 and 1, and recipients with GVHD grade 2-4.

The results were collated using a custom-built analysis and database system. Peak height data were translated into allele frequencies, and significance tests performed as described in the methodology section.
This system automatically extracted the strongest associated allele for each marker ( $2 \times 2$ ), and all markers associated by $2 x m$ analysis (result details see table). While all markers positive by $2 x m$ analysis also had at least one allele associated by $2 \times 2$ analyses; not all markers who carried an associated allele were also positive by $2 x m$ analysis.

In first screening analysis (tables 3.1, 3.2), 34 ( $0.79 \%$, donor) and 35 ( $0.81 \%$, recipient) markers were excluded because of technical failure in PCR or genotyping.

103 (2.38\%, donor) and 105 (2.43\%, recipient) markers were nonpolymorphic. This is an expected result as the microsatellite marker panel
used in this study contains microsatellites that are polymorphic for some, but not all populations.

## $1^{\text {st }}$ pooled DNA screening results:

In the donor pools analysis, 1016 alleles ( $2 \times 2$ test) and 624 MS markers ( 2 xm test) showed an association with acute GVHD grade 2-4, either as a risk or protective. In the recipient analysis, 931 alleles and 543 MS markers were associated.

All markers that were positive by $2 x m$ or $2 x 2$ analyses were typed again in the $2^{\text {nd }}$ screening step (tables 4.1, 4.2). Inclusion of markers positive only for $2 \times 2$ but not for $2 \times m$ analysis was a measure of additional sensitivity for the second screening step. Naturally, the first screening step contained many false positive associations:

- Statistically false positives, estimated as $5 \%$ of 4,321 markers (that would equal 216 markers in the $2 x m$ analysis) or $5 \%$ of 20,197 (donor analysis) or 20,132 (recipient analysis) alleles (which would lead to 1010 (donor analysis) and 1007 (recipient analysis) false positives in the $2 \times 2$ analyses).
- Errors introduced by DNA pooling process (e.g. variation in number of DNA copies per pool).
- Inherent artefacts of microsatellite typing (e.g. +A alleles, preferential amplification).


## Second pooled DNA screening (Confirmation Cohort)

The main purpose of the secons pooled DNA screening step was to eliminate false positive associations by independent confirmation. Following $2^{\text {nd }}$ pooled screening, identification of true and false positives was much more specific as the independent typing, in addition to $p$-value, introduces criteria which could be used to distinguish true and false positive associations:

- Association of the same allele within a marker
- Consistency of the odd's ratio ('risk', 'protective') of the same allele between the two screening steps
- Consistency of the microsatellite pattern and typing quality (as assessed by the peak image).

Tables 4.1 and 4.2 give the details of the $1^{\text {st }}$ and $2^{\text {nd }}$ screening steps separate for the donor GVHD 0-1 versus donor GVHD 2-4 analysis, and the recipient results accordingly. Results for $2 \times 2$ and $2 x m$ analysis were also separated.
All markers that showed a positive $2 x m$ or $2 \times 2$ result in first screening (1016 ( $23.51 \%$, donor analysis) and 931 ( $21.54 \%$, recipient analysis) were typed again in second screening, but analysed separately for $2 \times 2$ and $2 x m$ Chi-Square tests.

In second screening, 6 (donor analysis) and 10 (recipient analysis) markers showed a non-polymorphic results. These markers were typed again in all screening pools of $1^{\text {st }}$ and $2^{\text {nd }}$ result, using a new primer set. The non-polymorphic result was confirmed (hence, the initial polymorphic result in $1^{\text {st }}$ screening represented a false positive association). We also excluded 17 (donor analysis) and 13 (recipient analysis) markers for which we technically could not reproduce the positive association in first screening despite repeated attempts of PCR and typing (as described above). Except for those markers displaying non-polymorphism or PCR
failure, all markers had satisfactory allele allocations as described in section 2.12.3.

## $2^{\text {nd }}$ pooled DNA screening results:

In the donor analysis, 335 alleles (6.44\%) had a significant result by p -value (<0.05), while in the $2 x m$ analysis 178 markers (27.73\%) were significant. In recipients, 314 alleles (6.32\%) and 141 markers (25.97\%) were confirmed.

Determining consistency of associations across the two screenings

In the next step, false positive markers in the $2 \times 2$ analysis were excluded by identifying and selecting those markers that shared the same associated allele, and had an Odd's ratio that consistently pointed in the same direction (towards risk/protection).

When inspecting the results of positive markers that did not share the same most strongly associated allele within the marker, we noticed that many markers had several positive alleles. On inspection of the peak image we found that occasionally presumed artefacts represent the strongest allele, with a 'true' allele, which showed strongest allele association in the other screening, 'obscured'. Therefore we decided to determine all associated alleles in markers positive for $2 \times 2$ analysis and $2 x m$ analysis (as we assumed that if $2 x 2$ associations of the strongest allele within a marker would not result in $2 x m$-positivity, it would be unlikely that an allele with an even weaker association would have had a significant effect on risk/protection). Markers that would have a shared associated allele and be $2 x m$ positive were entered into the odd's ratio analysis.

## Pooled DNA screening results -

 same allele and Odd's ratio direction:Eventually, 97 (donor analysis) and 74 (recipient analysis) alleles would remain with a $p$-value of $<0.05$ for $2 \times 2$ analysis in both screenings, a shared allele and consistent odd's ratio direction; with 57 (donor) and 40 (recipient) markers by $2 x m$ analysis, accordingly.

| Donors |  | markers overall | \% | alleles $2 \times 2$ | \% | markers $2 \times 2$ | \% |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1st screen | tested | 4321 | 100 | 20197 | 100 | 4321 | 100 |
|  | positive | 1016 | 23.51 | 1016 | 5.03 | 642 | 14.86 |
|  | negative | 3175 | 73.32 | 19181 | 94.97 | 3548 | 82.11 |
|  | non-polymorph | 103 | 2.38 | N/A |  | 97 | 2.24 |
|  | failed | 34 | 0.79 | N/A |  | 34 | 0.79 |
|  | expected false pos 2 xm | 216 | 5 | 1009 | 5 | 216 | 5 |
|  | difference pos-false pos 2xm | 800 | 18.51 | 7 | 0.03 | 425 | 9.86 |
|  |  |  |  |  |  |  |  |
| 2nd screen | tested | 1016 | 100 | 5205 | 100 | 642 | 100 |
|  | positive | 335 | 32.97 | 335 | 6.44 | 178 | 27.73 |
|  | negative | 658 | 64.77 | 4870 | 93.56 | 442 | 68.69 |
|  | non-polymorph | 6 | 0.59 |  |  | 6 | 0.93 |
|  | failed | 17 | 1.67 |  |  | 17 | 2.65 |
|  | same allele as 1st screen allele |  |  | 125 |  |  |  |
|  | not same allele |  |  | 210 |  |  |  |
|  | - 2xm pos AND 2nd allele |  |  | 47 |  |  |  |
|  | sum same allele |  |  | 172 |  |  |  |
|  | same OR direction |  |  | 97 | 10.42 | 57 | 10.50 |
|  | expected false pos 2 xm |  |  | 51 | 5 | 32 | 5 |
|  | difference pos-false pos 2xm |  |  | 46 |  | 25 |  |

Table 4.1: Results of the pooled donor GVHD 0-1 v donor GVHD 2-4 analysis

| Recipient |  | markers overall | \% | $\begin{aligned} & \text { alleles } \\ & 2 \times 2 \\ & \hline \end{aligned}$ | \% | $\begin{aligned} & \text { markers } \\ & 2 \times 2 \\ & \hline \end{aligned}$ | \% |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1st screen | tested | 4321 | 100 | 20132 | 100 | 4321 | 100 |
|  | positive | 931 | 21.54 | 931 | 4.62 | 543 | 12.57 |
|  | negative | 3252 | 75.22 | 19201 | 95.38 | 3641 | 84.26 |
|  | non-polymorph | 105 | 2.43 | 103 | 0.51 | 105 | 2.43 |
|  | failed | 35 | 0.81 | 35 | 0.17 | 32 | 0.74 |
|  | expected false pos | 216 | 5 | 1006 | 5 | 216 | 5 |
|  | difference pos-false pos | 715 | 16.55 | -75 | -0.38 | 327 | 7.57 |
|  |  |  |  |  |  |  |  |
| 2nd screen | tested | 931 | 100 | 4969 | 100 | 543 | 100 |
|  | positive | 314 | 33.73 | 314 | 6.32 | 141 | 25.97 |
|  | negative | 594 | 63.80 | 4655 | 93.68 | 386 | 71.09 |
|  | non-polymorph | 10 | 1.07 | N/A |  | 10 | 1.84 |
|  | failed | 13 | 1.40 | N/A |  | 6 | 1.10 |
|  | same allele as 1st screen allele |  |  | 136 |  |  |  |
|  | not same allele |  |  | 208 |  |  |  |
|  | - 2xm pos AND 2nd allele |  |  | 27 |  |  |  |
|  | sum same allele |  |  | 163 |  |  |  |
|  | same OR direction |  |  | 74 | 7.95 | 40 | 7.37 |
|  | expected false pos $2 x \mathrm{~m}$ |  |  | 46.55 | 5 | 27.15 | 5 |
|  | difference pos-false pos 2xm |  |  | 27.45 |  | 12.85 |  |
|  |  |  |  |  |  |  |  |

Table 4.2: Results of the pooled recipient GVHD 0-1 v recipient GVHD 2-4 analysis

## Further steps of selecting associated microsatellite markers

The confirmation of the first screening results by independent did reduce, but not completely eliminate false positive associations. The number of associations found after analysis of the second screening step would still exceed the number of expected true positives (as compared to previous GWAS using this approach) as well as the resources allocated to individual genotyping. Therefore, within the $2 \times 2$ datasets we used a two-step selection system to identify strong association and high quality typing markers for preferential selection for individual genotyping, aiming to eliminate markers falsely positive for lower quality genotyping.
$1^{\text {st }}$ step:

- Selection by allele frequency (=frequency of a certain allele in the pool): Alleles with a consistent allele frequency of $>0.10$ higher were selected. Low frequency alleles had a lower fluorescent signal peak height (as peak signal height correlates with allele frequency in the pool; a low signal at the border of technical resolution and/or a low allele frequency at the border of statistical power were more likely to represent artefacts), and may represent new mutations within the microsatellite; and/or:
- $2 x m$ positive: An allele association also resulted in the marker being positive in the $2 x m$ analysis. This was more likely to the case in markers with smaller number of alleles (=number o alleles of a marker), increasing the statistical power of each allele.
$2^{\text {nd }}$ step:
- The fluorescent signal height was determined. The signal height depends on the number of allele copies in the sample, hence on the effectiveness of PCR. The genotyping process is calibrated to give accurate readings of fluorescent signals between 500-30,000 fu, i.e. the allele frequency distribution within one marker could expected to
be proportionally accurate, even with a variation of amount of DNA in the sample. Signals below or above this range are prone to distort the distribution of allele frequencies (i.e. an overall low signal may miss or misread low frequency alleles, while a very high signal may exaggerate the reading for high frequency alleles). A signal of 1000 fu or above (but $<30,000 \mathrm{fu}$ ) of the associated allele was regarded as of highest quality.
- Inspection of the microsatellite pattern: Consistency in the microsatellite pattern in all eight pools (allele number, sequence, peak height).


## Associated markers remaining following genotyping quality assessment:

This process resulted in a 'shortlist' of 48 ms markers (31 in the donor analysis, 17 in the recipient analysis) selected for individual genotyping (table 3.3).

### 4.2. Individual genotyping

### 4.2.1. Individual genotyping of the alleles found associated with GVHD grade 2-4 in the pooled screenings

Most artefacts introduced by pooled PCR and genotyping (as described above in methodology section) were readily identifiable by 'test-typing' on a small number of individual samples, therefore all remaining 48 markers were subjected to typing on 14 samples each that stem from a healthy Japanese control population, unrelated to this study. This step eliminated 9 (donor) and 2 ms markers (recipient) from further analysis due to pooling artefacts, copy number variation error or discovered non-polymorphism.

Eventually 19 (donor) and 11 (recipient) markers underwent individual genotyping on the full sample set (922 donors or recipients). Three (donors) and 4 (recipient) markers with weaker associations, despite passing the criteria for individual genotyping, were eventually not individually typed due to resource restrictions. The overview results of the individual genotyping are presented in table 4.3.

After applying Hardy-Weinberg Equilibrium tests for genotyping quality control, 10 MS markers were confirmed to have an association with Grade 2-4 acute GVHD that was consistent in both pooled DNA screenings and individual genotyping (tables 4.4-4.6) in univariate analyses. Associations with $p$-values that would withstand application of Bonferroni's correction for the total number of alleles in individual genotyping ( $n=123,2 \times 2$ analysis: corrected threshold for $p=0.05$ association: $p=0.0004065 ; p=0.1$ trend: $p=0.0008131$ ) and markers ( $\mathrm{n}=30,2 x m$ analysis: corrected threshold for $p=0.05$ association: $p=0.00166, p=0.1$ trend: $p=0.00333$ ), as well as those showing a trend with correction, were indicated.

Five further markers (D16S0452i, D5S1173i, D3S1225i, D14S0499i and AJ133269.1_180046) showed significant associations but failed the HWE for both case and control cohorts. From a genotyping quality control perspective these markers were therefore excluded from further analysis. Nevertheless, from a biological perspective failed HWE does not necessarily imply an invalid result. Due to the underlying malignant disease which is in part genetically determined, both the recipient as well as the HLA-matched donor population cannot be expected to reflect an allele distribution that would be expected in a 'healthy' population.

> Results after applying multiple testing correction statistics
> Four markers (recipient D5S424, donor D6S0035i, D1S0818i, D17S0219i), demonstrated associations by $2 \times m$ and/or $2 \times 2$ analyses that had $p$-values that held up against statistics for multiple testing correction, while one further marker (D6S0330i) showed a trend when Bonferroni's correction was applied. When including markers that failed the HWE test, one further marker (D16S04521i) would show an association.

| Target gene | MS identifier 2 | Donor/ Recipient | Test typing | Full individual typing Yes/No | outcome individual typing |
| :---: | :---: | :---: | :---: | :---: | :---: |
| SNRPN | D15S122 | D | passed | Y | not confirmed |
| AGPAT4 | D6S0330i | D | passed | Y | confirmed |
| PDE4B | D1S0716i | D | passed | N | Not tested |
| ALKBH1 | D14S594 | D | failed | N | - |
| TRAF7 | D16S0452i | D | passed | Y | failed HWE |
| NFKBIZ | $\text { DISO7_1 }_{4}$ | D | failed | N | - |
| TIAF1 | D17S0406i | D | passed | Y | artefact |
| ELTD1 | D1S0818i | D | passed | Y | confirmed |
| ITPKB | D1S1143i | D | passed | Y | not confirmed |
| MCM2 | D3S3607 | D | passed | Y | not confirmed |
| SMARCAL1 | D2S0809i | D | passed | Y | artefact |
| EDAR | D2S1281i | D | failed | N | - |
| CD86 | D3S1225i | D | passed | Y | failed HWE |
| FBXW7 | D4S0270i | D | passed | Y | not confirmed |
| C1QTNF2 | D5S403 | D | failed | N | - |
| MAPK14 | D6S0035i | D | passed | Y | confirmed |
| ETV1 | D7S0119i | D | passed | Y | not confirmed |
| HSPB1 | D7S1218i | D | passed | Y | not confirmed |
| DLG5 | D10S0603i | D | passed | N | not tested |
| TCF8 | D10S565 | D | passed | Y | not confirmed |
| CAV1 | $\begin{gathered} \text { AJ133269.1_180 } \\ 046 \end{gathered}$ | D | passed | Y | failed HWE |
| C1QBP | D17S0113i | D | failed | N | - |
| EIF4A3 | D17S0294i | D | passed | N | Not tested |
| DSCAM | D21S0184i | D | failed | N | - |
| ATF4 | D22S428 | D | failed | N | - |
| BTK | DXS0923i | D | failed | N | - |
| IL1RAPL2 | DXS0629i | D | passed | Y | confirmed |
| IL1RAPL2 | DXS0151i | D | passed | Y | confirmed |
| TGM3 | $\begin{gathered} \text { AL031678.2_901 } \\ 37 \end{gathered}$ | D | failed | N | - |
| SOCS3 | D17S0219i | D | passed | Y | confirmed |
| LTB | TNF C | D | passed | Y | trend only |
| F2RL1/S100Z | D5S424 | R | passed | Y | confirmed |
| HRK | D12S0781i | R | passed | Y | multiple alleles |
| MAP3K7 | D6S0738i | R | passed | N | not tested |
| NFKBIZ | $\begin{gathered} \text { DISO7_1000118 } \\ 4 \end{gathered}$ | R | failed | N | - |
| C1QA | D1S1655i | R | passed | Y | not confirmed |
| AKT3 | D1S1335i | R | passed | Y | confirmed |
| NMI | D2S1334i | R | passed | Y | not confirmed |
| EDAR | D2S1281i | R | failed | N | - |
| CSF2 | D5S1174i | R | passed | Y | non-polymorphic |
| IL7R | D5S1173i | R | passed | Y | failed HWE |
| RNASE6 | D14S0499i | R | passed | Y | failed HWE |
| MMP25 | D16S3082 | R | passed | Y | not confirmed |
| DDX42 | D17S0271i | R | passed | Y | confirmed |
| TRIM26 |  | R | passed | N | not tested |
| TBL1X | DXS0324i | R | passed | Y | confirmed |
| SSTR2 | $\begin{gathered} \hline \text { chr17.fa.07frz. } 7 \\ 8835314 \end{gathered}$ | R | passed | N | Not tested |
| ISG20 | D15S0049i | R | passed | N | Not tested |

Table 4.3 : overview results of individual genotyping

| Target Gene | Marker name (internal) | Database name | Donor/Patient | allele size | allele no | $\begin{aligned} & \text { 1st screen } \\ & 2 \times m \end{aligned}$ | 1st screen $2 \times 2$ p-value | $\begin{gathered} \text { 1st } \\ \text { screen } \end{gathered}$ OR | $\begin{aligned} & \text { 2nd screen } \\ & \text { 2xm } \end{aligned}$ | 2nd screen $2 \times 2$ p-value | $\begin{gathered} \text { 2nd } \\ \text { screen } \\ \text { OR } \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| F2RL1/S100Z | 0507E03 | D5S424 | P | 117.7 | 05 | 0.00465911 | 0.00123516 | 1.581714 | $2.37636 \mathrm{E}-09$ | $5.88101 \mathrm{E}-10$ | 2.29902 |
| MAPK14 | 0611B04 | D6S0035i | D | 373.2 | 02 | 0.0331471 | 0.0331471 | 0.688211 | 0.00163933 | 0.00163933 | 0.595028 |
| ELTD1 | 0111F11 | D1S0818i | D | 172 | 01 | 0.000310432 | $3.58175 \mathrm{E}-05$ | 1.902177 | 0.0167968 | 0.0167968 | 1.470116 |
| IL1RAPL2 | 2310B01 | DXS0151i | D | 466.6 | 04 | 0.000451 | 0.000451 | 1.783574 | 0.000105178 | 0.000105 | 1.864524 |
| SOCS3 | T002C05 | D17S0219i | D | 206 | 03 | 0.000157143 | $4.3431 \mathrm{E}-05$ | 0.42191 | 0.0111833 | 0.00396346 | 0.471879 |
| IL1RAPL2 | 2309D05 | DXS0629i | D | 393.4 | 03 | 0.151619 | 0.030286 | 1.45962 | 0.0201712 | 0.006152 | 1.568938 |
| TBL1X | 2309A04 | DXS0324i | P | 385.1 | 01 | 0.001843 | 0.001843 | 0.604196 | 0.0259134 | 0.025913 | 0.693579 |
| DDX42 | 1704G02 | D17S0271i | P | 222 | 01 | 0.0360375 | 0.0128895 | 0.676094 | 0.0397464 | 0.0397464 | 0.743281 |
| AGPAT4 | 0606E02 | D6S0330i | D | 166.6 | 07 | 0.000453657 | 0.000453657 | 1.616777 | 0.0123369 | 0.0123369 | 1.40784 |
| AKT3 | 0109C12 | D1S1335i | P | 90.3 | 03 | 0.00030278 | 0.000182015 | 1.637257 | 0.0111414 | 0.00302767 | 1.488903 |
| TRAF7 | 1601E07 | D16S04521i | D | 271 | 12 | 0.00002303 | 0.00002303 | 0.331837 | 0.0008224 | 0.00035560 | 0.535539 |
| CD86 | 0310G01 | D3S1225i | D | 97.4 | 03 | 0.000021968 | 0.000021968 | 0.370631 | 0.0268393 | 0.0268393 | 0.647246 |
| CAV1 | 137G11 | AJ133269.1_180046 | D | 384.1 | 05 | 0 | 0.000000087 | 4.635323 | 0.002575 | 0.0041277 | 2.163354 |
| RNASE6 | 1406B07 | D14S0499i | P | 359.9 | 02 | 0.00121386 | 0.000101171 | 1.760132 | 0.00000002 | 0.000000001 | 2.354878 |
| IL7R | 0508H03 | D5S1173i | P | 146.7 | 04 | 0.001375 | 0.00007793 | 0.634464 | 0.00000013 | 0.000000116 | 0.529155 |

Table 4.4: Genotyping results from both pooled screening steps of 15 microsatellite markers that showed an association in the individual genotyping.

| TargetGene | Marker name (internal) | Database name | Donor/Patient | $\begin{gathered} \text { allele } \\ \text { size (bp) } \end{gathered}$ | Associated allele no | $\begin{gathered} \text { aGVHD } \\ 01-24 \\ 2 \times m p= \end{gathered}$ | $\begin{gathered} \hline \text { aGVHD } \\ 01-242 \times 2 \\ p= \\ \hline \end{gathered}$ | $\begin{aligned} & \text { 2x2 Odds } \\ & \text { ratio } \end{aligned}$ | 95\% CI lower | 95\% CI higher | HWE decision |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| F2RL1/S100Z | 0507E03 | D5S424 | P | 117.7 | 05 | 0.0004* | 0.002017 | 1.338207 | 1.113486 | 1.608281 | ok |
| MAPK14 | 0611B04 | D6S0035i | D | 373.2 | 02 | 0.0004* | 0.00035* | 0.685984 | 0.558405 | 0.84271 | ok |
| ELTD1 | 0111F11 | D1S0818i | D | 172 | 01 | 0.0007* | 0.0000783* | 1.519239 | 1.242103 | 1.858208 | ok |
| IL1RAPL2 | 2310B01 | DXS0151i | D | 466.6 | 04 | 0.0066 | 0.007038 | 1.41669 | 1.125021 | 1.783976 | ok |
| SOCS3 | T002C05 | D17S0219i | D | 206 | 03 | 0.005 | 0.000275* | 0.418673 | 0.259705 | 0.674948 | ok |
| IL1RAPL2 | 2309D05 | DXS0629i | D | 393.4 | 03 | 0.016 | 0.001315 | 0.7778 | 2.698158 | 4.312269 | ok |
| TBL1X | 2309A04 | DXS0324i | P | 385.1 | 01 | 0.021 | 0.013253 | 0.753511 | 0.603937 | 0.940129 | ok |
| DDX42 | 1704G02 | D17S0271i | P | 222 | 01 | 0.0404 | 0.008597 | 0.71826 | 0.563124 | 0.916134 | ok |
| AGPAT4 | 0606E02 | D6S0330i | D | 166.6 | 07 | 0.071 | $0.00074 \dagger$ | 1.38941 | 1.150105 | 1.678508 | ok |
| AKT3 | 0109C12 | D1S1335i | P | 90.3 | 03 | 0.195 | 0.032222 | 1.226761 | 1.017861 | 1.478534 | ok |
| TRAF7 | 1601E07 | D16S04521i | D | 271 | 12 | 0* | 0.0002735* | 0.672402 | 0.54339 | 0.83204 | failed |
| CD86 | 0310G01 | D3S1225i | D | 97.4 | 03 | 0.029 | 0.0080907 | 0.75196 | 0.60978 | 0.92729 | failed |
| CAV1 | 137 G 11 | AJ133269.1_180046 | D | 384.1 | 05 | 0.354 | 0.0279587 | 1.391733 | 1.04506 | 1.85340 | failed |
| RNASE6 | 1406B07 | D14S0499i | P | 359.9 | 02 | 0.137 | 0.0076539 | 1.337914 | 1.08112 | 1.65570 | failed |
| IL7R | 0508H03 | D5S1173i | P | 146.7 | 04 | 0.0056 | 0.0122439 | 0.7072632 | 0.427571 | 0.98156 | failed |

Table 4.5: individual genotyping associations of microsatellite markers ( 2 xm ) and alleles ( $2 \times 2$ ) with aGVHD grade 2-4. $P$-values shaded dark (*) are significant against multiple testing correction; p-values shaded bright ( $\dagger$ ) show a trend. Included in this table are the five markers failing HWE testing, one of these showing an association.

| Target Gene | Marker | allele <br> no | $\begin{aligned} & \text { aGVHD } \\ & 01-24 p= \end{aligned}$ | total | $\begin{aligned} & \text { cases } \\ & \text { all } \\ & \hline \end{aligned}$ | $\begin{aligned} & \text { controls } \\ & \text { all } \end{aligned}$ | cases pos | cases neg | controls <br> pos | controls neg | Odds ratio | Iower Cl (95\%) | higher Cl (95\%) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| F2RL1/S100Z | D5S424 | 05 | 0.00202 | 1842 | 842 | 1000 | 446 | 396 | 457 | 543 | 1.338 | 1.1135 | 1.60828 |
| MAPK14 | D6S0035i | 02 | 0.00035 | 1824 | 832 | 992 | 207 | 625 | 323 | 669 | 0.686 | 0.5584 | 0.84271 |
| ELTD1 | D1S0818i | 01 | 0.000078 | 1832 | 842 | 990 | 619 | 223 | 634 | 347 | 1.519 | 1.2421 | 1.85821 |
| IL1RAPL2 | DXS0151i | 04 | 0.00704 | 1297 | 595 | 702 | 228 | 367 | 214 | 488 | 1.417 | 1.125 | 1.78398 |
| SOCS3 | D17S0219i | 03 | 0.00028 | 1826 | 838 | 988 | 24 | 814 | 65 | 923 | 0.419 | 0.2597 | 0.67495 |
| IL1RAPL2 | DXS0629i | 03 | 0.00132 | 1291 | 591 | 700 | 307 | 284 | 426 | 274 | 0.778 | 0.6234 | 0.97071 |
| TBL1X | DXS0324i | 01 | 0.01325 | 1271 | 585 | 686 | 264 | 321 | 358 | 328 | 0.754 | 0.6039 | 0.94013 |
| DDX42 | D17S0271i | 01 | 0.0086 | 1824 | 834 | 990 | 129 | 705 | 201 | 789 | 0.718 | 0.5631 | 0.91613 |
| AGPAT4 | D6S0330i | 07 | 0.00074 | 1828 | 838 | 990 | 360 | 478 | 348 | 642 | 1.389 | 1.1501 | 1.67851 |
| AKT3 | D1S1335i | 03 | 0.03222 | 1842 | 842 | 1000 | 364 | 478 | 383 | 617 | 1.227 | 1.0179 | 1.47853 |

Table 4.6: Allele numbers and Odds Ratio calculation for associated alleles from individual genotyping, illustrating the effect sizes of the associations.

### 4.2.2. HLA subgroup analysis of alleles

Mirroring the analysis of SNP markers in the pilot study, the effects of the alleles were also analysed in a subgroup of higher HLA matching. HLA matching was defined as high-resolution (allele level) match for HLA-A, B, C, DRB1 (i.e. 8/8 matching), with allowing for either a HLA-DQB1 or DPB1 only, and including 12/12 matches. Results of this subgroup analysis are presented in table 4.7. While some of the associations were limited to the HLA mismatched group only, others showed an effect on both degrees of matching, and some associations appeared to have a larger effect on the HLA-matched subgroup than on the mismatched one. Two markers had other alleles than the one identified by pooled screening associated with acute GVHD grade 2-4. In both cases, the markers had two main alleles only, hence could be analysed like a single nucleotide polymorphism (SNP) marker. While one allele of these markers indicated a GVHD risk, the corresponding 'opposite' allele would have a protective effect, as the OR of the associations showed (table 4.8.)

### 4.2.3. Genotype analysis with HLA subgroup analysis

An analysis of genotypes was also conducted where possible. As MS have many alleles, and therefore a large number of possible allele combinations forming a genotype, such analyses would have only be useful with a reasonable frequency of the genotype in question. A limit of an allele frequency of 0.1 or above was applied. Primarily the homozygosity and heterozygosity of the associated allele versus the remaining genotypes was investigated; and then all other genotypes that had a frequency of $10 \%$ or more were analysed.

Five markers showed an association with acute GVHD grade 2-4 of the homozygous genotype of the same associated allele (table 4.10); while four further markers showed associations of other genotypes with acute GVHD grade 2-4 (tables 4.11, 4.12). All of the latter four markers had only
two or three major alleles accounting for $>90 \%$ of the total allele frequency; the genotypic analysis showed that the genotype of the 'oppositional' allele had a stronger (and opposite) effect as compared to the allelic effect of the originally identified allele. One example was the above mentioned marker 0611B04 (D6S0035i): Whilst the allelic screening suggested that the minor allele 02 had a protective effect, it is in fact the major homozygous allele 01 genotype that constituted a risk of moderate-severe acute GVHD.

| marker info |  | all alleleic association |  |  |  |  | HLA mismatched |  |  | HLA matched |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| TargetGene | Marker name | Donor/Patient | allele no | $\begin{aligned} & \text { aGVHD 01-24 } \\ & 2 \times m p= \end{aligned}$ | $\begin{aligned} & \text { aGVHD 01-24 } \\ & 2 \times 2 p= \\ & \hline \end{aligned}$ | $\begin{aligned} & \hline 2 \times 2 \text { Odds } \\ & \text { ratio } \\ & \hline \end{aligned}$ | $\begin{aligned} & \text { aGVHD } \\ & 01-24 p= \\ & \hline \end{aligned}$ | Odds ratio | 95\% CI | $\begin{aligned} & \begin{array}{l} \text { aGVHD } \\ 01-24 p= \end{array} \\ & \hline \end{aligned}$ | Odds ratio | 95\% CI |
| F2RL1/S100Z | D5S424 | P | 05 | 0.0004 | 0.002017 | 1.338207 | 0.0107 | 1.349883 | 1.1-1.7 | 0.066 | 1.347273 | 0.9-1.8 |
| MAPK14 | D6S0035i | D | 02 | 0.0004 | 0.00035 | 0.685984 | 0.02457 | 0.74731 | 0.6-0.9 | 0.00273 | 0.579385 | 0.4-0.8 |
| ELTD1 | D1S0818i | D | 01 | 0.0007 | 0.0000783 | 1.519239 | 0.0011 | 1.52398 | 1.2-2.0 | 0.051 | 1.404808 | 1.0-2.0 |
| IL1RAPL2 | DXS0151i | D | 04 | 0.0066 | 0.007038 | 1.41669 | 0.071 | 1.30469 | 0.98-1.70 | 0.019 | 1.628809 | 1.1-2.4 |
| SOCS3 | D17S0219i | D | 03 | 0.005 | 0.000275 | 0.418673 | 0.000872 | 0.384298 | 0.2-0.7 | 0.115 | 0.480896 | 0.2-1.1 |
| IL1RAPL2 | DXS0629i | D | 03 | 0.016 | 0.001315 | 0.7778 | 0.06 | 0.768652 | 0.6-1.1 | 0.005 | 0.575585 | 0.4-0.85 |
| TBL1X | DXS0324i | P | 01 | 0.021 | 0.013253 | 0.753511 | 0.298 | 0.85887 | 0.7-1.1 | 0.009 | 0.598058 | 0.4-0.9 |
| DDX42 | D17S0271i | P | 01 | 0.0404 | 0.008597 | 0.71826 | 0.0316 | 0.713717 | 0.5-0.9 | 0.155 | 0.739331 | 0.5-1.1 |
| AGPAT4 | D6S0330i | D | 07 | 0.071 | 0.00074 | 1.38941 | 0.027 | 1.304192 | 1.0-1.6 | 0.00669 | 1.571115 | 1.1-2.1 |
| AKT3 | D1S1335i | P | 03 | 0.195 | 0.032222 | 1.226761 | 0.01824 | 1.323822 | 1.1-1.7 | 0.685 | 1.068627 | 0.8-1.5 |

Table 4.7: Association of alleles separated by degree of HLA matching. The effect of polymorphisms is either visible mainly in the HLA-matched subgroup (MAPK14, AGPAT4), or mainly in the HLA-mismatched subgroup (F2RL1, ELTD1, SOCS3, DDX42, AKT3).

| marker info |  | all alleleic association |  |  |  |  | HLA mismatched |  |  |  | HLA matched |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| TargetGene | Marker name | Donor/Patient | allele no | $\begin{aligned} & \text { aGVHD 01-24 } \\ & \begin{array}{l} 2 \times m p= \end{array} \\ & \hline \end{aligned}$ | $\begin{aligned} & \text { aGVHD 01- } \\ & 24 \geqslant \times 0 n= \end{aligned}$ | $\begin{array}{\|l\|} \hline 2 \times 2 \\ \text { Odds } \\ \text { ratio } \end{array}$ | allele | $\begin{array}{\|l\|} \hline \text { aGVHD } \\ 01-24 p= \\ \hline \end{array}$ | $\begin{aligned} & \text { Odds } \\ & \text { ratio } \\ & \hline \end{aligned}$ | $\begin{aligned} & \text { lower Cl } \\ & (95 \%) \\ & \hline \end{aligned}$ | aGVHD 01 $24 p=$ | $\begin{aligned} & \text { Odds } \\ & \text { ratio } \\ & \hline \end{aligned}$ | 95\% CI |
| F2RL1/S100Z | D5S424 | P | 05 | 0.0004 | 0.002017 | 1.338207 | 04 | 0.03319 | 0.76 | 0.6-1.0 | 0.000696 | 0.54 | 0.4-0.8 |
| MAPK14 | D6S0035i | D | 02 | 0.0004 | 0.00035 | 0.685984 | 01 | 0.0344 | 1.31 | 1.0-1.7 | 0.0066 | 1.64 | 1.2-2.3 |
| ELTD1 | D1S0818i | D | 01 | 0.0007 | 0.0000783 | 1.519239 |  |  |  |  |  |  |  |
| IL1RAPL2 | DXS0151i | D | 04 | 0.0066 | 0.007038 | 1.41669 |  |  |  |  |  |  |  |
| socs3 | D17S0219i | D | 03 | 0.005 | 0.000275 | 0.418673 |  |  |  |  |  |  |  |
| IL1RAPL2 | DXS0629i | D | 03 | 0.016 | 0.001315 | 0.7778 |  |  |  |  |  |  |  |
| TBL1X | DXS0324i | P | 01 | 0.021 | 0.013253 | 0.753511 |  |  |  |  |  |  |  |
| DDX42 | D17S0271i | P | 01 | 0.0404 | 0.008597 | 0.71826 |  |  |  |  |  |  |  |
| AGPAT4 | D6S0330i | D | 07 | 0.071 | 0.00074 | 1.38941 |  |  |  |  |  |  |  |
| AKT3 | D1S1335i | P | 03 | 0.195 | 0.032222 | 1.226761 |  |  |  |  |  |  |  |

Table 4.8: Association of alleles other than those identified by pooled DNA genotyping, with HLA matching subgroup analysis

| Target Gene | DS name | allele position | allele ID | P value aGVHD 01-24 association $p=$ | total | cases all | controls all | cases pos | cases neg | controls pos | controls neg | Odds ratio | $\begin{gathered} \text { lower } \\ \text { CI } \\ (95 \%) \end{gathered}$ | higher Cl (95\%) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| ELTD1 | D1S0818i | 172 | 01 | 0.0006 | 916 | 421 | 495 | 228 | 193 | 211 | 284 | 1.59 | 1.224 | 2.066 |
| IL1RAPL2 | DXS0151i | 466.6 | 04 | 0.0313 | 918 | 420 | 498 | 117 | 303 | 108 | 390 | 1.39 | 1.031 | 1.885 |
| TBL1X | DXS0324i | 385.1 | 01 | 0.0053 | 914 | 420 | 494 | 145 | 275 | 216 | 278 | 0.68 | 0.519 | 0.888 |
| AGPAT4 | D6S0330i | 166.6 | 07 | 0.0003 | 914 | 419 | 495 | 90 | 329 | 55 | 440 | 2.19 | 1.52 | 3.151 |
| AKT3 | D1S1335i | 90.3 | 03 | 0.0197 | 921 | 421 | 500 | 82 | 339 | 68 | 432 | 1.54 | 1.081 | 2.184 |

Table 4.9: Associations with moderate-to severe acute GVHD (grade 2-4) of homozygous genotypes of the same alleles as identified in the pooled and individual genotyping.

| marker info |  | all homozygous genotypes |  |  |  | HLA mismatched homozygous genotypes |  |  | HLA matched homozygous genotypes |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| TargetGene | Marker name | homozyg genotype | $\begin{aligned} & \text { aGVHD 01- } \\ & 24 p= \end{aligned}$ | Odds ratio | 95\% CI | $\begin{aligned} & \text { aGVHD 01- } \\ & 24 p= \\ & \hline \end{aligned}$ | Odds ratio | 95\% CI | $\begin{aligned} & \text { aGVHD 01- } \\ & 24 p= \\ & \hline \end{aligned}$ | Odds ratio | 95\% CI |
| F2RL1/S100Z | D5S424 | 0505 | 0.101 | 1.3 | 0.9-1.8 | 0.0817 | 1.41 | 0.9-2.1 | 0.69 | 1.12 | 0.7-1.9 |
| MAPK14 | D6S0035i | 0202 | 0.099 | 0.64 | 0.3-1.1 | 0.0737 | 0.52 | 0.3-1.0 | 1 | 0.93 | 0.4-2.1 |
| ELTD1 | D1S0818i | 0101 | 0.0005 | 1.59 | 1.2-2.1 | 0.0031 | 1.63 | 1.2-2.3 | 0.14 | 1.44 | 0.9-2.2 |
| IL1RAPL2 | DXS0151i | 0404 | 0.031 | 1.39 | 1.0-1.9 |  |  |  |  |  |  |
| SOCS3 | D17S0219i | 0303 | 0.52 | 0.59 | 0.1-2.4 | 0.22 | 0.21 | 0.1-1.8 | 1 | 0.74 | 0.1-8.2 |
| IL1RAPL2 | DXS0629i | 0303 | 0.02 | 0.73 | 0.6-0.9 |  |  |  |  |  |  |
| TBL1X | DXS0324i | 0101 | 0.0053 | 0.68 | 0.5-0.9 |  |  |  |  |  |  |
| DDX42 | D17S0271i | 0101 | 0.14 | 0.55 | 0.3-1.2 |  |  |  |  |  |  |
| AGPAT4 | D6S0330i | 0707 | 0.00025 | 2.19 | 1.5-3.2 | 0.0059 | 1.94 | 1.2-3.1 | 0.0008 | 2.77 | 1.5-5.1 |
| AKT3 | D1S1335i | 0303 | 0.019 | 1.54 | 1.1-2.2 | 0.0233 | 1.7 | 1.1-2.6 | 0.56 | 0.83 | 0.5-1.5 |

Table 4.10: homozygous genotype associations of the same allelic associations identified by pooled DNA genotyping, with analysis separate for HLA matched/mismatched subgroups.

| marker info |  | all homozygous genotypes |  |  |  | HLA mismatched homozygousgenotypes |  |  | HLA matched homozygous genotypes |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| TargetGene | Marker name | homozyg genotype | $\begin{aligned} & \text { aGVHD 01- } \\ & 24 p= \\ & \hline \end{aligned}$ | Odds ratio | 95\% CI | $\begin{aligned} & \text { aGVHD 01- } \\ & 24 p= \\ & \hline \end{aligned}$ | Odds ratio | 95\% CI | $\begin{aligned} & \text { aGVHD 01- } \\ & 24 p= \\ & \hline \end{aligned}$ | Odds ratio | 95\% CI |
| F2RL1/S100Z | D5S424 | 0404 | 0.000059 | 0.36 | 0.2-0.6 | 0.0067 | 0.44 | 0.2-0.8 | 0.00149 | 0.18 | 0.1-0.6 |
| MAPK14 | D6S0035i | 0101 | 0.00053 | 1.6 | 1.2-2.1 | 0.069 | 1.36 | 1.0-1.9 | 0.00093 | 2.18 | 1.4-3.4 |
| ELTD1 | D1S0818i |  |  |  |  |  |  |  |  |  |  |
| IL1RAPL2 | DXS0151i |  |  |  |  |  |  |  |  |  |  |
| SOCS3 | D17S0219i |  |  |  |  |  |  |  |  |  |  |
| IL1RAPL2 | DXS0629i |  |  |  |  |  |  |  |  |  |  |
| TBL1X | DXS0324i |  |  |  |  |  |  |  |  |  |  |
| DDX42 | D17S0271i |  |  |  |  |  |  |  |  |  |  |
| AGPAT4 | D6S0330i |  |  |  |  |  |  |  |  |  |  |
| AKT3 | D1S1335i |  |  |  |  |  |  |  |  |  |  |

Table 4.11: Genotypic associations other than those allelic associations identified by pooled DNA genotyping, with subgroup analysis for HLA matched/mismatched subgroup

| Target Gene | Marker name | allele position | allele ID | P value <br> aGVHD 01-24 <br> Association <br> $p=$ | total | cases all | controls all | $\begin{gathered} \text { cases } \\ \text { pos } \end{gathered}$ | $\begin{gathered} \text { cases } \\ \text { neg } \end{gathered}$ | $\begin{gathered} \text { controls } \\ \text { pos } \end{gathered}$ | controls neg | Odds ratio | $\begin{gathered} \text { lower } \\ \text { CI } \\ (95 \% \%) \end{gathered}$ | $\begin{gathered} \text { higher } \\ \text { CI } \\ (95 \%) \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $\begin{aligned} & \hline \text { F2RL1/ } \\ & \text { S100Z } \end{aligned}$ | D5S424 | 116 | 04 | 6E-05 | 921 | 421 | 500 | 20 | 401 | 61 | 439 | 0.36 | 0.21 | 0.605 |
| MAPK14 | D6S0035i | 369 | 01 | 5E-04 | 912 | 416 | 496 | 229 | 187 | 215 | 281 | 1.6 | 1.23 | 2.081 |
| IL1RAPL2 | DXS0629i | 397 | 04 | 0.001 | 916 | 419 | 497 | 159 | 260 | 139 | 358 | 1.58 | 1.19 | 2.08 |
| LTB | TNFC | 160 | 05 | 0.003 | 915 | 419 | 496 | 23 | 396 | 9 | 487 | 3.14 | 1.44 | 6.869 |

Table 4.12: Associated genotypes of other alleles than those identified through pooled screening and individual genotyping.

### 4.2.4. Analysis of MS marker associations on Chromosome $X$

Alleles from three MS markers (DXS0629i, DXS0324i, DXS0151i) were found to be associated with grade 2-4 acute GVHD. Analysis in the pooled screening was by counting overall alleles within the pool, correcting for the overall number of alleles (i.e. for males only one allele was counted, for females two). The analysis of individual genotyping was mirroring this approach in order to confirm the findings from pooled DNA screening.

Two of the alleles had a protective effect, while one indicated a risk for GVHD. One allele was recipient-intrinsic with a protective effect, whereas the two others derived from the donor, exerting a protective or risk effect. Two markers of these alleles were intronic to the same, very large gene IL1RAPL2.

An analysis separating the gender effects did show that the markers in the IL1RAPL2 gene had very similar effects on the recipient when coming from a female or male donor. The effect of the marker in TBL1X appeared to be mainly on the male recipient.

In the context of transplantation, many polymorphisms on chromosome X have been described as minor histocompatibility antigens (mHag). These are antigens outside the major histocompatibility complex (MHC) that can induce strong immunological responses leading to either graft rejection, GVHD or graft-versus-leukaemia effects. The analysis of markers on chromosome X in this respect is complex and beyond the scope of this study. Such analysis would require careful evaluation of confounding variables relating to donor/recipient sex, rejection and chronic GVHD as outcomes, subgroup analysis of the different female/male combinations of donor and recipient, as well as consideration of HLA matching.

|  |  |  | All |  |  |  | Female |  |  |  | Male |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Gene | Marker | allele no | $\begin{aligned} & p \text { for aGVHD } \\ & 01-24 p= \\ & \hline \end{aligned}$ | Odds ratio | lower CI (95\%) | higher $\mathbf{C l}$ (95\%) | $\begin{aligned} & \hline \text { p for } \\ & \text { aGVHD } \end{aligned}$ $01-24 p=$ | Odds ratio | $\begin{aligned} & \text { lower CI } \\ & \text { (95\%) } \end{aligned}$ | higher Cl (95\%) | $\begin{aligned} & \hline \text { p for } \\ & \text { aGVHD } \end{aligned}$ $01-24 p=$ | Odds ratio | $\begin{aligned} & \text { lower Cl } \\ & (95 \%) \\ & \hline \end{aligned}$ | higher Cl (95\%) |
| IL1RAPL2 | DXS0151i | 04 D | 0.00704 | 1.417 | 1.125 | 1.7839764 | 0.021 | 1.428118 | 1.05609 | 1.9311997 | 0.082 | 1.395542 | 0.9754874 | 1.9964762 |
| IL1RAPL2 | DXS0629i | 03 D | 0.00132 | 0.778 | 0.623 | 0.970705 | 0.018 | 0.70393 | 0.5264287 | 0.9412825 | 0.036 | 0.683761 | 0.4851211 | 0.9637359 |
| TBL1X | DXS0324i | 01 R | 0.01325 | 0.754 | 0.604 | 0.9401294 | 0.232 | 0.829365 | 0.618453 | 1.1122049 | 0.022 | 0.672497 | 0.4805425 | 0.9411286 |

Table 4.13: X-chromosomal markers associated with acute GVHD grade 2-4

### 4.2.5. Multivariate analysis

In order to understand which of the identified associations would be consistent when compared to other major variables in the dataset which we identified previously, multivariate analysis was conducted in STATA v 11 (performed by Dr Hirofumi Nakaoka). This was undertaken as backward multiple logistic regression, i.e. all variables showing a significant association in univariate analysis were include and eliminated in a stepwise fashion until no further improvement to the model could be achieved.

Variables included were recipient age, donor age, female into male transplant, diagnosis, use of total body irradiation, use of antithymoglobulin, use of cyclosporine A versus tacrolimus for GVHD prophylaxis, relapse and HLA matching (HLA-DQB1 or DPB1 mismatch and fully matched pairs only, versus all other grades of mismatching).

A single dataset containing all clinical variables and genotyping results was constructed. Samples for which we did not have all variable information or genotyping results were excluded (53 samples), therefore p-values for univariate analysis differ slightly from those reported in the tables above. Markers on the X-chromosome were not included.

Diagnosis (ALL > ANLL), donor age (older) and HLA mismatch were the strongest competing variables in multivariate analysis.

## Results of the multivariate analysis

Five markers (D6S0035i D17S0219i D1S0818i D6S0330i D5S424) showed associations in multivariate analysis that had effect sizes larger than any of the clinical variables, and are therefore independent predictors of moderate-severe GVHD.

|  | Univariate |  | $P$ value | Multivariate |  | $P$ value |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | OR | 95\% CI |  |
| Recipient age | 0.984 | 0.972-0.997 |  | 0.015 | 0.987 | 0.973-1.001 | 0.06 |
| Donor age | 1.018 | 1.000-1.035 | 0.045 | 1.021 | 1.003-1.040 | 0.023 |
| F-M transplant | 1.181 | 0.856-1.629 | 0.311 | 1.15 | 0.818-1.616 | 0.421 |
| Diagnosis (ANLL vs ALL) | 0.628 | 0.482-0.818 | $5.6 \times 10^{-4}$ | 0.644 | 0.485-0.857 | $2.5 \times 10^{-3}$ |
| TBI | 0.751 | 0.506-1.115 | 0.155 | 0.687 | 0.449-1.053 | 0.085 |
| ATG | 1.186 | 0.467-3.016 | 0.72 | 0.788 | 0.292-2.126 | 0.638 |
| Cya vs tac | 1.109 | 0.854-1.440 | 0.439 | 1.067 | 0.810-1.406 | 0.644 |
| Relapse | 0.69 | 0.508-0.939 | 0.018 | 0.727 | 0.528-1.001 | 0.051 |
| HLA match | 0.705 | 0.536-0.928 | 0.013 | 0.727 | 0.544-0.972 | 0.031 |
| D6S0035i (MAPK14) allele 2 | 0.67 | 0.541-0.829 | $2.3 \times 10^{-4}$ | 0.672 | 0.538-0.839 | $4.5 \times 10^{-4}$ |
| D17S0219i (SOCS3) allele 3 | 0.456 | 0.289-0.721 | $7.5 \times 10^{-4}$ | 0.426 | 0.264-0.685 | $4.3 \times 10^{-4}$ |
| D1S0818i (ELTD1) allele 1 | 1.49 | 1.218-1.822 | $1.0 \times 10^{-4}$ | 1.46 | 1.185-1.799 | $3.9 \times 10^{-4}$ |
| D6S0330i (AGPAT4) allele 7 | 1.376 | 1.141-1.659 | $8.4 \times 10^{-4}$ | 1.435 | 1.180-1.745 | $2.9 \times 10^{-4}$ |
| D5S424 (F2RL1) allele 5 | 1.495 | 1.212-1.838 | $1.4 \times 10^{-4}$ | 1.497 | 1.206-1.859 | $2.5 \times 10^{-4}$ |
| D17S0271i (DDX42) allele 1 | 0.719 | 0.564-0.918 | $8.0 \times 10^{-3}$ | 0.731 | 0.566-0.944 | 0.016 |
| D1S1335i (AKT3) allele 3 | 1.229 | 1.018-1.483 | 0.032 | 1.221 | 1.005-1.484 | 0.045 |
| TNFC Allele 5 | 1.238 | 0.977-1.568 | 0.077 | - | - | - |

Table 4.14: Multivariate analysis of microsatellite alleles associated with grade 2-4 acute GVHD.

### 4.3 Further exploration of a susceptibility region by SNP typing

Microsatellites as applied in this study 'represented' and identified a region of linkage disequilibrium to disease-associated genetic features like e.g. functional polymorphisms, assuming an average length of linkage disequilibrium of approximately 100 kb . Further work of investigation was therefore needed aiming to limit down in size the disease-associated locus, or even identify the underlying genetic variation that causes the disease association (see the more detailed discussion on this Topic in the discussion section).

As an example, a small exploration was undertaken into the MAPK14 locus (marker D6S0035i) using tag SNP identified through the HapMap project. The region on Chr 6 was searched for 50 kb on each side of the microsatellite (6:36,100.000-6:36,200.000), identifying 159 SNP in 6 haplotype blocks, of which 25 SNP were tag SNP. Focussing on the largest of the haplotype blocks, five SNP (rs6934216, rs851020, rs16884919, rs12530381, rs7760405) were selected and genotyped using TaqMan methodology. As not all of the 25 tag SNP were available for this platform, not all haplotypes would be captured, but haplotype analysis was attempted.

Three of these SNP markers showed association with grade 2-4 acute GVHD:

- rs851020 and rs12530381 (the latter is the closest to the microsatellite, and the coding region of MAPK14) both associated with allelic and genotypic risk (table 4.15). These markers had an almost identical allele and genotype distribution, and may have been linked.
- rs6934216 and rs851020 showed a protective allelic and genotypic trend towards association (table 4.15).

Due to the limited capture no haplotypes were derivable from these data. Further work with either SNP or microsatellites would be required to investigate associations at this locus.


Figure 4.2: Map of microsatellite and tagSNP positions in the intronic region of the MAPK14 gene. The shaded area of the gene indicates the exon. The large arrow indicates the position of the microsatellite, the small arrows show the position of selected SNP. Dark arrows indicate association of the marker with acute GVHD grade 2-4.

| marker | allele freq | allele freq | assoc <br> allele | $\mathrm{p}=$ | OR | OR 95\% CI |
| :--- | :--- | :--- | :--- | ---: | :--- | :--- |
| rs12530381 | $\mathrm{A}(73 \%)$ | $\mathrm{G}(27 \%)$ | A | 0.0013 | 1.407493 | $1.1-1.8$ |
| rs6934216 | $\mathrm{A}(9 \%)$ | $\mathrm{G}(91 \%)$ | G | 0.287 | 0.831325 | $0.6-1.1$ |
| rs851020 | $\mathrm{C}(72 \%)$ | $\mathrm{G}(28 \%)$ | C | 0.0035 | 1.362699 | $1.1-1.7$ |


| marker | genotype <br> freq | genotype <br> freq | genotype <br> freq | assoc <br> genotype | p= | OR | OR 95\% CI |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| rs12530381 | GG (0.08) | AA (0.53) | AG (0.39) | AA | 0.0043 | 1.584844 | $1.2-2.1$ |
| rs6934216 | GG (0.82) | AA (0.07) | AG (0.16) | GG | 0.08 | 0.732426 | $0.5-1.1$ |
| rs851020 | GG (0.10) | CC (0.53) | CG (0.37) | CC | 0.00014 | 1.667178 | $1.3-2.2$ |

Table 4.15: SNP allele and genotype associations of markers close to microsatellite D6S0035i

### 4.4. Genetic susceptibility regions for moderate-severe acute GVHD

### 4.4.1. Introduction

Each of the identified alleles or markers represented a region of approximately 100 kb of linkage disequilibrium (LD) with an associated genetic variation. Microsatellites in themselves are rarely directly disease causative, but linked to such a variation which could be a polymorphism, a mutation, deletion or duplication, or epigenetic trait.

The specific LD for any of the associated genetic loci in this study was not immediately known. The HapMap database gave information about the LD of SNP markers but not microsatellites. Previous studies have used well known association loci, placing a variety of SNP and MS markers around it (Koch et al., 2000, Ohashi and Tokunaga, 2003) or well characterized populations (Varilo et al., 2003), or mathematical models (Terwilliger et al., 2002, Shifman et al., 2003) to determine LD of MS markers. These studies found that the LD of MS markers extended beyond that of SNP markers (up to 400 kb , compared to up to 30 kb for SNP). Also, LD decreased with distance from the marker, even on the same haplotype block.

Therefore it was reasonable to assume a LD of 100 kb as a starting point for locus analysis, until detailed exploration of each locus by higher density typing with more MS or SNP markers indicated its specific LD.

Obviously, intronic MS markers with no other genes within a 100 kb range were prime candidates genes for further association testing. Six loci (AKT3, ELTD1, AGPAT4, MAPK14, IL1RAPL2 with 2 loci) fulfilled these criteria. All of these had been target genes, and two of these (ELTD1 and MAPK14) had marker as well as allele associations that were consistent even with application of multiple testing correction and in multivariate analysis. Two loci had intronic markers that were not within the target gene,
but coincidentally within other genes within 100 kb distance from the target gene (TBL1X $\rightarrow$ SHROOM2; SOCS3 $\rightarrow$ DNEL2), both had further genes within LD range. Finally, the loci F2RL1 and DDX42 were intronic in the target gene but had several other, non-targeted genes at close range. The following sections will describe each locus in turn. Information has been obtained from the gene cards website (http://www.genecards.org/) and additional sources, as referenced.

### 4.4.2. AKT3 locus

Chr 1:241718158-242080053
MS marker D1S1335i location: Chr1:241980312-241980412

The AKT3 (Protein Kinase B isoform 3, PKB) gene is a large gene, more than 360 kb long. In the pooled screening it was covered by 5 MS markers, of which only this one, which was also closest to the coding region, showed an association (Figure 4.3).
AKT3 has a broad effect on cell function, it is an important regulator of cell signalling in response to insulin and growth factors, it has a role in cell proliferation, differentiation, apoptosis, tumorigenesis as well as glycogen synthesis and glucose uptake (Somanath et al., 2006). AKT dysregulation, mainly studied in mice, leads to diseases like diabetes, cancer, cardiovascular and neurological disease (Hers et al., 2011). In Tlymphocytes, AKT3 has in important role in cell development and proliferation. AKT3 regulates glucose uptake, protein synthesis, and stimulates the E2F and forkhead transcription factors (Matthews and Cantrell, 2006). In a genome-wide gene expression study of GVHD, PKB expression was repressed in donor CD4 T-cells in chronic GVHD (Baron et al., 2007).


Figure 4.3: Genomic map of the AKT 3 gene and position of the associated microsatellite

### 4.4.3. ELTD1 locus

Chr 1:79128037-79279105
MS marker D1S0818i location: Chr 1:79149764-79149943

ELTD1 (EGF latrophillin and seven transmembrane containing 1) is part of the EGF-TM7 (Epidermal Growth Factor seven transmembrane) family. It has important funcions in leukocyte adhesion and neutrophil migration, and defects in this gene had consequences for innate and adaptive immunity (Yona and Stacey, 2010, Leemans et al., 2004). Genetic variation in this gene had been associated with parasitic susceptibility in cattle (Porto Neto et al., 2011) and subcutaneous fat thickness in humans (Lee et al., 2011).


Figure 4.4: Genomic map of the ELTD1 gene and position of the associated microsatellite

### 4.4.4. AGPAT4 locus

Chr 6: 161332749-161458407
MS marker D1S0818i location: Chr 6: 161511402-161511576

AGPAT4 (1-acylglycerol-3-phosphate O-acyltransferase 4) is a membrane enzyme that is involved in de novo phospholipid biosynthesis. The wider function of this protein is not known. Variations in this gene, however, had been associated with acute as well as chronic GVHD in a population from Finland (Turpeinen et al., 2009). This study found the donor SNP rs749013 associating with risk of acute as well as chronic GVHD, mirroring the finding of this study (donor allele of MS associated with risk of acute GVHD). The SNP marker is located approximately 50 kb upstream towards the exon, and in contrast to the Finnish population is non-polymorphic in Japanese.


Figure 4.5: Genomic map of the AGPAT4 gene and position of the associated microsatellite

### 4.4.5. MAPK14 locus

Chr 6: 36129769-36215820
MS marker D6S0035i location: Chr 6: 36178949-36179320

The gene for MAPK14 (Mitogen-activated Protein Kinase 14) is located on chromosome 6 close to, but outside the HLA region. The function of this gene has been explored extensively. MAPK14 responded to activation by environmental stress, pro-inflammatory cytokines, HSP70 and lipopolysaccharides (as part of the TLR pathway) (Kang et al., 2008, Lissauer et al., 2009, Mackay and Sallusto, 2006). It was a regulator of chronic inflammation in rheumatoid arthritis (Korb et al., 2006) and inflammatory bowel disease (Waetzig et al., 2002). MAPK14 had effects on the recruitment of immune cells to the colonic mucosa (Kang et al., 2010) and epithelia of the skin (Eckert et al., 2003). MAPK14 was a key element in the activation of the glucocorticoid kinase, which decreased susceptibility to cytotoxic drugs and promotes cell survival (Meng et al., 2005).

On the basis of the largely pro-inflammatory effects of MAPK14, inhibitors have been developed for the treatment of rheumatoid arthritis and other inflammatory conditions. While in vitro and animal models did show very promising results, a recent randomized controlled clinical trial had failed to show any long term benefit for patients with rheumatoid arthritis (Genovese et al., 2011), indicating that the role of MAPK14 is complex.

MAPK13, a splicing variant of MAPK14, is located close to this gene.


Figure 4.6: Genomic map of the MAPK14 gene and position of the associated microsatellite

### 4.4.6. IL1RAPL2 loci

Chr X: 103697652-104898478
MS marker DXS0629i location: Chr X: 103769044-103769435 (intronic) MS marker DXS0151i location: Chr X: 103689786-103690263

The IL1RAPL2 (interleukin 1 receptor accessory protein-like 2) gene is very large, more than 1000 kb , it was covered for this study by 10 MS markers. Only these two markers, which closely flank the coding region of IL1RAPL2, showed association (albeit in opposite directions).
This gene is part of the IL1 receptor family, which was the reason for inclusion in this study. It has, however, so far no documented role in the immune system. There is extensive literature of the association of IL1RAPL2 with cognitive impairment and mental retardation (Valnegri et al., 2011).


Figure 4.7: IL1RAPL2 gene locus on chromosome $X$ with two MS markers (left: DXS0151i, right: DXS0629i), flanking the coding region of the gene.

### 4.4.7. TBL1X locus

Chr X: 9391369-9647778
MS marker DXS0324i location: Chr X: 9722847-9723231

The target gene TBL1X (transducin (beta)-like 1 X-linked) is a further large X-chromosome gene. The marker flanked the coding region at a distance of $\sim 80 \mathrm{~kb}$, and was co-incidentally located intronically within the gene SHROOM2. A further gene, GPR143, was located within a 100 kb LD range of the marker between SHROOM2 and TBL1X (see Figure 4.8).

Little knowledge exists about the function of TBL1X. It had an essential role in transcription activation mediated by nuclear receptors (Glass and Ogawa, 2006). It recruited NFkB to its target for gene transcription and had a potential role in tumorigenesis (Ramadoss et al., 2011), and also regulated MYC gene expression, which is important for growth and expansion of somatic cells (Toropainen et al., 2010). Genetic polymorphisms in TBL1X had recently been linked to autism in males (Chung et al., 2011), but the mechanism remained unclear.

SHROOM2 (shroom family member 2) had a broad role in the morphogenesis of thickened epithelial shields during embryonal development (Lee et al., 2009), and regulated epithelial proliferation and angiogenesis (Farber et al., 2011).

GPR143 (G-protein coupled receptor 143) was involved in intracellular signal transduction, in particular the transfer of melanin. Mutations in GPR143 lead to variant forms of albinism and mental retardation. The protein expressed by GPR143 also represented a self or tumour antigen (Touloukian et al., 2003).


Figure 4.8: TBL1X gene locus on chromosome X . The marker is intronic to SHROOM2.The MS position is located outside TBL1X, but LD of the marker includes the coding regions of TBL1X and GPR143.

### 4.4.8. F2RL1 locus

Chr 5:76150610-76166896
MS marker D5S424 location: Chr 5: 76193683-76193804

F2RL1 (coagulation factor II (thrombin) receptor-like 1) was implicated in chronic responses associated with vessel inflammation and wound healing; stimulated activation of T-cells and neutrophils, promoted leukocyte rolling, adhesion and extravasation, increased capillary permeability and enhances production of cytokines. High F2RL1 expression in experimental intestinal radiation injury promoted inflammation and fibrosis. F2RL1 has been demonstrated on T-cells, where it triggered in an essential manner the IL6-secretion induced by thrombin, trypsin and tryptase (Li and $\mathrm{He}, 2006$ )

The location of the MS marker was intronic of the S100Z gene, the function of which is unclear. A genome-wide association study has found SNP polymorphism in this gene to associate with severity of ulcerative colitis (Festen et al., 2010). The S100 gene family was described to have wide-ranging roles in tumorigenesis, autoimmunity and innate immunity.


Figure 4.9: F2RL1 gene locus. The MS marker is located within S100Z, but its LD range includes F2RL1.

### 4.4.9. DDX42 locus

Chr 17: 59205299-59250409 MS marker D17S0271i location: Chr 17: 59224879-59225107

This marker was intronic of DDX42 (DEAD (Asp-Glu-Ala-Asp) box polypeptide 42) in a very gene dense region. Other genes within 100 kb LD region included CCDC47 (coiled-coil domain containing 47) which had an unknown function, FTSJ3 (FtsJ homolog 3 (E.coli)), PSMC5 (proteasome (prosome, macropain) 26 S subunit, ATPase 5) and SMARCD2 (SWI/SNF related, matrix associated, actin dependent regulator of chromatin, subfamily d, member 2).

DDX42 was involved in the survival of cells, counteracting the apoptotic effect of TP53BP2 (Uhlmann-Schiffler et al., 2009), while FTSJ3 was involved in ribosome synthesis (Morello et al., 2011).

PSMC5 is a functional element of the proteasome. An essential function of a modified proteasome, the immunoproteasome, was the processing of class I MHC peptides. It participated in transcriptional regulation since it has been shown to interact with the thyroid hormone receptor and retinoid X receptor-alpha.

SMARCD2 was involved in transcriptional activation and repression of selected genes by chromatin remodelling.


Figure 4.10: DDX42 gene locus. The marker is located in DDX42, but the genes CCDC47 and FTSJ3 are also within in LD range.

### 4.4.9. SOCS3 locus

Chr 17: 73864454-73867753
MS marker D17S0219i location: Chr 17: 73941484-73941691

This locus has a high gene density. The target gene, SOCS3 (suppressor of cytokine signalling 3) was just within LD range of the marker, which was intronic to DNEL2, also known as DNAH17 (dynein, axonemal, heavy chain 17). The other gene within range was PGS1 (phosphatidylglycerophosphate synthase 1).
DNAH17 is the force generating protein of respiratory tract ciliae and sperm flagellates. The function of PGS1 is unclear. SOCS3 is an important negative regulator of the janus kinase pathway, which is used by most cytokines for transcription activation (Tamiya et al., 2011), hence it had an important anti-inflammatory effect (Alexander and Hilton, 2004). In the animal model, SOCS3 had a suppressing role on the severity of GVHD (Hill et al., 2010).


Figure 4.11: The SOCS3 gene locus. SOCS3 is just at the border of the LD range of the marker, which locates to DNAH17 (DNEL2). PGS lies between SOCS3 and DNAH17.

## 5. DISCUSSION AND CONCLUSION

5.1 Strengths and limitations of the methodology
5.2 Discussion of results
5.3 Future
5.4 Conclusions

### 5.1. Strengths and limitations of the methodology

### 5.1.1. Introduction: Towards a high-quality gene association study

In the introduction to this study, existing literature on non-HLA gene polymorphisms associating with HSCT outcome was reviewed in a systematic way; the finding was that the majority of these studies have methodological quality issues, relating to study populations (small, heterogeneous), target gene loci (candidate gene selection) and statistical analysis (statistical power, application of multiple testing correction). There is now quite clear guidance on the design of high-quality genetic association studies(Colhoun et al., 2003, Gambaro et al., 2000, Lander and Schork, 1994, Schork, 1997, Rannala, 2001). The question is:

## To what extent did this study fulfil criteria of a high quality genetic association study?

The main areas pointed out by these authors include:

- Population: Genetic structure, confounding variables, case and control definitions, phenotype heterogeneity, outcome classification
- Statistics: Requirement for a pathophysiological link between gene and disease, failure to attribute results to chance (type I/II error), sample size, statistical power, independent confirmation, multiple testing
- Selection of target genes and markers

More recently, a review (McCarthy et al., 2008) of the methodology of several very large scale genome-wide association studies with SNP markers for common variant genetic traits highlighted the need for careful definitions of cases and controls, large sample sizes and replication study
in order to achieve robust results. Replication, even when using robust methodology, had often not being consistent; not necessarily indicating spurious results but the varying impact of clinical or environmental variables on a small effect size polymorphism. This phenomenon had been termed 'informative heterogeneity'.

### 5.1.2. Population

## Did the study population stem from a homogeneous genetic background?

It was pointed out that the studied population should stem from a homogenous genetic background without genetic admixture. All HSCT pairs entered into this study were genetically of Japanese origin. We knew from data of the HapMap Consortium (Consortium, 2005, Stranger et al., 2005) that genetic variation varies enormously between different populations. The Japanese population had a genetic structure of less, and better preserved haplotype blocks than Caucasians or Africans (Conrad et al., 2006, Gabriel et al., 2002). This was advantageous for the power of the study, as less MS markers were required for gene coverage (details of the genetic background were described in more detail in supplementary file 2.1).

As this approach was expected to yield robust results for a Japanese population, findings may be less applicable to other populations and would require confirmation. Allele frequencies and genotypes of MS as well as SNP markers, and associated genetic disease risks, vary amongst populations, some markers may be polymorphic and disease associated in one population and non-polymorphic in another. Nevertheless, there is substantial concordance of genetic variation across populations also, allowing for comparison of risk and large scale studies.

How well did this study address demographic or clinical confounders that affect acute GVHD severity?

Clear definition of cases, controls, risk factors and outcomes were essential. In Japan, since 1992 the consensus on the modified Glucksberg criteria for the diagnosis of GVHD (Glucksberg et al., 1974, Przepiorka et al., 1995, Rowlings et al., 1997) was used. As these contained subjective elements, and as several independent centres across Japan were applying these criteria, there were likely elements of intra- and inter-observer variability, which were not quantifiable for this population.
The criteria for selection of this study population did reduce confounding variables to some degree:

- Selection by diagnosis: Different underlying diagnoses carried variable risks for GVHD. CML and ALL in particular had an intrinsic GVHD risk, compared to other malignancies. Focusing on ALL and ANLL had reduced this variability. However, ALL and ANLL in itself had significantly different risks for GVHD, remaining a confounder.
- Recipient age: Recipient age >40 years per se was a risk factor for GVHD, and exclusion of patients above this age removed the effect of older age. Infants and young children with leukaemia also have a higher risk of GVHD, therefore exclusion of this age group reduced confounding. However, a minor effect towards GVHD risk remained in the age group below 10 years.
- Selection of unrelated bone marrow as graft source and myeloablative conditioning: These measures avoided effects on GVHD as an outcome by donor source (reduced GVHD risk with related donors), other stem cell source (higher, or lower GVHD risk with e.g. peripheral blood stem cell transplantation (PBSCT) or cord blood), conditioning (lower GVHD risk with reduced-intensity conditioning).
- The selection by diagnosis and age 'streamlined' some of the clinical confounders. Treatment for leukaemia was largely protocol-
driven, compared to other indications, hence variables such as previous chemotherapy, conditioning regimen and GVHD prophylaxis were very similar.

As previous data and sample collections were often small, there was little room for considering clinical risk factors for GVHD or other HSCT outcomes, although these were well established (Loren et al., 2006, Randolph et al., 2004, Perez-Simon et al., 2005, Kollman et al., 2001). In this situation, multivariate analysis was applied to correct for the clinical confounders. Despite the selection criteria applied, this study had remaining confounding variables that required multivariate analysis:

- Donor age >30 years
- HLA mismatching
- GVHD prophylaxis with Cyclosporin A or tacrolimus
- Minor effects of recipient age, conditioning regimen

The prevention, diagnosis and management of GVHD had changed over this time period. Although all patients had myeloablative conditioning, this varied with underlying disease and staging. While the prophylactic regimen in Japan consisted initially of cyclosporine A, methotrexate and steroids, a change from cyclosporine A to tacrolimus reduced the incidence of GVHD (Hara et al., 2007). The JMDP database did not provide detailed information on dosage of methotrexate and steroids in the earlier HSCT.
ATG and T-cell depletion, in contrast to Europe or North America, had little role in Japan as it was found to dramatically increase the relapse rate of leukaemia. A few of the more recent transplants in this study cohort used in addition newer agents such as mycophenolate mofetil. Similarly, the management of GVHD over these 12 years had evolved - improved diagnostics, better control of immunosuppression, better control of concurrent infections through improved diagnostic tests and newer
antibiotics, better nursing care and monitoring, and new, more powerful treatments of GVHD had gradually changed the characteristics of GVHD. Most of these characteristics were not captured in the dataset as such information was not prospectively collected at the time.

The application of two separate time frames by this study was likely to provide some correction for this type of confounding, but may also have had effects on the sensitivity of the study:

- Associations consistent across both subsequent time frames would indicate some degree of independence from the effects that changes in supportive transplant practice, factors that were not recorded in the dataset, over time would have had on outcome.
- The disadvantage of this approach was that it ruled out all associations that did not have that degree of consistency, i.e. associations that evolved in the cohort of the second screening alone would have went undetected. This was likely to have a negative effect on sensitivity of the study. The findings indicate associations that were valid consistently over a long period of time, but may not necessarily represent the strongest effects on GVHD
- Despite the effort of correcting for genetic and clinical confounders, significant variables affect outcome, as the multivariate analysis demonstrated.
- The effects of HLA matching and mismatching could not effectively been adjusted between the cohorts. This study was able to adjust HLA match or mismatch by pairing of samples from the first screening cohort with those from the second cohort at the HLA locus or locus combination, but not by serogroups or even alleles. More recent research from the Japanese registry has shown that different allele mismatches at the same HLA locus may have risk as well as protection effects, depending on the specific allele (see below).


## What was the effect of HLA matching and mismatching on GVHD?

Because HLA matching was known as a significant risk factor for GVHD (Morishima et al., 2002, Morishima et al., 2007, Kawase et al., 2009, Oh et al., 2005, Sasazuki et al., 1998, Kawase et al., 2007), and possible to control in a study setting, most previous studies in the 1990's and 2000's used higher HLA-matched HSCT pairs (e.g. 8/8, or even 12/12 matched) or related HSCT for genetic association studies. This measure was thought to reduce genetic confounding caused by HLA mismatching. The disadvantage of this approach was that data from these studies were meaningful to only a small subset of the HSCT population in clinical practice, and that other clinical confounders could not be addressed due to sample size issues.

Consequences for this study of the decision not to select samples by HLA matching or mismatching:

- Not restricting inclusion for degree of HLA matching made this study population more representative of a HSCT population seen in clinical practice.
- The large proportion of HLA mismatched pairs would allow for an analysis of the effects of non-HLA gene polymorphisms in an HLA mismatched population.
- The proportion of HLA matched HSCT pairs was large enough to permit subgroup analysis.
- The adjustment of the degree of HLA matching of the confirmatory cohort ( $2^{\text {nd }}$ screening) to that of the discovery cohort ( $1^{\text {st }}$ screening), in order to achieve a similar degree of genetic confounding, has introduced stratification. The JMDP registry population between 2001 and 2005 had an overall higher degree of HLA matching than the population from 1993-2000. Hence, the HLA matching of the confirmatory cohort was not representative of the degree of HLA matching of the Japanese registry cohort during that time; mismatching was over-represented. This was reflected in the GVHD
prevalence, which was higher in the confirmatory cohort than in the discovery cohort.


## How could the selection study be modified to minimize confounding and stratification?

As HSCT is rapidly expanding, larger registry cohorts may become available for research in the near future. Important steps to reduce clinical and genetic confounding would include:

- Reducing the time frame of sampling (e.g. 5 years), to reduce confounding that stems from development in supportive therapy.
- Use of most recent transplants - reflect more the current clinical practice
- Aim for higher cohort size to increase statistical power
- Focus on single large diagnostic groups (eg. ALL, AML separate) rather than combined, as these carry in themselves different risks for outcomes
- Rather than by time frames, the study cohort could be divided by other important variables, such as degree of HLA matching. This requires more detailed understanding of the risk of HLA mismatching by e.g. high-risk allele mismatches or HLA haplotypes. Such analysis would be very valuable in directly comparing the competing risks of HLA mismatch and non-HLA gene polymorphisms. In addition, this analysis would give an insight into the non-HLA immunogenetics of HLA mismatched HSCT, an area that it under-researched.
- As SNP gene-chip GWAS typing is becoming more readily available, it would be useful to conduct this approach in a parallel study mirroring the same set up. This would provide a complementary perspective on the genetic variation in the same study population, and facilitate the fine-mapping of associated microsatellite loci (SNP or SNP haplotype associations within LD

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range of associated MS could limit down the area requiring sequencing).
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## SUMMARY: Study population

- The study population stemmed from a genetically relatively homogeneous background
- Case and control definition followed standard practice of GVHD classification.
- Extensive efforts were made to identify and address confounding variables. The effect of confounding variables could be reduced by careful evaluation in a larger stem population and selection of a study population.
- Nevertheless, substantial clinical confounding remained and required multivariate analyses.
- Selection carried a risk of stratification for new confounders, which needed to be identified (e.g. HLA matching, see above).
- Limited availability of large study cohorts and rapid evolution of the field of HSCT made effective control of confounding variables difficult.
- Careful selection of a population that would be relevant in clinical practice rather than just of research interest (e.g. understanding pathobiology) may facilitate clinical application of study findings (e.g. risk stratification).


### 5.1.3. Reasoning for choice of scale of approach

## Which genes to focus on in relation to GVHD - all, some, or few?

The findings from the literature appraisal indicated that a larger scale, more systematic approach to candidate gene and marker selection was required to better understand the effect of non-HLA gene polymorphisms on GVHD. The initial decision was on the extent of cover of the genome - ranging from a selected candidate gene approach to whole genome scanning. Table 5.1 lays out the advantages and disadvantages of three approaches to gene selection, representing three degrees of indiscrimination: A candidate gene approach, where a small number of genes believed to have a high likelihood of being involved in the pathogenesis of acute GVHD (highly discriminate); a targeted genomic approach (e.g. focusing on the immune system as a whole, less discriminatory), and a genome wide scanning approach (indiscriminate).

The decision for a targeted genome scanning approach, which includes the genes of the entire immune system in a broader sense ('Immunogenome'), supplemented by genes relevant for GVHD (previously associated genes and other strong candidates) and leukaemia (e.g. susceptibility genes, drug metabolism genes), was based on aspects of feasibility and study quality:

- Previous data from genome-wide association studies indicated that immune response associated conditions tend to have genetic associations within the immune system genes (Zhernakova et al., 2009) - as GVHD is a result of immune system dysregulation, focusing on immune system genes should have yielded a higher positive predictive value for associations.
- Reduced number of markers reduced the need for multiple testing statistics - resulting in higher power
- Feasible in the available time (3 years) and with available resources
- Reasonable trade-off between number of available samples, resulting statistical power of cohorts, multiple testing statistics

|  | Candidate approach | Targeted genome scanning | Whole genome scanning |
| :---: | :---: | :---: | :---: |
| Advantages | - Simple to perform <br> - Simple statistics <br> - Cheap <br> - Powerful with small sample size | - Systematic approach to biological meaningful selection of genes <br> - Statistically robust with reasonable sample size, positive predictive value | - Indiscriminate approach to genes <br> - Reflects full genetic contribution to disease |
| Disadvantages | - Likely to miss important associations | - May miss important associations outside targeted system | - Requires large sample size <br> - Issues with statistical power, multiple testing <br> - Expensive <br> - Time consuming |

Table 5.1: Advantages and disadvantages of different approaches to gene selection

SUMMARY - Scale of approach

- A targeted genome scanning (indiscriminate selection of all immune system genes, 'immunogenome') provided the best trade-off between the competing factors of gene selection, statistical power and available resources.
- Main disadvantage was that important associations outside the selected panel may not be detected.


### 5.1.4. Reasoning for choice of marker type and marker number, and pooled/individual typing approach

## Single Nucleotide Polymorphisms or microsatellite markers? <br> What were the consequences for study design by choosing microsatellite markers?

Modern high-throughput genetic screening approaches mainly use two different types of genetic markers:

- Single Nucleotide Polymorphisms (SNP): A SNP is a genetic variation where a single nucleotide within the genome sequence is altered. SNP in coding sequences may alter gene expression and cause a change in biology, or may link to a causative variant.
- Microsatellites (MS): MS are short sequence repeats, commonly as a variable number of di, tri, or tetra tandem base repeats. Microsatellite markers often, but not always stand in linkage desequilibrium (LD) with SNP on the same haplotype block.

SNP and MS both have their specific advantages and disadvantages (see table 5.2) (Bahram and Inoko, 2007, Jorgenson and Witte, 2007).

At the time this study was developed (2005-2006), new miniaturized, standardised and automated genome-wide SNP typing platforms were evolving, using hundreds of thousands of SNP markers on a single sample. During this time there was still an ongoing debate about which markers to include in these assays, sample size and statistical power, typing quality, and processing of the vast amount of data. In addition, these systems were still very expensive, especially for larger scale studies (table 5.3 shows a cost comparison of a SNP based and MS based approach for 2007). The advantages of such an approach were obvious: The methodology became technically ever simpler, reproducible, faster and cheaper, individual
sample data were retained and allowed for multiple outcome measures. Initial studies had limited numbers of samples (in the hundreds), and statistical power was limited, due to the enormous multiple testing burden. In order to detect the small effect sizes of common variants, now sample sizes of tens of thousands have been used (Vercelli and Martinez, 2006, Anonymus, 2007). Such large numbers still remained a challenge for many conditions, including HSCT where the largest collections of data and sample have just reached (CIBMTR/NMDP) or are approaching (JMDP) the 10,000' mark. Providing consistent case, control and phenotype definition was still a major problem.

There were clearly advantages of the MS approach in this setting. The method of genome-wide scanning with MS markers was pioneered and brought to a high standard by a Japanese group (Prof Inoko, Tokai University), with the first study published in 2005 (Tamiya et al., 2005) (summarised in Figure 5.1). This approach used almost 30,000 MS markers spread throughout the genome at high density (charting at 100 kb with overlap), and sequential screening in three steps on pooled DNA. More than ten genome wide studies using this approach have since been published.

MS markers have a larger LD range - therefore less markers were required to provide genome coverage, as compared to SNP, which gave MS an advantage with regards to power, sample size and multiple testing correction. The haplotype block structure of the Japanese population was highly preserved - allowing the full exploitation of the large LD of MS markers. Our selection of markers achieved a very high density of coverage for the target genes - 97\% of target genes had either two flanking MS markers, or at least one, within the projected LD range of 100 kb. (the genome-wide MS panel would provide cover to $\sim 90 \%$ of the euchromatic region of the genome). On the other hand, true LD range of MS markers was not known, therefore the LD may have been shorter or longer, creating either gaps or extended coverage. In contrast to SNP arrays, which were so densely packed that individual markers could not
have been regarded as independent, MS loci could be regarded as independent and gave a clearer distinction of haplotype blocks.

By evolution, SNP and MS were often closely linked, with a MS indicating mutation or causative SNP polymorphism within its LD. By mathematical models, SNP markers, which have only two alleles, were more resistant to mutations as compared to MS, which could have 2-20 or even more alleles. A MS marker therefore may have 'mutated away' and lost its linkage; while on the other hand it may have indicated an evolutionary more recent genetic risk variation that SNP may not had captured (Oka et al., 2012, Hiruma et al., 2011). From this model, MS and SNP approaches were complimentary to each other by having a large area of overlap, and each additional aspects of genomic variation which the other approach did not cover.

Limitations of using a microsatellite-based approach include existing gaps in cover, and the fact that the pooled approach would allow for allelic, rather than genotypic association.

- This study had a very good cover of target genes (almost 90\% full cover, and a further 7\% partial cover. However, there are gene regions within the genome which have no suitable natural microsatellites, or have genetic variation (e.g. gene duplication) that make microsatellite typing inherently difficult.
- High-throughput MS typing required pooling of DNA, which lead to loss of individual genotypic information in the screening stages, relying on allele frequency differences between pools alone. Allele frequency differences often but not always reflect genotypic risks, and important associations may have been missed that way. Additionally, at individual level a pooled allele frequency difference indicating a protective effect may translate at individual genotypic analysis into a risk, and vice versa. Hence, at the individual genotyping stage the finding of an associated allele should ideally
backed by identifying a genotypic association. This, however, is not always possible given the large number of genotypes for microsatellites resulting from the large number of alleles. For some markers, nevertheless, this study was able to confirm the association of homozygous genotypes.

|  | SNP | MS |
| :---: | :---: | :---: |
| Advantages | - Allows for individual genotyping and data analysis <br> - High-throughput platforms available <br> - Technically more simple <br> - Lower mutation rate more stable over time, but misses recently evolved genetic variation | - Wider linkage desequilibrium <br> - Requires less markers than SNP for same coverage (~30,000 for whole genome) <br> - Statistically more powerful allows for lower sample size with same coverage <br> - more alleles - more informative <br> - Clear definition of haplotypes <br> - Cheaper to perform than SNP |
| Disadvantages | - Short LD range - many gaps, may miss epigentic variation <br> - Unclear definition of haplotype blocks ('virtual' haplotype definition) may miss important associations <br> - Large number of markers required for coverage (>500,000 for whole genome) <br> - Multiple testing requires large sample size for sufficient power <br> - Expensive <br> - Time consuming <br> - Requires high-density SNP typing of candidate regions | - Higher mutation rate - may miss older SNP associations <br> - No high-throughput platform available - requires pooled DNA approach <br> - Technically more demanding <br> - Requires high-density SNP typing of candidate regions <br> - Artifacts introduced by DNA pooling <br> - DNA pooling allows for single outcome measure only at screening stage |

Table 5.2: Advantages and disadvantages of large scale genomic approaches using SNP and MS

|  | SNP | MS <br> (individual) | MS (pooled) |
| :--- | :--- | :--- | :---: |
| Estimated no <br> of markers | 70,000 | 4,000 | 4,000 |
| Estimated <br> assay costs | $\$ 600,000$ | $\$ 8,000,000$ | $\$ 70,000$ |
| Estimated <br> time | 3 years | $>10$ years | 3 years |

Table 5.3: Estimation of assay costs and time requirements for a targeted immunogenome scanning study, based on $n=1000$ sample pairs (time point March 2007) in US \$
*this includes costs for 3 pooled screening steps and individual genotyping of identified candidate gene regions with approximately 100 SNP.

## Approach to genome-wide scanning (Tamiya et.al.)

- A simple case-control study design of a cohort of $n=375$ patients and an equal number of controls
- Highly accurate DNA pooling, constructing 3 pools each for cases and controls, each containing $n=125$ individuals
- A phased three-step genomic screening on pooled DNA. A panel of 27,039 MS markers was tested on the first set of case and control pools, markers found associated ( $\mathrm{n}=2,748$ ) were tested on the second pool set. Markers still remaining positive ( $\mathrm{n}=372$ ) were subjected to typing in the third screening step, which still left n=133 MS markers associated. Chi Square and Fisher's exact test for $2 \times 2$ and $2 \times m$ tables, with a significance level of $p=0,05$ were used to establish the association after each screen
- The remaining $n=133$ markers were individually genotyped on the combined set of $n=375$ cases and control pairs, to eliminate errors potentially caused by the pooling process. Of these, $n=47$ still remained associated.
- Of these $\mathrm{n}=47$ associated gene regions, $\mathrm{n}=7$ regions were selected for fine mapping with SNP, based on high allele frequency and high degree of significance. Fine mapping was performed on a further independent cohort of $n=565$ cases and control pairs.
- Statistical power estimation indicated that the power to detect allele frequencies $<0.25$ is limited given the sample size per pool ( $n=125$ )
- Based on data on the pooling method, and data from test markers typed individually and in the created pools, difference in allele frequency was calculated to be $<4 \%$.
- As the number of multiple comparisons in this screening is $\mathrm{n}+1$, pseudopositive markers were calculated as $\mathrm{n}=1352$ (first screening), $\mathrm{n}=257$ (second screening) and $\mathrm{n}=25$ (third screening).
- Pritchard's method was used on a set of 69 randomly selected markers to verify the absence of stratification.

Figure 5.1: Methodological summary of a genome-wide scanning approach with MS markers

SUMMARY - marker choice and consequences for study design

Microsatellite markers have inherent advantages over SNP markers:

- They are more polymorphic and therefore more informative.
- Because their LD range is wider, for a given genomic region less MS markers are required than SNP to provide the same cover, giving the MS approach a statistical advantage.

Choosing a MS marker based approach has consequences for study design:

- The variability in PCR and typing requirements means that there are no commercial high-throughput platforms available.
- High throughput can be achieved by DNA pooling, which is technically complex.


### 5.1.5. Technical aspects

What were the technical challenges of the study design, and how were they addressed?

The microsatellite approach, however, had some drawbacks, mainly for technical reasons, some of which were discussed in the first reported genome-wide association study using this approach (Tamiya et al., 2005). This study was derived from this approach and shared its methodology, marker set and analysis tools.
Due to their larger variability and resulting variable length of the marker amplicon, MS markers did require variable PCR conditions which hinder automatization. The genome wide study approach mentioned above (Tamiya et al., 2005) had therefore selected preferentially those markers that had similar amplicon sizes and PCR conditions. However, $3-5 \%$ of markers frequently failed in genome-wide scanning, requiring reprocessing in order to keep the fail rate below $2 \%$. These circumstances made the development of standardized, fast and cheap assays difficult, and ruled out a large-scale, high throughput approach on individual samples.
There was a focus on MS markers that had 3-6 major alleles, which was a trade-off between technical conditions, LD, statistical power and controlling the false-positive error rate. Smaller numbers of alleles (e.g. 2) reduced the informative content, while markers with larger numbers of alleles could be highly informative and indicate rarer variants but had a poor statistical power.
Another source of technical difficulties was the requirement for a highthroughput platform, as the individual sample PCR and typing of several thousand MS markers would have been prohibitive from a time and cost perspective. Pooling of DNA was applied, and the method for this was refined to a degree that the SD for allele frequency differences between pools and individual genotyping could be kept near +/-1\%. However, the pooling process required expertise and time, had size limitations (the volume of the PCR reaction limits the maximal pool size to $\sim 1000$
individuals) and produced some inherent artefacts that required individual inspection and allele selection of each set of pools per marker. Some algorithms were developed to semi-automate this inspection step and take the element of judgement out of this process (Schnack et al., 2004, Perlin et al., 1995, Miller and Yuan, 1997, Matsumoto et al., 2004, Olejniczak and Krzyzosiak, 2006). A further disadvantage of DNA pooling was that the capability to analyze individual information was lost. Samples had to be pooled towards a single outcome, which restricted the extent of the analysis.

SUMMARY - technical challenges of genomic screening with microsatellites

- Due to their high polymorphism, MS markers have varying PCR and typing conditions.
- Markers with 3-6 alleles are preferentially selected, representing a trade-off between marker informative content and statistical power.
- DNA pooling requires time and expertise, and has drawbacks such as PCR and typing errors, and the loss of individual data and analysis.


## What was the technical validity of this study?

## Typing errors

It was already mentioned in the results section that the MS marker panel for this study was re-plated from the genome-wide marker panel plates which were sorted according to PCR typing conditions; therefore the higher number of initial PCR and typing error was expected, and corrected successfully (overall $1.1 \%$ (donor) and $1.0 \%$ (recipient) of the $4,321 \mathrm{MS}$ markers eventually failed PCR and typing over the two screening steps.

Was the pooled DNA representative of the true allele frequencies?

The quality of the DNA pools, as assessed by a test marker, was in keeping with the previously reported studies, with no significant allele frequency differences between pools and individual typing. The mean allele frequency difference was below $1 \%$ in all pools (which was lower than the $2 \%$ reported previously (Tamiya et al., 2005)).
This marker, however, represented only a spot measure of an 'ideal' marker under optimal PCR and typing conditions. To gain a better understanding of the consistency of DNA pool quality the pooled and individual allele frequencies for the 30 MS markers that were typed individually were reviewed (data not shown). Again, a very high concordance was found between pooled and individual typing.

How were pooling/typing artefacts identified and addressed?

The methodology was successful in identifying spurious associations due to artefacts. Of the eight MS markers that were not confirmed in individual typing, six had additional low frequency alleles that were not detected in pooled typing, and rendered the pooled allele association non-significant. Only two markers had an over-estimation of the associated allele in the pool resulting in an association that could not be confirmed in individual typing. This represented a known MS PCR error (preferential amplification
of the shorter-repeat alleles) which was exaggerated through the pooling process. Other artefacts that became apparent only in individual genotyping resulting in non-confirmation of the pooled association included mistaking a starter peak for an allele, non-polymorphism, and multiple alleles due to copy number variation.

How were false positive associations addressed?

The two screening steps of this study had, compared to the initial publication of this methodology, a higher rate of false positive association (table 5.4). This table shows also a projection of this study based on the data of Tamiya et.al.(Tamiya et al., 2005) Tamiya et.al.'s study had seen a reduction in marker number by approximately $90 \%$ in the first and second pooled screening step, and around $60 \%$ in the third pooled screening. In this study a reduction by approximately $75 \%$ and $65 \%$ was observed in the pooled screenings. Only the third screening step (which was not a pooled screening in this study, but a selection by allele identity and OR direction) reduced the number of associations close to the number projected.

There were differences between the study by Tamiya et.al. and this study that could explain these findings. Tamiya et.al. had a smaller pool size ( $\mathrm{n}=125$ cases/controls), so the statistical power to detect low frequency alleles was much reduced, and the investigators would consequently not include alleles of a frequency of $20 \%$ or lower. The pool sizes of this study were larger, which increased statistical power and sensitivity of low frequency allele detection. In addition, in order to capture all potential associations even if of low frequency, alleles of a frequency of $5 \%$ and above were included. This allele frequency was within the technical resolution of the pooled DNA approach, but at the border or below the statistical power for the sample size. Analysis of the $2^{\text {nd }}$ pooled screening showed indeed that $48 \%$ (donor) and $51 \%$ (recipient) of associations had allele frequencies below $15 \%$.
Tamiya et.al. did conduct the three pooled screening steps with selection of identified alleles and markers by p-value only (without correction for multiple testing), and selected by OR direction only after the third
screening. This study has shown that using a reduced number of MS markers for a targeted screening, a two-step pooled screening approach was sufficient.

|  | $\begin{aligned} & \hline \text { Tamiya } \\ & (2005) \end{aligned}$ |  |  | This study projected |  |  | This study Donors |  |  | This study Recipients |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Markers total | Markers positive | \% | Markers total | Markers positive | \% | Markers total | Markers positive | \% | Markers total | Markers positive | \% |
| 1st screen | 27039 | 2847 | 10.53 | 4321 | 455 | 10.53 | 4321 | 1016 | 23.51 | 4321 | 931 | 21.55 |
| 2nd screen | 2847 | 372 | 1.38 | 455 | 60 | 1.4 | 1016 | 335 | 7.75 | 931 | 314 | 7.27 |
| 3rd screen | 372 | 133 | 0.49 | 60 | 22 | 0.5 | 335 | 31 | 0.72 | 314 | 17 | 0.39 |
| Indiv typing | 133 | 47 | 0.17 | 22 | 7 | 0.17 | 31 | 6 | 0.14 | 17 | 4 | 0.09 |

Table 5.4: Comparison of rates of positive association between the original genome-wide study (Tamiya 2005), the projection for and the actual results for this study. The rate of positives in the first and second screens is higher than in the genome-wide study and the projection, but reaches very similar rates at the entry and after individual genotyping.

SUMMARY - Internal quality controls point to a high degree of technical validity of this study:

- PCR and genotyping errors were followed up and largely eliminated
- Artefacts were actively sought and resolved/excluded
- Prospective typing of a test marker showed a high degree of concordance between allele frequencies in the pools and by individual genotyping, in keeping with previous studies using the same approach.
- Retrospective analysis of individually typed associated markers confirmed the high concordance
- A larger number than expected of false positive associations was anticipated and effective measures taken to separate false positives from true positives


### 5.1.6. Data analysis

## Statistical aspects: Are the analyses valid?

The targeted genomic approach had advantages over genome-wide approaches with regards to statistical power. Investigation of the immune system was achieved with around $15 \%$ of the number of markers needed for a genome-wide scan. The selection of disease-relevant genes provided a higher positive predictive value and immediately gave a pathophysiological link between gene and disease, thus increasing the likelihood that the association were true.

Aspects of type I and II errors were already discussed in the previous section 'technical aspects'. Knowing that non-HLA associations with GVHD were likely to be of small effect size, for this study a high degree of sensitivity was deliberately chosen for the two pooled DNA screening steps (by including allele frequencies at the level of technical resolution, rather than at the level of statistical power; and non-application of multiple testing correction), accepting a large number of false positive associations. Independent confirmation in a second screening cohort was sought to confirm or refute the associations of the first screening cohort, and provided a powerful tool for further selection (allele identification, OR in the same direction). This approach did indeed reduce the number of false positive associations efficiently to levels that were projected, and identified a number of consistent associations. Most of these associations had effect sizes close to the limit of detection given the statistical power of this study; hence this was a successful strategy. It was, nevertheless, not possible to establish which falsely negative associations may have been missed, as a validation study of the approach was beyond the remit of this project. On the other hand, several publications of studies having used the pooled DNA microsatellite based approach have been successful in identifying new associations, and most of these studies confirmed the findings of previous SNP studies, i.e. on rheumatoid arthritis (Tamiya et al., 2005),
hypertension (Yatsu et al., 2007), Asthma (Hui et al., 2008), adult height (Kimura et al., 2008), anorexia nervosa (Nakabayashi et al., 2009), complications after radiotherapy for cancer (Michikawa et al., 2010), psoriasis (Hiruma et al., 2011) and macular degeneration (Meguro et al., 2012). The genetic risk for Beh et's disease was explored using the identical genome-wide marker set independently in two populations (Japanese, Korean), resulting in the identification of HLA-B51 as a genetic risk (Meguro et al., 2010, Horie et al., 2012). Multiple correction statistics (Bonferroni's correction) were applied for the individual genotyping step only, still resulting in four MS markers being associated with grade 2-4 acute GVHD.

Nevertheless, detection of effect sizes much below an OR=1.5 was very limited due to the small number of samples (for a genomic screening approach). Low frequency alleles caused a large number of false positive association, hence these were not replicated very well which may have statistical power as well as technical reasons.

SUMMARY - data analyses

This study's approach was deliberately sensitive by:

- using technical resolution of the scan (rather than statistical power resolution)
- non-application of multiple testing correction through a twostep screening process.
- Provision of specificity by independent confirmation of associations that were true but had a low effect size.

Ten such loci were identified, while the number of false positive associations were effectively reduced.

The statistical power given the pool and cohort sizes, allele frequencies and number of tests was moderate - associations of alleles with an $O R<1.3-1.5$ may not be reliably detected.

SUMMARY - methodological strengths and limitations I

Strengths:

This study fulfilled criteria for a high-quality genetic association study:

- Population of genetically homogeneous background
- Attempt to control genetic, demographic and clinical confounders
- Discovery/independent confirmation study design
- Large scale, indiscriminate gene targets

Technical strengths:

- Microsatellite markers - wide LD, informative
- control of typing errors
- high quality DNA pooling
- Control of artefacts
- Control of false positive associations


## Statistical strengths:

- Sensitive and specific for small effect size associations

SUMMARY - methodological strengths and limitations II

## Limitations:

The design of study cohorts for genetic association studies with HSCT outcomes is difficult:

- Control of confounding variables still requires multivariate analyses
- Cohort design by selection may induce stratification and new confounders
- HLA matching adjusted between cohorts, but not representative of registry

Technical limitations:

- Targeted genomic approach may miss important associations
- MS markers may not capture all genetic variation at a locus
- MS requires pooled DNA approach - loss of individual information, potential of typing errors
- Lack of validation study for this approach - scope of false negative markers unknown

Statistical limitations:

- Sample size provides limited statistical power - associations with Odd's ratio <1.5 not well represented, HLA subgroup analysis not very powerful


### 5.2 Discussion of results

### 5.2.1. Hypothesis and objectives

The hypothesis of this study stated: "Allele frequency differences of microsatellite markers are associated with moderate-severe acute GVHD".

The finding of 10 microsatellite loci associated with moderate-severe acute GVHD through a methodology that complied with many requirements for a high-quality genetic association study, with valid technical and statistical results, confirmed this hypothesis; rejecting the null hypothesis: "Allele frequency differences of microsatellite markers are not associated with moderate-severe acute GVHD."

The assumption for this hypothesis was that GVHD is in part a complex genetic trait, and that common allele polymorphism of non-HLA genes in the patient and donor genomes contributed to the development of GVHD. It was also assumed that such non-HLA risk alleles had an effect size that reached that of certain HLA mismatches, hence could be consistent despite variation in clinical and genetic risk factors over time. The objective of this study, the demonstration of the existence of such polymorphisms through a targeted genome scanning with MS markers, was therefore achieved.

### 4.2.2. Roles and functions of identified associations

## Exploratory study

Before embarking on a large scale genomic approach, the study population was explored by a smaller scale pilot study in order to establish whether identification of small size non-HLA polymorphism would be feasible in this population. For this purpose 41 SNP markers, all
of these stemming from previous candidate gene studies, were genotyped using TaqMan® (Applied Biosystems) assays in a confirm/refute approach (Harkensee et al., 2012).

The study was capable of confirming previously reported SNP associations: IL2-330 as a risk for development of chronic GVHD, TNF1031 as a risk for severe acute GVHD, and CTLA4-CT60 as protective against acute GVHD. These findings add credibility to the previous results, and confirmed the capability of the study population to demonstrate such non-HLA polymorphisms in a consistent manner. The associated IL2 and TNF genotypes represent high-producer variants of these cytokines that have been extensively studied in the context of GVHD. Both have essential roles in inducing and maintaining GVHD (see introduction section above). CTLA4 is an important second signal for Tcell activation, and the findings of these studies are in keeping with previous results.

## Pooled DNA screening and confirmation with MS markers

The identified genomic loci associated with a MS marker represented an area of LD within which presumably a genetic variant that had a causative role in grade 2-4 acute GVHD is located. Further work, using a variety of methods that include higher density mapping with MS or SNP, or sequencing, would be required to determine the true LD and detect such genetic variants. The following description of roles and functions of genes within the LD ranges of the associated markers is therefore assumptive; the causative genetic variants still remained to be found.

The findings from this work underline the notion that modulation of the antigen recognition and subsequent T-cell activation by non-HLA genes may occur during any of the three stages in the pathophysiology of GVHD. The microsatellite loci identified by this study put new pathways onto the map of GVHD pathobiology. Presuming that LD is strongest close to the MS location, genes that are in close range or are an isolated gene within the LD range have a higher likelihood to be a player in GVHD.

In this study, the MAPK14 (key player of proinflammatory response within the TLR pathway) and ELTD1 (a membrane protein closely involved in leukocyte adhesion and migration through endothelia) loci not only clearly have important roles in immune function, but also strong associations withstanding Bonferroni' correction. AGPAT4, another isolated gene within the MS markers LD, has an unknown function but SNP polymorphisms had been associated with GVHD severity before (Turpeinen et al., 2009). Strikingly, that study found a donor risk associated with a genetic variant, as did the microsatellite in this study. The associated SNP in the study by Turpeinen et.al. is polymorphic in the Finnish as well as Caucasian population, however, not in the Japanese. AGPAT4 is a transcriptional variant of the AGPAT1 locus which is situated in the MHC Class III region on chromosome 6, and is expressed uniformly in most tissues. AGPAT4 is highly polymorphic, and the linkage it may have to other MHC genes has not been studied sufficiently. However, ectopic expression of AGPAT in cytokine responsive cell lines may lead to enhanced expression of TNF when stimulated with IL1 $\beta$, suggesting the gene could have a role in immunoregulation (Leung, 2001). The finding that the AGPAT4 locus shows association in a consistent pattern across two genetically very diverse population makes this gene a very strong candidate for further exploration and, hopefully, clinical application.

Some loci have more than one gene within the LD range. Target gene F2RL1 is a clotting factor with well known immunomodulatory properties, like regulation of chronic inflammation in blood vessels, leukocyte rolling, adhesion and extravasation, and activation of T cells and neutrophils. The other gene at this locus, S100Z, is a known risk gene for inflammatory bowel disease. SOCS3 is attributed with an important negative feedback loop of cytokine secretion, which is a potent driver of GVHD.

Other loci implicate apoptosis pathways (DDX42) or broader metabolic and regulatory pathways (e.g. SHROOM2 - epithelial generation and regeneration, TBL1X - NFkB recruitment for gene transcription, PSMC5 - proteasome function, SMARCD2 - transcriptional activator).

The loci on the X-chromosome (IL1RAPL2 and TBL1X) are of particular interest, as they could represent potential minor histocompatibility antigens (mHag). In fact, the MS DXS0151i showed a protective effect for non-female to male HSCT ( $p=0.03$ ) with regards to acute GVHD grade 24. The prevalence of rejection was too small $(n=20)$ to yield robust results, hence further study is required to clarify the role of this marker.

## How should the results of these studies be followed up? <br> How can the findings from these studies be applied in clinical practice?

Exploratory study (further details see supplementary file 5.1):

- A confirmatory study on a larger cohort with more clearly defined HLA matching and clinical subgroups (this study, with participation of the author, of SNP markers associated with HSCT outcomes from previous studies in Japan, is currently under way).
- A full typing of all SNP in both screening and confirmation cohort would be desirable to understand the issue of reproducibility better (there are no plans to conduct this at present).
- Functional data: the possibility of correlating gene expression (e.g. of IL2 and TNF) with clinical phenotypes should be explored, prospective observation of recipients with genotype, expression profiles and clinical phenotype
- If this would show a difference in GVHD outcome between cases and controls (i.e. confirm that these genotypes would indeed predict risk), an interventional clinical trial could be undertaken.

MS-based pooled DNA scanning study:

- Confirmation of all MS markers in a separate cohort would be desirable
- Associated MS loci should be explored further in order to identify underlying causative genetic variants. This requires fine mapping with SNP and MS markers in the first instance to limit the size of the susceptibility area (practically, mapping the haplotype block of the MS marker). Sequencing can then be applied to identify functional gene variations. (JMDP and Tokai University have agreed to perform this investigation for the MAPK14 locus lead by the author using newly designed MS markers, tagSNP and next generation sequencing).
- All identified causative genetic variants should be confirmed independently, functional data gained and prospective clinical application sought, as described for the exploratory study.
- The microsatellite locus in the MAPK14 gene could be a treatment target: This gene is known to promote inflammatory responses, and specific MAPK14 inhibitors have been developed and trialled in conditions like rheumatoid arthritis. The effects were short lasting with no overall benefit after a 6 month treatment period. The reasons for the short duration of the effect and how to overcome this are not currently known, nevertheless, MAPK14 inhibitors potentially could have a future role in prevention or treatment of acute GVHD.
- It would be conceivable that the information of genetic risk/protective loci gained by this study could be used to intensify or relax GVHD prophylaxis regimen, based on a predictive score. This information could also serve in the selection of donors, should there be a choice available and should GVHD be a particular risk to be avoided. It would be essential to test such hypotheses in a prospective manner on a contemporaneous HSCT population.
- The identified MS loci suggest many pathways not implicated with GVHD before - there is a wide scope for further genetic and functional studies.
- Further study of the X-chromosome loci - could these associations represent mHags?

SUMMARY - Discussion of results

- Demonstrating the existence of non-HLA MS susceptibility regions for acute GVHD risk confirms the hypothesis of this study
- These susceptibility regions contain genes that implicate several new pathways with the pathophysiology of acute GVHD
- All findings should be confirmed in a further independent cohort
- Further exploration of MS loci include fine mapping or sequencing
- Prospective clinical evaluation of risk genotype, gene expression profiling and clinical phenotype is required before application in clinical practice
- The TNF-1031 and IL2-330 SNP could be applicable for prospective clinical observation
- The MAPK14 locus brings potentially a clinical application. The pathophysiological role in inflammatory diseases is reasonably well understood, a treatment exists and has been used in a clinical trial. MAPK14/p38 inhibitors could have a future role in GVHD prophylaxis and treatment.


### 5.2.3. Implication of the results for study design and methodology

In the SNP-based pilot study and the MS-based immunogenome screening alike, results pointed to themes that had so far not been well addressed in previous studies:

- The effect size of associated markers remained small
- Reproducibility of previous associations was low
- Construction of appropriate study cohorts remained a challenge

The objectives of the design of these two studies were to apply stringent methodology that would eliminate spurious results, but also allow a high degree of sensitivity in order to pick up low effect size associations. These studies were two of few who considered control of confounding variables by actively selecting cases to provide more homogeneity.
The screening step of the exploration study identified several SNP genotypes associating with HSCT outcomes, some of these confirming previous reports, with appropriate significance level and effect size. It was surprising that only a very small number of these replicated independently in the confirmatory cohort. The observation was that associations disappeared or even reversed.
These findings were mirrored in the MS-based study. This study included all genes that were reported as being associated with acute GVHD in the literature. Forty MS markers linking to 25 previously associated gene loci were positive in the first screening (table 5.5). None of these, with the exception of the TNF locus which showed a trend, was replicated.

## What could have been the reasons for this lack of replication?

- Despite the effort of reducing confounders, such as demographic (recipient age), clinical (diagnosis, HSCT source, conditioning) and genetic (homogeneous ethnic background, subgroup analysis of HLA matching) ones, this study population had still known confounders
which were significant in multivariate analysis (donor age, HLA matching). There were probably unknown confounders which may not have been captured in the dataset.
- The adjustment of the degree of HLA matching of the confirmatory cohort to that of the screening cohort. As the HSCT's of the earlier time frame (1993-2000) were more mismatched due to lack of better donors, the confirmation cohort represented the same degree of mismatching, although the overall HSCT population in Japan already experienced better HLA matching. As a result, prevalence of GVHD in the confirmatory cohort was slightly higher than in the overall HSCT population in Japan from the same time frame.
- Allocating HSCT into two distinct time frames (1993-2000 versus 20012005). This was introduced as an additional measure to make consistency more robust, by making associations independent of changes that occurred over the 13 years of development of HSCT. To some degree it could be expected that this measure reduced unknown confounding variables, but it was likely to have reduced the power of confirming associations from the screening cohort. This means that some of the positive associations in the screening step of both studies may well not have been spurious, but not confirmed because of competing confounders.
- Statistical power of these studies was limited - if the effect size in the confirmatory cohort was below the level of adequate statistical power, it may have escaped detection.

Most of the previous studies in the field of non-HLA genetics tried to control genetic confounding by selecting HSCT pairs that were either HLA matched related or unrelated HSCT. The thinking behind this strategy was that HLA mismatching had strong effects on HSCT outcome while non-HLA genetic variants had small effects. In order to show these small effects, HLA mismatching would have to be reduced, otherwise smaller effect size associations could not be detected because these would not be visible in the 'noise' of genetic confounding.

These studies have observed that indeed some non-HLA polymorphisms showed effects mainly or only in the HLA-matched subgroup (e.g. IL2330 and CTLA4-CT60 SNP in the exploratory study; F2RL1, AGPAT4 loci in the MS-based study). However, it was remarkable that some effects were clearly stronger in the HLA mismatched subgroups, with weak or absent effects in the HLA matched subgroups (e.g. TNF-1031 SNP in the exploratory study, ELTD1, SOCS3, DDX42 and AKT3 loci in the MS-based study). As the statistical power was only moderate, future confirmatory studies are certainly required to confirm this finding.
If confirmed, the finding that different degrees of HLA matching or mismatching involves different genetic risk loci, could expand the current knowledge of the pathophysiology of GVHD. Although the T-cell mediated alloreaction remained in the centre of the pathophysiological process, modulating mechanisms that were responsible for the severity of GVHD could be very different depending on the degree of HLA matching. It could be postulated that GVHD in fact involves very diverse pathomechanisms according to the HSCT setting (e.g. related/unrelated donor, cord HSCT, presence of minor histocompatibility antigens, preceding tissue damage through infection or chemotherapy, etc.), and therefore would require diverse strategies of prevention and treatment.

The literature review undertaken for these studies showed that small effect sizes and poor reproducibility are notorious for non-HLA polymorphisms associating with HSCT outcome. This study tried to overcome these problems by using stricter criteria for cohort inclusion and independent confirmation, thus improving the study quality. Despite this, low effect sizes and poor reproducibility persisted.

## What are the methodological lessons learned from this study? What should future genetic association studies in the field of HSCT consider?

- Studies should be adequately powered: Most associations are expected to be of low effect (i.e. OR <2), therefore the size of the cohorts should be able to provide statistical power within this range.
- An independent confirmatory cohort from the same ethnic and clinical background would be essential.
- Construction of cohorts with careful exploration and measures of control for demographic, clinical and confounders. Such confounders include: Age of donor and recipient, sex of donor and recipient (demographic), type and source of transplant, conditioning regimen, GVHD prophylaxis, etc (clinical), ethnic background, HLA matching, KIR, mHag (genetic). The expanding registries and application of HSCT may soon be able to provide adequate numbers of subjects for such approaches.
- Expanding the scope of investigated populations - 'from the bench to the bedside'. The research question should not just focus on understanding the mechanisms of GVHD, but translate into clinical practice (i.e. studying small subgroups like HLA-matched HSCT may have helped clarifying some of the non-HLA genetic risk, but was of little relevance in clinical practice because the results refer to a small and diverse population). Selecting cohorts that share clinical characteristics like diagnosis, type of HSCT etc. makes results more relevant to clinical practice and facilitates translation.

| Genetic loci previously associated - positive in $1^{\text {st }}$ screening |  | Within 'top 100 ' associations by p value |
| :---: | :---: | :---: |
| ABCB1 | IL6 | IFNG |
| CCL2 | IL7R |  |
| CCR5 | ITGA4 |  |
| CTLA4 | KIR |  |
| CXCR3 | MTHFR |  |
| ESR | PTPN22 |  |
| FCGR3 | TGFB |  |
| GSST1 | TGFBR |  |
| HSPA1L | TLR3 |  |
| IFNG | TLR4 |  |
| IL1 | TNF |  |
| IL1RN | TNFRSF1B |  |
| IL2 |  |  |

Table 5.5: Gene loci associated with GVHD in previous studies showing an association in the $1^{\text {st }}$ screening step. None of these were replicated in the $2^{\text {nd }}$ screening step, except TNF which showed a trend.

SUMMARY - methodological implications for future studies

- Although study power and cohort selection had an impact on results, lack of reproducibility and small effect sizes of associated genotypes is a common theme in non-HLA gene association studies
- Associations differ between HLA-matched and HLAmismatched subgroups, indicating that non-HLA gene polymorphisms may have variable effects in different HSCT settings
- Future genetic association studies need to consider adequate statistical power, stringent study designs and careful cohort construction.
- The perspective, aim and objectives of future studies should also be on clinical questions


### 5.3. Future

Despite the plethora of data from genome-wide association studies, very few of these, SNP or MS based alike, have actually been able to identify causative genetic variants. The epigenetic function of the genome just has begun of being understood, projects such as ENCODE provide a deeper insight on how genes are regulated (Consortium, 2007). According to this research, gene function could be regulated by structural (mutations, polymorphisms, etc) or functional (epigenetic) elements, the latter ones may not necessarily comply with the limitations of haplotype blocks or LD. For example, remote regulatory elements may cause an association signal, but the regulated gene could be far away outside the LD range (Consortium, 2007).

The scope of genetic disease association research is rapidly expanding. Genome-wide, high-throughput approaches with SNP or MS markers have rapidly advanced knowledge about the influence of genetic polymorphisms on health and disease, and have enhanced understanding of underlying pathomechanisms.

Despite these advances, common genetic variables only explain a very small fraction of the total genetic risk (2-3\%), the missing proportion has been referred to as the 'dark matter' of genetic risk (Maher, 2008). SNPbased genome-wide association studies of common genetic variants may not be best suited to explore missing genetic risk due to the high threshold of significance, which makes small effect size association escape detection. The MS approach using a multi-step confirmation design without application of multiple testing correction statistics at this stage, maybe somewhat more sensitive, but no direct comparison studies have been performed. Apart from small effect size, there are several reasons for this lack of detecting larger proportions of genetic risk (Maher, 2008):

- Associated marker in LD with causative variant - loss of effect size
- Rare alleles that may not be captured by scanning for common variants
- Copy Number Variations (CNV) - which are not picked up well by current genome-wide approaches (Stefansson et al., 2008, Consortium, 2008)
- Transcriptional control of a gene locus by several other genes which are not necessarily in LD (Brem et al., 2002)
- Epigenetic effects (Waterland and Jirtle, 2003)

There are several ways how more knowledge about the 'dark matter' could be established. Fine mapping of regions within LD of associated markers, using SNP, MS or sequencing, could reveal causative variations. Most recently, sequencing of selected genomic regions or the whole genome came within reach of broad application with the advent of faster and cheaper sequencing technology (next generation sequencing). Next generation sequencing (NGS) works by high-throughput, parallel sequencing of overlapping short stretches of genome (100-250 bp), which are annotated by computational methods (Bentley et al., 2009). NGS is capable of detecting rare variants that escape detection by genome-wide studies using SNP or MS markers; and is very effective in detecting new microsatellites fur further investigation (Santana et al., 2009, Zalapa et al., 2012).
Future efforts are aiming to integrate data from large scale genomic and gene transcription or expression research (Hansen, 2008). A few microarray studies have already been undertaken measuring protein signatures of GVHD in urine (Weissinger and Dickinson, 2009, Kaiser et al., 2004) or gene transcription in blood (Buzzeo et al., 2008, Krijanovski et al., 2007, Paczesny et al., 2009b). These studies, especially if capable of discriminating profiles in different HSCT settings, could greatly contribute to the understanding of GVHD pathophysiology.

The future of genetic research into the causes of GVHD is likely see new approaches:

- larger scale marker-based genomic studies, using SNP or MS markers
- integration of genetic and functional data, dissection of GVHD pathophysiology
- Whole genome sequencing approaches is likely to identify further, and rarer genetic variants associating with GVHD, but have not yet been conducted.


### 5.4. Conclusions

This study has demonstrated that a MS-based, pooled DNA scanning methodology, derived from a genome-wide scanning approach and for the first time applied in an HSCT setting, was capable of identifying nonHLA genetic risk loci for the development of moderate-severe acute GVHD.

The expected low effect size of associations suggested an approach that was robust, powerful and sensitive. This study did show that a microsatellite-based approach had some inherent advantages (i.e. more informative markers, a study design of a multi-step screening) over similar SNP-based approaches, but also some disadvantages (e.g. the requirement for DNA pools, lack of high-throughput platforms).

With regards to translating the study design, overall this study went much further to control confounding variables than previous studies, but some confounders remained. Due to the nature of rapidly evolving progress in this field, robust cohort design is difficult. The choice of genetic marker type, study design with independent confirmation, and selection of a population from an ethnically homogeneous background with attempt of controlling confounders complied with well laid out requirements for a high-quality genetic association study. A larger number of subjects, providing the study with a stronger statistical power, could potentially have led to a larger number of susceptibility loci identified. From a technical perspective, extensive quality controls had ensured adequate quality of pooled DNA and interpretation of typing signals. False positive markers had effectively been eliminated, despite a deliberately 'sensitive' approach by using technical resolution of DNA pools as a threshold for inclusion, and non-application of multiple testing correction whilst building on confirmation in an independent cohort to rule out false positives.

This approach confirmed three previous SNP associations (IL2-330, TNF-1031 and CTLA4-CT60) in an exploration study, and ten new target gene microsatellite loci (F2RL1, MAPK14, ELTD1, IL1RAPL2 (x2),

SOCS3, TBL1X, DDX42, AGPAT4 and AKT3) in MS-based pooled DNA approach. All of these loci should be confirmed in a further independent cohort. Some of these loci, e.g. the SNP from the exploration study which have known high-expression genotypes, are close to potential prospective observation and application in clinical practice as predictors of risk. The MAPK14 locus was confirmed in this study by SNP typing, was already one of the better understood with regards to pathophysiology (it was involved in pro-inflammatory responses). A specific treatment (p38 inhibitor) already exists that has been trialled in humans. Hence, the way to clinical application could be promising. An associated MS in the AGPAT4 locus has confirmed the finding of association in this locus in a genetically diverse population, rendering this locus a strong candidate for further exploration.

Some observations in these studies raised new questions and hypotheses. This study demonstrated that small effect size associations with HSCT outcome did exist and could be consistent, but most associations from the screening step were not reproducible, even with this more stringent study design. Heterogeneity of confounders, hence cohort construction, was a likely cause for the lack of reproducibility. Future studies should consider more the issues of statistical power, study and cohort design.

Another important observation was the discrepancy of effect markers had in HLA matched or mismatched subgroups. With the focus mainly on HLA-matched study cohorts, previous studies may have missed associations that are predominant in HLA-mismatched subgroups only. It may be possible that the pathogenesis of GVHD involves different genes in different degrees of HLA matching - a hypothesis that would have to be proven by future studies.
The availability of ever larger HSCT registries for research is likely to facilitate larger scale investigations that are likely to overcome the methodological problems of previous studies (i.e. statistical power), including this one. The future will probably see larger scale genomic
approaches (e.g. SNP, MS or sequencing based) and integration with gene expression, elucidating the pathophysiology of GVHD and identifying new targets for clinical application.

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## APPENDICES

Appendix 2.1: Characteristics of the study population
Appendix 2.2: Procedure for custom-design of MS markers
Appendix 2.3: Estimation of DNA requirements for the pooled screening steps
Appendix 2.4: Calibration of pipettes for DNA pooling
Appendix 2.5: PCR mixtures for test typing
Appendix 3.1: Information of SNP assays used in the exploration study
Appendix 4.1: Genotyping errors and their resolution

## Appendix 2.1

## Characteristics of the study population

|  | Screening | Confirmation | Difference $p=$ |
| :---: | :---: | :---: | :---: |
| Demographic |  |  |  |
| Recipients | 460 | 462 |  |
| Recipient gender Male | 269 (58.48\%) | 289 (62.55\%) | $\mathrm{n} / \mathrm{s}$ |
| Recipient gender Female | 191 (41.52\%) | 173 (37.45\%) | $\mathrm{n} / \mathrm{s}$ |
| Donor gender Male | 267 (58.04\%) | 278 (60.17\%) | $\mathrm{n} / \mathrm{s}$ |
| Donor gender Female | 193 (41.96\%) | 182 (39.39\%) | $\mathrm{n} / \mathrm{s}$ |
| Female donor to Male recipient transplant | 102 (22.17\%) | 84 (18.18\%) | $\mathrm{n} / \mathrm{s}$ |
| Recipient age range | 4-40 y | $4-40 \mathrm{y}$ | $\mathrm{n} / \mathrm{s}$ |
| Recipient age mean | 21.7 y | 24.1 y | $\mathrm{n} / \mathrm{s}$ |
| Donor age range | 20-70y | $19-51 \mathrm{y}$ | $\mathrm{n} / \mathrm{s}$ |
| Donor age mean | 34 y | 34.3 y | $\mathrm{n} / \mathrm{s}$ |
| Clinical |  |  |  |
| Diagnosis Acute lymphoblastic leukaemia | 260 (56.52\%) | 254 (54.98\%) | $\mathrm{n} / \mathrm{s}$ |
| Diagnosis Acute non-ALL | 200 (43.48\%) | 208 (45.02\%) | $\mathrm{n} / \mathrm{s}$ |
| High risk leukaemia | 279 (60.65\%) | 246 (53.25\%) | <0.1 |
| HLA matching - 12/12 loci | 41 (8.91\%) | 37 (8\%) | $\mathrm{n} / \mathrm{s}$ |
| HLA matching - 12/12 and 10/10 loci | 160 (34.78\%) | 166 (35.93\%) | $\mathrm{n} / \mathrm{s}$ |
| HLA matching - GVHD risk mismatches | 220 (47.83\%) | 229 (49.57\%) | $\mathrm{n} / \mathrm{s}$ |
| Conditioning - Cyclophosphamide + total body irradiation (TBI) | 334 (72.61\%) | 322 (69.67\%) | $\mathrm{n} / \mathrm{s}$ |
| Conditioning - Busulphan/Cyclophosphamide or Busulphan based | 53 (11.52\%) | 47 (10.17\%) | $\mathrm{n} / \mathrm{s}$ |
| Conditioning - TBI based | 73 (15.87\%) | 83 (17.97\%) | $\mathrm{n} / \mathrm{s}$ |
| Conditioning - other | 0 | 10 (2.16\%) | $\mathrm{n} / \mathrm{s}$ |
| GVHD prophylaxis - Cyclosporin A based | 279 (60.65\%) | 154 (33.33\%) | <0.05 |
| GVHD prophylaxis - Tacrolimus based | 177 (34.48\%) | 305 (66.02\%) | <0.05 |
| GVHD prophylaxis - other | 4 (0.87\%) | 3 (0.65\%) | $\mathrm{n} / \mathrm{s}$ |
| Outcome |  |  |  |
| Acute GVHD grade 0 | 124 (26.96\%) | 124 (26.84\%) | $\mathrm{n} / \mathrm{s}$ |
| Acute GVHD grade 1 | 153 (33.26\%) | 99 (21.42\%) | <0.05 |
| Acute GVHD grade 2 | 105 (22.83\%) | 143 (30.95\%) | <0.05 |
| Acute GVHD grade 3 | 50 (10.87\%) | 72 (15.58\%) | <0.05 |
| Acute GVHD grade 4 | 28 (6.09\%) | 24 (5.19\%) | $\mathrm{n} / \mathrm{s}$ |
| Chronic GVHD - none | 244 (53.04\%) | 242 (52.38\%) | $\mathrm{n} / \mathrm{s}$ |
| Chronic GVHD - limited disease | 71 (15.43\%) | 63 (13.64\%) | $\mathrm{n} / \mathrm{s}$ |
| Chronic GVHD - extensive disease | 95 (20.65\%) | 106 (22.94\%) | $\mathrm{n} / \mathrm{s}$ |
| Chronic GVHD - unknown | 50 (10.86\%) | 49 (10.6\%) | $\mathrm{n} / \mathrm{s}$ |
| Relapse | 115 (25\%) | 110 (23.81\%) | $\mathrm{n} / \mathrm{s}$ |
| Survival - 100 days | 395 (86.9\%) | 403 (87.23\%) | $\mathrm{n} / \mathrm{s}$ |
| Survival - 1 year | 306 (66.52\%) | 312 (67.53\%) | $\mathrm{n} / \mathrm{s}$ |
| Survival - 3 years | 245 (53.26\%) | 258 (55.84\%) | $\mathrm{n} / \mathrm{s}$ |

Appendix 2.1: Population characteristics. $p$ refers to statistically significant differences between the screening and confirmation cohorts.

## Appendix 2.2:

## Procedure for custom-design of MS markers

## 1. Procedure for custom-design of MS markers

Some target genes were located in gene regions that did not have adequate cover with MS markers from the genome-wide panel. Therefore, for 59 genes MS markers had to be custom-designed following the procedure below:

- Identification of target gene genomic location on the NCBI or GeneCard databases
- Retrieving the genomic sequence, plus an additional sequence of 50 kb to both sides of the target gene, from the UCSC genome browser website (http://genome.ucsc.edu/), using the 'Gene Sorter' function
- Importing the retrieved sequence into the 'Blat' function of the same website, checking for multiple sequence locations, applying settings to identify SNP and possible amplicons
- Importing the same sequence into Sputnik (http://cbi.labri.ubordeaux.fr/outils/Pise/sputnik.html) to identify microsatellite repeats within the obtained $\sim 100 \mathrm{~kb}$ sequence.
- Selection of an appropriate candidate microsatellite, criteria: as many repeats as possible, preferably 3-4 base repeats, uninterrupted repeat sequence.
- Importing the same sequence into geneview, location of the identified microsatellite, cut and paste the microsatellite with 1000 base pairs flanking on each side.
- Import of obtained sequence into primer express, primer search using the settings: Melting temperature (Tm) $56-58^{\circ} \mathrm{C}$, amplicon length max 400 base pairs.
- If suitable primer pairs not found: Trying of a different microsatellite from the Sputnik output
- If suitable primer pair found: Checking of primer pair with the 'In silico PCR' function on the UCSC website to ensure uniqueness of primer pair for the intended target.
- For $n=8$ loci it was not possible to design appropriate primers. Reasons were either a complete absence of suitable microsatellites within the target regions, or extensive duplication of highly similar sequence (e.g. within the FCGR gene cluster) disabling identification of a unique sequence.


## Appendix 2.3:

## Estimation of DNA requirements for the pooled screening steps

## 1. Estimation of DNA amounts required for single genotyping

The essential DNA requirements for conducting pooled DNA genotyping are as outlined in table 1.

| Screen | Cohort | No. <br> markers | Amount <br> for <br> $\mathbf{c o n c} \mathbf{6}$ <br> $\mathbf{n g} / \boldsymbol{\mu l}$ | Amount <br> for <br> $\mathbf{c o n c} \mathbf{8}$ <br> $\mathbf{n g} / \boldsymbol{\mu}$ | No. <br> samples | Total <br> DNA at 6 <br> $\mathbf{n g} / \boldsymbol{\mu l}$ | Total <br> DNA at 8 <br> $\mathbf{n g} / \boldsymbol{\mu l}$ |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 1 | Pat 0-1 | 4000 | 48 | 64 | 281 | 683.27 | 911.03 |
| 1 | Pat 2-4 | 4000 | 48 | 64 | 195 | 984.61 | 1312.82 |
| 1 | Don 0-1 | 4000 | 48 | 64 | 281 | 683.27 | 911.03 |
| 1 | Don 2-4 | 4000 | 48 | 64 | 195 | 984.61 | 1312.82 |
| 2 | Pat 0-1 | 500 | 48 | 64 | 128 | 187.50 | 250.00 |
| 2 | Pat 2-4 | 500 | 48 | 64 | 111 | 216.21 | 288.28 |
| 2 | Don 0-1 | 500 | 48 | 64 | 128 | 187.50 | 250.00 |
| 2 | Don 2-4 | 500 | 48 | 64 | 111 | 216.21 | 288.28 |

Table 1: Basic DNA requirement estimates

## 2. Additional DNA amounts required for processing, further typing

To calculate the total amount required for the first screening step, an extra one third for possibly necessary repeat experiments was added, and, depending on sample concentration, an extra 100-500 ng of DNA required for the pooling process (repeated concentration measurements). Therefore final requirements for the first and second pooled screenings were as outlined in table 2.

| Screen | Cohort | High <br> conc | Low conc |
| :---: | :---: | :---: | :---: |
| 1 | Pat 0-1 | 1714 | 1314 |
| 1 | Pat 2-4 | 2249 | 1849 |
| 1 | Don 0-1 | 1714 | 1314 |
| 1 | Don 2-4 | 2249 | 1849 |
| 2 | Pat 0-1 | 833 | 433 |
| 2 | Pat 2-4 | 884 | 484 |
| 2 | Don 0-1 | 833 | 433 |
| 2 | Don 2-4 | 884 | 484 |

## Table 2: Corrected DNA requirements including allocations for pipetting/pooling, in ng amount.

Following the first and second pooled screening steps, around 100 markers were expected to show a significant association. These markers would then be individually genotyped on the entire cohort. As the amount of DNA required for one MS marker typing was 1-2 ng/sample, this would require a further 200 ng of DNA, plus a $15 \%$ margin for repeat experiments.

Only the markers which showed a consistently significant association after the two pooled screenings and the individual genotyping confirmation step would be subjected to SNP 'scanning' of the 100 kb region, with approximately 50 SNP. Using a TaqMan® assay, this required 1-2 ng of DNA per SNP studied (hence, 100 ng ), plus a margin of $15 \%$ - approximately 120 ng of DNA.

Separately, the plan was to study all SNP associated with HSCT outcomes by individual genotyping using a TaqMan® assay. These were approximately 150 SNP, including a margin of $15 \%$ requiring 350 ng of DNA.

After summing up all these requirements, a further $15 \%$ of DNA amount was added for pipetting and measurement variability.

In summary, maximum DNA requirements for the entire study were as in table 3.

| Usage | Entry $1^{\text {st }}$ Screen, ng | Entry $2^{\text {nd }}$ Screen, ng |
| :---: | :---: | :---: |
| First Screen | 2250 | N/A |
| Second Screen | N/A | 890 |
| Individ MS typing | 230 | 230 |
| SNP screen | 120 | 120 |
| SNP previous | 350 | 350 |
| Intermed sum | 2950 | 1590 |
| Add for pipet/measure | 450 | 260 |
| Total | 3400 | 1850 |

Table 3: Estimated total DNA amounts required for entire study

## Appendix 2.4: <br> Calibration of pipettes for DNA pooling

The process of pooling requires a very high degree of accuracy. Therefore for all pipetting of pools only calibrated pipettes with a fixed volume are used (i.e. the pipette is dedicated to this process for the duration of the study, the volume on the pipette is not changed). For calibration, a set of at least 3 pipettes is tested, and the one is chosen that has the lowest volume variation (which has to be $<1 \%$ ). Only original pipette tips of the maker are used. The aim was to have fixed pipette volumes for all standard procedure volumes. Volumes of individual samples had to be pipetted with multiple fixed pipettes if necessary. For all pooling into intermediate or large pools fixed volume pipettes with the exact volume required were 'customcalibrated'.

## Procedure of pipette calibration

## Preparation

- A set of five 2 ml plastic flipcap tubes is prepared for each pipette
- Each tube is weighed three times on a high sensitivity scale (nanograms, Mettler® Toledo) at standard conditions (stable lab table and temperature)

Practical procedure

- Each of the five tubes is pipetted with the target volume by a standard technique:
- Attachment of the tip by a single, firm but gentle movement in vertical direction
- Gentle, slow aspiration with 1-2 seconds wait after completion of aspiration
- Keeping pipette vertical at all times of aspiration, transfer, dispensing
- Slow dispensing of sample into tube, wait for 1-2 seconds with tip inside fluid before removing
- Each filled tube is then weighed three times
- Calculation of mean, standard deviation, standard error and variance on an excel template spreadsheet.

| Pipette <br> volume <br> range | Nominal <br> volume | Pipetted <br> volume | SD | Variance | Upper <br> error \% | Lower <br> error \% |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| $2-20$ | 20 | 19.7 | 0.00006 | 0.00000003 | 0.4 | -0.44 |
| $10-100$ | 100 | 98.9 | 0.0001 | 0.00000002 | 0.12 | -0.24 |
| $100-1000$ | 380 | 378 | 0.0009 | 0.0000009 | 0.35 | -0.22 |
| $20-200$ | 95 | 94.5 | 0.0001 | 0.00000003 | 0.24 | -0.28 |
| $20-200$ | 87 | 86.8 | 0.00007 | 0.000000005 | 0.09 | -0.06 |

Table 1: Examples of pipette tests. The pipette that delivered the volume closest to the aimed volume was chosen, if the error was within a range of + - $1 \%$.

## Appendix 2.5:

## PCR mixtures for test typing

High amount DNA

| DNA <br> $\mathrm{amt} /$ well <br> ng | 93.18677 |
| :--- | ---: |
| Marker <br> typing <br> capacity | 6000 |

Medium amount DNA

| DNA <br> $\mathrm{amt} /$ well <br> ng | 62.12451 |
| :--- | ---: |
| Marker <br> typing <br> capacity | 9000 |

Low amount DNA

| DNA <br> amt/well <br> ng | 46.59338 |
| :--- | ---: |
| Marker <br> typing <br> capacity | 12,000 |


| ingredient | vol <br> $\boldsymbol{\mu l} /$ well |
| :--- | ---: |
| dH2O | 0 |
| $10 x$ uffer | 2 |
| 2.0 mM <br> dNTP | 2.5 |
| Primer mix | 2 |
| AmpliTaqG | 0.1 |
| DNA | 13.4 |
| total vol <br> ( $\boldsymbol{\mu l}$ ) | $\mathbf{2 0}$ |


| ingredient | vol <br> $\boldsymbol{\mu l} /$ well |
| :--- | ---: |
| dH2O | 4.2 |
| 10xbuffer | 2 |
| 2.0mM <br> dNTP | 2.5 |
| Primer mix | 2 |
| AmpliTaqG | 0.1 |
| DNA | 9.2 |
| total vol <br> $(\boldsymbol{\mu l})$ | $\mathbf{2 0}$ |


| ingredient | vol <br> $\boldsymbol{\mu l} /$ well |
| :--- | ---: |
| dH2O | 6.5 |
| 10xbuffer | 2 |
| 2.0 mM <br> dNTP | 2.5 |
| Primer mix | 2 |
| AmpliTaqG | 0.1 |
| DNA | 6.9 |
| total vol <br> $(\boldsymbol{\mu l})$ | $\mathbf{2 0}$ |

Table 3.11: PCR mixtures for three options of DNA amount in the PCR procedure, as applied for MS marker testing

## Appendix 3.1:

## Information of SNP assays used in the

exploration study

Primers for Luminex Genotyping of IL10 SNP

| PR0 BE | oligonuc kotide nam e | sequence ( 5 ' --> 3') |
| :---: | :---: | :---: |
| L10P comm on probe | L10CR-2 | TTTTTTTTTTTTTTTTTTTTCAGACTACTCTTACCCA |
| -1082A probe | 1082AS-4 | TTTTTTTTTTTTTTTTTTTTCTGTTCCCCTTCCCAAAGA |
| -1082G probe | 1082GS-3 | TTTTTTTTTTTTTTTTTTTTTTCCCCTCCCAAAG |
| -819C probe | 819CR-2 | TTTTTTTTTTTTTTTTTTTTAGGTGATGTAACATCTCTGTGC |
| -819T probe | 819TS-2 | TTTTTTTTTTTTTTTTTTTTGCACAGAGATATTACATCACCT |
| -592A probe | 592AR-8 | TTTTTTTTTTTTTTTTTTTTCCGCCTGTACTGTAGG |
| -592C probe | 592CR-2 | TTTTTTTTTTTTTTTTTTTTCGCCTGTCCTGTAGGAA |


| PRMER | oligonuc kotide nam e | sequence $\left(5^{\prime}-->3^{\prime}\right)$ |
| :--- | :--- | :--- |
| $\mathbb{L} 10$ P fonw ard prim er | $\mathbb{L} 10-$ F2 | CAAATCCAAGACAACACTACTAAGGC |
| $\mathbb{L} 10$ P reverse prim er | $\mathbb{L} 10-$ R2 | GGCTAAATATCCTCAAAGTTCCCAAG |

## TaqMan Assays

| Assay ID | Context Sequence | Design Strand | Category ID | Group ID | Gene Symbol | NCBI Gene Reference | Cytogenetic Band | SNP Type | Location on NCBI Assembly |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| C__15944115_20 | TATCTAGCTATATGATTGT GAGTTA[A/G]CTTCTTAAAT CTTCTATGACTCAGT | Forward | Chr12 | D12S313 | IFNG |  | 12q15a | INTERGENIC/UNK NOWN | 66841278 |
| C__15820717_10 | GTCAGCCTGTGGGGTAAC TTGGTCC[A/G]TGGGATTT CCCCTAAAAAGGTAGCC | Forward | Chr16 | D16S3136 | NOD2 | NM_022162.1 | 16q12.1c | INTRON | 49291470 |
| C__ 7514871_10 | GGAAGCAAAGGAGAAGCT GAGAAGA[C/T]GAAGGAAA AGTCAGGGTCTGGAGGG | Reverse | Chr6 | D6S276 | TNF;LTA;LTB |  | 6p21.33a | INTERGENIC/UNK NOWN | 31650287 |
| C___9077561_20 | AATGGAAAATCCCAGAAAT TCTCCC[A/G]TTTGGATCCC ACCTTCTCCATCCCA | Forward | Chr1 | D1S484 | FCGR2A | NM_021642.2 | 1q23.3a | MIS-SENSE MUTATION | 159746369 |
| C__25651063_10 | CATTGCATTCTTGACAGAT TCTCTT[A/G]TTGCCTTAAA AAGAATCACTGGCCT | Forward | Chr16 | D16S3136 | NOD2 | NM_022162.1 | 16q12.1c | INTRON | 49322877 |
| C___9546481_20 | GATTTTTACATATGAGCCT TCAATG[A/G]TGTTGCCTG GTTACTATTATTAAAG | Reverse | Chr2 | D2S160 | IL1A | NM_000575.3 | 2q13d | UTR 5 | 113259431 |
| C__34029672_10 | GAGCTTCTGCAAAGTGGA AGAATAC[C/T]GCTTGGCC CTAACTCCTCACCCCAA | Reverse | Chr16 | D16S3136 | NOD2 | NM_022162.1 | 16q12.1c | INTRON | 49305205 |
| C___1384440_10 | GTGCCACCCATTTATTGGG GAAAAG[C/T]CCTAAAAGG GGAAGTGGGGAAGGGA | Reverse | Chr16 | D16S3136 | NOD2 | NM_022162.1 | 16q12.1c | INTRON | 49310316 |
| C___8861232_20 | GTGGCCATCCCTGGGAAT GCAAGCA[G/T]GGATGCAG TCTGCACGTCCACGTCC | Forward | Chr1 | D1S2667 | TNFRSF1B | NM_001066.2 | 1p36.22a | MIS-SENSE MUTATION | 12175542 |
| C__16049347_10 | TTGCTGTGTTTGTAATTCA GGTAAA[C/T]CTATTTTCTG TAAAGCAGGCATGAT | Reverse | Chr8 | D8S285 | CCL4 |  | 8q12.1c | INTERGENIC/UNK NOWN | 59833385 |
| C__15859930_10 | AGTAACTCAGAAAATTTTC TTTGTC[C/A]TAAAACTACA CTGAACATGTGAATA | Forward | Chr4 | D4S402 | IL2 |  | 4q27d | INTERGENIC/UNK NOWN | 123597430 |
| C___1839943_10 | TACCTTGGGTGCTGTTCTC TGCCTC[G/A]GGAGCTCTC TGTCAATTGCAGGAGC | Forward | Chr2 | D2S160 | IL1B |  | 2q13d | INTERGENIC/UNK NOWN | 113311338 |


| C__1202883_20 | gaAAAGCTGCGTGATGAT GAAATCG[G/A]CTCCCGCA GACACCTTCTCCTTCAA | Forward | Chr1 | D1S2667 | CLCN6;MTHF <br> R;C1orf167 | NM_005957.3 | 1p36.22a | MIS-SENSE <br> MUTATION | 11778965 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| C__8708473_10 | GAGGAGGGGGCAACAGGA CACCTGA[A/G]GGATGGAA GGGTCAGGAGGCAGACA | Forward | Chr19 | D19S220 | $\begin{aligned} & \text { TMEM91;TGF } \\ & \text { B1;B9D2 } \end{aligned}$ |  | 19q13.2c | INTERGENIC/UNK NOWN | 46552136 |
| C___9578811_10 | TCATATGGTTAACTGTCCA TTCCAG[A/G]AACGTCTGT GAGCCTCTCATGTTGC | Forward | Chr10 | D10S1765 | FAS |  | 10q23.31b | INTERGENIC/UNK NOWN | 90739943 |
| C__15820716_10 | TTGCTCTTGACTCTTGGCA GGAAAC[A/G]TACAACTCTT TCTTTCTTCTTTTCT | Reverse | Chr16 | D16S3136 | NOD2 | NM_022162.1 | 16q12.1c | INTRON | 49291534 |
| C___3296043_10 | TCTTCACCACTATTTGGGA TATAAC[A/G]TGGGTTAACA CAGACATAGCAGTCC | Reverse | Chr2 | D2S117 | CTLA4 |  | 2q33.2a | INTERGENIC/UNK NOWN | 204447164 |
| C__11722137_10 | TTTTCAAACAAGAAGTAGT TTTTCA[C/T]CAAACAATGT CTCTTATGTAATTCA | Reverse | Chr9 | D9S1776 | TLR4 | NM_138554.2 | 9q33.1c | INTRON | 119512585 |
| C__15873886_20 | ATGTCAGAGACGGAGACG AGGCAAC[A/C]GGACCGTG GAGGAGAAAAATAGAAA | Forward | Chr19 | D19S220 | B9D2;TGFB1 | NM_000660.3 | 19q13.2c | INTRON | 46548726 |
| C__1882528_10 | GGAGGCGGATGCTGTGAG TTCCCAG[G/T]TCTTGGCT GTTGTCTGAGAGGGGGT | Forward | Chr10 | D10S591 | IL15RA | $\begin{aligned} & \text { NM_002189.2 } \\ & \text {,NM_172200. } \\ & 1 \end{aligned}$ | 10p15.1b | MIS-SENSE MUTATION | 6042374 |
| C__2213785_10 | TTTCTAGCCGCCAAGTGGA GAACAG[C/G]TTGGAGCGG TGCGCCGGGCTTAGCG | Forward | Chr22 | D22S539 | SLC2A11;MIF |  | 22q11.23a | INTERGENIC/UNK NOWN | 22566392 |
| C__1272348_10 | CTGTTCCTATTCAGCCATC TTGGCT[C/T]GGGACCAGA GAACTTCGTATTTCTT | Reverse | Chr1 | D1S198 | IL23R |  | 1p31.3a | INTERGENIC/UNK NOWN | 67421048 |
| C__7504226_10 | CCATATACCTGAAAGATCT GATGAA[A/G]CCCAGCGTG TTTTTAAAAGTTCGAA | Forward | Chr3 | D3S1558 | CD86 | $\begin{aligned} & \text { NM_006889.3 } \\ & \text {,NM_175862. } \\ & 3 \end{aligned}$ | 3q13.33c | MIS-SENSE MUTATION | 123321009 |
| C___3052613_1_ | GTGATGATAGGGTTACACA TCTGCT[C/T]CAATTCCTTT CTCTTATGATCAAAC | Forward | Chr6 | D6S276 | $\begin{aligned} & \text { HSPA1A;HSP } \\ & \text { A1L;LSM2 } \end{aligned}$ | NM_005527.3 | 6p21.33a | MIS-SENSE MUTATION | 31885925 |
| C__2215707_10 | GGCCCAGAAGACCCCCCT CGGAATC[A/G]GAGCAGGG AGGATGGGGAGTGTGAG | Reverse | Chr6 | D6S276 | LTB;LTA;TNF |  | 6p21.33a | INTERGENIC/UNK NOWN | 31651080 |
| C__1384434_10 | TGGCCTTTGGAAGGGGCA TTTCTGA[A/T]TAAGATCTG GGCCGCTCTCCGCTGG | Reverse | Chr16 | D16S3136 | NOD2 | NM_022162.1 | 16q12.1c | INTRON | 49317048 |
| C__3017467_10 | TGTTTTATTTAAGCCTCAC AAGGGT[A/G]TAGTGTGAC TACACTGTTTCTTAAC | Forward | Chr16 | D16S3136 | NOD2 | NM_022162.1 | 16q12.1c | INTRON | 49294157 |


| C__15873887_10 | ATTGTATGGTTTGTGTTCT TCTATC[C/T]TTCAGGGACC ATCTAGGTGGACCTT | Reverse | Chr19 | D19S220 | TGFB1;B9D2 | NM_000660.3 | 19q13.2c | INTRON | 46545926 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| C__7514879_10 | GAGGCAATAGGTTTTGAG GGGCATG[A/G]GGACGGG GTTCAGCCTCCAGGGTCC | Reverse | Chr6 | D6S276 | LTB;LTA;TNF |  | 6p21.33a | INTERGENIC/UNK NOWN | 31651010 |
| C___2404008_10 | TGGACAGGCGGTCCTGGA TGGCCTC[A/G]ATCAGCGC GGCGTCCTGCACCCCAG | Forward | Chr12 | D12S85 | $V D R$ | $\begin{aligned} & \text { NM_000376.2 } \\ & \text {,NM_0010175 } \\ & 35.1 \end{aligned}$ | 12q13.11c | $\begin{aligned} & \text { SILENT } \\ & \text { MUTATION } \end{aligned}$ | 46525024 |
| C___2415786_20 | GCACAAGGCTCAGCTGAA CCTGGCT[A/G]CCAGGACC TGGCCCTGCACTCTCCT | Reverse | Chr2 | D2S117 | CTLA4 | $\begin{aligned} & \text { NM_0010376 } \\ & 31.1, N M \_005 \\ & 214.3 \\ & \hline \end{aligned}$ | 2q33.2a | MIS-SENSE MUTATION | 204440959 |
| C__11918223_10 | GTCGAGTATGGGGACCCC CCCTTAA[C/T]GAAGACAG GGCCATGTAGAGGGCCC | Reverse | Chr6 | D6S276 | LTB;LTA;TNF |  | 6p21.33a | INTERGENIC/UNK NOWN | 31650461 |
| C___3017459_20 | ACATTTCTCTTGGCTTCCT GGTGCG[G/T]GCCAAAGGT GTCGTGCCAGGGAGTA | Forward | Chr16 | D16S3136 | NOD2 | NM_022162.1 | 16q12.1c | SILENT MUTATION | 49303084 |
| C__31784020_10 | TCTATTAAGGTAGACCACC TCTCCC[C/T]TTTTTTTTTTT TCAAACAAGAAGTA | Forward | Chr9 | D9S1776 | TLR4 | NM_138554.2 | 9q33.1c | INTRON | 119512551 |
| C___2415784_10 | CTTATCTCTCTCTAGACCT TCTTGG[C/T]TAAGAAACCA TGTAGTTTGTATGAA | Forward | Chr2 | D2S117 | CTLA4 | $\begin{aligned} & \hline \text { NM_0010376 } \\ & 31.1, N M \_005 \\ & 214.3 \\ & \hline \end{aligned}$ | 2q33.2a | INTRON | 204441833 |
| C__30031638_10 | CCTGAATTCTCAGTAACTT AGAAGT[A/C]ATTTCTAATG ATTCCGGCTGGGCAC | Forward | Chr19 | D19S220 | TGFB1;B9D2 | NM_000660.3 | 19q13.2c | INTRON | 46543349 |
| RS1264457-GA |  | Forward |  |  | HLA-E |  |  |  |  |
| RS1800795-GC |  | Reverse |  |  | HLA-E |  |  |  |  |

## Appendix 4.1:

## Genotyping errors and their resolution

## 1. $1^{\text {st }}$ screening $P C R$ and genotyping - technical results (all pools combined)

The initial round of PCR and genotyping had an error rate of $11.36 \%$. This was expected as there is some variation in PCR conditions of markers on the panel for the genome-wide association studies. In that set-up, these variations were addressed by grouping markers requiring the same conditions on the same plate. As the selection of markers for this study disrupted this order, markers with different typing conditions were on the same plate. The main causes of error were absence of peak signal (4.34\%) and low peak signal (5.08\%), indicating either PCR failure or high dilution/low concentration of PCR product. High peak signal, size standard errors, data collection failure and mechanical errors of PCR and genotyping accounted for the remaining $2.06 \%$ of errors.

### 1.1. Strategy for resolving error samples ( $1^{\text {st }}$ step):

- Samples with an absent peak signal were re-typed using a x10 dilution.
- Samples with a low peak signal were re-typed using a x20 dilution.
- Samples with size standard errors, data collection failure or mechanical errors were re-typed using a $\times 40$ dilution.
- Samples with high peak signals were subjected to a dilution sequence of $x 80, x 200, x 400$ and $x 800$

Results of the $1^{\text {st }}$ step of error sample resolution:
All of the high peak/size standard/data collection/mechanical error samples had adequate results. Most of the low peak signal and some of the absent
peak signal sample errors were resolved, but $5.20 \%$ of samples remained with absent/low peak signal error. We presumed that the reason is primary PCR failure.

### 1.2. Strategy for resolving error samples ( $2^{\text {nd }}$ step):

- Repeat PCR of all samples with absent/low peak signal
- DNA purification of PCR product in persisting low peak signal samples

Results of the $2^{\text {nd }}$ step of error sample resolution:
A further $2.53 \%$ of samples were resolved, remaining $2.67 \%$ with persistently absent/low peak signal. Possible reasons include degradation of primers (primers for studies in this institution stem from a master primer set on plates which are defrosted and re-frozen each time a primer aliquot is taken) or inadequate PCR conditions.

### 1.3. Strategy for resolving error samples ( $3^{\text {rd }}$ step):

- Re-PCR with primers from a separate primer stock
- If error persists: purchase of fresh primer, modification of PCR conditions (extension to 35 cycles, increasing concentration of dNTP, change of annealing temperature), re-typing after PCR product purification

Results of the $3^{\text {rd }}$ step of error sample resolution:
Only $0.8 \%$ of samples remained as error. At this point, no further attempts of resolution were undertaken - the additional information gained would not have stood in relation to the time and effort to achieve it.

## 2. Error resolution for the second pooled DNA screening

Error resolution of the second pooled screening followed the same principles.
Eventually, $1.55 \%$ of markers were not reproducible and excluded (1.67\% in the donor screen, $1.3 \%$ in the recipient screen).

## Supplementary Material

A data disc containing the supplementary material is attached to the back of this page.

## Supplementary File 1.1

## List of 248 gene association studies: Associations of non-HLA gene polymorphisms with HSCT outcomes

## Last updated: 15 April 2012

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## Supplementary file 2.1:

## JMDP registry analysis and study cohort design

## 1. Genetic background of the Japanese population

It has been pointed out that the studied population should stem from a homogenous genetic background without genetic admixture. It was known from data of the HapMap Consortium (Consortium, 2005, Stranger et al., 2005) that genetic variation varies enormously between different populations, more so the closer the population was to the evolutionary African population, and less so the further the population had moved away from Africa. The Japanese, as the furthest East of all Asian populations, had a genetic structure of less, and better preserved haplotype blocks than Caucasians or Africans (Conrad et al., 2006, Gabriel et al., 2002). While the European and North American History was characterized by large shifts and admixing of populations (e.g. migration within Europe, immigration and displacement of Caucasians, Africans and Asians to North America); due its geography Japan had experienced fewer admixtures. The Japanese population is genetically closely related to that of South East and North East Asia, areas on the Asian mainland from which the Japanese archipelago was initially and repeatedly colonised, and to which it maintained close relations through history (Omoto and Saitou, 1997, Nanta, 2008, Hanihara, 1991).
According to their records, all HSCT donors and recipients in this study were of Japanese origin, hence genetic admixture is very low. As this population had larger size and a smaller number of preserved haplotype blocks, with fewer haplotypes per block, the linkage disequilibrium range of a selected marker could expected to be wider as compared to other populations, therefore less markers would be required to give the same LD range coverage as for a more genetically diverse population. This had implications for statistical power (potentially, the population size required could be set lower, and a smaller number of markers required less statistical correction for
multiple testing).
Difference of genetic variation between populations has been an area of intense research in the HSCT field. For example, the lower incidence of GVHD in the Japanese population could be attributed in part to differences in HLA alleles amongst the Japanese and other population, and subsequent HLA matching for HSCT (Petersdorf et al., 2007, Morishima et al., 2007a). With growing HSCT registries, there has been a move towards studying populations from a more homogenous ethnic background. Non-HLA gene polymorphisms previously associated with GVHD also showed considerable variation of allele and genotype distribution (e.g. IL10 (Middleton et al.), IL6 (Visentainer et al., 2008), NOD2 (Tanabe et al., 2011)).

## 2. Cohort Selection

### 2.1. Analysis of clinical and genetic risk factors in a large, unselected HSCT population

Clinical risk factors for acute GVHD have been documented in studies from the JMDP registries or other unrelated HSCT studies in Japan. These included donor-recipient gender mismatch, patient age, donor age, diagnosis (acute lymphoblastic leukaemia(ALL) versus chronic myeloid leukaemia (CML)), GVHD prophylaxis (cyclosporine versus tacrolimus), relapse risk, antithymoglobulin (ATG), conditioning regimen (total body irradiation (TBI) versus non-TBI), stem cell source (bone marrow transplant (BMT) versus peripheral blood stem cell transplant (PBSCT)) and blood group (ABO) mismatch (Kawase et al., 2007, Morishima et al., 2007a, Hara et al., 2007, Kimura et al., 2008). Multivariate analyses in Japanese genetic association studies had confirmed the significance of these risk factors. Studies in other populations identified Donor and recipient cytomegaly virus (CMV) status, intensity of conditioning regimens, malignant disease stage, diagnosis, unrelated versus related HSCT, early transplants (before 2000), older donor age, older patient age, high nucleated cell count, stem cell source, ethnicity, gender mismatch, donor parity and ABO mismatch as further clinical risk factors (Hill et al., 1997, Socie et al., 2001, Aschan, 2007, Chaidos et al., 2007, Wojnar et al., 2006, Svennilson et al., 2003, Nash et al., 1992). For the Japanese population, more recent work identified KIR and KIRL mismatches, HLA haplotypes and minor histocompatibility antigens ( mHag ) as genetic risk factors.(Yabe et al., 2008, Morishima et al., 2010, Kawase et al., 2008, Ogawa et al., 2008). Eventually, it had to be assumed that many more clinical and genetic risk factors exist of which we have little evidence, owed to the restricted nature of data collections.

In order to identify risk factors for moderate-severe GVHD (grade 2-4), a large cohort ( $n=2469$ HSCT pairs) from the JMDP registry, representing a majority of unrelated donor HSCT in Japan between 1993-2000, was investigated for
significant associations by both log rank test and Fisher's exact test, and binary logistic regression.
The baseline characteristics of this population were summarised in table 1.
Risk factors for grade 2-4 GVHD considered included recipient age, recipient gender, donor age, donor gender, donor-recipient gender mismatch (female donor to male recipient), diagnosis, ABO mismatch, conditioning regimen, cyclophosphamide dose, total body irradiation dose, number of nucleated cells in graft, GVHD prophylaxis, antithymocyte globulin, T-cell depletion and HLA matching.
Table 2 shows the associations of clinical risk factors with grade 2-4 acute GVHD in univariate analysis, using Fisher's Exact Test with 95\% confidence intervals for Odds Ratio. Donor T-cell depletion (TcD) and the use of antithymocyte globulin (ATG) to eradicate recipient T-cells and 5-locus HLA matching confered the strongest effects on the prevention of acute GVHD. Certain diagnosis in itself, or by their population and treatment characteristics, carried a higher risk of acute GVHD. Donor age $>30$ years almost matched the effect size of HLA mismatch. A conditioning regimen consisting of standard dose cyclophosphamide and total body irradiation had the lowest risk of acute GVHD. Cyclophosphamide in itself modulated the recipient immune response in a GVHD protective way, even at high dose. TBI at standard doses has the lowest GVHD risk, but as part of conditioning (TBI versus no TBI) had no impact on GVHD risk. No significant effect on GVHD risk was found with recipient or donor gender, female into male mismatch, AB0 matching, graft nucleated cell count, or recipient age. Recipient age $>40$ years, however, showed a tendency towards a higher risk of grade 2-4 GVHD. The multivariate analysis for this unselected population is shown in table 3. Taken all risk factors from the univariate analysis into account, seven of these remained significant, relating to demographics (donor age $>30 y$, high risk diagnosis (ALL, CML)), conditioning (other conditioning regimen than CyTBI, cyclophosphamide dose none/low versus standard/high) and GVHD prophylaxis (cyclosporine A versus tacrolimus, no ATG versus ATG, HLA mismatching).

| Category | Factor | Frequency | Percent |
| :---: | :---: | :---: | :---: |
| Recipient age | 0-10y | 406 | 16.4 |
|  | 11-20y | 542 | 22.0 |
|  | 21-30y | 600 | 24.3 |
|  | 31-40y | 453 | 18.3 |
|  | 41-50y | 421 | 17.1 |
|  | 51-60y | 47 | 1.9 |
| Recipient gender | Female | 994 | 40.3 |
|  | Male | 1475 | 59.7 |
| Donor age | 0-10y | 0 | . 0 |
|  | 11-20y | 20 | . 8 |
|  | 21-30y | 895 | 36.2 |
|  | 31-40y | 941 | 38.1 |
|  | 41-50y | 602 | 24.4 |
|  | 51-60y | 9 | . 4 |
|  | 61-70y | 1 | . 0 |
|  | unknown | 1 | . 0 |
| Donor gender | Female | 974 | 39.4 |
|  | Male | 1495 | 60.6 |
| Female-male gender mismatch | no F-M mismatch | 1950 | 79.0 |
|  | F-M mismatch | 518 | 21.0 |
| Diagnosis | unknown | 7 | . 3 |
|  | ALL (acute lymphoblastic leukaemia) | 653 | 26.4 |
|  | ANLL (acute non-lymphoblastic leukaemia) | 617 | 25.0 |
|  | CML (chronic myeloid leukaemia) | 643 | 26.0 |
|  | HD (Hodgkin's disease) | 52 | 2.1 |
|  | ID (primary immunodeficiency) | 17 | . 7 |
|  | LPD (lymphoproliferative disease) | 1 | . 0 |
|  | MDS (myelodysplastic syndrome) | 201 | 8.1 |
|  | MF (myelofibosis) | 2 | . 1 |
|  | MM (multiple myeloma) | 2 | . 1 |
|  | NHL (non-Hodgkin lymphoma) | 96 | 3.9 |
|  | SAA (severe aplastic anaemia) | 178 | 7.2 |
| ABO matching | matched | 1248 | 50.5 |
|  | minor mismatch | 526 | 21.3 |
|  | major mismatch | 620 | 25.1 |
|  | minor \& major mismatch | 55 | 2.2 |
|  | unknown | 20 | . 8 |
| Conditioning | No of different regimens | 86 |  |
|  | Busulphan+Cyclophosphamide | 241 | 9.8 |
|  | Busulphan+Cyclophosphamide+other | 145 | 5.9 |
|  | Cyclophosphamide+other | 53 | 2.1 |
|  | Cyclophosphamide+total body irradiation | 639 | 25.9 |


|  | Cyclophosphamide+total body irradiation+other | 90 | 3.6 |
| :---: | :---: | :---: | :---: |
|  | Cyclophosphamide+total body irradiation+busulphan | 251 | 10.2 |
|  | Cyclophosphamide+total body irradiation+Ara-C | 504 | 20.4 |
|  | Cyclophosphamide+total body irradiation+VP16213 | 220 | 8.9 |
|  | other | 59 | 2.4 |
|  | Total body irradiation+other | 267 | 10.8 |
| Cyclophosphamide dose | none | 295 | 11.9 |
|  | <120 | 576 | 23.3 |
|  | 120 (standard) | 1262 | 51.1 |
|  | >120 | 312 | 12.6 |
|  | unknown | 24 | 1.0 |
| Total Body Irradiation | none | 486 | 19.7 |
|  | $<1200$ | 336 | 13.6 |
|  | 1200 | 1485 | 60.1 |
|  | >1200 | 149 | 6.0 |
|  | unknown | 13 | . 5 |
| No of nucleated cells | <251 | 509 | 20.6 |
|  | 251-400 | 1437 | 58.2 |
|  | >400 | 503 | 20.4 |
|  | unknown | 20 | . 8 |
| GVHD prophylaxis | No of different regimens | 21 |  |
|  | Cyclosporin A - based | 1818 | 73.6 |
|  | Tacrolimus - based | 641 | 26.0 |
|  | Other | 10 | . 4 |
| Antithymoglobulin | unknown | 4 | . 2 |
|  | no ATG | 2279 | 92.3 |
|  | ATG | 186 | 7.5 |
| T-cell depletion | unknown | 4 | . 2 |
|  | no T-cell depletion | 2448 | 99.1 |
|  | T-cell depletion | 17 | . 7 |
| HLA matching | 5-locus matched | 925 | 37.5 |
|  | mismatched | 1544 | 62.5 |

Table 1: Baseline characteristics of an unselected HSCT population
( $n=2469$ ) from the JMDP registry

| Category | comparison | $p$-value | case <br> all no | control all no | case <br> pos | case <br> neg | control pos | $\begin{gathered} \text { control } \\ \text { neg } \\ \hline \end{gathered}$ | Odds <br> Ratio | Confidence interval | confidence interval | comment |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| GVHD prophylaxis | No TcD v TcD | 0.039 | 1444 | 968 | 13 | 1431 | 3 | 965 | 2.9222 | 0.83051 | 10.282 | No TcD higher risk |
| GVHD prophylaxis | No ATG v ATG | 0.002 | 1447 | 968 | 128 | 1316 | 54 | 914 | 1.64629 | 1.18468 | 2.28777 | ATG protective |
| HLA matching | mismatch v 5 locus match | <0.0001 | 968 | 1447 | 653 | 315 | 837 | 610 | 1.5108 | 1.27436 | 1.79111 | HLA mismatch higher risk |
| Diagnosis | high risk (ALL+CML) v low risk (other) | <0.0001 | 968 | 1447 | 566 | 402 | 703 | 744 | 1.49007 | 1.2644 | 1.75603 | ALL, CML risk |
| GVHD prophylaxis | Cyclosporin v tacrolimus | <0.0001 | 968 | 1447 | 751 | 214 | 1032 | 411 | 1.39762 | 1.15566 | 1.69023 | CyA higher risk |
| Donor age | >30y v <30y | <0.0001 | 968 | 1447 | 652 | 316 | 863 | 584 | 1.39625 | 1.17735 | 1.65585 | higher age-higher risk |
| Conditioning | Other v CyTBI | 0.002 | 968 | 1447 | 218 | 750 | 408 | 1039 | 1.35098 | 1.11792 | 1.63263 | Non-CyTBI higher risk |
| Conditioning | Cy nollow dose v standard/high dose | 0.001 | 956 | 1436 | 964 | 472 | 576 | 380 | 1.3474 | 1.13674 | 1.59709 | no/low have higher risk |
| Conditioning | TBI standard dose v lower/higher dose | 0.023 | 964 | 1438 | 797 | 167 | 1135 | 303 | 1.27406 | 1.03325 | 1.57098 | standard dose protective |
| Donor age | >40y v <40y | 0.021 | 968 | 1447 | 265 | 703 | 335 | 1112 | 1.25127 | 1.03809 | 1.50822 | higher age-higher risk |
| Recipient gender | Male v Female | 0.062 | 968 | 1447 | 600 | 368 | 842 | 605 | 1.17151 | 0.99173 | 1.38388 |  |
| Donor gender | Male v Female | 0.14 | 968 | 1447 | 605 | 363 | 856 | 590 | 1.14875 | 0.97195 | 1.35772 |  |
| ABO matching | matched $v$ mismatched | 0.156 | 962 | 1434 | 474 | 488 | 749 | 685 | 0.88832 | 0.75437 | 1.04604 |  |
| No of nucleated cells | high v low | 0.215 | 954 | 1441 | 85 | 869 | 108 | 1333 | 1.20727 | 0.89727 | 1.62438 |  |
| Donor-Recipient gender mismatch | Female into Male v other | 0.495 | 968 | 1446 | 197 | 771 | 311 | 1135 | 0.9325 | 0.76304 | 1.13958 |  |
| Recipient age | >40y v <40y | 1 | 968 | 1447 | 179 | 789 | 267 | 1180 | 1.00264 | 0.81297 | 1.23657 |  |
| Conditioning | TBI v non-TBI | 1 | 964 | 1438 | 774 | 190 | 1155 | 283 | 0.99814 | 0.81302 | 1.22542 |  |

Table 2: Univariate analysis of clinical risk factors in an unselected cohort of Japanese HSCT from the JMDP register (1993-2000)

|  | B | SE | Wald | df | Sig. | Exp(B) | $\begin{gathered} 95.0 \% \mathrm{Cl} \text { for } \\ \operatorname{Exp}(\mathrm{B}) \end{gathered}$ |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  |  |  | Lower | Upper |
| No ATG v ATG | -. 500 | . 147 | 11.604 | 1 | . 001 | 1.648 | 1.236 | 2.197 |
| HLA mismatch v 5locus match | -. 413 | . 071 | 34.048 | 1 | . 000 | 1.511 | 1.315 | 1.736 |
| Cyclosporin A v Tacrolimus | . 347 | . 080 | 18.812 | 1 | . 000 | 1.415 | 1.210 | 1.656 |
| No CyTBI v CyTBI | -. 241 | . 084 | 8.244 | 1 | . 004 | 1.271 | 1.079 | 1.498 |
| High risk diagnosis (ALL+CML) v other | . 236 | . 068 | 12.080 | 1 | . 001 | 1.266 | 1.108 | 1.447 |
| $\begin{aligned} & \text { Donor age }>30 y \text { v } \\ & <30 y \end{aligned}$ | . 009 | . 003 | 6.862 | 1 | . 009 | 1.009 | 1.002 | 1.016 |
| Cyclophosphamide non/low v standard/high | -. 007 | . 003 | 4.992 | 1 | . 025 | 1.007 | 1.001 | 1.013 |
| TBI v no TBI | . 152 | . 092 | 2.769 | 1 | . 096 | 1.164 | . 973 | 1.393 |
| $\begin{aligned} & \text { Donor age }>40 \mathrm{y} v \\ & <40 \mathrm{y} \end{aligned}$ | . 004 | . 003 | 1.870 | 1 | . 172 | 1.004 | . 998 | 1.009 |
| TBI dose low/high v standard | -. 009 | . 008 | 1.546 | 1 | . 214 | . 991 | . 976 | 1.005 |
| TcD v no TCD | -. 672 | . 581 | 1.339 | 1 | . 247 | . 511 | . 164 | 1.594 |
| ABO matched v mismatched | . 072 | . 066 | 1.190 | 1 | . 275 | 1.075 | . 944 | 1.225 |
| Nucleated cell count low v high | . 122 | . 117 | 1.086 | 1 | . 297 | 1.129 | . 898 | 1.419 |
| Female into male gender mismatch | -. 110 | . 139 | . 625 | 1 | . 429 | . 896 | . 682 | 1.177 |
| Recipient gender | -. 061 | . 088 | .474 | 1 | . 491 | . 941 | . 792 | 1.119 |
| Donor gender | -. 072 | . 107 | . 455 | 1 | . 500 | . 930 | . 754 | 1.147 |
| Recipient age $>40 y$ v $<40 y$ | -. 001 | . 002 | . 135 | 1 | . 713 | . 999 | . 995 | 1.003 |

Table 3: Multivariate analysis (Cox regression) of clinical risk factors in an unselected Japanese HSCT cohort from the JMDP registry, 1993-2000

### 2.2. Designing a study cohort accounting for clinical risk factors

A large proportion of HSCT in Japan by JMDP between 1993 and 2000 included one or more statistically significant risk factors for moderate-severe acute GVHD (grade 2-4), (see table 4). This outcome was chosen because GVHD was one of the most commonly reported HSCT outcomes in the literature. The challenge was to devise a model for cohort selection that eliminated the strongest effects of these clinical risk factors, while preserving a sample size that would provide adequate statistical power. Knowing that the effect of HLA mismatching has an OR of approximately 1.5, it was aimed for a sample size that would undercut this OR with regards to statistical power. It was estimated that for a single outcome a sample size of approximately $\mathrm{n}=500$ in each of the two screening steps (or 1000 altogether) would be required to achieve an OR of 1.3 for an allele frequency of 0.2 .
I order to reduce the effect of HLA mismatching, a higher degree HLAmatched subgroup was analysed as a comparison (mismatched only for either one HLA-DQB1 or HLA-DPB1 locus, largely representing an 8/8 match of HLA A, B, C, DR) and non- ATG group as a control to compare variability and significance of the clinical risk factors in the different models.

GVHD prophylaxis with either cyclosporine A or tacrolimus was not applied for cohort selection as this would have introduced a strong time bias (cyclosporine A was largely replaced by tacrolimus in the late 1990's when registry studies showed that this measure significantly reduced the incidence of acute GVHD). Donor age >30 years has a strong effect, but such limitation would have been practically unrealistic with regards to sample size and power. These data showed that a standard dose Cyclophosphamide/TBI regimen carried the lowest risk of acute GVHD, but this regimen was in practice not suitable for all patients, especially high-risk ones. TBI in itself was known to induce a higher risk for acute GVHD; however, in this analysis TBI showed only a trend in this direction. In summary, the data on the different conditioning regimen and their dosages indicate that a TBI versus non-TBI analysis carried a large number of confounders.

Table 5 gives an overview of the models devised. Model 1 followed strictly the order of the effect size of the multivariate analysis. The second selection step (ATG given - HLA mismatched removed) already reduced the sample size below target ( 870 HSCT pairs). This is the model used as a control to study variability, significance and effect sizes of risk factors in the other models.

In Model 2, the focus was on the conditioning regimen (ATG given conditioning other than Cyclophosphamide/TBI), reaching a sample size of 609 HSCT pairs, which would have been unfeasible.

Model 3 concentrated on the underlying diagnosis (excluding ALL and CML ATG given included), resulting in a sample size of 1054 HSCT pairs. A diagnosis-driven approach appeared the most feasible. However, concerns about this approach included the bundling of a multitude of diagnoses in the low-risk group (including malignancies and non-malignancies), versus two very different diagnoses in the high risk group (ALL, and CML - the latter one almost abandoned as a HSCT indication). Hence, a cohort selecting the two large, relevant acute leukaemia groups (ALL, ANLL) was included in the following comparison (Model 4).
In model 1, donor age $>30$, cyclosporine-based GVHD prophylaxis, nonCy/TBI conditioning regimens and TBI remained the most important clinical confounders (table 6), despite high degree of HLA matching. Model 3 (table 7) favoured a multitude of low-risk diagnosis over full HLA matching, resulting in a wider variety of clinical confounders, which amongst donor age $>30$ years and cyclosporine-based GVHD prophylaxis also included noncyclophosphamide/TBI conditioning regimens, cyclophosphamide dose, TBI dose and HLA mismatching.

In contrast, model 4 (table 8) displayed cyclosporine GVHD prophylaxis, donor age and HLA mismatching as the strongest confounders, while aspects of conditioning regimens and ABO mismatch had a borderline significant role. As this model provided a sample size of adequate statistical power, it was chosen as the preferred model for cohort design. At this point, it was chosen to exclude HSCT pairs of a recipient age $<4$ years and $>40$ years. Leukaemia in infants and small children had very different causes and outcomes compared to the leukaemia of older children and adults. Also, an analysis of age groups (5 year intervals) showed a significant increase in risk of acute

GVHD grade 2-4 with recipient age $>40$ (data not shown here). This reduced the number of eligible sample pairs to $n=1000$.

| risk factor | frequency <br> unselected |
| :--- | ---: |
| No ATG | $92.50 \%$ |
| HLA mismatch | $62.50 \%$ |
| CyA GVHd <br> prophylaxis | $73.60 \%$ |
| Non-CyTBI regimen | $74.10 \%$ |
| High risk diagnosis | $54.40 \%$ |
| Donor age >30y | $63 \%$ |
| Cy dose none/low | $36.30 \%$ |
| TBI | $80.30 \%$ |

Table 4: Proportion of clinical risk factors in unselected HSCT population from the JMDP registry, 1993-2000

| Model | 1 | ATG given removed | HLA mismatched removed | high risk diagnosis removed | donor age $>30 y$ removed | Cy dose none/low removed | TBI removed |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Remaining no | 2469 | 2279 | 870 | 416 | 170 | 107 | 25 |
| Model | 2 | ATG given removed | Non-CyTBI removed | high risk diagnosis removed | donor age $>30 y$ removed | Cy dose none/low removed | TBI removed |
| Remaining no | 2469 | 2279 | 609 | 311 | 118 | 93 | 0 |
| Model | 3 | high risk diagnosis removed | ATG given removed | Non-CyTBI removed | donor age $>30 y$ removed | Cy dose none/low removed | TBI removed |
| Remaining no | 2469 | 1173 | 1054 | 311 | 118 | 93 | 0 |
| Model | 4 | ALL/ANLL selected | ATG given removed | Non-CyTBI removed | $\begin{aligned} & \text { donor age } \\ & >30 \mathrm{y} \\ & \text { removed } \end{aligned}$ | Cy dose none/low removed | TBI removed |
|  | 2469 | 1270 | 1209 | 273 | 99 | 82 | 0 |

Table 5: Modelling of hypothetic study cohorts. Model 1 represents a 5-locus HLA matched and ATG removed cohort, Model $\mathbf{2}$ is selected for homogeneity of conditioning regimens, while models 3 and 4 were selected by diagnosis (model 3: non-high risk diagnosis, model 4: ALL and ANLL selected). Model 4 was chosen for cohort design, as it provides the best statistical power.

| Variables in the Equation |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | log rank univariate |  |  |  |  |  |  | 95.0\% Cl for Exp(B) |  |
|  |  | B | SE | Wald | df | Sig. | Exp(B) | Lower | Upper |
| Cyclosporin v tacrolimus | 0.003 | . 468 | . 148 | 10.047 | 1 | . 002 | 1.597 | 1.196 | 2.134 |
| Donor age <30 v >30 | 0.001 | . 016 | . 006 | 6.912 | 1 | . 009 | 1.016 | 1.004 | 1.029 |
| Cyclophosphamide+TBI v other | 0.021 | -. 387 | . 151 | 6.586 | 1 | . 010 | 679 | 505 | . 913 |
| TBI v noTBI | 0.778 | . 319 | . 162 | 3.884 | 1 | . 049 | 1.375 | 1.002 | 1.889 |
| Cyclophosphamide no/low v standard/high dose | 0.104 | -. 008 | . 006 | 2.045 | 1 | . 153 | . 992 | . 981 | 1.003 |
| recipient age <40 v >40 | 0.368 | . 003 | . 003 | . 843 | 1 | . 359 | 1.003 | 996 | 1.010 |
| donor age <40 v >40 | 0.06 | . 003 | . 004 | . 473 | 1 | . 492 | 1.003 | . 994 | 1.012 |
| abo matched v mismatched | 0.62 | . 068 | . 119 | . 330 | 1 | . 566 | 1.071 | . 848 | 1.353 |
| Female donor into male recip v other | 0.977 | -. 082 | . 148 | . 309 | 1 | . 578 | 921 | .689 | 1.231 |
| high risk doagnosis v other | 0.148 | . 040 | . 119 | . 113 | 1 | . 737 | 1.041 | . 824 | 1.315 |
| T-cell depletion v no T cell depletion | 0.94 | -. 178 | 1.012 | . 031 | 1 | . 860 | . 837 | 115 | 6.083 |
| ATG given v no ATG | NA |  |  | . | $0^{\text {a }}$ | . |  |  |  |
| 5-locus HLA match v mismatch | NA |  |  | . | $0^{\text {a }}$ | . |  |  |  |

Table 6: Multivariate analysis (binary logistic regression) of model 1

| Variables in the Equation |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | log rank univariate | B | SE | Wald | df | Sig. | $\operatorname{Exp}(\mathrm{B})$ | 95.0\% CI for Exp(B) |  |
|  |  |  |  |  |  |  |  | Lower | Upper |
| Donor age <30 v >30 | 0.0001 | . 021 | . 006 | 14.678 | 1 | . 000 | 1.021 | 1.010 | 1.032 |
| Cyclosporin v tacrolimus | 0.00001 | . 483 | . 126 | 14.631 | 1 | . 000 | 1.621 | 1.265 | 2.075 |
| Cyclophosphamide+TBI v other | 0.001 | -. 343 | . 135 | 6.467 | 1 | . 011 | . 710 | . 545 | . 924 |
| 5-locus HLA match v mismatch | 0.013 | -. 254 | . 111 | 5.192 | 1 | . 023 | . 776 | . 624 | . 965 |
| Cyclophosphamide no/low v standard/high dose | 0.001 | . 248 | . 113 | 4.842 | 1 | . 028 | 1.282 | 1.027 | 1.599 |
| TBI dose standard v none/high | 0.009 | -. 024 | . 012 | 4.095 | 1 | . 043 | . 976 | . 954 | . 999 |
| TBI v notBI | 0.798 | . 218 | . 142 | 2.350 | 1 | 125 | 1.244 | . 941 | 1.645 |
| Female donor into male recip v other | 0.376 | . 098 | . 137 | . 507 | 1 | . 476 | 1.103 | . 842 | 1.444 |
| recipient age <40 v >40 | 0.575 | -. 001 | . 003 | . 166 | 1 | . 684 | . 999 | . 993 | 1.005 |
| donor age <40 v >40 | 0.029 | . 001 | . 004 | . 084 | 1 | . 772 | 1.001 | . 993 | 1.009 |
| T-cell depletion $v$ no $T$ cell depletion | 0.846 | -. 147 | 1.011 | . 021 | 1 | . 885 | . 863 | . 119 | 6.265 |
| abo matched v mismatched | 0.621 | . 009 | . 107 | . 008 | 1 | . 929 | 1.010 | . 819 | 1.245 |
| high risk doagnosis v other | NA |  |  |  | $0^{\text {a }}$ |  |  |  |  |
| ATG given v no ATG | NA |  |  |  | $0^{\text {a }}$ |  |  |  |  |

Table 7: multivariate analysis (binary logistic regression) of model 3

|  | log rank univariate | B | SE | Wald | df | Sig. | Exp(B) | 95.0\% CI for Exp(B) |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  |  |  |  | Lower | Upper |
| 5-locus HLA match v mismatch | 0.0001 | -3.879 ×10-1 | . 099 | 15.224 | 1 | . 000 | . 678 | . 558 | . 824 |
| Cyclosporin v tacrolimus | 0.01 | . 308 | . 111 | 7.694 | 1 | . 006 | 1.361 | 1.095 | 1.691 |
| Donor age <30 v >30 | 0.0001 | . 013 | . 005 | 7.309 | 1 | . 007 | 1.013 | 1.004 | 1.023 |
| Cyclophosphamide no/low v standard/high dose | 0.009 | -8.952 $\times 10-3$ | . 004 | 4.135 | 1 | . 042 | . 991 | . 983 | 1.000 |
| abo matched v mismatched | 0.024 | . 192 | . 095 | 4.119 | 1 | . 042 | 1.212 | 1.007 | 1.460 |
| donor age <40 v >40 | 0.0001 | . 007 | . 004 | 3.944 | 1 | . 047 | 1.007 | 1.000 | 1.014 |
| high risk doagnosis v other | 0.042 | . 167 | . 097 | 2.950 | 1 | . 086 | 1.181 | . 977 | 1.428 |
| Cyclophosphamide+TBI v other | 0.046 | -2.150 x10-1 | . 126 | 2.906 | 1 | . 088 | . 807 | . 630 | 1.033 |
| Female donor into male recip v other | 0.6 | -6.642 $\times 10-2$ | 118 | . 320 | 1 | . 572 | . 936 | . 743 | 1.178 |
| recipient age <40 v >40 | 0.382 | -1.168 $\times 10-3$ | . 003 | . 132 | 1 | . 717 | . 999 | . 993 | 1.005 |
| TBI v notBl | 0.785 | -2.637 ×10-2 | . 147 | . 032 | 1 | . 858 | . 974 | . 730 | 1.299 |
| T-cell depletion v no T cell depletion | 0.188 | -1.009 $\times 10-1$ | 117.788 | . 007 | 1 | . 932 | . 000 | $\begin{array}{r} 2.269 \times 10- \\ 105 \end{array}$ | $7.551 \times 10+95$ |
| TBI dose standard v none/high | 0.976 | . 000 | . 011 | . 000 | 1 | . 986 | 1.000 | . 979 | 1.022 |
| ATG given v no ATG | NA |  |  |  | $0^{\text {a }}$ |  |  |  |  |

Table 8: multivariate analysis (binary logisitc regression) of model 4. This model was chosen as the preferred model for cohort design.

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Power calculation

The power of association testing was calculated based on following $2 \times 2$ contingency tables.
Given a microsatellite loci with $k$ alleles, we created $2 \times 2$ contingency tables (table 3.1) for each indiv chi-squared test or Fisher's exact test.
Power for each pool was calculated, changing combinations of following parameters:

- Odds ratio: 1.5, 2.0, and 2.5
- Marker allele frequency: $0.03,0.05,0.10$, and 0.30
- Type I error rate: 0.05

Power was calculated by using the software 'PS: Power and Sample Size Calculation'.
The statistical power was calculated in the total cells as the power to detect association in successiv association tests in pool 1 and pool 2 are significant ( $p<0.05$ ). Therefore, Power (Total) = Power(Pool 1) * Power(Pool 2)
The reason why we calculated the total power described above is that we select microsatellite mark successive pools 1 and 2 as candidates for next screening.


Fig 1: Statistical power for $O R=1.5$


Fig 2: Statistical powe
jidual allele, which can be analyzed using a
'e pools 1 and 2 . In other words, both of the
<ers that confer statistical significance in

|  | Marker allele |  |
| :--- | :---: | :---: |
|  | $\mathrm{M}_{1}$ | $\mathrm{M}_{2} \sim \mathrm{M}_{\mathrm{k}}$ |
| Grade 2- <br> 4 GVDH <br> Grade 0- <br> 1 GVDH | a | b |

## Chi Square table for power calculation.

3 r for $\mathrm{OR}=2.0$


Fig 3: Statistical power for $\mathrm{OR}=\mathbf{2 . 5}$
a~d: Allele counts


## Supplementary file 2.3:

## Construction of final study cohorts and their characteristics

## 1. Initial assessment of DNA sample number, quantity and concentration

At this point it was decided to assess the actual availability of DNA samples for the chosen cohort model. The first step was the identification of selected samples from the database and the sample collection, stored in different freezers at $-70^{\circ} \mathrm{C}$.

The Japan marrow Donor Programme (JMDP) provided two large data files: One contained the clinical data, with a unique pair number (UPN) and a DNA sample number as identifiers. The second file contained the UPN, the DNA sample number, and a new DNA Bank number for each sample which was established recently when the JMDP DNA collections from different time periods were integrated into one system. Samples were in cardboard boxes in simple numerical order, representing the timely order of the transplantations.

As an initial step, a data file of the intended cohort linking all sample information and clinical information together was created. From this data file lists were extracted to enable the identification of the targeted samples in the boxes. Original DNA samples of the intended cohorts were then obtained from the original collection and sorted into a separate set of boxes. Samples were separated into donors/recipients, ALL/ANLL, grades of GVHD 0-4, and finally in numerical order.

At the same instance, the total volume of each sample was estimated by comparing its volume to standard test volumes pipetted in $50 \mu$ intervals (50$800 \mu \mathrm{l})$ into a set of identical test tubes. This later enabled an estimation of the total amount of DNA per original sample tube in $\mathrm{ng} / \mu \mathrm{g}$ (see below).

Missing and depleted samples were identified by simple visual inspection, and listed accordingly. Of the initial $n=1000$ pairs, $n=112$ were completely (both pair partners) or partially (one pair partner) depleted and therefore excluded. $N=543$ pairs had at least some DNA and were therefore extracted from the collection for further exploration. N=345 pairs had been transferred to Tokyo University (Research Group Professor Ogawa) to be included in a separate study. These samples were mostly fully HLA matched. Enquiries with Professor Ogawa's team revealed that of the $n=345$ sample pairs, $n=74$ pairs were also depleted and excluded.
This meaned that a maximum of only $n=814$ samples would be available for this study, with at that point an unknown amount of DNA, and at that point uncertainty when a larger proportion of fully HLA matched pairs would become available from Tokyo University.

As time was constraint, a feasibility report explored the available options.

### 1.1. Sample Availability and Study Scenario Feasibility

Three different study scenarios, reflecting a spectrum between a genome wide scanning study and an individual genotyping study have been assessed for feasibility from a sample availability point of view (table 1). From the experience of previous microsatellite and SNP studies in the Tokai University laboratory, the required amount of DNA was estimated for these scenarios (table 2).

| 1 | Genome wide scanning using approximately 30.000 MS markers |
| :--- | :--- |
| 2 | Limited genome scanning of immune regulatory genes, approximately <br> 3000 MS markers |
| 3 | Individual genotyping study using approximately 100 MS and SNP <br> markers |

Table 1: Scenarios for feasibility assessment

| Study type | Minimum DNA <br> in microgr |
| :--- | :--- |
| Genome wide, 30.000 MS | 30 |
| Limited scanning, 3000 MS | 5 |
| Individual genotyping, $100 \mathrm{MS} /$ SNP | 1 |

Table 2: Estimates of DNA amount required

### 1.2. Sample Concentration and volumes

Samples available at Tokai University. All available samples of pairs from the initial ALL and ANLL cohorts (age stratified 4-40 years) were identified from different freezers at Tokai University. Available and unavailable samples were marked in a list, and the volume of the available samples was estimated using a simple model.

The DNA concentration of 1086 (543 pairs) available samples was measured using the PICO Green method (described in the methods section). Total amount was calculated multiplying concentration with estimated volume.

Tokyo University samples. Professor Ogawa kindly provided a table with concentration and quality data of all samples from this cohort he used in his study. These data did not contain any total volume estimations, therefore at this stage these have been estimated applying data available from the samples at Tokai University. We estimated samples with a concentration $>5 \mathrm{ng} / \mu \mathrm{l}$ to correspond with a total amount of $>1 \mu$, a concentration of $>10 \mathrm{ng} / \mu \mathrm{g}$ with a total amount of $>5 \mu \mathrm{~g}$, and a concentration of $>50 \mathrm{ng} / \mu \mathrm{l}$ with a total amount of $>30 \mu$, accordingly.

Summary of sample availability: Table 3 summarises the sample availability for the different study scenarios. The data for Tokai University samples are accurate, while the Tokyo University data are estimates for the reasons explained above.

| Scenario | Samples available at Tokai | HLA <br> matched $8 / 8$ | Sample <br> s <br> availabl <br> e at <br> Tokyo | HLA matche d 8/8 | Total (HLA matched) | Pairs require d for scenari 0 | + / - |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Genome wide | 44 | 15 (34\%) | 41 | $\begin{aligned} & \hline 27 \\ & (66 \%) \end{aligned}$ | $\begin{aligned} & \hline 85 \\ & (42,49 \%) \end{aligned}$ | 900+ | -815 |
| Limited genome | 407 | $\begin{aligned} & 81 \\ & (19.9 \%) \end{aligned}$ | 271 | $\begin{aligned} & 195 \\ & (72 \%) \end{aligned}$ | $\begin{aligned} & \hline 678 \\ & (276,40 \%) \end{aligned}$ | $\begin{aligned} & 600+1 \\ & 900+ \end{aligned}$ | $\begin{aligned} & +781 \\ & -222 \end{aligned}$ |
| Individual typing | 543 | 120 (22\%) | 271 | $\begin{aligned} & \hline 195 \\ & (72 \%) \end{aligned}$ | $\begin{aligned} & \hline 814 \\ & (315,38 \%) \end{aligned}$ | 500+ | $\begin{aligned} & +31 \\ & 4 \end{aligned}$ |

Table 3: Summary of expected sample availability for different study approaches

### 1.3. Conclusions

Genome wide scanning. This scenario was not feasible given the very limited number of pairs having a sufficient amount of DNA. Even extrapolated to the entire study population (2469 pairs), which would be completely unselected, no more than 330 pairs would have had a sufficient amount of DNA.

Limited genome scanning. This approach was a feasible option if samples from Tokyo University were included, and if the study would only implicate two screening steps. Application to JMDP for further sample access was required.

Individual Genotyping. This was also a feasible option. The cohort size of over 800 may even allow for some further selection. Although even with the samples available at Tokai University this could be feasible, the proportion of 8/8 HLA matched pairs is low for a Japanese population (20\%). A better approach would be to include the matched pairs from Tokyo University.

The options were discussed between the team at Tokai University and JMDP in October 2007. All partners were keen on undertaking a genomic screening study, rather than a candidate gene association study. It was agreed to combine samples from Tokai and Tokyo Universities for a first screening step on pooled DNA, using microsatellite markers, on the selected cohort that was proposed. Access to further samples was approved, and samples and dataset prepared. The first screening would encompass HSCT between 1993 and 2000, while the second cohort would include those between 2001 and 2005.

## 2. Application of selection criteria for construction of a discovery and a confirmatory cohort

HSCT pairs for the first cohort (time frame 1993-2000, $n=460$ ) were selected on the basis of criteria for model 4, and DNA availability. Criteria included:

- Acute leukaemia (ALL or ANLL)
- Myeloablative conditioning
- T-cell replete
- Full bone marrow HSCT
- Recipient age 4-40 years
- DNA availability for both donor and recipient sample of $5 \mu \mathrm{~g}$.

The second cohort ( $\mathrm{n}=462$ ) was selected by the same criteria. In order to reduce confounding by different grades of HLA mismatching, samples were paired for HLA matching between first and second cohort. In practice, for each of the 48 allele mismatch combinations, an equivalent was chosen from the 2001-2005 stem population.

All donor-recipient pairs were HLA-typed retrospectively to allele level at six loci (HLA-A, HLA-B, HLA-C, HLA-DRB1, HLA-DQB1, HLA-DPB1). The distribution of HLA matching of the confirmatory cohort was adjusted to that of the screening cohort by matching each sample of the screening cohort with a confirmatory cohort sample of the same HLA class or HLA class combination according to the previous literature (Sasazuki et al., 1998, Morishima et al., 2002) and our own analyses of risk matches/mismatches within this study population (data not shown).

Table 4 shows the demographic and clinical characteristics of the selected cohorts. There was no statistically significant difference between the cohorts in the baseline demographic criteria. Tables 5 and 6 specify the degree of HLA matching and mismatching. For reasons of comparison, we have used the NMDP/CIBMTR classification of HLA matching (Weisdorf et al., 2008). According to this classification, 357 HSCT pairs had an 8/8 (HLA A, B, C, DRB1) high-resolution allele match, 331 (35.9\%) were partially matched (1
mismatch within these HLA loci), and 234 (25.4\%) were mismatched (two or more mismatches within these HLA loci). Considering the HLA DQ and DP loci also, only 78 HSCT pairs (8.5\%) had a 12/12 allele match. In Japanese, HLA A, B, and C mismatches were associated with risk of acute GVHD. HLA C mismatches, however, had a protective effect on relapse (whilst HLA A, C and B mismatches associate with a risk of death) (Sasazuki et al., 1998, Morishima et al., 2002, Morishima et al., 2007b). More recent research had focused on specific allele mismatches, rather than mismatches in loci, aiming to identify non-permissive mismatches for acute GVHD (Kawase et al., 2007) or protective mismatches against relapse (Kawase et al., 2009), as well as risk HLA haplotypes for GVHD(Morishima et al., 2010).

Multivariate analysis of the finally selected combined cohorts ( $n=922$ HSCT pairs, table 7) showed that diagnosis, recipient age and HLA mismatch remain the most significant confounding variables. ALL had a higher risk for moderate-severe acute GVHD than ANLL. Recipient age below 10 years was carrying a higher GVHD risk. 8/8 loci HLA match is protective against GVHD. Relapse and major ABO mismatch still showed trends towards risk.

|  | Screening | Confirmation | Difference $p=$ |
| :---: | :---: | :---: | :---: |
| Demographic |  |  |  |
| Recipients | 460 | 462 |  |
| Recipient gender Male | 269 (58.48\%) | 289 (62.55\%) | $\mathrm{n} / \mathrm{s}$ |
| Recipient gender Female | 191 (41.52\%) | 173 (37.45\%) | $\mathrm{n} / \mathrm{s}$ |
| Donor gender Male | 267 (58.04\%) | 278 (60.17\%) | $\mathrm{n} / \mathrm{s}$ |
| Donor gender Female | 193 (41.96\%) | 182 (39.39\%) | $\mathrm{n} / \mathrm{s}$ |
| Female donor to Male recipient transplant | 102 (22.17\%) | 84 (18.18\%) | $\mathrm{n} / \mathrm{s}$ |
| Recipient age range | 4-40 y | $4-40 \mathrm{y}$ | $\mathrm{n} / \mathrm{s}$ |
| Recipient age mean | 21.7 y | 24.1 y | $\mathrm{n} / \mathrm{s}$ |
| Donor age range | 20-70y | $19-51$ y | $\mathrm{n} / \mathrm{s}$ |
| Donor age mean | 34 y | 34.3 y | $\mathrm{n} / \mathrm{s}$ |
| Clinical |  |  |  |
| Diagnosis Acute lymphoblastic leukaemia | 260 (56.52\%) | 254 (54.98\%) | $\mathrm{n} / \mathrm{s}$ |
| Diagnosis Acute non-ALL | 200 (43.48\%) | 208 (45.02\%) | $\mathrm{n} / \mathrm{s}$ |
| High risk leukaemia | 279 (60.65\%) | 246 (53.25\%) | <0.1 |
| HLA matching - 12/12 loci | 41 (8.91\%) | 37 (8\%) | $\mathrm{n} / \mathrm{s}$ |
| HLA matching - 12/12 and 10/10 loci | 160 (34.78\%) | 166 (35.93\%) | $\mathrm{n} / \mathrm{s}$ |
| HLA matching - GVHD risk mismatches | 220 (47.83\%) | 229 (49.57\%) | $\mathrm{n} / \mathrm{s}$ |
| Conditioning - Cyclophosphamide + total body irradiation (TBI) | 334 (72.61\%) | 322 (69.67\%) | $\mathrm{n} / \mathrm{s}$ |
| Conditioning - <br> Busulphan/Cyclophosphamide or Busulphan based | 53 (11.52\%) | 47 (10.17\%) | $\mathrm{n} / \mathrm{s}$ |
| Conditioning - TBI based | 73 (15.87\%) | 83 (17.97\%) | $\mathrm{n} / \mathrm{s}$ |
| Conditioning - other | 0 | 10 (2.16\%) | $\mathrm{n} / \mathrm{s}$ |
| GVHD prophylaxis - Cyclosporin A based | 279 (60.65\%) | 154 (33.33\%) | <0.05 |
| GVHD prophylaxis - Tacrolimus based | 177 (34.48\%) | 305 (66.02\%) | <0.05 |
| GVHD prophylaxis - other | 4 (0.87\%) | 3 (0.65\%) | $\mathrm{n} / \mathrm{s}$ |
| Outcome |  |  |  |
| Acute GVHD grade 0 | 124 (26.96\%) | 124 (26.84\%) | $\mathrm{n} / \mathrm{s}$ |
| Acute GVHD grade 1 | 153 (33.26\%) | 99 (21.42\%) | <0.05 |
| Acute GVHD grade 2 | 105 (22.83\%) | 143 (30.95\%) | <0.05 |
| Acute GVHD grade 3 | 50 (10.87\%) | 72 (15.58\%) | <0.05 |
| Acute GVHD grade 4 | 28 (6.09\%) | 24 (5.19\%) | $\mathrm{n} / \mathrm{s}$ |
| Chronic GVHD - none | 244 (53.04\%) | 242 (52.38\%) | $\mathrm{n} / \mathrm{s}$ |
| Chronic GVHD - limited disease | 71 (15.43\%) | 63 (13.64\%) | $\mathrm{n} / \mathrm{s}$ |
| Chronic GVHD - extensive disease | 95 (20.65\%) | 106 (22.94\%) | $\mathrm{n} / \mathrm{s}$ |
| Chronic GVHD - unknown | 50 (10.86\%) | 49 (10.6\%) | $\mathrm{n} / \mathrm{s}$ |
| Relapse | 115 (25\%) | 110 (23.81\%) | $\mathrm{n} / \mathrm{s}$ |
| Survival - 100 days | 395 (86.9\%) | 403 (87.23\%) | $\mathrm{n} / \mathrm{s}$ |
| Survival - 1 year | 306 (66.52\%) | 312 (67.53\%) | $\mathrm{n} / \mathrm{s}$ |
| Survival - 3 years | 245 (53.26\%) | 258 (55.84\%) | $\mathrm{n} / \mathrm{s}$ |

Table 4: Population characteristics. $p$ refers to statistically significant differences between the screening and confirmation cohorts.

|  | alleles mismatched | $\mathrm{n}=$ |  |
| :---: | :---: | :---: | :---: |
| $\begin{aligned} & \text { Matched 8/8 } \\ & \mathrm{n}=357 \text { (38.7\%) } \end{aligned}$ | HLA-DQ-DP | 30 | 3.3 |
|  | HLA-DQ | 6 | 0.7 |
|  | HLA-DP | 243 | 26.4 |
|  | Fully matched | 78 | 8.5 |
| Partially mismatchedn=331 (35.9\%) | HLA-DR | 1 | 0.1 |
|  | HLA-C-DQ | 3 | 0.3 |
|  | HLA-C-DP | 92 | 10 |
|  | HLA-A | 11 | 1.2 |
|  | HLA-C | 25 | 2.7 |
|  | HLA-DR-DQ-DP | 104 | 11.3 |
|  | HLA-DR-DQ | 18 | 2 |
|  | HLA-DR-DP | 17 | 1.8 |
|  | HLA-A-DQ | 1 | 0.1 |
|  | HLA-A-DP | 41 | 4.4 |
|  | HLA-B-DQ | 1 | 0.1 |
|  | HLA-B-DP | 3 | 0.3 |
|  | HLA-C-DQ-DP | 11 | 1.2 |
|  | HLA-B-DQ-DP | 2 | 0.2 |
|  | HLA-A-DQ-DP | 1 | 0.1 |
| Mismatched$\mathrm{n}=234 \text { (25.4\%) }$ | HLA-C-DR | 6 | 0.7 |
|  | HLA-A-DR | 2 | 0.2 |
|  | HLA-C-DR-DQ | 11 | 1.2 |
|  | HLA-C-DR-DP | 14 | 1.5 |
|  | HLA-B-DR-DQ | 1 | 0.1 |
|  | HLA-A-B-C-DR-DQ-DP | 4 | 0.4 |
|  | HLA-A-B-C-DQ-DP | 2 | 0.2 |
|  | HLA-A-B-C-DP | 8 | 0.9 |
|  | HLA-A-B-C | 2 | 0.2 |
|  | HLA-A-B-DR-DQ-DP | 2 | 0.2 |
|  | HLA-A-C-DR-DQ-DP | 22 | 2.4 |
|  | HLA-B-C-DR-DQ-DP | 2 | 0.2 |
|  | HLA-A-C-DR-DQ | 3 | 0.3 |
|  | HLA-A-C-DR-DP | 5 | 0.5 |
|  | HLA-A-C-DQ-DP | 3 | 0.3 |
|  | HLA-B-C-DR-DP | 1 | 0.1 |
|  | HLA-B-C-DQ-DP | 1 | 0.1 |
|  | HLA-A-B-DP | 1 | 0.1 |
|  | HLA-A-C-DQ | 2 | 0.2 |
|  | HLA-A-C-DP | 22 | 2.4 |
|  | HLA-B-C-DP | 21 | 2.3 |
|  | HLA-A-B | 1 | 0.1 |
|  | HLA-A-C | 10 | 1.1 |
|  | HLA-B-C | 2 | 0.2 |
|  | HLA-A-DR-DQ-DP | 9 | 1 |
|  | HLA-B-DR_DQ_DP | 6 | 0.7 |
|  | HLA-C-DR-DQ-DP | 68 | 7.4 |
|  | HLA-A-DR-DQ | 2 | 0.2 |
|  | HLA-A-DR-DP | 1 | 0.1 |
| Total |  | 922 | 100 |

Table 5: Detailed HLA mismatch. All matches and mismatches are based on high-resolution allele typing of the HLA A, B, C, DRB1, DQB1 and DPB1 loci. The classification as outlined by NMDPICIBMTR (Weisdorf et al., 2008), which focuses on the HLA A, B, C and DRB1 loci only, without consideration of HLA DQB1 or DPB1. Matched $8 / 8$ refers to allele match at the loci HLA A, B, C and DRB1. Partially matched HSCT allows for one allele mismatch within this group, whereas mismatched HSCT has two or more allele mismatches within HLA A, B, C and DRB1. 340 HSCT pairs (36.9\%) had an HLA C mismatch.

| mismatch group | group total $\mathrm{n}=$ | $\%$ |
| :--- | ---: | ---: |
| 3 HLA I +3 HLA II | 4 | 0.433839 |
| 3 HLA I +2 HLA II | 2 | 0.21692 |
| 3 HLA I + HLA II | 8 | 0.867679 |
| 3 HLA I + 0 HLA II | 2 | 0.21692 |
| 2 HLA I +3 HLA II | 26 | 2.819957 |
| 2 HLA I +2 HLA II | 13 | 1.409978 |
| 2 HLA I +1 HLA II | 46 | 4.989154 |
| 2 HLA I +0 HLA II | 13 | 1.409978 |
| 1 HLA I +3 HLA II | 83 | 9.002169 |
| 1 HLA I +2 HLA II | 43 | 4.663774 |
| 1 HLA I +1 HLA II | 149 | 16.16052 |
| 1 HLA I +0 HLA II | 36 | 3.904555 |
| 3 HLA II | 104 | 11.27983 |
| 2 HLA II | 65 | 7.049892 |
| 1 HLA II | 250 | 27.11497 |
| fully matched | 78 | 8.45987 |
| Total | 922 | 100 |

Table 6: Summary of HLA mismatch by HLA class I and II.

|  | B |  | S.E. | Wald |  |  | df | Sig. | $\operatorname{Exp}(\mathrm{B})$ | 95\% C.I.for EXP(B) |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  |  |  |  |  |  | Lower | Upper |
| ALL v ANLL |  | 0.427 |  | 0.143 |  | 8.92683 | 1 | 0.0028101 | 1.533921 | 1.158557 | 2.0309 |
| Recipient age group <10y |  | -0.151 |  | 0.069 |  | 4.718277 | 1 | 0.0298436 | 0.859 | 0.748918 | 0.985262 |
| 8/8 HLA match |  | -0.303 |  | 0.145 |  | 4.324559 | 1 | 0.0375661 | 0.738402 | 0.55483 | 0.982711 |
| Relapse |  | 0.307 |  | 0.163 |  | 3.553663 | 1 | 0.0594142 | 1.360325 | 0.987856 | 1.873232 |
| ABO major mismatch |  | 0.134 |  | 0.079 |  | 2.859926 | 1 | 0.0908117 | 1.143999 | 0.978841 | 1.337022 |
| TBI given |  | -0.346 |  | 0.215 |  | 2.595402 | 1 | 0.1071743 | 0.706824 | 0.463431 | 1.078048 |
| Female to male transplant |  | -0.458 |  | 0.288 |  | 2.531509 | 1 | 0.1115935 | 0.632177 | 0.359336 | 1.112182 |
| Donor age > 30 |  | -0.229 |  | 0.159 |  | 2.060951 | 1 | 0.1511158 | 1.257433 | 0.919734 | 1.719125 |
| Recipient sex |  | 0.24 |  | 0.189 |  | 1.605145 | 1 | 0.2051755 | 1.271931 | 0.876715 | 1.845307 |
| CyA GVHD prophylaxis |  | 0.09855 |  | 0.138 |  | 0.505388 | 1 | 0.477142 | 1.103564 | 0.841017 | 1.448073 |
| High risk leukaemia |  | 0.797 |  | 1.166 |  | 0.467833 | 1 | 0.4939857 | 2.220418 | 0.225799 | 21.83475 |
| Non-Cy-TBI conditioning |  | 0.0796 |  | 0.156 |  | 0.259843 | 1 | 0.6102281 | 1.082876 | 0.797305 | 1.47073 |
| Donor age >40y |  | 0.024 |  | 1.442 |  | 0.000283 | 1 | 0.9865867 | 1.024548 | 0.060624 | 17.315 |
| Donor sex |  | -41.663 |  | 56841.8 |  | 5.37E-07 | 1 | 0.9994152 | 8.05E-19 | 0 |  |

Table 7: Multivariate analysis of risk factors for grade 2-4 acute GVHD in the finally chosen combined study cohort.

| Gene Symbol | status |
| :---: | :---: |
| 3.8-1 | included |
| 3.8-1.2 | included |
| 3.8-1.3 | included |
| 3.8-1.4 | included |
| 3.8-1.5 | included |
| A2M | excluded - no marker |
| A2ML1 | included |
| A4GALT | included |
| AATK | included |
| ABCA1 | included |
| ABCB1 | included |
| ABCC1 | included |
| ABCC11 | included |
| ABCC4 | included |
| ABCF1 | included |
| ABCG2 | included |
| ABO | included |
| ACACA | included |
| ACE | included |
| ACE2 | included |
| ACHE | included - new MS design |
| ACOT11 | included |
| ACOT8 | included |
| ACTA1 | included |
| ACTB | included |
| ACTC1 | included |
| ACTG1 | included |
| ACTL7B | included |
| ACTN1 | included |
| ACTN2 | included |
| ACTN3 | included |
| ADA | included |
| ADAM10 | included |
| ADAM12 | included |
| ADAM17 | included |
| ADAM8 | included |
| ADAMTS13 | included |
| ADAT2 | included |
| ADD1 | included |
| ADIPOQ | excluded - no marker |
| ADK | included |
| ADM | included |
| ADORA1 | included |
| ADORA2 | included |
| ADORA3 | included |
| AGER | included |
| AGPAT1 | included |
| AGPAT4 | included |
| AGPS | included |
| AGTR1 | included |
| AGTR2 | included |
| AGTRL1 | included |
| AICDA | included |
| AIF1 | included |
| AIFM1 | included |


| AIM2 | included |
| :---: | :---: |
| AIP | included |
| AIRE | included |
| AKAP12 | included |
| AKAP13 | included |
| AKAP7 | included |
| AKT1 | included |
| AKT2 | included |
| AKT3 | included |
| AKTIP | included - new MS design |
| ALAS2 | included |
| ALCAM | included |
| ALK | included |
| ALKBH1 | included |
| ALKBH2 | included |
| ALKBH3 | included |
| ALKBH4 | included |
| ALKBH5 | included |
| ALKBH6 | included |
| ALKBH7 | included |
| ALKBH8 | included |
| ALOX12 | included |
| ALOX12B | included |
| ALOX15 | included |
| ALOX15B | included |
| ALOX5 | included |
| ALOX5AP | included |
| ANGPT1 | included |
| ANGPT2 | included |
| ANGPTL1 | included |
| ANK1 | included |
| ANK2 | included |
| ANK3 | included |
| ANKDD1A | included |
| ANKRD6 | included |
| ANP32B | included |
| ANPEP | included |
| ANXA1 | included |
| ANXA2 | included |
| ANXA5 | included |
| APAF1 | included |
| APBB1IP | included |
| APC | included |
| APEX1 | included |
| API5 | included |
| APLN | included |
| APOL6 | included |
| APOM | included |
| APS | included |
| AQP1 | included |
| AR | included |
| AREG | included |
| ARG1 | included |
| ARHGDIA | included |
| ARHGDIB | included |
| ARID1B | included |


| ARL6IP5 | included |
| :---: | :---: |
| ARRB1 | included |
| ARRB2 | included |
| ART1 | included |
| ART4 | included |
| ARTN | included |
| ARTS1 | included |
| ASB1 | included |
| ASCL1 | included |
| ASPM | included |
| ATBF1 | included |
| ATF1 | included |
| ATF3 | included |
| ATF4 | included |
| ATF5 | included |
| ATG5 | included |
| ATM | included |
| ATP10A | included |
| ATP1B3 | included |
| ATP6V1G2 | included |
| ATXN1 | included |
| AVEN | included |
| AXIN1 | included |
| AXIN2 | included |
| AZGP1 | included |
| AZU1 | included |
| B2M | included |
| B3GALNT1 | included |
| B3GALT4 | included |
| B3GAT1 | included |
| B3GNT3 | included |
| BAALC | included |
| BAD | included |
| BAG1 | included |
| BAG2 | included |
| BAG3 | included |
| BAG4 | included |
| BAG5 | included |
| BAGE | included |
| BAGE2 | included |
| BAGE3 | included |
| BAGE4 | included |
| BAGE5 | included |
| BAI3 | included |
| BAIAP2L1 | included |
| BAK1 | included |
| BANK1 | included |
| BAT1 | included |
| BAT2 | included |
| BAT2, BAT2 GT, BAT2 included |  |
| BAT3 | included |
| BAT4 | included |
| BAT5 | included |
| BAX | included |
| BAZ1A | included |
| BBC3 | included |


| BCAM | included |
| :---: | :---: |
| BCAP31 | included |
| BCAS2 | included |
| BCKDHB | included |
| BCL10 | included |
| BCL2 | included |
| BCL2A1 | included |
| BCL2L1 | included |
| BCL2L10 | included |
| BCL2L11 | included |
| BCL2L12 | included |
| BCL2L13 | included |
| BCL2L14 | included |
| BCL2L2 | included |
| BCL3 | included |
| BCR | included |
| BDKRB1 | included |
| BDKRB2 | included |
| BDNF | included |
| BFAR | excluded - no marker |
| BGN | included |
| BID | included |
| BIK | included |
| BIRC2 | included |
| BIRC3 | included |
| BIRC4 | included |
| BIRC5 | included |
| BIRC6 | included |
| BIRC7 | included |
| BIRC8 | included |
| BLK | included |
| BLM | included |
| BLNK | included |
| BLR1 | included |
| BLVRB | included |
| BMI-1 | included |
| BMP2 | included |
| BMP3 | included |
| BMP4 | included |
| BMP5 | included |
| BMP6 | included |
| BMP7 | included |
| BMPR1A | included |
| BMPR1B | included |
| BMPR2 | included |
| BMX | included |
| BNIP1 | included |
| BNIP2 | included |
| BNIP3 | included |
| BNIP3L | included |
| BNIP3P | excluded - no marker |
| BNIPL | included |
| BOK | included |
| BPI | included |
| BRCA2 | included |
| BRD2 | included |


| BRD8 | included |
| :---: | :---: |
| BRDG1 | included |
| BRF1 | included |
| BSG | included |
| BST1 | included |
| BST2 | included |
| BTAF | included |
| BTBD9 | included |
| BTC | included |
| BTG1 | included |
| BTG3 | included |
| BTK | included |
| BTLA | included |
| BTN3A1 | included |
| BTNL2 | included |
| BTRC | included |
| BUB3 | included |
| BXDC1 | included |
| C10orf26 | included |
| C19orf10 | included |
| C1QA | included |
| C1QB | included |
| C1QBP | included |
| C1QG | included |
| C1QL1 | included |
| C1QL2 | included |
| C1QL3 | included |
| C1QL4 | included |
| C1QR | included |
| C1QTNF2 | included |
| C1QTNF3 | included |
| C1QTNF4 | included |
| C1QTNF5 | included |
| C1QTNF6 | included |
| C1QTNF7 | included |
| C1R | excluded - no marker |
| C1RL | included |
| C1S | excluded - no marker |
| C2 | excluded - no marker |
| C2orf47 | included |
| C3 | included |
| C3AR1 | included |
| C4A | excluded - no marker |
| C4B | excluded - no marker |
| C4BPA | included |
| C4BPB | included |
| C5 | included |
| C5R1 | included |
| C6 | included |
| C6orf10 | included |
| C6orf12 | included |
| C6orf123 | included |
| C6orf134 | included |
| C6orf136 | included |
| C6orf138 | included |
| C6orf15 | included |


| C6orf174,KIAA0408 | included |
| :---: | :---: |
| C6orf18 | included |
| C6orf204 | included |
| C6orf205 | included |
| C6orf21 | included |
| C6orf25 | included |
| C6orf27 | included |
| C6orf47 | included |
| C6orf48 | included |
| C6orf65 | included |
| C6orf91 | included |
| C7 | included |
| C8A | included |
| C8B | included |
| C8G | included |
| C9 | included |
| CABIN1 | included |
| CADM1 | included |
| CADM2 | included |
| CADM3 | included |
| CADM4 | included |
| CALR | included |
| CAMK2D | included |
| CAMK4 | included |
| CAMP | excluded - no marker |
| CANX | included |
| CARD10 | included |
| CARD11 | included |
| CARD14 | included |
| CARD6 | included |
| CARD8 | included |
| CARD9 | excluded - no marker |
| CARM1 | included |
| CASP1 | included |
| CASP10 | included |
| CASP12 | included |
| CASP14 | included |
| CASP2 | included |
| CASP3 | excluded - no marker |
| CASP4 | included |
| CASP5 | included |
| CASP7 | included |
| CASP8 | included |
| CASP8AP2 | included |
| CASP9 | included |
| CAT | included |
| CAV1 | included |
| CBFA2T2 | included |
| CBFB | included |
| CBL | excluded - no marker |
| CBLB | included |
| CCBP2 | included |
| CCL1 | included |
| CCL11 | included |
| CCL13 | included |
| CCL14 | included |


| CCL15 | included |
| :---: | :---: |
| CCL16 | included |
| CCL17 | included |
| CCL18 | included |
| CCL19 | included |
| CCL2 | included |
| CCL20 | included |
| CCL21 | included |
| CCL22 | included |
| CCL23 | included |
| CCL24 | included |
| CCL25 | included |
| CCL26 | included |
| CCL27 | included |
| CCL28 | included |
| CCL3 | included |
| CCL3L1 | excluded - no marker |
| CCL3L3 | excluded - no marker |
| CCL4 | included |
| CCL4L1 | excluded - no marker |
| CCL4L2 | excluded - no marker |
| CCL5 | included |
| CCL7 | included |
| CCL8 | included |
| CCNA1 | included |
| CCNA2 | included |
| CCNB1 | included |
| CCNB1IP1 | included |
| CCNB2 | included |
| CCNB3 | included |
| CCNC | excluded - no marker |
| CCND1 | included |
| CCND2 | included |
| CCND3 | included |
| CCNE1 | included |
| CCNE2 | included |
| CCNG1 | included |
| CCNH | included |
| CCR1 | included |
| CCR10 | included |
| CCR2 | included |
| CCR3 | included |
| CCR4 | included |
| CCR5 | included |
| CCR6 | included |
| CCR7 | included |
| CCR8 | included |
| CCR9 | included |
| CCRL1 | included |
| CCRL2 | included |
| CCRN4L | included |
| CD109 | included |
| CD139 | excluded - unknown location |
| CD14 | included |
| CD151 | included |
| CD160 | included |


| CD163 | excluded - no marker |
| :---: | :---: |
| CD164 | included |
| CD164L1 | included |
| CD177 | included |
| CD19 | excluded - no marker |
| CD1A | included |
| CD1B | included |
| CD1C | included |
| CD1D | included |
| CD1E | included |
| CD2 | included |
| CD200 | included |
| CD200R1 | included |
| CD200R2 | included |
| CD207 | included |
| CD209 | included |
| CD22 | included |
| CD24 | included |
| CD244 | included |
| CD245 | excluded - unknown location |
| CD274 | included |
| CD276 | included |
| CD28 | included |
| CD2AP | included |
| CD2BP2 | included |
| CD300A | included |
| CD300C | included |
| CD300E | included |
| CD300LB | included |
| CD300LF | included |
| CD302 | included |
| CD320 | excluded - no marker |
| CD33 | included |
| CD33L3 | included |
| CD34 | included |
| CD36 | included |
| CD37 | included |
| CD38 | included |
| CD3D | included |
| CD3E | included |
| CD3EAP | included |
| CD3G | included |
| CD3Z | included |
| CD4 | included |
| CD40LG | included |
| CD44 | included |
| CD47 | included |
| CD48 | included |
| CD5 | included |
| CD53 | included |
| CD55 | included |
| CD58 | included |
| CD59 | included |
| CD5L | included |
| CD6 | excluded - no marker |
| CD63 | included |


| CD65 | excluded - unknown location |
| :---: | :---: |
| CD68 | included |
| CD69 | included |
| CD7 | included |
| Cd72 | included |
| CD74 | included |
| CD79A | included |
| CD79B | included |
| CD80 | included |
| CD81 | included |
| CD82 | included |
| CD83 | included |
| CD84 | included |
| CD86 | included |
| CD8A | included |
| CD8B1 | included |
| CD8BP | excluded - no marker |
| CD9 | included |
| CD96 | included |
| CD97 | included |
| CD99 | excluded - no marker |
| CD99L2 | included |
| CDA | included |
| CDC2 | included |
| CDC20 | included |
| CDC25A | excluded - no marker |
| CDC25B | included |
| CDC37 | included |
| CDC42 | excluded - no marker |
| CDC42EP5 | included |
| CDCP1 | included |
| CDH1 | included |
| CDH2 | included |
| CDH5 | included |
| CDK10 | excluded - no marker |
| CDK2 | included |
| CDK3 | excluded - no marker |
| CDK4 | included |
| CDK5 | included |
| CDK6 | included |
| CDK7 | included |
| CDK8 | included |
| CDK9 | included |
| CDKAL1 | included |
| CDKN1A | included |
| CDKN1B | excluded - no marker |
| CDKN1C | included |
| CDKN2A | included |
| CDKN2B | included |
| CDKN2C | included |
| CDKN2D | included |
| CDKN3 | included |
| CDO1 | included |
| CDSN | included |
| CDw12 | excluded - unknown location |
| CDW52 | included |


| CDW93 | included |
| :---: | :---: |
| CDX2 | included |
| CEACAM1 | included |
| CEACAM3 | included |
| CEACAM5 | included |
| CEACAM6 | included |
| CEACAM8 | included |
| CEBPA | included |
| CEBPE | included |
| CEBPG | included |
| CENPF | included |
| CENPM | included |
| CERK | included |
| CES1 | included |
| CFB | excluded - no marker |
| CFD | included |
| CFDP1 | included |
| CFH | included |
| CFHR1 | included |
| CFHR2 | included |
| CFHR3 | included |
| CFHR4 | included |
| CFHR5 | included |
| CFI | included |
| CFL1 | included |
| CFL2 | included |
| CFLAR | included |
| CFP | included |
| CGA | included |
| CGB | included |
| CHAF1B | included |
| CHERP | included |
| CHES1 | included |
| CHI3L2 | included |
| CHLI | included |
| CHMP1B | included |
| CHMP2A | included |
| CHMP2B | included |
| CHMP4A | included |
| CHMP4B | included |
| CHMP4C | included |
| CHMP5 | included |
| CHMP6 | included |
| CHMP7 | included |
| CHRAC1 | included |
| CHUK | included |
| CIDEA | excluded - no marker |
| CIITA | included |
| CIR | included |
| CISH | excluded - no marker |
| CITED2 | included |
| CKLF | included |
| CKS1B | included |
| CKS2 | included |
| CLASP1 | included |
| CLASP2 | included |


| CLC | included |
| :---: | :---: |
| CLCF1 | included |
| CLDN23 | included |
| CLDN3 | included |
| CLEC12A | included |
| CLEC1A | included |
| CLEC1B | included |
| CLEC2B | included |
| CLEC3A | included |
| CLEC4a | included |
| CLEC4C | included |
| CLEC4D | included |
| CLEC4E | included |
| CLEC4M | included |
| CLEC5A | included |
| CLEC6A | excluded - no marker |
| CLEC7A | included |
| CLECSF10A | included |
| CLIC1 | included |
| CLIP1 | included |
| CLIP2 | included |
| CLK1 | included |
| CLU | included |
| CMKLR1 | included |
| CMTM1 | included |
| CMTM2 | included |
| CMTM3 | included |
| CMTM4 | included |
| CMTM5 | included |
| CMTM6 | included |
| CMTM7 | included |
| CMTM8 | included |
| CNTN2 | included |
| COL11A2 | included |
| COL14A1 | included |
| COL1A1 | included |
| COL1A2 | included |
| COL21A1 | included |
| COL3A1 | included |
| COL4A1 | included |
| COL4A2 | included |
| COL4A3 | included |
| COL4A3BP | included |
| COL4A4 | included |
| COL4A5 | included |
| COL4A6 | included |
| COL5A1 | included |
| COL5A2 | included |
| COL5A3 | included |
| COL6A1 | included |
| COL6A2 | included |
| COL6A3 | included |
| COLEC12 | included |
| COP1 | included |
| CORO1A | excluded - no marker |
| COX2 | included |


| CPE | included |
| :---: | :---: |
| CPS1 | included |
| CR1 | included |
| CR1L | included |
| CR2 | included |
| CRADD | included |
| CREB1 | included |
| CREBBP | included |
| CREBL1 | excluded - no marker |
| CREM | included |
| CRF1R | included |
| CRF2R | included |
| CRH | included |
| CRHR2 | included |
| CRKL | included |
| CRLF1 | included |
| CRLF2 | excluded - no marker |
| CRLF3 | included |
| CRP | included |
| CSDA | included |
| CSF1 | included |
| CSF1R | included |
| CSF2 | included |
| CSF2RA | excluded - no marker |
| CSF2RB | included |
| CSF3 | included |
| CSF3R | included |
| CSH1 | included |
| CSK | included |
| CSNK1A1 | included |
| CSNK2B | included |
| CSPG2 | included |
| CST1 | included |
| CTDSP2 | included |
| CTDSPL | included |
| CTF1 | included |
| CTGF | included |
| CTLA4 | included |
| CTNNB1 | included |
| CTNS | included |
| CTPS | included |
| CTSA | included |
| CTSB | included |
| CTSD | included |
| CTSF | included |
| CTSG | included |
| CTSH | included |
| CTSK | included |
| CTSL1 | included |
| CTSS | included |
| CTSZ | included |
| CUGBP2 | included |
| CUL4A | included |
| CX3CL1 | included |
| CX3CR1 | included |
| CXCL1 | included |


| CXCL10 | included |
| :---: | :---: |
| CXCL11 | included |
| CXCL12 | included |
| CXCL13 | included |
| CXCL14 | included |
| CXCL16 | included |
| CXCL2 | included |
| CXCL3 | included |
| CXCL5 | included |
| CXCL6 | included |
| CXCL9 | included |
| CXCR3 | included |
| CXCR4 | included |
| CXCR6 | included |
| CXCR7 | included |
| CYBA | included |
| CYBB | included |
| CYCS | included |
| CYP11A1 | included |
| CYP11B1 | included |
| CYP11B2 | included |
| CYP17A1 | included |
| CYP19A1 | included |
| CYP1A1 | included |
| CYP1A2 | included |
| CYP1B1 | included |
| CYP20A1 | included |
| CYP21A2 | excluded - no marker |
| CYP26A1 | included |
| CYP26B1 | included |
| CYP26C1 | included |
| CYP27A1 | included |
| CYP27B1 | included |
| CYP2A13 | included |
| CYP2A6 | included |
| CYP2A7 | included |
| CYP2B | included |
| CYP2B6 | included |
| CYP2C18 | included |
| CYP2C19 | included - new MS design |
| CYP2C8 | included |
| CYP2C9 | included - new MS design |
| CYP2D6 | included |
| CYP2E1 | included |
| CYP2F1 | included |
| CYP2J2 | included |
| CYP2R1 | included |
| CYP2S1 | included |
| CYP2U1 | included |
| CYP2W1 | included |
| CYP39A1 | included |
| CYP3A11 | included |
| CYP3A3 | included |
| CYP3A43 | included |
| CYP3A5 | included |
| CYP3A7 | included |


| CYP46A1 | included |
| :---: | :---: |
| CYP4B1 | included |
| CYP4F11 | included |
| CYP4F12 | included |
| CYP4F2 | included |
| CYP4F3 | included |
| CYP4F8 | included |
| CYP4V2 | included |
| CYP4X1 | included |
| CYP4Z1 | included |
| CYP51A1 | included |
| CYP7A1 | included |
| CYP7B1 | included |
| CYP8B1 | included |
| CYSLTR1 | included |
| CYSLTR2 | included |
| CYTL1 | included |
| D6S2723E | included |
| DAAM2 | included |
| DAD1 | included |
| DAG1 | included |
| DAP | included |
| DAPK1 | included |
| DAPK2 | included |
| DAPK3 | excluded - no marker |
| DARC | included |
| DAXX | included |
| DCAL1 | included |
| DCD | included |
| DCLRE1C | included |
| DCN | included |
| DCTN2 | included |
| DDAH1 | included |
| DDAH2 | included |
| DDR1 | included |
| DDX1 | included |
| DDX10 | included |
| DDX11 | included |
| DDX12 | included |
| DDX17 | included |
| DDX18 | included |
| DDX19A | included |
| DDX19B | included |
| DDX20 | included |
| DDX21 | included |
| DDX23 | included |
| DDX24 | included |
| DDX25 | included |
| DDX26B | included |
| DDX27 | excluded - no marker |
| DDX28 | included |
| DDX31 | included |
| DDX39 | included |
| DDX3X | included |
| DDX3Y | included - new MS design |
| DDX4 | included |


| DDX41 | included |
| :---: | :---: |
| DDX42 | included |
| DDX43 | included |
| DDX46 | included |
| DDX47 | included |
| DDX49 | included |
| DDX5 | included |
| DDX50 | included |
| DDX51 | excluded - no marker |
| DDX52 | included |
| DDX53 | included |
| DDX54 | included |
| DDX55 | included |
| DDX56 | included |
| DDX58 | included |
| DDX59 | included |
| DDX6 | included |
| DEDD | included |
| DEDD2 | included |
| DEFA1 | included |
| DEFA1A3 | included |
| DEFA3 | included |
| DEFA4 | included |
| DEFA5 | included |
| DEFA6 | included |
| DEFB1 | included |
| DEFB4 | excluded - no marker |
| DFB103A | excluded - no marker |
| DFB103B | excluded - no marker |
| DFB104A | excluded - no marker |
| DFB104B | excluded - no marker |
| DFB105A | excluded - no marker |
| DFB106A | excluded - no marker |
| DFB106B | excluded - no marker |
| DFB107A | excluded - no marker |
| DFB107B | included |
| DFB108B | included |
| DFB110 | included |
| DFB111 | included |
| DFB112 | included |
| DFB113 | included |
| DFB114 | included |
| DFB115 | included |
| DFB116 | included |
| DFB118 | excluded - no marker |
| DFB119 | excluded - no marker |
| DFB121 | excluded - no marker |
| DFB122 | included |
| DFB123 | included |
| DFB124 | included |
| DFB125 | included |
| DFB126 | included |
| DFB127 | included |
| DFB128 | included |
| DFB129 | included |
| DFB130 | included |


| DFB131 | included |
| :---: | :---: |
| DFB133 | excluded - unknown location |
| DFB134 | included |
| DFB136 | included |
| DFB137 | included |
| DFFA | included |
| DFFB | included |
| DGCR2 | included |
| DGK alpha | included |
| DGKB | included |
| DHFR | included |
| DHH | included |
| DHX15 | included |
| DHX16 | included |
| DHX29 | included |
| DHX30 | included |
| DHX32 | included |
| DHX33 | included |
| DHX34 | included |
| DHX35 | included |
| DHX36 | included |
| DHX37 | included |
| DHX38 | included |
| DHX40 | included |
| DHX57 | included |
| DHX8 | included |
| DHX9 | included |
| DIABLO | included |
| DIAPH2 | included |
| DIDO1 | included |
| DIP | included |
| DKC1 | included |
| DLG5 | included |
| DLL1 | included |
| DLL4 | included |
| DLX3 | included |
| DMBT1 | included |
| DNAH8 | included |
| DNAM1 | included |
| DNASE1 | included |
| DNM2 | included |
| DNTT | included |
| DOCK2 | included |
| DOK1 | included |
| DOK2 | included |
| DOM3Z | excluded - no marker |
| DPCR1 | included |
| DPP4 | included |
| DQX1 | included |
| DRG1 | included |
| DRG2 | included |
| DSCAM | included |
| DSP | included |
| DSS1 | included |
| DTX1 | included |
| DUSP1 | included |


| DUSP10 | included |
| :---: | :---: |
| DUSP11 | included |
| DUSP12 | included |
| DUSP13 | included |
| DUSP14 | included |
| DUSP15 | included |
| DUSP16 | included |
| DUSP18 | included |
| DUSP19 | included |
| DUSP2 | included |
| DUSP21 | included |
| DUSP22 | included |
| DUSP23 | included |
| DUSP26 | included |
| DUSP28 | included |
| DUSP3 | included |
| DUSP4 | included |
| DUSP5 | included |
| DUSP6 | included |
| DUSP7 | included |
| DUSP8 | excluded - no marker |
| DUSP9 | included |
| DVL1 | included |
| DYRK1A | included |
| DYRK1B | included |
| DYRK2 | included |
| DYRK3 | included |
| DYRK4 | included |
| E2F4 | included |
| EBF | included |
| EBF2 | included |
| EBI2 | included |
| EBI3 | included |
| ECGF1 | included |
| ECSIT | included |
| EDA | included |
| EDAR | included |
| EDARADD | included |
| EDG1 | included |
| EDN1 | included |
| EDN2 | excluded - no marker |
| EDN3 | included |
| EEA1 | included |
| EEF1A1 | included |
| EEF1A2 | included |
| EEF2 | included |
| EFHC1 | included |
| EFNA1 | included |
| EGF1 | included |
| EGFL11 | included |
| EGFL8 | included |
| EGFR | included |
| EGFTM7 | included |
| EGR1 | included |
| EGR2 | included |
| EGR3 | included |


| EHMT2 | excluded - no marker |
| :---: | :---: |
| EIF4A3 | included |
| EIF4G2 | excluded - no marker |
| ELA1 | included |
| ELA2 | included |
| ELA2A | included |
| ELA2B | included |
| ELA3A | excluded - no marker |
| ELA3B | excluded - no marker |
| ELF3 | included |
| ELK4 | included |
| ELMO1 | included |
| EMP3 | included |
| EMR1 | included |
| EMR2 | included |
| EMR3 | included |
| ENC1 | included |
| ENG | included |
| ENPEP | included |
| ENPP3 | included |
| ENSG00000179038 | included |
| ENSG00000204345 | included |
| ENTPD1 | included |
| EOMES | included |
| EP300 | included |
| EPB41L2 | included |
| EPHA1 | included |
| EPHA7 | included |
| EPHB1 | included |
| EPHB2 | included |
| EPHX1 | included |
| EPO | included |
| EPOR | included |
| EPX | included |
| ERBB2 | included |
| ERC2 | included |
| ERCC1 | included |
| ERCC2 | included |
| ERCC5 | included |
| EREG | included |
| ERG | included |
| ERGIC2 | included |
| ERMAP | included |
| ESR1 | included |
| ESR2 | included |
| ESRRA | included |
| ESRRB | included |
| ESRRG | included |
| ETF1P1 | included |
| ETV1 | included |
| ETV6 | included |
| EVI1 | included |
| EXO1 | included |
| EZH2 | included |
| F2 | included |
| F2R | included |


| F2RL1 | included |
| :---: | :---: |
| F2RL2 | included |
| F3 | included |
| F5 | included |
| F8 | included |
| FABP3 | included |
| FABP4 | included |
| FABP5 | included |
| FADD | included |
| FAF1 | included |
| FAIM3 | included |
| FAM120B | included |
| FAT10 | included |
| FBXW7 | included |
| FCAMR | included |
| FCAR | included |
| FCER1A | included |
| FCER1G | included |
| FCER2 | included |
| FCGR1A | included |
| FCGR1B | included - new MS design |
| FCGR1C | excluded - unknown location |
| FCGR2A | included - new MS design |
| FCGR2B | included |
| FCGR2C | included |
| FCGR3A | included |
| FCGR3B | included |
| FCGRT | included - new MS design |
| FCN1 | included |
| FCN2 | included |
| FCN3 | excluded - no marker |
| FCRL1 | included |
| FCRL2 | included |
| FCRL3 | included |
| FCRL4 | included |
| FCRL5 | included |
| FCRL6 | included |
| FCRLA | included |
| FCRLB | included |
| FEEL-2 | included |
| FEN1 | included |
| FES | included |
| FGA | included |
| FGB | included |
| FGC | included |
| FGF1 | included |
| FGF10 | included |
| FGF11 | included |
| FGF12 | included |
| FGF13 | included |
| FGF14 | included |
| FGF16 | included |
| FGF17 | included |
| FGF18 | excluded - no marker |
| FGF19 | included |
| FGF2 | included |


| FGF20 | included |
| :---: | :---: |
| FGF21 | included |
| FGF22 | included |
| FGF23 | included |
| FGF3 | included |
| FGF4 | included |
| FGF5 | included |
| FGF6 | included |
| FGF7 | included |
| FGF8 | included |
| FGF9 | included |
| FGFR1 | included |
| FGFR2 | included |
| FGFR3 | included |
| FGFR4 | included |
| FGG | included |
| FGL2 | included |
| FGR | included |
| FIGF | included |
| FKBP10 | included |
| FKBP11 | included |
| FKBP14 | excluded - no marker |
| FKBP15 | included |
| FKBP1A | included |
| FKBP1AC | included |
| FKBP1B | included |
| FKBP2 | included |
| FKBP3 | included |
| FKBP4 | included |
| FKBP5 | included |
| FKBP8 | included |
| FKBP9 | included |
| FKBP9L | included |
| FKBPL | included |
| FLJ20105 | included |
| FLJ43763 | included |
| FLJ45422 | included |
| FLJ46831 | included |
| FLOT1 | included |
| FLT1 | included |
| FLT3 | included |
| FLT3LG | included |
| FLT4 | included |
| FMOD | included |
| FN1 | included |
| FOS | included |
| FOSB | included |
| FOSL1 | included |
| FOXA1 | included |
| FOXA2 | included |
| FOXA3 | included |
| FOXB1 | included |
| FOXB2 | included |
| FOXC1 | included |
| FOXC2 | included |
| FOXD1 | included |


| FOXD2 | included |
| :---: | :---: |
| FOXD3 | included |
| FOXD4 | excluded - no marker |
| FOXE1 | included |
| FOXE3 | included |
| FOXF1 | included |
| FOXF2 | included |
| FOXG1B | included |
| FOXG1C | included |
| FOXH1 | excluded - no marker |
| FOXI2 | included |
| FOXJ1 | included |
| FOXJ2 | included |
| FOXJ3 | included |
| FOXK1 | included |
| FOXK2 | included |
| FOXL1 | included |
| FOXL2 | included |
| FOXN1 | included |
| FOXN2 | included |
| FOXN4 | included |
| FOXO1A | included |
| F0X01B | included |
| FOXO3A | included |
| FOXO3B | included |
| FOXP1 | included |
| FOXP2 | included |
| FOXP3 | included |
| FOXP4 | included |
| FOXQ1 | included |
| FOXR1 | included |
| FOXR2 | excluded - no marker |
| FPR1 | included |
| FPRL1 | included |
| FRAP1 | included |
| FRK | included |
| FRZB | included |
| FSHR | included |
| FURIN | included |
| FUT1 | included |
| FUT3 | included |
| FUT4 | included |
| FYB | included |
| FYN | included |
| FZD10 | included |
| FZD4 | included |
| FZD9 | included |
| G6PD | included |
| GAB2 | included |
| GADD45B | included |
| GADD45G | included |
| GALC | included |
| GAPDH | included |
| GAS1 | included |
| GAS2 | included |
| GATA1 | included |


| GATA2 | included |
| :---: | :---: |
| GATA3 | included |
| GATA4 | included |
| GATA5 | included |
| GATA6 | included |
| GBA | excluded - no marker |
| GBP1 | included |
| GBP2 | included |
| GBP3 | included |
| GBP5 | included |
| GCA | included |
| GCG | included |
| GCK | included |
| GCLC | included |
| GCLM | included |
| GCNT2 | included |
| GDF15 | included |
| GEM | included |
| GFRAL | included |
| GGT1 | included |
| GGT2 | included |
| GH1 | included |
| GH2 | included |
| GHR | included |
| GINS2 | included |
| GLI1 | included |
| GLI2 | included |
| GLI3 | included |
| GMDS | included |
| GNA13 | included |
| GNAI1 | included |
| GNAI2 | included |
| GNAI3 | included |
| GNAL | included |
| GNB3 | included |
| GNL1 | included |
| GNL2 | included |
| GNLY | included |
| GNRH1 | included |
| GNRHR | included |
| GP1BA | included |
| GP1BB | included |
| GP5 | included |
| GP9 | included |
| GPATCH3 | included |
| GPHA2 | included |
| GPHB5 | included |
| GPNMB | included |
| GPR107 | included |
| GPR109B | included |
| GPR132 | included |
| GPR4 | included |
| GPR44 | included |
| GPR56 | included |
| GPR65 | included |
| GPR68 | included |


| GPS2 | included |
| :---: | :---: |
| GPSM3 | included |
| GPX1 | included |
| GPX2 | included |
| GPX3 | included |
| GPX4 | excluded - no marker |
| GRAIL | included |
| GRAP2 | included |
| GRB10 | included |
| GRB2 | included |
| GRIK1 | included |
| GRIK2 | included |
| GSK3A | included |
| GSK3B | included |
| GSR | included |
| GSST1 | included |
| GSTA1 | included |
| GSTM1 | included |
| GSTP1 | included |
| GTF2A1 | included |
| GTF2A2 | included |
| GTF2B | included |
| GTF2E1 | included |
| GTF2E2 | included |
| GTF2F1 | included |
| GTF2F2 | included |
| GTF2F2L | included |
| GTF2H1 | included |
| GTF2H2 | excluded - no marker |
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| GTF2H4 | included |
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| GTF2IRD2B | included |
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| GTF3C3 | included |
| GTF3C4 | included |
| GTF3C5 | included |
| GTF3C6 | included |
| GUSB | included |
| GYPA | included |
| GYPB | included |
| GYPC | included |
| GYPE | excluded - no marker |
| GZMA | included |
| GZMB | included - new MS design |
| GZMH | excluded - no marker |
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| GZMM | included |
| H2AFX | included |
| H2AFZ | included |
| HACE1 | included |


| HAMP | included |
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| HAVCR2 | included |
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| HCG4P8 | included |
| HCG4P9 | excluded - unknown location |
| HCG5P8 | included |
| HCG8 | excluded - unknown location |
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| HCG9P5 | included |
| HCGVIII-2 | included |
| HCK | included |
| HCP5 | included |
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| HDAC4 | included |
| HDAC5 | included |
| HDAC6 | included |
| HDAC7A | included |
| HDAC8 | included |
| HDAC9 | included |
| HDC | included |
| HDGFL1 | included |
| HECA | included |


| HERC5 | included |
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| HEXA | included |
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| HGF | included |
| HISPPD2A | included |
| HIST2H2AA4 | excluded - no marker |
| HIST2H4A | excluded - no marker |
| HIST3H2A | included |
| HIVEP2 | included |
| HLA-16 | included |
| HLA-21 | included |
| HLA-75 | included |
| HLA-80 | included |
| HLA-90 | included |
| HLA-A | included |
| HLA-B | excluded - no marker |
| HLABC-CA | included |
| HLA-C | excluded - no marker |
| HLA-DMA | included |
| HLA-DMB | included |
| HLA-DOA | included |
| HLA-DOB | included |
| HLA-DPA1 | included |
| HLA-DPA3 | excluded - unknown location |
| HLA-DPB1 | included |
| HLA-DPB2 | included |
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| HLA-DQA2 | included |
| HLA-DQB1 | included - new MS design |
| HLA-DQB2 | included |
| HLA-DQB3 | excluded - unknown location |
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| HLA-DRB1 | included |
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| HLA-DRB9 | included |
| HLA-E | included |
| HLA-F | included |
| HLA-G | included |
| HLA-H | included |
| HLA-J | included |
| HLA-K | included |
| HLA-L | included |
| HLA-N | excluded - unknown location |
| HLA-S | excluded - unknown location |
| HLA-X | excluded - unknown location |
| HLA-Z | excluded - unknown location |
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| HLTF | included |
| HLX1 | included |
| HM13 | included |
| HMGB1 | included |


| HMGCR | included |
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| HMHB1 | included |
| HMMR | included |
| HMOX1 | included |
| HMOX2 | included |
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| HNF4A | included |
| HNF4G | included |
| HNMT | included |
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| HOXB4 | included |
| HOXC11 | included |
| HOXD10 | included |
| HPA | included |
| HPGD | included |
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| HRH1 | included |
| HRH2 | included |
| HRH3 | included |
| HRH4 | included |
| HRK | included |
| HS3ST1 | included |
| HSCT | included |
| HSD17B8 | included |
| HSD3B1 | included |
| HSH2D | included |
| HSP B8 | included |
| HSP90AA1 | included |
| HSP90AB1 | included |
| HSP90B1 | included |
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| HSPA1B | included |
| HSPA1L | included |
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| HSPA5 | included |
| HSPB1 | included |
| HSPD1 | included |
| HSPG2 | included |
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| HTR1B | included |
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| HTR2B | included |
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| IBD2 | included |
| IBD3 | included |
| IBD5 | included |
| ICAM1 | included |
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| ICAM4 | included |
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| ICEBERG | included |
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| IFI16 | included |
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| IFI30 | included |
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| IFIT1L | included |
| IFIT2 | included |
| IFIT3 | included |
| IFIT5 | included |
| IFITM1 | included |
| IFITM4P | included |
| IFNA1 | included |
| IFNA10 | included |
| IFNA13 | included |
| IFNA14 | included |
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| IFNA2 | included |
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| IFNA7 | included |
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| IFNB1 | included |
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| IFNE1 | included |
| IFNG | included - new MS design |
| IFNGR1 | included |
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| IFRD1 | included |
| IFRG28 | included |
| IGF1 | included |
| IGF1R | included |
| IGF2 | included |
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| IGFBP2 | included |
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| IGJ | included |
| IGKV1-12 | included |
| IGLL1 | included |
| IGSF1 | included |
| IGSF10 | included |
| IGSF11 | included |
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| IGSF22 | included |
| IGSF3 | included |
| IGSF5 | included |
| IGSF6 | included |
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| IGSF9B | included |
| IHH | included |
| IKBKAP | included |
| IKBKB | included |
| IKBKE | included |
| IKBKG | included |
| IKZF1 | included |
| IL10 | included - new MS design |
| IL10RA | included |
| IL10RB | included |
| IL11 | included |
| IL11RA | included |
| IL12A | included |
| IL12B | included |
| IL12RB1 | included |
| IL12RB2 | included |
| IL13 | included |
| IL13RA1 | included |
| IL13RA2 | included |
| IL15 | included |
| IL15RA | included |
| IL15RB | excluded - unknown location |
| IL16 | included |
| IL17A | included |
| IL17B | included |
| IL17C | included |
| IL17D | included |
| IL17F | included |
| IL17RA | included |
| IL17RB | included |
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| IL17RD | included |
| IL17RE | included |
| IL18 | included |
| IL18BP | included |
| IL18R1 | included |
| IL18RAP | included |
| IL19 | included |
| IL1A | included |
| IL1B | included - new MS design |
| IL1F10 | included |
| IL1F5 | included |
| IL1F7 | included - new MS design |
| IL1F8 | included |
| IL1F9 | included |
| IL1R1 | included |
| IL1R2 | included |
| IL1RAP | included |
| IL1RAPL1 | included |


| IL1RAPL2 | included |
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| IL1RL2 | included |
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| IL20RA | included |
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| IL21 | included |
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| IL23A | included |
| IL23R | included |
| IL24 | included |
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| IL26 | included |
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| IL2RB | included |
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| IL3 | included |
| IL31 | included |
| IL31RA | included |
| IL32 | included |
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| IL4 | included |
| IL4I1 | included |
| IL4R | included |
| IL5 | included |
| IL5RA | included |
| IL6 | included |
| IL6R | included - new MS design |
| IL6RL1 | included |
| IL6ST | included |
| IL6STP | included |
| IL7 | included |
| IL7R | included |
| IL8 | included - new MS design |
| IL8RA | included |
| IL8RB | included |
| IL9 | included |
| IL9R | excluded - no marker |
| ILF2 | included |
| ILF3 | included |
| ILK | included |
| INCA | included |
| INDO | included |
| INHA | included |


| INHBA | included |
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| INHBB | included |
| INHBC | included |
| INHBE | included |
| INPP5D | included |
| INSR | included |
| IRAK1 | included |
| IRAK1BP1 | included |
| IRAK2 | included |
| IRAK3 | included |
| IRAK4 | included |
| IRF1 | included |
| IRF2 | included |
| IRF3 | included |
| IRF4 | included |
| IRF5 | included |
| IRF6 | included |
| IRF7 | excluded - no marker |
| IRF8 | included |
| IRGC | included |
| IRGM | included |
| ISG15 | included |
| ISG20 | included |
| ISGF3G | included |
| ITCH | included |
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| ITGA10 | included |
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| ITGA9 | included |
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| ITGAM | included - new MS design |
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| ITGAX | included |
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| ITGB2 | included |
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| ITK | included |


| ITPKB | included |
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| JUN | included |
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| JUND | included |
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| KDR | included |
| KEAP1 | included |
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| KHDRBS1 | included |
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| KIAA0020 | included |
| KIAA1949 | included |
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| KIR2DL1 | included |
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| KIR3DL2 | included |
| KIR3DL3 | included |
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| KIR3DS1 | included |
| KIR3DX1 | included |
| KIT | included |
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| KLF11 | included |
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| KLHL32 | included |
| KLRA1 | included |
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| KLRD1 | included |
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| KLRG2 | included |
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| KRT15 | included |
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| KRT34 | included |
| KRT35 | included |
| KRT5 | included |
| KRT6A | included |
| KRT8 | included |
| L1CAM | included |
| LAG3 | included |
| LAIR1 | included |
| LAIR2 | included |
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| LAMA4 | included |
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| LAMC1 | excluded - no marker |
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| LAMC3 | included |
| LAMP1 | included |
| LAMP2 | included |
| LAMP3 | included |
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| LAT2 | included |
| LATS2 | included |
| LAX1 | included |
| LBP | included |
| LCK | included |
| LCN2 | included |
| LCP2 | included |
| LCT | included |
| LDLR | included |
| LEAP2 | included |
| LECT1 | included |
| LECT2 | included |
| LEDGF | included |
| LENG8 | included |
| LEP | included |
| LEPR | included |
| LGALS1 | excluded - no marker |
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| LGALS14 | included |
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| LGALS2 | included |
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| LGALS3BP | included |
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| LGALS7 | included |
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| LGMN | included |
| LGP2 | included |
| LHB | included |
| LHCGR | included |
| LHFPL2 | included |
| LIF | included |
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| LOC441792 | included |
| LOC643962 | included |
| LOC645740 | included |
| LOC646702 | included |
| LOC728195 | excluded - no marker |
| LPC2 | included |
| LPO | excluded - no marker |
| LRDD | excluded - no marker |
| LRP1 | excluded - no marker |
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| LRRC23 | included |
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| LST1 | included |
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| LTA4H | included |
| LTB | included |
| LTB4DH | included |


| LTB4R2 | included |
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| LTBP3 | included |
| LTBR | included |
| LTC4S | included |
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| LY6G5B | included |
| LY6G5C | included |
| LY6G6C | included |
| LY6G6D | included |
| LY6G6E | included |
| LY75 | included |
| LY86 | included |
| LY9 | included |
| LY96 | included |
| LYG2 | included |
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| LYN | included |
| LYPLA2P1 | included |
| LYST | included |
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| MAD2LI | included |
| MADCAM1 | included |
| MADD | included |
| MAF | included |
| MALT1 | included |
| MAML2 | included |
| MAN1A1 | included |
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| MAP2K2 | included |
| MAP2K3 | included |
| MAP2K4 | included |
| MAP2K5 | included |
| MAP2K6 | included |
| MAP2K7 | included |
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| MAP3K11 | included |
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| MAP3K2 | included |
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| MAP3K7 | included |
| MAP3K7IP1 | included |
| MAP3K7IP2 | included |
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| MAP4K1 | included |
| MAP4K2 | included |


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| MARCH7 | included |
| MARCO | included |
| MASP1 | included |
| MASP2 | included |
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| MC3R | included |
| MC4R | included |
| MC5R | included |
| MCAM | included |
| MCCD1 | included |
| MCL1 | included |
| MCM2 | included |
| MCM6 | included |
| MCP | included |
| MCRS1 | included |
| MDC1 | included |
| MDFIC | included |
| MDM2 | included |
| MDM4 | included |
| MEF2D | included |
| MEFV | included |
| MELK | included |
| MERTK | included |
| MET | included |
| MFI2 | included |
| MGA | included |
| MGMT | included |
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| MICB | included |
| MICC | included |
| MICD | included |
| MICE | included |


| MICF | included |
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| MICG | included |
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| MITF | included |
| MKLN1 | included |
| MLLT7 | included |
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| MMP21 | included |
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| MMP23B | included |
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| MMP26 | included |
| MMP27 | included |
| MMP28 | included |
| MMP3 | included |
| MMP7 | included |
| MMP8 | included |
| MMP9 | included |
| MN1 | included |
| MOAP1 | included |
| MOG | included |
| MOXD1 | included |
| MPG | included |
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| MPO | included - new MS design |
| MPS1 | included |
| MPZL1 | included |
| MR1 | included |
| MRC1 | included |
| MRC1L1 | included |
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| MRE11A | included |
| MRPL28 | included |
| MRPS18B | included |
| MS4A1 | included |
| MS4A3 | included |
| MS4A5 | included |
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| MT1F | included |
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| MT1G | included |
| MT1X | included |
| MT2A | included |
| MT3 | included |
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| MTMR11 | included |
| MTRR | included |
| MUC1 | included |
| MVP | included |
| MX1 | included |
| MX2 | included |
| MXD1 | included |
| MYC | included |
| MYCL1 | included |
| MYCN | included |
| MyD88 | included - new MS design |
| MYH11 | included |
| MYH2 | included |
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| MYO1G | included |
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| NCAPH | included |
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| NCOA7 | included |
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| NCR3 | included |
| NDUFA2 | included |
| NDUFS3 | included |
| NEDD9 | included |
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| NFATC2 | included |
| NFATC2IP | excluded - no marker |
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| NFE2L1 | included |
| NFIL3 | included |


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| NFKBIB | included |
| NFKBIE | included |
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| NFYA | included |
| NFYB | included |
| NFYC | included |
| NGFB | included |
| NGFR | included |
| NHLH1 | included |
| NID1 | included |
| NKAIN2 | included |
| NKIRAS1 | included |
| NKTR | included |
| NLRC3 | included |
| NLRC4 | included |
| NLRC5 | included |
| NLRP1 | included |
| NLRP10 | included |
| NLRP11 | included |
| NLRP12 | included |
| NLRP13 | included |
| NLRP14 | included |
| NLRP2 | included |
| NLRP2P | included |
| NLRP3 | included |
| NLRP3P | included |
| NLRP4 | included |
| NLRP5 | included |
| NLRP6 | included |
| NLRP7 | included |
| NLRP8 | included |
| NLRP9 | included |
| NLRP9P | included |
| NLRX1 | excluded - no marker |
| NME1 | included |
| NMI | included |
| NOD1 | included |
| NOD2 | included |
| NOL3 | included |
| NOS1 | included |
| NOS2A | included |
| NOS3 | included |
| NOSIP | included |
| NOTCH1 | included |
| NOTCH2 | included |
| NOTCH3 | included |
| NOTCH4 | included - new MS design |
| NOV | included |
| NOX1 | included |
| NOX3 | included |
| NOX4 | included |


| NOXA1 | included - new MS design |
| :---: | :---: |
| NOXO1 | included |
| NPM1 | excluded - no marker |
| NPPA | included |
| NPPB | included |
| NPTN | included |
| NPY | included |
| NPY2R | included |
| NQO1 | included |
| NR0B1 | included |
| NR0B2 | included |
| NR1D1 | included |
| NR1D2 | included |
| NR1H2 | included |
| NR1H3 | included |
| NR1H4 | included |
| NR112 | included |
| NR113 | included |
| NR2C1 | included |
| NR2C2 | included |
| NR2D1 | included |
| NR2E1 | included |
| NR2E3 | included |
| NR2F1 | included |
| NR2F2 | included |
| NR2F6 | included |
| NR3C1 | included |
| NR3C2 | included |
| NR4A1 | excluded - no marker |
| NR4A2 | included |
| NR4A3 | included |
| NR5A1 | included |
| NR5A2 | included |
| NR6A1 | included |
| NRAS | included |
| NRF1 | included |
| NRG2 | included |
| NRG3 | included |
| NRG4 | included |
| NRM | included |
| NRP1 | included |
| NSMAF | included |
| NT5E | included |
| NUMA1 | included |
| NUSAP1 | included |
| NXF1 | included |
| OAS1 | included |
| OAS2 | excluded - no marker |
| OAS3 | included |
| OASL | included |
| OFCC1 | included |
| OLIG2 | included |
| OLR1 | included |
| OPMR1 | included |
| OPRK1 | included |
| OPRL1 | included |


| ORM1 | included |
| :---: | :---: |
| ORM2 | included |
| OSCAR | included |
| OSM | included |
| P2RX1 | included |
| P2RX2 | included |
| P2RX3 | included |
| P2RX4 | included |
| P2RX5 | included |
| P2RX6 | included |
| P2RX7 | included |
| P2RY1 | included |
| P2RY11 | included |
| P2RY13 | included |
| P2RY14 | included |
| P2RY2 | included |
| P2RY4 | included |
| P2RY5 | included |
| P2RY6 | included |
| P5-04 | included |
| P5-05 | included |
| P5-07 | included |
| P5-09 | included |
| P5-11 | included |
| PACRG | included |
| PACSIN1 | included |
| PADI4 | included |
| PAFAH1B1 | included |
| PAFAH1B2 | excluded - no marker |
| PAFAH1B3 | included |
| PAFAH2 | included |
| PAG1 | included |
| PAK1 | included |
| PAK2 | included |
| PAK3 | included |
| PARK2 | included |
| PARP1 | included |
| PARVG | included |
| PAWR | included |
| PBEF1 | included |
| PBX2 | included |
| PCDHB16 | included |
| PCDHB5 | included |
| PCMT1 | included |
| PCOLN3 | excluded - no marker |
| PDCD1 | included |
| PDCD10 | included |
| PDCD1LG2 | included |
| PDCD2 | included |
| PDCD5 | included |
| PDCD6 | included |
| PDCD6IP | included |
| PDCD7 | included |
| PDE10A | included |
| PDE4A | included |
| PDE4B | included |


| PDE4C | included |
| :---: | :---: |
| PDE4D | included |
| PDGFA | excluded - no marker |
| PDGFB | included |
| PDGFRA | included |
| PDGFRB | included |
| PDIA3 | included - new MS design |
| PDK1 | included |
| PDK2 | included |
| PDRG1 | included |
| PDXK | included |
| PEA15 | excluded - no marker |
| PECAM1 | included |
| PELI1 | included |
| PELI2 | included |
| PELI3 | included |
| PEX6 | included |
| PF4 | included |
| PF4V1 | included |
| PFDN6 | included |
| PFN1 | included |
| PGDS | included |
| PGLYRP1 | included |
| PGLYRP2 | included |
| PGLYRP3 | included |
| PGLYRP4 | included |
| PGR | included |
| PHACTR1 | included |
| PHACTR2 | included |
| PHLDA2 | excluded - no marker |
| PI3 | included |
| PI4K2B | included |
| PIAS4 | included |
| PIGF | included |
| PIK3C2A | included |
| PIK3C2B | included |
| PIK3C2G | included |
| PIK3C3 | included |
| PIK3CA | included |
| PIK3CB | included |
| PIK3CD | included |
| PIK3CG | included |
| PIK3R1 | included |
| PIK3R2 | included |
| PIK3R3 | included |
| PIK3R4 | included |
| PIK3R5 | included |
| PIK4CA | excluded - no marker |
| PIK4CB | included |
| PILB | included |
| PILRA | included |
| PIM1 | included |
| PIN1 | included |
| PINX1 | included |
| PKD1 | included |
| PKD2 | included |


| PKD3 | included |
| :---: | :---: |
| PKHD1 | included |
| PLA1A | included |
| PLA2G10 | excluded - no marker |
| PLA2G2D | included |
| PLA2G6 | included |
| PLA2G7 | included |
| PLA2R1 | included |
| PLAA | included |
| PLAU | included |
| PLAUR | included |
| PLCB2 | included |
| PLCB3 | included |
| PLCG1 | included |
| PLCL2 | included |
| PLEC1 | excluded - no marker |
| PLEKHB1 | included |
| PLEKHH2 | included |
| PLK1 | included |
| PLK3 | included |
| PLK4 | included |
| PLXNA1 | included |
| PLXNB1 | included |
| PLXNC1 | included |
| PLXND1 | included |
| PMS2L3 | included |
| POLA2 | included |
| POLD3 | included |
| POLDIP3 | included |
| POLE3 | included |
| POMC | included |
| POP1 | included |
| POT1 | included |
| POU2AF1 | included |
| POU2F3 | included |
| POU4F1 | included |
| POU5F1 | included |
| PPAP2B | included |
| PPARA | included |
| PPARD | included |
| PPARG | included |
| PPBP | included |
| PPIA | included |
| PPIAL | included |
| PPIAL4 | included |
| PPIB | included |
| PPIC | included |
| PPID | included |
| PPIE | included |
| PPIF | included |
| PPIG | included |
| PPIH | included |
| PPIL1 | included |
| PPIL2 | included |
| PPIL3 | included |
| PPIL4 | included |


| PPIL5 | included |
| :---: | :---: |
| PPIL6 | included |
| PPIP9 | included |
| PPP1R10 | included |
| PPP1R11 | included |
| PPP1R16A | included |
| PPP1R16B | excluded - no marker |
| PPP1R2P1 | included |
| PPP2R4 | included |
| PPP3CA | included |
| PPP3CB | included |
| PPP3CC | included |
| PPP3R1 | included |
| PPP3R2 | included |
| PPT2 | included |
| PRAME | included |
| PRDM1 | included |
| PRDX4 | included |
| PRDX6 | included |
| PRF1 | included |
| PRG1 | included |
| PRG2 | included |
| PRG4 | included |
| PRKAA2 | included |
| PRKACA | included |
| PRKACB | included |
| PRKACG | included |
| PRKCA | included |
| PRKCB1 | included |
| PRKCD | included |
| PRKCG | included |
| PRKCH | included |
| PRKCI | included |
| PRKCQ | included |
| PRKCZ | included |
| PRKD1 | included |
| PRKDC | included |
| PRKRIR | included |
| PRL | included |
| PRLR | included |
| PRM3 | included |
| PRMT1 | included |
| PRMT2 | included |
| PRMT3 | included |
| PRMT5 | included |
| PRMT6 | included |
| PRMT7 | included |
| PRMT8 | included |
| PRNP | included |
| PROC | included |
| PROCR | included |
| PROM1 | included |
| Protein S | included |
| PRPF40A | included |
| PRR3 | included |
| PRRT1 | included |


| PRSS16 | included |
| :---: | :---: |
| PRTN3 | included |
| PSCD1 | included |
| PSCDBP | included |
| PSG1 | included |
| PSMB1 | included |
| PSMB10 | included |
| PSMB5 | included |
| PSMB6 | included |
| PSMB7 | included |
| PSMB8 | included |
| PSMB9 | included |
| PSMC6 | included |
| PSME1 | included |
| PSME2 | included |
| PSME3 | included |
| PSMF1 | included |
| PSORS1C1 | included |
| PSORS1C2 | included |
| PSTPIP1 | included |
| PTAFR | included |
| PTCH1 | included |
| PTDSR | included |
| PTDSS1 | included |
| PTEN | included |
| PTGDR | included |
| PTGDS | included |
| PTGER1 | included |
| PTGER2 | included |
| PTGER3 | included |
| PTGER4 | included |
| PTGES | included |
| PTGES2 | included |
| PTGFR | included |
| PTGFRN | included |
| PTGIR | excluded - no marker |
| PTGIS | included |
| PTGS1 | included |
| PTGS2 | included |
| PTH | included |
| PTHLH | included |
| PTHR1 | included |
| PTK 2B | included |
| PTK2 | included |
| PTK7 | included |
| PTP4A1 | included |
| PTP4A2 | included |
| PTP4A3 | included |
| PTPMT1 | included |
| PTPN1 | included |
| PTPN11 | included |
| PTPN12 | included |
| PTPN13 | included |
| PTPN14 | included |
| PTPN18 | included |
| PTPN2 | included |


| PTPN20A | excluded - no marker |
| :---: | :---: |
| PTPN20B | excluded - no marker |
| PTPN20C | included |
| PTPN21 | included |
| PTPN22 | included |
| PTPN23 | included |
| PTPN3 | included |
| PTPN4 | included |
| PTPN5 | included |
| PTPN6 | included |
| PTPN7 | included |
| PTPN9 | included |
| PTPRA | included |
| PTPRB | included |
| PTPRC | included |
| PTPRCAP | included |
| PTPRD | included |
| PTPRE | included |
| PTPRF | included |
| PTPRG | included |
| PTPRH | included |
| PTPRJ | included |
| PTPRK | included |
| PTPRM | included |
| PTPRN | included |
| PTPRN2 | included |
| PTPRO | included |
| PTPRQ | included |
| PTPRR | included |
| PTPRS | included |
| PTPRT | included |
| PTPRU | included |
| PTPRV | included |
| PTPRZ1 | included |
| PTX3 | included |
| PVR | included |
| PVRL1 | included |
| PVRL2 | included |
| PVRL3 | included |
| PXDN | included |
| PXMP3 | included |
| PYCARD | excluded - no marker |
| PYDC1 | excluded - no marker |
| QSCN6 | included |
| RAB19 | included |
| RAB3D | included |
| RAC1 | included |
| RAC2 | included |
| RAC3 | excluded - no marker |
| RAD23A | included |
| RAD50 | included |
| RAD51 | included - new MS design |
| RAD9A | included |
| RAD9B | included |
| RAE1 | included |
| RAET1E | included |


| RAF1 | excluded - no marker |
| :---: | :---: |
| RAG1 | included |
| RAG2 | included |
| RAGE | included |
| RALBP1 | included |
| RAN | included |
| RANBP2 | included |
| RANP1 | included |
| RAP1A | included |
| RAPGEF1 | included |
| RARA | included |
| RARB | included |
| RARG | included |
| RARRES2 | included |
| RARRES3 | included |
| RASA1 | included |
| RASGRP1 | included |
| RASGRP2 | included |
| RASGRP3 | included |
| RASSF5 | included |
| RAX | included |
| RBPSUH | included |
| RDBP | excluded - no marker |
| RDX | included |
| REL | included |
| RELA | included |
| RELB | included |
| RELN | included |
| RFC1 | included |
| RFX1 | included |
| RFX2 | included |
| RFX3 | included |
| RFX4 | included |
| RFX5 | included |
| RFXANK | included |
| RFXAP | included |
| RFXDC1 | included |
| RGL2 | included |
| RGS13 | included |
| RHAG | included |
| RHBDL2 | included |
| RHCE | included |
| RHD | included |
| RHOA | included |
| RHOC | included |
| RHOD | included |
| RHOH | included |
| RING1 | included |
| RIPK1 | included |
| RIPK2 | included |
| RIPK3 | included |
| RNASE6 | included |
| RNASE7 | included |
| RNASEH2A | included |
| RND2 | excluded - no marker |
| RNF39 | included |


| RNF4 | included |
| :---: | :---: |
| RNF5 | included |
| RNF7 | included |
| RNMT | included |
| ROCK1 | included |
| ROCK2 | included |
| RORA | included |
| RORB | included |
| RORC | included |
| RPA3 | included |
| RPL23AP1 | included |
| RPL32P1 | included |
| RPL7AP7 | included |
| RPLP1 | included |
| RPP21 | included |
| RPS18 | included |
| RPS4X | included |
| RPS4Y1 | included - new MS design |
| RPS6KA2 | included |
| RRM1 | included |
| RSAD2 | included |
| RUNX1 | included |
| RUNX1T1 | included |
| RUNX2 | included |
| RUNX3 | included |
| RXRA | excluded - no marker |
| RXRB | included |
| RXRG | included |
| S100A11 | included |
| S100A12 | included |
| S100A4 | included |
| S100A6 | included |
| S100A8 | included |
| S100A9 | included |
| SAA1 | included |
| SAA2 | included |
| SAA3P | included |
| SAMHD1 | included |
| SAP18 | included |
| SARM1 | included |
| SATB1 | excluded - no marker |
| SCAP1 | included |
| SCARA3 | included |
| SCARA5 | included |
| SCARB1 | included |
| SCARB2 | included |
| SCARF1 | included |
| SCARF2 | included |
| SCGB3A1 | included |
| SCL11A1 | included |
| SCMH1 | included |
| SCML2 | included |
| SCYE1 | included |
| SDC1 | included |
| SDC2 | included |
| SDC3 | included |


| SDC4 | included |
| :---: | :---: |
| SDCBP | included |
| SDF2 | excluded - no marker |
| SDF2L1 | included |
| SEC61A1 | included |
| SEC61A2 | included |
| SEC61B | included |
| SECTM1 | included |
| SELE | included |
| SELL | included |
| SELP | included |
| SELPLG | included |
| SEMA3E | included |
| SEMA4D | included |
| SEMA7A | included |
| SEMG1 | included |
| SEPHS2 | included |
| SEPT2 | included |
| SERPINA2 | included |
| SERPINB2 | included |
| SERPINB8 | included |
| SERPINB9 | included |
| SERPINC1 | included |
| SERPINE1 | included |
| SERPING1 | included |
| SET | included |
| SFRS2IP | included |
| SFTBA1B | excluded - no marker |
| SFTPA1 | included |
| SFTPB | included |
| SFTPD | included |
| SGK | included |
| SH2D1A | included |
| SH2D1B | included |
| SH2D2A | included |
| SH3KBP1 | included |
| SHB | included |
| SHH | included |
| SHMT1 | excluded - no marker |
| SHMT2 | included - new MS design |
| SIGIRR | included - new MS design |
| SIGLEC1 | included |
| SIGLEC10 | included |
| SIGLEC5 | included |
| SIGLEC6 | included |
| SIGLEC7 | included |
| SIGLEC8 | included |
| SIGLEC9 | included |
| SILV | included |
| SIM1 | included |
| SIPA1 | included |
| SIRPA | included |
| SIRPB1 | included |
| SIRPB2 | included |
| SIT1 | included |
| SIVA | included |


| SKAP1 | included |
| :---: | :---: |
| SKIV2L | excluded - no marker |
| SKP2 | included |
| SLAMF1 | included |
| SLAMF6 | included |
| SLAMF7 | included |
| SLAMF8 | included |
| SLAMF9 | included |
| SLC14A1 | included |
| SLC19A1 | included - new MS design |
| SLC1A5 | included |
| SLC22A1 | included |
| SLC22A4 | included |
| SLC22A5 | included |
| SLC25A19 | included |
| SLC39A7 | included |
| SLC3A2 | included |
| SLC40A1 | included |
| SLC44A1 | included |
| SLC44A4 | excluded - no marker |
| SLC4A1 | included |
| SLC6A4 | included |
| SLC7A5 | included |
| SLC9A1 | included |
| SLC9A2 | included |
| SLC9A3 | included |
| SLC9A4 | included |
| SLC9A5 | included |
| SLC9A6 | included |
| SLC9A7 | included |
| SLC9A8 | included |
| SLC9A9 | included |
| SLPI | included |
| SMAD1 | included |
| SMAD3 | included |
| SMAD7 | included |
| SMARCA1 | included |
| SMARCA2 | included |
| SMARCA4 | included |
| SMARCA5 | included |
| SMARCAL1 | included |
| SMC3 | included |
| SMG6 | included |
| SMO | included |
| SNFT | included |
| SNRP70 | included |
| SNRPN | included |
| SNX9 | included |
| SOBP | included |
| SOCS1 | included |
| SOCS2 | included |
| SOCS3 | included |
| SOCS4 | included |
| SOCS5 | included |
| SOCS6 | included |
| SOCS6/CBLN2 | included |


| SOCS7 | excluded - no marker |
| :---: | :---: |
| SOD1 | included - new MS design |
| SOD2 | included |
| SOD3 | included |
| SORL1 | included |
| SOS1 | included |
| SOX13 | included |
| SP1 | included |
| SP110 | included |
| SP3 | included |
| SPEN | included |
| SPHK1 | included |
| SPHK2 | included |
| SPI1 | included |
| SPN | included |
| SPP1 | included |
| SQSTM1 | included |
| SRC | included |
| SREBF1 | included |
| SRPK1 | included |
| SRPK2 | included |
| SRPK3 | excluded - no marker |
| SST | included |
| SSTR1 | included |
| SSTR2 | included |
| ST3GAL5 | included |
| ST6GAL1 | included |
| ST8SIA6 | included |
| STAB1 | included |
| STAT1 | included |
| STAT2 | included |
| STAT3 | included |
| STAT4 | included |
| STAT5A | included |
| STAT5B | included |
| STAT6 | included |
| STIL | included |
| STK17A | included |
| STK17B | included |
| STK19 | excluded - no marker |
| STK3 | included |
| STK38 | included |
| STK4 | included |
| STT3B | included |
| STX11 | included |
| STXBP5 | included |
| STYX | included |
| SULT1C2 | included |
| SUMO1 | excluded - no marker |
| SUV39H1 | included |
| SVEP1/MUSK | included |
| SYK | included |
| SYN3/LARGE | included |
| SYNE1 | included |
| SYT7 | included |
| SYTL1 | excluded - no marker |


| Tac1 | included |
| :---: | :---: |
| TACC1 | included |
| TACSTD1 | included |
| TAF13 | included |
| TAF9 | included |
| TAL1 | included |
| TAL2 | included |
| TA-NFKBH | included |
| TANK | included |
| TAP1 | included |
| TAP2 | included |
| TAPBP | included |
| TBK1 | excluded - no marker |
| TBL1X | included |
| TBL1XR1 | included |
| TBL1Y | included |
| TBP | included |
| TBX21 | included |
| TBXAR2 | included |
| TBXAS1 | included |
| TCAM2 | included |
| TCD@ | included |
| TCF12 | included |
| TCF19 | included |
| TCF3 | included |
| TCF4 | included |
| TCF7 | excluded - no marker |
| TCF8 | included |
| TCIRG1 | included |
| TCN2 | included |
| TEC | included |
| TEK | included |
| TEP1 | included |
| TERC | included |
| TERT | included |
| TFAP2C | included |
| TFB1M | included |
| TFDP2 | included |
| TFF1 | excluded - no marker |
| TFF2 | excluded - no marker |
| TFF3 | included |
| TFRC | included |
| TGFA | included |
| TGFB1 | included - new MS design |
| TGFB2 | included |
| TGFB3 | included |
| TGFBR1 | included |
| TGFBR2 | included |
| TGFBR3 | included |
| TGIF1 | included |
| TGM3 | included |
| THBD | included |
| THBS1 | included |
| THOC1 | included |
| THPO | included |
| THRA | included |


| THRB | included |
| :---: | :---: |
| THY1 | included |
| TIAF1 | included |
| TICAM1 | included |
| TIMELESS | included |
| TIMP1 | included |
| TIMP2 | included |
| TIMP3 | included |
| TIMP4 | included |
| TIRAP | included |
| TIRG1 | included |
| TLE3 | included |
| TLN1 | included |
| TLR1 | included |
| TLR10 | included - new MS design |
| TLR11 | excluded - unknown location |
| TLR12 | excluded - unknown location |
| TLR13 | excluded - unknown location |
| TLR2 | included |
| TLR3 | included |
| TLR4 | included |
| TLR5 | included |
| TLR6 | included |
| TLR7 | included |
| TLR8 | included |
| TLR9 | included |
| TM7SF4 | included |
| TMC8 | included |
| TMEM142A | excluded - no marker |
| TMEM158 | included |
| TMEM37 | included |
| TMPO | included |
| TMPRSS11D | included |
| TMSB4X | included |
| TMSB4Y | included |
| TNC | included |
| TNF | included |
| TNFa | included |
| TNFAIP3 | included |
| TNFb | included |
| TNFd | included |
| TNFRSF10A | included - new MS design |
| TNFRSF10B | included |
| TNFRSF10C | included - new MS design |
| TNFRSF10D | included - new MS design |
| TNFRSF11A | included |
| TNFRSF11B | included |
| TNFRSF12A | included |
| TNFRSF13B | included |
| TNFRSF13C | included |
| TNFRSF14 | included - new MS design |
| TNFRSF17 | included - new MS design |
| TNFRSF18 | included - new MS design |
| TNFRSF19 | included |
| TNFRSF19L | included |
| TNFRSF1A | included |


| TNFRSF1B | included |
| :---: | :---: |
| TNFRSF21 | included |
| TNFRSF25 | included |
| TNFRSF4 | excluded - no marker |
| TNFRSF5 | included |
| TNFRSF6 | included |
| TNFRSF6B | included |
| TNFRSF7 | included |
| TNFRSF8 | included |
| TNFRSF9 | included |
| TNFSF10 | included |
| TNFSF11 | included |
| TNFSF12 | included |
| TNFSF12-13 | included |
| TNFSF13 | included |
| TNFSF13B | included |
| TNFSF14 | included |
| TNFSF15 | included |
| TNFSF18 | included |
| TNFSF4 | included |
| TNFSF6 | included |
| TNFSF7 | included |
| TNFSF8 | included |
| TNFSF9 | included |
| TNN | included |
| TNR | included |
| TNXB | excluded - no marker |
| TOLLIP | excluded - no marker |
| TOP2A | included |
| TOR3A | included |
| TP35 | included |
| TP73L | included |
| TPMT | included |
| TPT1 | included |
| TRA@ | included |
| TRADD | included |
| TRAF1 | included |
| TRAF2 | included |
| TRAF3 | included |
| TRAF3IP1 | included |
| TRAF4 | excluded - no marker |
| TRAF5 | included |
| TRAF6 | included |
| TRAF7 | included |
| TRAM1 | included |
| TRAM2 | included |
| TRAT1 | included |
| TRB@ | included |
| TREM1 | included |
| TREM2 | included |
| TREML1 | included |
| TREML2 | included |
| TREML3 | included |
| TREML4 | included |
| TRERF1 | included |
| TRG@ | included |


| TRH | included |
| :---: | :---: |
| TRHR | included |
| TRIM10 | included |
| TRIM15 | included |
| TRIM22 | included |
| TRIM25 | included |
| TRIM26 | included |
| TRIM31 | included |
| TRIM39 | included |
| TRIM40 | included |
| TRIM59 | included |
| TRPM2 | included |
| TRPV1 | included |
| TRPV2 | included |
| TSC22D3 | included |
| TSHB | included |
| TSHR | included |
| TSLP | included |
| TSPAN7 | included |
| TSPYL2 | included |
| TTRAP | included |
| TTYH1 | included |
| TUBA1 | included |
| TUBA1A | included |
| TUBB | included |
| TWIST1 | included |
| TXK | included |
| TXN | included |
| TXNDC | included |
| TXNRD1 | included |
| TYK2 | included |
| TYMS | included |
| TYR | included |
| TYROBP | included |
| UBB | excluded - no marker |
| UBC | included |
| UBD | included |
| UBE2D1 | included |
| UBE2L6 | included |
| UBE2N | included |
| UBE2V1 | included |
| UCN | included |
| UGCG | included |
| UGT1A1 | included |
| UGT1A9 | included |
| UGT2B17 | included |
| UGT2B28 | included |
| UGT2B7 | included |
| ULBP1 | included |
| ULBP2 | included |
| ULBP3 | included |
| UNC5CL | included |
| UNC84B | included |
| UNC93B1 | included |
| UNG | included |
| Unknown | included |


| USF1 | included |
| :---: | :---: |
| USF2 | included |
| USP18 | excluded - no marker |
| USP9Y | included |
| UTRN | included |
| UTY | included |
| UVRAG | included |
| VARS | included |
| VARSL | included |
| VASP | included |
| VAV1 | included |
| VAV2 | included |
| VAV3 | included |
| VCAM1 | included |
| VDR | included |
| VEGFA | included |
| VEGFB | included |
| VEGFC | included |
| VIM | included |
| VIP | included |
| VISA | included |
| VNN1 | included |
| VPREB1 | included |
| VPS24 | included |
| VPS52 | included |
| VRK2 | included |
| VTCN1 | included |
| VTN | included |
| VWF | included |
| WAS | included |
| WASF1 | included |
| WASF3 | included |
| WDR46 | included |
| WFDC12 | included |
| WIPF1 | included |
| WNT1 | included |
| WNT3 | included |
| WNT4 | included |
| WNT5A | included |
| WRNIP1 | included |
| WT1 | excluded - no marker |
| WTAP | included |
| XBP1 | included |
| XCL1 | included |
| XCL2 | included |
| XCR1 | included |
| XDH | included |
| XG | included |
| XK | included |
| XPA | included |
| XPC | included |
| XRCC1 | included |
| XRCC3 | included |
| XRCC5 | included |
| YES1 | included |
| YWHA2 | included |

YWHAQ
YY1
ZAP70
ZBTB12
ZBTB22
ZBTB32
ZBTB7
ZFAND3
ZFP36
ZNF192
ZNF193
ZNF3
ZNF451
ZNRD1
included
included
included
excluded - no marker
included
included
excluded - no marker
included
included
included
included
included
included
included

| Gene Symbol | Aliases | Gene location (genecard) | Chromosome |
| :---: | :---: | :---: | :---: |
| TNFRSF18 | GITR | Chromosome 1:1,128,751-1,131,952 | 1 |
| DVL1 | DSH | Chromosome 1:1,260,521-1,274,623 | 1 |
| MMP23B |  | Chromosome 1:1,557,337-1,623,109 | 1 |
| PRKCZ | PKC§ | Chromosome 1:1,971,769-2,106,694 | 1 |
| DFFA |  | Chromosome 1:10,439,166-10,455,200 | 1 |
| VCAM1 | CD106 | Chromosome 1:100,957,885-100,977,189 | 1 |
| EDG1 | sphingosine-1-phosphate | Chromosome 1:101,475,032-101,479,662 | 1 |
| PRMT6 | PRMT6 | Chromosome 1:107,400,824-107,403,439 | 1 |
| VAV3 |  | Chromosome 1:107,915,305-108,309,108 | 1 |
| VAV3 |  | Chromosome 1:107,915,305-108,309,108 | 1 |
| VAV3 |  | Chromosome 1:107,915,305-108,309,108 | 1 |
| VAV3 |  | Chromosome 1:107,915,305-108,309,108 | 1 |
| TAF13 | POLII | Chromosome 1:109,406,644-109,420,147 | 1 |
| GNAI3 | G protein alpha i3 | Chromosome 1:109,892,824-109,938,498 | 1 |
| GNAI3 | G protein alpha i3 | Chromosome 1:109,892,824-109,938,498 | 1 |
| MASP2 | MAp19 | Chromosome 1:11,009,167-11,029,877 | 1 |
| MASP2 | MAp19 | Chromosome 1:11,009,167-11,029,877 | 1 |
| FRAP1 | mTOR | Chromosome 1:11,089,179-11,245,176 | 1 |
| FRAP1 | mTOR | Chromosome 1:11,089,179-11,245,176 | 1 |
| FRAP1 | mTOR | Chromosome 1:11,089,179-11,245,176 | 1 |
| MTHFR |  | Chromosome 1:11,768,367-11,788,702 | 1 |
| NPPA | ANP | Chromosome 1:11,828,353-11,830,989 | 1 |
| CSF1 | M-CSF | Chromosome 1:110,254,778-110,275,144 | 1 |
| CD53 |  | Chromosome 1:111,215,344-111,244,081 | 1 |
| CD53 |  | Chromosome 1:111,215,344-111,244,081 | 1 |
| CHI3L2 | Chi3I3 | Chromosome 1:111,571,804-111,587,585 | 1 |
| ADORA3 | Adenosin receptor 3 | Chromosome 1:111,827,493-111,908,107 | 1 |
| ADORA3 | Adenosin receptor 3 | Chromosome 1:111,827,493-111,908,107 | 1 |
| RAP1A | Rap-1 | Chromosome 1:111,886,363-112,060,836 | 1 |
| RHOC | RhoC | Chromosome 1:113,045,251-113,051,579 | 1 |
| PTPN22 |  | Chromosome 1:114,092,981-114,215,904 | 1 |
| PTPN22 |  | Chromosome 1:114,092,981-114,215,904 | 1 |
| BCAS2 |  | Chromosome 1:114,911,701-114,925,788 | 1 |
| NRAS | Ras | Chromosome 1:115,048,613-115,102,147 | 1 |
| TSHB | TSHB | Chromosome 1:115,373,938-115,378,464 | 1 |
| NGFB | NGFB | Chromosome 1:115,630,060-115,682,380 | 1 |
| CD58 | LFA-3 | Chromosome 1:116,858,680-116,915,184 | 1 |
| IGSF3 |  | Chromosome 1:116,918,554-117,011,898 | 1 |
| CD2 | LFA-2 | Chromosome 1:117,098,530-117,113,374 | 1 |
| PTGFRN | CD315, prostaglandin rec | Chromosome 1:117,254,202-117,334,503 | 1 |
| PTGFRN | CD315, prostaglandin rec | Chromosome 1:117,254,202-117,334,503 | 1 |
| VTCN1 |  | Chromosome 1:117,487,732-117,555,079 | 1 |
| VTCN1 |  | Chromosome 1:117,487,732-117,555,079 | 1 |
| HSD3B1 | 3 beta hydroxysteroid deh | Chromosome 1:119,851,356-119,859,200) | 1 |
| TNFRSF8 | CD30 | Chromosome 1:12,046,021-12,126,851 | 1 |
| TNFRSF1B | CD120b | Chromosome 1:12,149,647-12,191,872 | 1 |
| NOTCH2 |  | Chromosome 1:120,255,699-120,413,799 | 1 |
| FCGR1B |  | Chromosome 1:120,728,502-120,737,460 | 1 |
| FCGR1B |  | Chromosome 1:120,728,502-120,737,460 | 1 |
| ITGA10 |  | Chromosome 1:144,236,248-144,255,225 | 1 |
| CD160 |  | Chromosome 1:144,407,155-144,426,971 | 1 |
| PPIAL4 |  | Chromosome 1:146,418,535-146,422,374 | 1 |
| FCGR1A | CD64 | Chromosome 1:146,567,361-146,577,147 | 1 |
| MTMR11 | CRA | Chromosome 1:148,167,168-148,175,396 | 1 |
| MCL1 |  | Chromosome 1:148,813,658-148,818,760 | 1 |


| CTSS | Cathepsin S | Chromosome 1:148,969,175-149,005,057 | 1 |
| :---: | :---: | :---: | :---: |
| CTSK | cathepsin K | Chromosome 1:149,035,311-149,047,436 | 1 |
| CTSK | cathepsin K | Chromosome 1:149,035,311-149,047,436 | 1 |
| BNIPL |  | Chromosome 1:149,275,670-149,286,700 | 1 |
| PIK4CB |  | Chromosome 1:149,531,037-149,566,815 | 1 |
| ELA2A |  | Chromosome 1:15,655,811-15,690,482 | 1 |
| ELA2B |  | Chromosome 1:15,655,811-15,690,482 | 1 |
| RORC | RORg, NR1F3 | Chromosome 1:150,039,364-150,070,972 | 1 |
| S100A11 | S100a11 | Chromosome 1:150,271,606-150,276,135 | 1 |
| PGLYRP3 | PGRP-Ia | Chromosome 1:151,536,962-151,549,818 | 1 |
| S100A12 | S100A12 | Chromosome 1:151,612,808-151,614,749 | 1 |
| S100A6 | S100a6 | Chromosome 1:151,773,699-151,775,344 | 1 |
| S100A4 | S100a4 | Chromosome 1:151,782,713-151,789,236 | 1 |
| ILF2 |  | Chromosome 1:151,900,905-151,910,148 | 1 |
| ILF2 |  | Chromosome 1:151,900,905-151,910,148 | 1 |
| MPS1 |  | Chromosome 1:152,229,853-152,231,250 | 1 |
| MPS1 |  | Chromosome 1:152,229,853-152,231,250 | 1 |
| IL6R | CD126 | Chromosome 1:152,644,293-152,708,550 | 1 |
| IL6R | CD126 | Chromosome 1:152,644,293-152,708,550 | 1 |
| CKS1B |  | Chromosome 1:153,213,753-153,218,348 | 1 |
| EFNA1 | Ephrin A1 | Chromosome 1:153,366,560-153,374,010 | 1 |
| MUC1 | MUC1 | Chromosome 1:153,424,924-153,429,330 | 1 |
| MEF2D |  | Chromosome 1:154,700,143-154,737,244 | 1 |
| SH2D2A | RIBP | Chromosome 1:155,042,659-155,053,270 | 1 |
| SH2D2A | RIBP | Chromosome 1:155,042,659-155,053,270 | 1 |
| FCRL5 | CD307 | Chromosome 1:155,749,791-155,788,934 | 1 |
| FCRL4 |  | Chromosome 1:155,810,163-155,834,494 | 1 |
| FCRL2 |  | Chromosome 1:155,982,145-156,013,546 | 1 |
| FCRL2 |  | Chromosome 1:155,982,145-156,013,546 | 1 |
| CD1D |  | Chromosome 1:156,416,361-156,421,310 | 1 |
| CD1C |  | Chromosome 1:156,526,200-156,530,044 | 1 |
| IFI16 | Ifi204 | Chromosome 1:157,236,382-157,291,569 | 1 |
| IFI16 | Ifi204 | Chromosome 1:157,236,382-157,291,569 | 1 |
| DARC | CD234, Duffy blood group | Chromosome 1:157,408,023-157,442,914 | 1 |
| FCER1A |  | Chromosome 1:157,526,128-157,544,638 | 1 |
| CRP |  | Chromosome 1:157,948,703-157,951,003 | 1 |
| DUSP23 |  | Chromosome 1:158,017,346-158,018,957 | 1 |
| IGSF9 |  | Chromosome 1:158,163,453-158,182,010 | 1 |
| IGSF8 | CD316 | Chromosome 1:158,327,754-158,335,103 | 1 |
| NHLH1 | HEN1 | Chromosome 1:158,603,481-158,609,262 | 1 |
| SLAMF6 | NTBA | Chromosome 1:158,721,444-158,759,676 | 1 |
| CD48 |  | Chromosome 1:158,915,160-158,948,265 | 1 |
| LY9 | CD229 | Chromosome 1:159,032,552-159,064,669 | 1 |
| JAM1 | CD321, JAM-A, F11R | Chromosome 1:159,231,625-159,275,404 | 1 |
| JAM1 | CD321, JAM-A, F11R | Chromosome 1:159,231,625-159,275,404 | 1 |
| FCER1G |  | Chromosome 1:159,451,693-159,457,113 | 1 |
| NR113 | NR113 | Chromosome 1:159,466,079-159,474,590 | 1 |
| FCGR2A | CD32 | Chromosome 1:159,741,844-159,755,984 | 1 |
| FCGR3B | CD16b | Chromosome 1:159,859,610-159,867,620 | 1 |
| SPEN | SHARP | Chromosome 1:16,046,946-16,139,542 | 1 |
| SH2D1B | EAT2b | Chromosome 1:160,631,680-160,648,552 | 1 |
| RXRG | NR2B3 | Chromosome 1:163,636,778-163,681,057 | 1 |
| CD3Z | CD247, CD3 zeta chain | Chromosome 1:165,666,501-165,754,471 | 1 |
| CD3Z | CD247, CD3 zeta chain | Chromosome 1:165,666,501-165,754,471 | 1 |
| MPZL1 | concanavalin A receptor | Chromosome 1:165,957,832-166,026,684 | 1 |
| XCL2 |  | Chromosome 1:166,776,626-166,779,859 | 1 |


| XCL1 |  | Chromosome 1:166,812,335-166,817,939 | 1 |
| :---: | :---: | :---: | :---: |
| F5 | Factor V | Chromosome 1:167,750,028-167,822,450 | 1 |
| SELP | CD62P | Chromosome 1:167,824,661-167,866,031 | 1 |
| SELP | CD62P | Chromosome 1:167,824,661-167,866,031 | 1 |
| SELL | L-Selectin, CD62L | Chromosome 1:167,926,432-167,947,463 | 1 |
| PADI4 |  | Chromosome 1:17,507,277-17,563,086 | 1 |
| TNFSF6 | CD178, FASL | Chromosome 1:170,894,777-170,902,637 | 1 |
| TNFSF6 | CD178, FASL | Chromosome 1:170,894,777-170,902,637 | 1 |
| TNFSF18 | GITRL | Chromosome 1:171,275,723-171,286,679 | 1 |
| TNFSF18 | GITRL | Chromosome 1:171,275,723-171,286,679 | 1 |
| TNFSF4 | CD252, OX40L | Chromosome 1:171,419,493-171,443,094 | 1 |
| TNFSF4 | CD252, OX40L | Chromosome 1:171,419,493-171,443,094 | 1 |
| PRDX6 |  | Chromosome 1:171,713,028-171,724,569 | 1 |
| SERPINC1 | Antithrombin | Chromosome 1:172,139,562-172,153,139 | 1 |
| TNN | tenascin | Chromosome 1:173,303,617-173,383,825 | 1 |
| TNR | tenascin | Chromosome 1:173,558,558-173,979,529 | 1 |
| TNR | tenascin | Chromosome 1:173,558,558-173,979,529 | 1 |
| TNR | tenascin | Chromosome 1:173,558,558-173,979,529 | 1 |
| TNR | tenascin | Chromosome 1:173,558,558-173,979,529 | 1 |
| TNR | tenascin | Chromosome 1:173,558,558-173,979,529 | 1 |
| TNR | tenascin | Chromosome 1:173,558,558-173,979,529 | 1 |
| ANGPTL1 | Angioarrestin | Chromosome 1:177,085,293-177,106,838 | 1 |
| TOR3A | ADIR | Chromosome 1:177,317,735-177,333,653 | 1 |
| TOR3A | ADIR | Chromosome 1:177,317,735-177,333,653 | 1 |
| QSCN6 |  | Chromosome 1:178,390,591-178,439,788 | 1 |
| QSCN6 |  | Chromosome 1:178,390,591-178,439,788 | 1 |
| MR1 | MR1=HLALS | Chromosome 1:179,269,762-179,292,312 | 1 |
| IGSF21 |  | Chromosome 1:18,306,827-18,577,563 | 1 |
| IGSF21 |  | Chromosome 1:18,306,827-18,577,563 | 1 |
| IGSF21 |  | Chromosome 1:18,306,827-18,577,563 | 1 |
| IGSF21 |  | Chromosome 1:18,306,827-18,577,563 | 1 |
| DHX9 |  | Chromosome 1:181,075,127-181,123,510 | 1 |
| DHX9 |  | Chromosome 1:181,075,127-181,123,510 | 1 |
| LAMC2 |  | Chromosome 1:181,422,022-181,480,662 | 1 |
| LAMC2 |  | Chromosome 1:181,422,022-181,480,662 | 1 |
| NCF2 | neutrophil cytosolic factor | Chromosome 1:181,791,320-181,826,634 | 1 |
| PRG4 | MSF | Chromosome 1:184,532,034-184,550,317 | 1 |
| PTGES2 |  | Chromosome 1:184,907,546-184,916,179 | 1 |
| RGS13 |  | Chromosome 1:190,871,905-190,896,059 | 1 |
| RGS13 |  | Chromosome 1:190,871,905-190,896,059 | 1 |
| CFH |  | Chromosome 1:194,887,631-194,983,257 | 1 |
| CFH |  | Chromosome 1:194,887,631-194,983,257 | 1 |
| CFHR4 |  | Chromosome 1:195,010,571-195,154,386 | 1 |
| CFHR2 |  | Chromosome 1:195,179,520-195,194,979 | 1 |
| PTPRC | CD45 | Chromosome 1:196,874,424-196,993,035 | 1 |
| NR5A2 | LRH-1 | Chromosome 1:198,263,353-198,413,175 | 1 |
| DDX59 |  | Chromosome 1:198,859,647-198,905,749 | 1 |
| DDX59 |  | Chromosome 1:198,859,647-198,905,749 | 1 |
| TNFRSF14 | CD270, LIGHTR, HVEM | Chromosome 1:2,479,150-2,486,613 | 1 |
| TNFRSF14 | CD270, LIGHTR, HVEM | Chromosome 1:2,479,150-2,486,613 | 1 |
| PLA2G2D | Phospholipase | Chromosome 1:20,311,019-20,318,637 | 1 |
| PLA2G2D | Phospholipase | Chromosome 1:20,311,019-20,318,637 | 1 |
| PLA2G2D | Phospholipase | Chromosome 1:20,311,019-20,318,637 | 1 |
| CDA | CDD | Chromosome 1:20,788,028-20,817,988 | 1 |
| CDA |  | Chromosome 1:20,788,028-20,817,988 | 1 |
| ELF3 | ERT | Chromosome 1:200,243,696-200,252,939 | 1 |


| PTPN7 |  | Chromosome 1:200,382,764-200,397,332 | 1 |
| :---: | :---: | :---: | :---: |
| PTPRV |  | Chromosome 1:200,403,802-200,425,104 | 1 |
| ADORA1 | Adenosin receptor 1 | Chromosome 1:201,326,405-201,403,156 | 1 |
| ADORA1 | Adenosin receptor 1 | Chromosome 1:201,326,405-201,403,156 | 1 |
| ADORA1 | Adenosin receptor 1 | Chromosome 1:201,326,405-201,403,156 | 1 |
| FMOD | Fibromodulin | Chromosome 1:201,576,375-201,587,240 | 1 |
| LAX1 |  | Chromosome 1:202,000,957-202,012,123 | 1 |
| SOX13 | SOX13 (SRY box 13) | Chromosome 1:202,308,866-202,363,494 | 1 |
| SOX13 | SOX13 (SRY box 13) | Chromosome 1:202,308,866-202,363,494 | 1 |
| PIK3C2B |  | Chromosome 1:202,658,379-202,726,175 | 1 |
| MDM4 |  | Chromosome 1:202,752,134-202,793,871 | 1 |
| CNTN2 |  | Chromosome 1:203,278,963-203,313,761 | 1 |
| CNTN2 |  | Chromosome 1:203,278,963-203,313,761 | 1 |
| ELK4 |  | Chromosome 1:203,833,330-203,868,623 | 1 |
| IKBKE | IKKepsilon, IKKi | Chromosome 1:204,710,414-204,736,846 | 1 |
| DYRK3 |  | Chromosome 1:204,875,504-204,924,381 | 1 |
| MAPKAPK2 |  | Chromosome 1:204,924,912-204,974,251 | 1 |
| IL10 | Interleukin 10 | Chromosome 1:205,007,570-205,012,462 | 1 |
| IL19 |  | Chromosome 1:205,038,838-205,082,949 | 1 |
| IL20 |  | Chromosome 1:205,105,322-205,109,191 | 1 |
| IL24 |  | Chromosome 1:205,137,411-205,144,107 | 1 |
| C4BPB |  | Chromosome 1:205,328,810-205,339,961 | 1 |
| C4BPA |  | Chromosome 1:205,344,230-205,384,940 | 1 |
| CD55 | CD55, DAF | Chromosome 1:205,561,476-205,600,934 | 1 |
| CR2 | CD21 | Chromosome 1:205,694,198-205,729,863 | 1 |
| CR1 |  | Chromosome 1:205,736,096-205,881,733 | 1 |
| CR1 |  | Chromosome 1:205,736,096-205,881,733 | 1 |
| CR1 |  | Chromosome 1:205,736,096-205,881,733 | 1 |
| MCP | CD46 | Chromosome 1:205,992,025-206,035,481 | 1 |
| CD34 |  | Chromosome 1:206,116,942-206,151,370 | 1 |
| CD34 |  | Chromosome 1:206,116,942-206,151,370 | 1 |
| CD34 |  | Chromosome 1:206,116,942-206,151,370 | 1 |
| LAMB3 |  | Chromosome 1:207,854,838-207,892,443 | 1 |
| LAMB3 |  | Chromosome 1:207,854,838-207,892,443 | 1 |
| LAMB3 |  | Chromosome 1:207,854,838-207,892,443 | 1 |
| IRF6 |  | Chromosome 1:208,025,659-208,046,102 | 1 |
| TRAF5 |  | Chromosome 1:209,566,580-209,614,911 | 1 |
| TRAF5 |  | Chromosome 1:209,566,580-209,614,911 | 1 |
| ATF3 | ATF3 | Chromosome 1:210,805,374-210,860,742 | 1 |
| ATF3 | ATF3 | Chromosome 1:210,805,374-210,860,742 | 1 |
| PTPN14 |  | Chromosome 1:212,597,474-212,791,265 | 1 |
| PTPN14 |  | Chromosome 1:212,597,474-212,791,265 | 1 |
| CENPF | Centromere protein F | Chromosome 1:212,843,155-212,904,537 | 1 |
| ESRRG | NR3B3 | Chromosome 1:214,743,211-215,377,720 | 1 |
| ESRRG | NR3B3 | Chromosome 1:214,743,211-215,377,720 | 1 |
| ESRRG | NR3B3 | Chromosome 1:214,743,211-215,377,720 | 1 |
| ESRRG | NR3B3 | Chromosome 1:214,743,211-215,377,720 | 1 |
| ESRRG | NR3B3 | Chromosome 1:214,743,211-215,377,720 | 1 |
| ESRRG | NR3B3 | Chromosome 1:214,743,211-215,377,720 | 1 |
| ESRRG | NR3B3 | Chromosome 1:214,743,211-215,377,720 | 1 |
| TGFB2 |  | Chromosome 1:216,586,200-216,684,584 | 1 |
| HLX1 | HIx | Chromosome 1:219,119,366-219,125,022 | 1 |
| HLX1 | HIX | Chromosome 1:219,119,366-219,125,022 | 1 |
| DUSP10 | MKP-5 | Chromosome 1:219,941,389-219,982,141 | 1 |
| DUSP10 | MKP-5 | Chromosome 1:219,941,389-219,982,141 | 1 |
| DUSP10 | MKP-5 | Chromosome 1:219,941,389-219,982,141 | 1 |


| HSPG2 | Perlecan | Chromosome 1:22,021,324-22,136,377 | 1 |
| :---: | :---: | :---: | :---: |
| WNT4 |  | Chromosome 1:22,318,177-22,342,197 | 1 |
| C1QA |  | Chromosome 1:22,835,705-22,838,762 | 1 |
| C1QA |  | Chromosome 1:22,835,705-22,838,762 | 1 |
| C1QA |  | Chromosome 1:22,835,705-22,838,762 | 1 |
| EPHB2 |  | Chromosome 1:22,910,045-23,114,405 | 1 |
| TLR5 | CD285 | Chromosome 1:221,350,270-221,383,247 | 1 |
| EPHX1 |  | Chromosome 1:224,064,459-224,099,884 | 1 |
| EPHX1 |  | Chromosome 1:224,064,459-224,099,884 | 1 |
| PARP1 |  | Chromosome 1:224,615,015-224,662,414 | 1 |
| PARP1 |  | Chromosome 1:224,615,015-224,662,414 | 1 |
| ITPKB |  | Chromosome 1:224,886,014-224,993,647 | 1 |
| ITPKB |  | Chromosome 1:224,886,014-224,993,647 | 1 |
| HIST3H2A | Histone 3 | Chromosome 1:226,711,303-226,712,197 | 1 |
| ACTA1 | F-actin | Chromosome 1:227,633,615-227,636,468 | 1 |
| ACTA1 | F-actin | Chromosome 1:227,633,615-227,636,468 | 1 |
| LYST |  | Chromosome 1:233,890,964-234,113,563 | 1 |
| LYST |  | Chromosome 1:233,890,964-234,113,563 | 1 |
| LYST |  | Chromosome 1:233,890,964-234,113,563 | 1 |
| LYST |  | Chromosome 1:233,890,964-234,113,563 | 1 |
| NID1 | entactin | Chromosome 1:234,205,753-234,303,706 | 1 |
| NID1 | entactin | Chromosome 1:234,205,753-234,303,706 | 1 |
| EDARADD |  | Chromosome 1:234,624,303-234,714,649 | 1 |
| EDARADD |  | Chromosome 1:234,624,303-234,714,649 | 1 |
| EDARADD |  | Chromosome 1:234,624,303-234,714,649 | 1 |
| ACTN2 | alpha actinin 2 | Chromosome 1:234,916,422-234,994,554 | 1 |
| IL22RA1 |  | Chromosome 1:24,318,848-24,342,198 | 1 |
| IL22RA1 |  | Chromosome 1:24,318,848-24,342,198 | 1 |
| IL22RA1 |  | Chromosome 1:24,318,848-24,342,198 | 1 |
| EXO1 |  | Chromosome 1:240,078,105-240,119,864 | 1 |
| AKT3 | PKB | Chromosome 1:241,718,158-242,080,053 | 1 |
| AKT3 | PKB | Chromosome 1:241,718,158-242,080,053 | 1 |
| AKT3 | PKB | Chromosome 1:241,718,158-242,080,053 | 1 |
| AKT3 | PKB | Chromosome 1:241,718,158-242,080,053 | 1 |
| AKT3 | PKB | Chromosome 1:241,718,158-242,080,053 | 1 |
| NLRP3 | NALP3, CIAS1, PYPAF1, | Chromosome 1:245,647,974-245,679,033 | 1 |
| NLRP3 | NALP3, CIAS1, PYPAF1, | Chromosome 1:245,647,974-245,679,033 | 1 |
| RUNX3 |  | Chromosome 1:25,098,596-25,164,062 | 1 |
| RUNX3 |  | Chromosome 1:25,098,596-25,164,062 | 1 |
| RHD | RhD antigen, CD240D | Chromosome 1:25,471,568-25,529,523 | 1 |
| RHCE | CD240CE | Chromosome 1:25,561,327-25,629,270 | 1 |
| PAFAH2 | PAF, platelet activating fa | Chromosome 1:26,158,845-26,197,235 | 1 |
| PAFAH2 | PAF, platelet activating fa | Chromosome 1:26,158,845-26,197,235 | 1 |
| CDW52 | CD52, CAMPATH | Chromosome 1:26,516,998-26,519,601 | 1 |
| GPATCH3 | NR0B2 | Chromosome 1:27,089,567-27,099,549 | 1 |
| NROB2 | NR0B2 | Chromosome 1:27,110,566-27,113,047 | 1 |
| SLC9A1 | NHE-1 | Chromosome 1:27,297,893-27,366,059 | 1 |
| SLC9A1 | NHE-1 | Chromosome 1:27,297,893-27,366,059 | 1 |
| FGR |  | Chromosome 1:27,811,162-27,834,375 | 1 |
| FGR |  | Chromosome 1:27,811,162-27,834,375 | 1 |
| PTAFR |  | Chromosome 1:28,346,264-28,392,971 | 1 |
| PTAFR |  | Chromosome 1:28,346,264-28,392,971 | 1 |
| PTPRU |  | Chromosome 1:29,435,611-29,525,899 | 1 |
| PTPRU |  | Chromosome 1:29,435,611-29,525,899 | 1 |
| DFFB |  | Chromosome 1:3,763,705-3,791,853 | 1 |
| SDC3 | Syndecan | Chromosome 1:31,114,901-31,166,301 | 1 |


| SDC3 | Syndecan | Chromosome 1:31,114,901-31,166,301 | 1 |
| :---: | :---: | :---: | :---: |
| FABP3 | FABP3 | Chromosome 1:31,610,687-31,618,510 | 1 |
| FABP3 | FABP3 | Chromosome 1:31,610,687-31,618,510 | 1 |
| PTP4A2 |  | Chromosome 1:32,144,609-32,176,578 | 1 |
| PTP4A2 |  | Chromosome 1:32,144,609-32,176,578 | 1 |
| KHDRBS1 |  | Chromosome 1:32,252,017-32,299,037 | 1 |
| LCK |  | Chromosome 1:32,489,480-32,524,353 | 1 |
| LCK |  | Chromosome 1:32,489,480-32,524,353 | 1 |
| HDAC1 |  | Chromosome 1:32,530,274-32,571,823 | 1 |
| CSF3R | CD114 | Chromosome 1:36,704,231-36,721,466 | 1 |
| CSF3R | CD114 | Chromosome 1:36,704,231-36,721,466 | 1 |
| GNL2 | Ngp | Chromosome 1:37,805,004-37,834,109 | 1 |
| PPIE |  | Chromosome 1:39,977,117-40,002,173 | 1 |
| PPIE |  | Chromosome 1:39,977,117-40,002,173 | 1 |
| MYCL1 | I-myc | Chromosome 1:40,133,685-40,140,274 | 1 |
| NFYC |  | Chromosome 1:40,929,829-41,009,864 | 1 |
| NFYC |  | Chromosome 1:40,929,829-41,009,864 | 1 |
| CTPS | CTP synthase | Chromosome 1:41,217,951-41,250,815 | 1 |
| SCMH1 | Scmh1 | Chromosome 1:41,265,461-41,480,375 | 1 |
| SCMH1 | Scmh1 | Chromosome 1:41,265,461-41,480,375 | 1 |
| FOXJ3 |  | Chromosome 1:42,414,797-42,574,135 | 1 |
| FOXJ3 |  | Chromosome 1:42,414,797-42,574,135 | 1 |
| PPIH |  | Chromosome 1:42,896,635-42,915,016 | 1 |
| ERMAP |  | Chromosome 1:43,055,363-43,083,247 | 1 |
| ERMAP |  | Chromosome 1:43,055,363-43,083,247 | 1 |
| MPL | CD110, F36VMpl, thromb | Chromosome 1:43,576,062-43,592,722 | 1 |
| CDC20 |  | Chromosome 1:43,597,213-43,601,461 | 1 |
| PTPRF |  | Chromosome 1:43,769,134-43,861,924 | 1 |
| PTPRF |  | Chromosome 1:43,769,134-43,861,924 | 1 |
| ARTN | ARTN | Chromosome 1:44,171,579-44,175,499 | 1 |
| PLK3 |  | Chromosome 1:45,038,623-45,049,479 | 1 |
| PIK3R3 |  | Chromosome 1:46,278,399-46,371,054 | 1 |
| CYP4B1 |  | Chromosome 1:47,037,305-47,057,672 | 1 |
| CYP4B1 |  | Chromosome 1:47,037,305-47,057,672 | 1 |
| CYP4X1 |  | Chromosome 1:47,261,827-47,289,010 | 1 |
| TAL1 | SCL=TAL1 | Chromosome 1:47,454,550-47,469,974 | 1 |
| TAL1 | SCL=TAL1 | Chromosome 1:47,454,550-47,469,974 | 1 |
| FOXE3 |  | Chromosome 1:47,654,331-47,656,311 | 1 |
| FOXD2 |  | Chromosome 1:47,674,276-47,678,950 | 1 |
| FAF1 |  | Chromosome 1:50,677,738-51,198,524 | 1 |
| FAF1 |  | Chromosome 1:50,677,738-51,198,524 | 1 |
| FAF1 |  | Chromosome 1:50,677,738-51,198,524 | 1 |
| FAF1 |  | Chromosome 1:50,677,738-51,198,524 | 1 |
| FAF1 |  | Chromosome 1:50,677,738-51,198,524 | 1 |
| FAF1 |  | Chromosome 1:50,677,738-51,198,524 | 1 |
| PPAP2B |  | Chromosome 1:56,732,527-56,817,845 | 1 |
| PPAP2B |  | Chromosome 1:56,732,527-56,817,845 | 1 |
| PPAP2B |  | Chromosome 1:56,732,527-56,817,845 | 1 |
| PRKAA2 | AMPK | Chromosome 1:56,883,583-56,953,596 | 1 |
| C8A |  | Chromosome 1:57,093,065-57,156,482 | 1 |
| JUN | AP-1 | Chromosome 1:59,019,048-59,022,587 | 1 |
| TNFRSF25 | TRAMP | Chromosome 1:6,443,798-6,502,708 | 1 |
| TNFRSF25 | TRAMP | Chromosome 1:6,443,798-6,502,708 | 1 |
| CYP2J2 |  | Chromosome 1:60,131,568-60,165,050 | 1 |
| FOXD3 |  | Chromosome 1:63,561,300-63,563,385 | 1 |
| JAK1 |  | Chromosome 1:65,071,500-65,204,775 | 1 |


| JAK1 |  | Chromosome 1:65,071,500-65,204,775 | 1 |
| :---: | :---: | :---: | :---: |
| JAK1 |  | Chromosome 1:65,071,500-65,204,775 | 1 |
| LEPR | CD295, leptin receptor | Chromosome 1:65,658,858-65,879,830 | 1 |
| LEPR | CD295, leptin receptor | Chromosome 1:65,658,858-65,879,830 | 1 |
| LEPR | CD295, leptin receptor | Chromosome 1:65,658,858-65,879,830 | 1 |
| PDE4B | Phosphodiesterases | Chromosome 1:66,030,781-66,612,850 | 1 |
| PDE4B | Phosphodiesterases | Chromosome 1:66,030,781-66,612,850 | 1 |
| PDE4B | Phosphodiesterases | Chromosome 1:66,030,781-66,612,850 | 1 |
| PDE4B | Phosphodiesterases | Chromosome 1:66,030,781-66,612,850 | 1 |
| PDE4B | Phosphodiesterases | Chromosome 1:66,030,781-66,612,850 | 1 |
| IL23R |  | Chromosome 1:67,404,671-67,498,250 | 1 |
| IL23R |  | Chromosome 1:67,404,671-67,498,250 | 1 |
| IL12RB2 |  | Chromosome 1:67,545,635-67,635,171 | 1 |
| TNFRSF9 | CDw137, 4-1BB | Chromosome 1:7,902,494-7,923,513 | 1 |
| TNFRSF9 | CDw137, 4-1BB | Chromosome 1:7,902,494-7,923,513 | 1 |
| PTGER3 | EP3 | Chromosome 1:71,090,624-71,286,079 | 1 |
| PTGER3 | EP3 | Chromosome 1:71,090,624-71,286,079 | 1 |
| PTGER3 | EP3 | Chromosome 1:71,090,624-71,286,079 | 1 |
| PTGER3 | EP3 | Chromosome 1:71,090,624-71,286,079 | 1 |
| PTGFR |  | Chromosome 1:78,542,156-78,778,974 | 1 |
| PTGFR |  | Chromosome 1:78,542,156-78,778,974 | 1 |
| PTGFR |  | Chromosome 1:78,542,156-78,778,974 | 1 |
| IFI44 |  | Chromosome 1:78,888,104-78,902,351 | 1 |
| IFI44 |  | Chromosome 1:78,888,104-78,902,351 | 1 |
| EGFTM7 |  | Chromosome 1:79,128,037-79,279,105 | 1 |
| EGFTM7 |  | Chromosome 1:79,128,037-79,279,105 | 1 |
| PRKACB |  | Chromosome 1:84,316,329-84,476,769 | 1 |
| PRKACB |  | Chromosome 1:84,316,329-84,476,769 | 1 |
| BCL10 |  | Chromosome 1:85,504,519-85,516,359 | 1 |
| BCL10 |  | Chromosome 1:85,504,519-85,516,359 | 1 |
| DDAH1 |  | Chromosome 1:85,556,756-85,703,415 | 1 |
| DDAH1 |  | Chromosome 1:85,556,756-85,703,415 | 1 |
| GTF2B | TFIIB | Chromosome 1:89,091,203-89,129,889 | 1 |
| GBP3 |  | Chromosome 1:89,244,948-89,261,132 | 1 |
| GBP2 |  | Chromosome 1:89,344,403-89,414,311 | 1 |
| GBP5 |  | Chromosome 1:89,498,853-89,511,119 | 1 |
| PIK3CD |  | Chromosome 1:9,634,390-9,711,564 | 1 |
| TGFBR3 | Betaglycan | Chromosome 1:91,918,488-92,144,147 | 1 |
| TGFBR3 | Betaglycan | Chromosome 1:91,918,488-92,144,147 | 1 |
| TGFBR3 | Betaglycan | Chromosome 1:91,918,488-92,144,147 | 1 |
| ISG15 | G1P2, ISRE, UCRP | Chromosome 1:938,666-939,783 | 1 |
| GCLM |  | Chromosome 1:94,123,349-94,147,600 | 1 |
| F3 | CD142, coag factor 3, tiss | Chromosome 1:94,767,369-94,779,944 | 1 |
| CHUK | IKK1 | Chromosome 10:101,899,841-101,979,366 | 10 |
| CHUK | IKK1 | Chromosome 10:101,899,841-101,979,366 | 10 |
| BTRC | Beta-TRCP | Chromosome 10:103,103,810-103,307,068 | 10 |
| BTRC | Beta-TRCP | Chromosome 10:103,103,810-103,307,068 | 10 |
| BTRC | Beta-TRCP | Chromosome 10:103,103,810-103,307,068 | 10 |
| FGF8 | FGF8 | Chromosome 10:103,519,877-103,525,817 | 10 |
| NFKB2 | p100, p52 | Chromosome 10:104,144,320-104,152,271 | 10 |
| C10orf26 | OPAL1 | Chromosome 10:104,525,996-104,566,011 | 10 |
| CYP17A1 | 17 alpha hydroxylase | Chromosome 10:104,580,278-104,587,280 | 10 |
| CUGBP2 |  | Chromosome 10:11,087,290-11,418,680 | 10 |
| CUGBP2 |  | Chromosome 10:11,087,290-11,418,680 | 10 |
| CUGBP2 |  | Chromosome 10:11,087,290-11,418,680 | 10 |
| CUGBP2 |  | Chromosome 10:11,087,290-11,418,680 | 10 |


| CUGBP2 |  | Chromosome 10:11,087,290-11,418,680 | 10 |
| :---: | :---: | :---: | :---: |
| CUGBP2 |  | Chromosome 10:11,087,290-11,418,680 | 10 |
| DUSP5 | VH3 | Chromosome 10:112,247,586-112,261,292 | 10 |
| SMC3 | BAM11 | Chromosome 10:112,317,439-112,354,384 | 10 |
| SMC3 | BAM11 | Chromosome 10:112,317,439-112,354,384 | 10 |
| CASP7 |  | Chromosome 10:115,428,925-115,480,654 | 10 |
| SEC61A2 |  | Chromosome 10:12,211,642-12,251,966 | 10 |
| SEC61A2 |  | Chromosome 10:12,211,642-12,251,966 | 10 |
| BAG3 |  | Chromosome 10:121,400,872-121,427,321 | 10 |
| BAG3 |  | Chromosome 10:121,400,872-121,427,321 | 10 |
| FGFR2 | KGF/FGF receptor, CD33 | Chromosome 10:122,473,377-123,347,962 | 10 |
| FGFR2 | KGF/FGF receptor, CD33 | Chromosome 10:122,473,377-123,347,962 | 10 |
| FGFR2 | KGF/FGF receptor, CD33 | Chromosome 10:122,473,377-123,347,962 | 10 |
| FGFR2 | KGF/FGF receptor, CD33 | Chromosome 10:122,473,377-123,347,962 | 10 |
| FGFR2 | KGF/FGF receptor, CD33 | Chromosome 10:122,473,377-123,347,962 | 10 |
| FGFR2 | KGF/FGF receptor, CD33 | Chromosome 10:122,473,377-123,347,962 | 10 |
| FGFR2 | KGF/FGF receptor, CD33 | Chromosome 10:122,473,377-123,347,962 | 10 |
| FGFR2 | KGF/FGF receptor, CD33 | Chromosome 10:122,473,377-123,347,962 | 10 |
| FGFR2 | KGF/FGF receptor, CD33 | Chromosome 10:122,473,377-123,347,962 | 10 |
| FGFR2 | KGF/FGF receptor, CD33 | Chromosome 10:122,473,377-123,347,962 | 10 |
| FGFR2 | KGF/FGF receptor, CD33 | Chromosome 10:122,473,377-123,347,962 | 10 |
| FGFR2 | KGF/FGF receptor, CD33 | Chromosome 10:122,473,377-123,347,962 | 10 |
| FGFR2 | KGF/FGF receptor, CD33 | Chromosome 10:122,473,377-123,347,962 | 10 |
| DMBT1 | gp340 | Chromosome 10:124,310,171-124,393,242 | 10 |
| DMBT1 | gp340 | Chromosome 10:124,310,171-124,393,242 | 10 |
| BUB3 |  | Chromosome 10:124,903,783-124,914,876 | 10 |
| BUB3 |  | Chromosome 10:124,903,783-124,914,876 | 10 |
| MMP21 |  | Chromosome 10:127,445,012-127,454,380 | 10 |
| DHX32 |  | Chromosome 10:127,514,896-127,575,017 | 10 |
| ADAM12 | metalloproteinase desine | Chromosome 10:127,690,940-128,067,055 | 10 |
| ADAM12 | metalloproteinase desine | Chromosome 10:127,690,940-128,067,055 | 10 |
| ADAM12 | metalloproteinase desined | Chromosome 10:127,690,940-128,067,055 | 10 |
| ADAM12 | metalloproteinase desined | Chromosome 10:127,690,940-128,067,055 | 10 |
| FOXI2 |  | Chromosome 10:129,425,504-129,429,440 | 10 |
| PTPRE |  | Chromosome 10:129,595,315-129,774,155 | 10 |
| PTPRE |  | Chromosome 10:129,595,315-129,774,155 | 10 |
| MGMT |  | Chromosome 10:131,155,456-131,455,358 | 10 |
| MGMT |  | Chromosome 10:131,155,456-131,455,358 | 10 |
| MGMT |  | Chromosome 10:131,155,456-131,455,358 | 10 |
| BNIP3 |  | Chromosome 10:133,631,181-133,645,450 | 10 |
| ADAM8 | CD156a | Chromosome 10:134,925,898-134,940,362 | 10 |
| CYP2E1 |  | Chromosome 10:135,190,857-135,224,714 | 10 |
| DCLRE1C |  | Chromosome 10:14,979,364-15,036,437 | 10 |
| DCLRE1C |  | Chromosome 10:14,979,364-15,036,437 | 10 |
| ITGA8 |  | Chromosome 10:15,595,954-15,802,130 | 10 |
| ITGA8 |  | Chromosome 10:15,595,954-15,802,130 | 10 |
| ITGA8 |  | Chromosome 10:15,595,954-15,802,130 | 10 |
| ITGA8 |  | Chromosome 10:15,595,954-15,802,130 | 10 |
| C1QL3 |  | Chromosome 10:16,595,748-16,604,010 | 10 |
| C1QL3 |  | Chromosome 10:16,595,748-16,604,010 | 10 |
| VIM | Vimentin | Chromosome 10:17,311,283-17,319,598 | 10 |
| MRC1L1 | Mannose receptor | Chromosome 10:17,891,368-17,993,184 | 10 |
| MRC1 | CD206, Mannose recepto | Chromosome 10:18,138,358-18,240,097 | 10 |
| BMI-1 | BMI-1 | Chromosome 10:22,650,146-22,660,194 | 10 |
| APBB1IP | RIAM, PEL1 | Chromosome 10:26,767,138-26,896,738 | 10 |
| APBB1IP | RIAM, PEL1 | Chromosome 10:26,767,138-26,896,738 | 10 |


| APBB1IP | RIAM，PEL1 | Chromosome 10：26，767，138－26，896，738 | 10 |
| :---: | :---: | :---: | :---: |
| MAP3K8 | TPL2 | Chromosome 10：30，762，872－30，790，768 | 10 |
| MAP3K8 | TPL2 | Chromosome 10：30，762，872－30，790，768 | 10 |
| TCF8 |  | Chromosome 10：31，647，430－31，858，748 | 10 |
| TCF8 |  | Chromosome 10：31，647，430－31，858，748 | 10 |
| TCF8 |  | Chromosome 10：31，647，430－31，858，748 | 10 |
| TCF8 |  | Chromosome 10：31，647，430－31，858，748 | 10 |
| TCF8 |  | Chromosome 10：31，647，430－31，858，748 | 10 |
| TCF8 |  | Chromosome 10：31，647，430－31，858，748 | 10 |
| ITGB1 | CD29 | Chromosome 10：33，229，326－33，287，204 | 10 |
| ITGB1 | CD29 | Chromosome 10：33，229，326－33，287，204 | 10 |
| NRP1 | CD304，BDCA4，Neuropil | Chromosome 10：33，506，426－33，665，196 | 10 |
| NRP1 | CD304，BDCA4，Neuropil | Chromosome 10：33，506，426－33，665，196 | 10 |
| CREM | ICER | Chromosome 10：35，455，807－35，541，892 | 10 |
| CREM | ICER | Chromosome 10：35，455，807－35，541，892 | 10 |
| CREM | ICER | Chromosome 10：35，455，807－35，541，892 | 10 |
| CXCL12 | SDF1 | Chromosome 10：44，185，611－44，200，548 | 10 |
| ALOX5 | 5－LO | Chromosome 10：45，189，635－45，261，571 | 10 |
| ALOX5 | 5－LO | Chromosome 10：45，189，635－45，261，571 | 10 |
| PTPN20C |  | Chromosome 10：48，926，216－49，033，022 | 10 |
| MAPK8 | JNK | Chromosome 10：49，184，739－49，317，409 | 10 |
| MAPK8 | JNK | Chromosome 10：49，184，739－49，317，409 | 10 |
| MBL2 |  | Chromosome 10：54，195，146－54，201，466 | 10 |
| UBE2D1 | UBCH5 | Chromosome 10：59，764，745－59，800，515 | 10 |
| UBE2D1 | UBCH5 | Chromosome 10：59，764，745－59，800，515 | 10 |
| IL15RA |  | Chromosome 10：6，034，340－6，060，156 | 10 |
| IL2RA |  | Chromosome 10：6，092，658－6，144，294 | 10 |
| PRKCQ | PKC入 | Chromosome 10：6，509，111－6，662，269 | 10 |
| PRKCQ | PKC入 | Chromosome 10：6，509，111－6，662，269 | 10 |
| PRKCQ | PKC入 | Chromosome 10：6，509，111－6，662，269 | 10 |
| ANK3 |  | Chromosome 10：61，458，165－61，819，494 | 10 |
| ANK3 |  | Chromosome 10：61，458，165－61，819，494 | 10 |
| ANK3 |  | Chromosome 10：61，458，165－61，819，494 | 10 |
| ANK3 |  | Chromosome 10：61，458，165－61，819，494 | 10 |
| ANK3 |  | Chromosome 10：61，458，165－61，819，494 | 10 |
| CDC2 | CDK1 | Chromosome 10：62，205，690－62，224，616 | 10 |
| EGR2 |  | Chromosome 10：64，241，762－64，246，133 | 10 |
| EGR2 |  | Chromosome 10：64，241，762－64，246，133 | 10 |
| DDX50 |  | Chromosome 10：70，331，040－70，376，609 | 10 |
| PRG1 | serglycin | Chromosome 10：70，517，834－70，534，573 | 10 |
| PRF1 | Perforin | Chromosome 10：72，027，110－72，032，521 | 10 |
| PPP3CB | calcineurin | Chromosome 10：74，866，192－74，925，765 | 10 |
| PPP3CB | calcineurin | Chromosome 10：74，866，192－74，925，765 | 10 |
| PLAU | Plasminogen activator | Chromosome 10：75，340，896－75，347，261 | 10 |
| ADK | adenosine kinase | Chromosome 10：75，580，971－76，139，067 | 10 |
| ADK | adenosine kinase | Chromosome 10：75，580，971－76，139，067 | 10 |
| ADK | adenosine kinase | Chromosome 10：75，580，971－76，139，067 | 10 |
| ADK | adenosine kinase | Chromosome 10：75，580，971－76，139，067 | 10 |
| ADK | adenosine kinase | Chromosome 10：75，580，971－76，139，067 | 10 |
| ADK | adenosine kinase | Chromosome 10：75，580，971－76，139，067 | 10 |
| DUSP13 |  | Chromosome 10：76，524，196－76，538，976 | 10 |
| DLG5 |  | Chromosome 10：79，220，557－79，356，384 | 10 |
| DLG5 |  | Chromosome 10：79，220，557－79，356，384 | 10 |
| GATA3 |  | Chromosome 10：8，136，662－8，157，170 | 10 |
| GATA3 |  | Chromosome 10：8，136，662－8，157，170 | 10 |
| PPIF |  | Chromosome 10：80，777，226－80，785，096 | 10 |


| PPIF |  | Chromosome 10:80,777,226-80,785,096 | 10 |
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| SFTPD | SP-D | Chromosome 10:81,687,476-81,698,841 | 10 |
| NRG3 | NRG3 | Chromosome 10:83,624,786-84,736,913 | 10 |
| NRG3 | NRG3 | Chromosome 10:83,624,786-84,736,913 | 10 |
| NRG3 | NRG3 | Chromosome 10:83,624,786-84,736,913 | 10 |
| NRG3 | NRG3 | Chromosome 10:83,624,786-84,736,913 | 10 |
| NRG3 | NRG3 | Chromosome 10:83,624,786-84,736,913 | 10 |
| NRG3 | NRG3 | Chromosome 10:83,624,786-84,736,913 | 10 |
| NRG3 | NRG3 | Chromosome 10:83,624,786-84,736,913 | 10 |
| NRG3 | NRG3 | Chromosome 10:83,624,786-84,736,913 | 10 |
| NRG3 | NRG3 | Chromosome 10:83,624,786-84,736,913 | 10 |
| NRG3 | NRG3 | Chromosome 10:83,624,786-84,736,913 | 10 |
| NRG3 | NRG3 | Chromosome 10:83,624,786-84,736,913 | 10 |
| NRG3 | NRG3 | Chromosome 10:83,624,786-84,736,913 | 10 |
| NRG3 | NRG3 | Chromosome 10:83,624,786-84,736,913 | 10 |
| NRG3 | NRG3 | Chromosome 10:83,624,786-84,736,913 | 10 |
| BMPR1A | CD292 | Chromosome 10:88,506,376-88,674,925 | 10 |
| PTEN |  | Chromosome 10:89,612,850-89,721,667 | 10 |
| PTEN |  | Chromosome 10:89,612,850-89,721,667 | 10 |
| TNFRSF6 | FAS, CD95 | Chromosome 10:90,739,206-90,765,522 | 10 |
| LIPA |  | Chromosome 10:90,963,306-91,164,294 | 10 |
| LIPA |  | Chromosome 10:90,963,306-91,164,294 | 10 |
| LIPA |  | Chromosome 10:90,963,306-91,164,294 | 10 |
| CYP26C1 |  | Chromosome 10:94,811,011-94,818,444 | 10 |
| CYP2C18 |  | Chromosome 10:96,433,368-96,485,937 | 10 |
| CYP2C19 |  | Chromosome 10:96,512,371-96,603,007 | 10 |
| CYP2C19 |  | Chromosome 10:96,512,371-96,603,007 | 10 |
| CYP2C9 |  | Chromosome 10:96,688,418-96,739,137 | 10 |
| CYP2C8 |  | Chromosome 10:96,786,519-96,819,244 | 10 |
| ENTPD1 | CD39 | Chromosome 10:97,461,526-97,619,442 | 10 |
| ENTPD1 | CD39 | Chromosome 10:97,461,526-97,619,442 | 10 |
| ENTPD1 | CD39 | Chromosome 10:97,461,526-97,619,442 | 10 |
| BLNK |  | Chromosome 10:97,941,445-98,021,316 | 10 |
| DNTT |  | Chromosome 10:98,054,075-98,088,311 | 10 |
| DNTT |  | Chromosome 10:98,054,075-98,088,311 | 10 |
| CTSD | cathepsin D | Chromosome 11:1,730,558-1,741,798 | 11 |
| CTSD | cathepsin D | Chromosome 11:1,730,558-1,741,798 | 11 |
| LSP1 |  | Chromosome 11:1,830,776-1,870,069 | 11 |
| ADM |  | Chromosome 11:10,283,172-10,285,499 | 11 |
| PGR | NR3C3 | Chromosome 11:100,414,313-100,506,465 | 11 |
| PGR | NR3C3 | Chromosome 11:100,414,313-100,506,465 | 11 |
| BIRC3 |  | Chromosome 11:101,693,404-101,713,675 | 11 |
| MMP7 |  | Chromosome 11:101,896,449-101,906,688 | 11 |
| MMP20 |  | Chromosome 11:101,952,776-102,001,273 | 11 |
| MMP27 |  | Chromosome 11:102,067,625-102,081,678 | 11 |
| MMP10 |  | Chromosome 11:102,146,444-102,156,569 | 11 |
| MMP3 |  | Chromosome 11:102,211,738-102,219,552 | 11 |
| MMP13 |  | Chromosome 11:102,318,934-102,331,672 | 11 |
| MMP13 |  | Chromosome 11:102,318,934-102,331,672 | 11 |
| CASP12 | Caspase 12 | Chromosome 11:104,261,876-104,274,607 | 11 |
| CASP4 |  | Chromosome 11:104,318,804-104,345,373 | 11 |
| CASP5 |  | Chromosome 11:104,370,180-104,384,909 | 11 |
| ICEBERG |  | Chromosome 11:104,513,879-104,515,663 | 11 |
| ICEBERG |  | Chromosome 11:104,513,879-104,515,663 | 11 |
| ALKBH8 |  | Chromosome 11:106,878,664-106,941,637 | 11 |
| ALKBH8 |  | Chromosome 11:106,878,664-106,941,637 | 11 |


| ATM | Ataxia teleangiectasia mu | Chromosome 11:107,598,769-107,745,036 | 11 |
| :---: | :---: | :---: | :---: |
| ATM | Ataxia teleangiectasia my | Chromosome 11:107,598,769-107,745,036 | 11 |
| ATM | Ataxia teleangiectasia my | Chromosome 11:107,598,769-107,745,036 | 11 |
| ATM | Ataxia teleangiectasia my | Chromosome 11:107,598,769-107,745,036 | 11 |
| DDX10 |  | Chromosome 11:108,041,014-108,316,866 | 11 |
| DDX10 |  | Chromosome 11:108,041,014-108,316,866 | 11 |
| DDX10 |  | Chromosome 11:108,041,014-108,316,866 | 11 |
| RDX |  | Chromosome 11:109,605,376-109,672,647 | 11 |
| RDX |  | Chromosome 11:109,605,376-109,672,647 | 11 |
| POU2AF1 |  | Chromosome 11:110,728,190-110,755,627 | 11 |
| POU2AF1 |  | Chromosome 11:110,728,190-110,755,627 | 11 |
| IL18 |  | Chromosome 11:111,519,186-111,540,050 | 11 |
| IL18 |  | Chromosome 11:111,519,186-111,540,050 | 11 |
| NCAM1 | CD56 | Chromosome 11:112,337,368-112,653,781 | 11 |
| NCAM1 | CD56 | Chromosome 11:112,337,368-112,653,781 | 11 |
| NCAM1 | CD56 | Chromosome 11:112,337,368-112,653,781 | 11 |
| CADM1 |  | Chromosome 11:114,550,227-114,880,325 | 11 |
| CADM1 |  | Chromosome 11:114,550,227-114,880,325 | 11 |
| CADM1 |  | Chromosome 11:114,550,227-114,880,325 | 11 |
| CADM1 |  | Chromosome 11:114,550,227-114,880,325 | 11 |
| CADM1 |  | Chromosome 11:114,550,227-114,880,325 | 11 |
| CADM1 |  | Chromosome 11:114,550,227-114,880,325 | 11 |
| IL10RA | CDw210a, IL10R1 | Chromosome 11:117,362,319-117,377,404 | 11 |
| IL10RA | CDw210a, IL10R1 | Chromosome 11:117,362,319-117,377,404 | 11 |
| CD3E | CD3 | Chromosome 11:117,680,662-117,692,100 | 11 |
| CD3D | CD3 | Chromosome 11:117,710,475-117,718,669 | 11 |
| DDX6 |  | Chromosome 11:118,125,623-118,167,082 | 11 |
| DDX6 |  | Chromosome 11:118,125,623-118,167,082 | 11 |
| BLR1 | CD185, CXCR5 | Chromosome 11:118,259,777-118,272,181 | 11 |
| BLR1 | CD185, CXCR5 | Chromosome 11:118,259,777-118,272,181 | 11 |
| MCAM | CD146 | Chromosome 11:118,684,444-118,693,050 | 11 |
| PVRL1 | CD111 | Chromosome 11:119,014,018-119,104,645 | 11 |
| PVRL1 | CD111 | Chromosome 11:119,014,018-119,104,645 | 11 |
| POU2F3 |  | Chromosome 11:119,616,256-119,695,863 | 11 |
| POU2F3 |  | Chromosome 11:119,616,256-119,695,863 | 11 |
| POU2F3 |  | Chromosome 11:119,616,256-119,695,863 | 11 |
| SORL1 |  | Chromosome 11:120,828,130-121,005,621 | 11 |
| SORL1 |  | Chromosome 11:120,828,130-121,005,621 | 11 |
| DDX25 |  | Chromosome 11:125,279,550-125,298,215 | 11 |
| DDX25 |  | Chromosome 11:125,279,550-125,298,215 | 11 |
| TIRAP | Mal | Chromosome 11:125,658,192-125,672,683 | 11 |
| TIRAP | Mal | Chromosome 11:125,658,192-125,672,683 | 11 |
| PTH |  | Chromosome 11:13,470,177-13,474,143 | 11 |
| IGSF9B |  | Chromosome 11:133,290,395-133,327,321 | 11 |
| IGSF9B |  | Chromosome 11:133,290,395-133,327,321 | 11 |
| IGSF9B |  | Chromosome 11:133,290,395-133,327,321 | 11 |
| JAM3 | CD323 | Chromosome 11:133,444,030-133,526,861 | 11 |
| JAM3 | CD323 | Chromosome 11:133,444,030-133,526,861 | 11 |
| B3GAT1 | CD57 | Chromosome 11:133,753,608-133,787,022 | 11 |
| CYP2R1 |  | Chromosome 11:14,856,131-14,870,327 | 11 |
| PIK3C2A |  | Chromosome 11:17,067,861-17,147,864 | 11 |
| PIK3C2A |  | Chromosome 11:17,067,861-17,147,864 | 11 |
| SAA3P | Saa3 | Chromosome 11:18,090,596-18,094,695 | 11 |
| SAA2 |  | Chromosome 11:18,223,365-18,226,758 | 11 |
| SAA2 |  | Chromosome 11:18,223,365-18,226,758 | 11 |
| GTF2H1 |  | Chromosome 11:18,300,719-18,345,153 | 11 |


| IGSF22 |  | Chromosome 11:18,682,435-18,704,353 | 11 |
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| PTPN5 |  | Chromosome 11:18,706,051-18,769,965 | 11 |
| IGF2 | IGF2 | Chromosome 11:2,106,918-2,125,616 | 11 |
| CD81 | TAPA1 | Chromosome 11:2,355,096-2,375,225 | 11 |
| CD81 | TAPA1 | Chromosome 11:2,355,096-2,375,225 | 11 |
| CDKN1C |  | Chromosome 11:2,861,019-2,863,577 | 11 |
| PRMT3 | PRMT3 | Chromosome 11:20,365,679-20,487,404 | 11 |
| PRMT3 | PRMT3 | Chromosome 11:20,365,679-20,487,404 | 11 |
| GAS2 | GAS2 | Chromosome 11:22,646,230-22,791,123 | 11 |
| GAS2 | GAS2 | Chromosome 11:22,646,230-22,791,123 | 11 |
| GAS2 | GAS2 | Chromosome 11:22,646,230-22,791,123 | 11 |
| NLRP6 | NALP6, PYPAF5, PAN3 | Chromosome 11:268,570-275,304 | 11 |
| NLRP6 | NALP6, PYPAF5, PAN3 | Chromosome 11:268,570-275,304 | 11 |
| BDNF | BDNF | Chromosome 11:27,633,016-27,699,872 | 11 |
| BDNF | BDNF | Chromosome 11:27,633,016-27,699,872 | 11 |
| ART1 | CHAT1, ADP ribosyl trans | Chromosome 11:3,622,937-3,642,222 | 11 |
| CD59 | protectin, MIRL | Chromosome 11:33,681,132-33,714,600 | 11 |
| CD59 | protectin, MIRL | Chromosome 11:33,681,132-33,714,600 | 11 |
| CAT | Catalase | Chromosome 11:34,417,054-34,450,183 | 11 |
| CD44 |  | Chromosome 11:35,116,993-35,210,525 | 11 |
| CD44 |  | Chromosome 11:35,116,993-35,210,525 | 11 |
| CD44 |  | Chromosome 11:35,116,993-35,210,525 | 11 |
| TRAF6 |  | Chromosome 11:36,467,299-36,488,398 | 11 |
| RAG1 |  | Chromosome 11:36,546,139-36,557,877 | 11 |
| SIGIRR |  | Chromosome 11:395,716-407,397 | 11 |
| SIGIRR |  | Chromosome 11:395,716-407,397 | 11 |
| MMP26 |  | Chromosome 11:4,745,076-4,970,235 | 11 |
| MMP26 |  | Chromosome 11:4,745,076-4,970,235 | 11 |
| MMP26 |  | Chromosome 11:4,745,076-4,970,235 | 11 |
| MMP26 |  | Chromosome 11:4,745,076-4,970,235 | 11 |
| API5 |  | Chromosome 11:43,290,109-43,322,655 | 11 |
| ALKBH3 |  | Chromosome 11:43,858,971-43,898,392 | 11 |
| CD82 | CD82, KAI1 | Chromosome 11:44,543,717-44,597,915 | 11 |
| CD82 | CD82, KAI1 | Chromosome 11:44,543,717-44,597,915 | 11 |
| MAPK8IP1 |  | Chromosome 11:45,863,778-45,884,592 | 11 |
| MAPK8IP1 |  | Chromosome 11:45,863,778-45,884,592 | 11 |
| F2 | Alpha-Thrombin, prothron | Chromosome 11:46,697,331-46,717,631 | 11 |
| F2 | Alpha-Thrombin, prothron | Chromosome 11:46,697,331-46,717,631 | 11 |
| NR1H3 | NR1H3 | Chromosome 11:47,227,083-47,246,972 | 11 |
| NR1H3 | NR1H3 | Chromosome 11:47,227,083-47,246,972 | 11 |
| NDUFS3 | NADH dehydrogenase | Chromosome 11:47,543,464-47,562,690 | 11 |
| NDUFS3 | NADH dehydrogenase | Chromosome 11:47,543,464-47,562,690 | 11 |
| C1QTNF4 |  | Chromosome 11:47,567,792-47,580,516 | 11 |
| PTPRJ | CD148 | Chromosome 11:47,958,689-48,146,246 | 11 |
| PTPRJ | CD148 | Chromosome 11:47,958,689-48,146,246 | 11 |
| HBD | Hbb-b1 | Chromosome 11:5,203,270-5,212,454 | 11 |
| HBD | Hbb-b1 | Chromosome 11:5,203,270-5,212,454 | 11 |
| TRIM22 |  | Chromosome 11:5,667,495-5,688,669 | 11 |
| TRIM22 |  | Chromosome 11:5,667,495-5,688,669 | 11 |
| TRIM22 |  | Chromosome 11:5,667,495-5,688,669 | 11 |
| AGTRL1 | apelin receptor | Chromosome 11:56,757,630-56,761,489 | 11 |
| P2RX3 | P2X3 | Chromosome 11:56,862,525-56,894,125 | 11 |
| PRG2 |  | Chromosome 11:56,910,832-56,914,706 | 11 |
| UBE2L6 | Ubch8 | Chromosome 11:57,075,705-57,092,333 | 11 |
| SERPING1 |  | Chromosome 11:57,121,603-57,138,902 | 11 |
| SERPING1 |  | Chromosome 11:57,121,603-57,138,902 | 11 |


| MS4A3 |  | Chromosome 11:59,580,677-59,595,164 | 11 |
| :---: | :---: | :---: | :---: |
| MS4A5 |  | Chromosome 11:59,953,638-59,971,841 | 11 |
| ILK | Integrin linked kinase | Chromosome 11:6,581,540-6,588,677 | 11 |
| ILK | Integrin linked kinase | Chromosome 11:6,581,540-6,588,677 | 11 |
| ILK | Integrin linked kinase | Chromosome 11:6,581,540-6,588,677 | 11 |
| GPR44 | CRTH2 | Chromosome 11:60,374,983-60,380,020 | 11 |
| CD5 | Leu1 | Chromosome 11:60,626,543-60,651,900 | 11 |
| SYT7 | SytVII | Chromosome 11:61,039,361-61,104,874 | 11 |
| SYT7 | SytVII | Chromosome 11:61,039,361-61,104,874 | 11 |
| FEN1 | ok | Chromosome 11:61,316,726-61,321,286 | 11 |
| NXF1 | TAP | Chromosome 11:62,316,219-62,329,529 | 11 |
| SLC3A2 | CD98 | Chromosome 11:62,380,094-62,412,929 | 11 |
| LGALS12 | galectin12 | Chromosome 11:63,030,132-63,040,815 | 11 |
| VEGFB |  | Chromosome 11:63,758,646-63,762,835 | 11 |
| FKBP2 | Calcineurin | Chromosome 11:63,764,989-63,768,262 | 11 |
| ESRRA | NR3B1 | Chromosome 11:63,829,616-63,840,786 | 11 |
| RASGRP2 | CalDAG-GEFI | Chromosome 11:64,250,959-64,269,504 | 11 |
| GPHA2 | GPHA2 | Chromosome 11:64,458,519-64,459,936 | 11 |
| GPHA2 | GPHA2 | Chromosome 11:64,458,519-64,459,936 | 11 |
| POLA2 |  | Chromosome 11:64,786,006-64,821,664 | 11 |
| LTBP3 |  | Chromosome 11:65,062,850-65,082,275 | 11 |
| LTBP3 |  | Chromosome 11:65,062,850-65,082,275 | 11 |
| RELA | p65 | Chromosome 11:65,177,649-65,186,959 | 11 |
| CFL1 | cofilin | Chromosome 11:65,378,858-65,383,462 | 11 |
| CFL1 | cofilin | Chromosome 11:65,378,858-65,383,462 | 11 |
| CD164L1 | CD248, endosialin, TEM1 | Chromosome 11:65,838,534-65,841,091 | 11 |
| PELI3 | Pellino3 | Chromosome 11:65,990,974-66,001,382 | 11 |
| ACTN3 | alpha actinin 3 | Chromosome 11:66,070,967-66,087,373 | 11 |
| CTSF | Cathepsin F | Chromosome 11:66,087,511-66,092,623 | 11 |
| RHOD | RHOD | Chromosome 11:66,580,897-66,596,060 | 11 |
| RHOD | RHOD | Chromosome 11:66,580,897-66,596,060 | 11 |
| CLCF1 | cardiotrophin-like cytokin¢ | Chromosome 11:66,888,215-66,897,782 | 11 |
| CLCF1 | cardiotrophin-like cytokin¢ | Chromosome 11:66,888,215-66,897,782 | 11 |
| AIP |  | Chromosome 11:67,007,097-67,015,150 | 11 |
| GSTP1 |  | Chromosome 11:67,107,862-67,110,701 | 11 |
| UNC93B1 |  | Chromosome 11:67,515,151-67,528,169 | 11 |
| TCIRG1 | V-ATPase | Chromosome 11:67,563,059-67,574,942 | 11 |
| LRP5 |  | Chromosome 11:67,836,674-67,973,317 | 11 |
| LRP5 |  | Chromosome 11:67,836,674-67,973,317 | 11 |
| LRP5 |  | Chromosome 11:67,836,674-67,973,317 | 11 |
| LRP5 |  | Chromosome 11:67,836,674-67,973,317 | 11 |
| LRP5 |  | Chromosome 11:67,836,674-67,973,317 | 11 |
| CCND1 | cyclinD1 | Chromosome 11:69,165,054-69,178,423 | 11 |
| FGF4 | FGF4 | Chromosome 11:69,296,978-69,299,352 | 11 |
| FADD | MC159 | Chromosome 11:69,726,917-69,731,144 | 11 |
| NLRP14 | NALP14, LRR, NOD5, G | Chromosome 11:7,016,373-7,049,333 | 11 |
| NLRP14 | NALP14, LRR, NOD5, G | Chromosome 11:7,016,373-7,049,333 | 11 |
| NLRP10 | NALP10, PYNOD, PAN5, | Chromosome 11:7,937,547-7,941,780 | 11 |
| NLRP10 | NALP10, PYNOD, PAN5, | Chromosome 11:7,937,547-7,941,780 | 11 |
| DFB108B |  | Chromosome 11:71,221,894-71,226,256 | 11 |
| IL18BP |  | Chromosome 11:71,387,587-71,394,409 | 11 |
| P2RY2 | P2Y2 | Chromosome 11:72,606,992-72,625,045 | 11 |
| P2RY2 | P2Y2 | Chromosome 11:72,606,992-72,625,045 | 11 |
| P2RY2 | P2Y2 | Chromosome 11:72,606,992-72,625,045 | 11 |
| TNFRSF19L |  | Chromosome 11:72,765,053-72,786,167 | 11 |
| PLEKHB1 |  | Chromosome 11:73,023,592-73,051,512 | 11 |


| POLD3 |  | Chromosome 11:73,981,277-74,031,413 | 11 |
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| ARRB1 | Beta arrestin 1 | Chromosome 11:74,654,130-74,740,521 | 11 |
| ARRB1 | Beta arrestin 1 | Chromosome 11:74,654,130-74,740,521 | 11 |
| UVRAG |  | Chromosome 11:75,203,923-75,531,342 | 11 |
| UVRAG |  | Chromosome 11:75,203,923-75,531,342 | 11 |
| UVRAG |  | Chromosome 11:75,203,923-75,531,342 | 11 |
| UVRAG |  | Chromosome 11:75,203,923-75,531,342 | 11 |
| PRKRIR | PKR | Chromosome 11:75,738,652-75,769,528 | 11 |
| PRKRIR | PKR | Chromosome 11:75,738,652-75,769,528 | 11 |
| PAK1 | PAK | Chromosome 11:76,710,708-76,862,581 | 11 |
| PAK1 | PAK | Chromosome 11:76,710,708-76,862,581 | 11 |
| GAB2 |  | Chromosome 11:77,603,990-77,806,414 | 11 |
| GAB2 |  | Chromosome 11:77,603,990-77,806,414 | 11 |
| GAB2 |  | Chromosome 11:77,603,990-77,806,414 | 11 |
| CD151 |  | Chromosome 11:822,952-828,835 | 11 |
| FZD4 | CD344, FZ4 | Chromosome 11:86,334,369-86,344,081 | 11 |
| FZD4 | CD344, FZ4 | Chromosome 11:86,334,369-86,344,081 | 11 |
| TYR |  | Chromosome 11:88,550,268-88,668,474 | 11 |
| TYR |  | Chromosome 11:88,550,268-88,668,474 | 11 |
| TYR |  | Chromosome 11:88,550,268-88,668,474 | 11 |
| NOX4 |  | Chromosome 11:88,699,160-88,864,301 | 11 |
| MRE11A |  | Chromosome 11:93,790,114-93,866,688 | 11 |
| MRE11A |  | Chromosome 11:93,790,114-93,866,688 | 11 |
| CLEC12A | MICL | Chromosome 12:10,015,281-10,029,461 | 12 |
| CLEC12A | MICL | Chromosome 12:10,015,281-10,029,461 | 12 |
| CLEC1A |  | Chromosome 12:10,113,421-10,142,872 | 12 |
| OLR1 | SCARE1, LOX1, LDL rec | Chromosome 12:10,202,167-10,216,004 | 12 |
| OLR1 | SCARE1, LOX1, LDL rec | Chromosome 12:10,202,167-10,216,004 | 12 |
| KLRD1 | CD94/NKG2A | Chromosome 12:10,351,684-10,359,983 | 12 |
| KLRC3 |  | Chromosome 12:10,456,181-10,464,461 | 12 |
| KLRA1 | Ly49 | Chromosome 12:10,633,039-10,643,431 | 12 |
| CSDA |  | Chromosome 12:10,742,955-10,767,171 | 12 |
| IGF1 | IGF1 | Chromosome 12:101,313,806-101,398,471 | 12 |
| IGF1 | IGF1 | Chromosome 12:101,313,806-101,398,471 | 12 |
| ASCL1 | MASH1 | Chromosome 12:101,875,594-101,878,421 | 12 |
| ASCL1 | MASH1 | Chromosome 12:101,875,594-101,878,421 | 12 |
| FEEL-2 | FEEL-2 | Chromosome 12:102,505,181-102,684,635 | 12 |
| FEEL-2 | FEEL-2 | Chromosome 12:102,505,181-102,684,635 | 12 |
| FEEL-2 | FEEL-2 | Chromosome 12:102,505,181-102,684,635 | 12 |
| FEEL-2 | FEEL-2 | Chromosome 12:102,505,181-102,684,635 | 12 |
| FEEL-2 | FEEL-2 | Chromosome 12:102,505,181-102,684,635 | 12 |
| HSP90B1 | gp96 phox | Chromosome 12:102,848,290-102,865,833 | 12 |
| NFYB |  | Chromosome 12:103,034,988-103,056,170 | 12 |
| NFYB |  | Chromosome 12:103,034,988-103,056,170 | 12 |
| TXNRD1 | TrxR alpha | Chromosome 12:103,204,857-103,268,192 | 12 |
| RFX4 | RFX4 | Chromosome 12:105,501,163-105,680,711 | 12 |
| RFX4 | RFX4 | Chromosome 12:105,501,163-105,680,711 | 12 |
| RFX4 | RFX4 | Chromosome 12:105,501,163-105,680,711 | 12 |
| CMKLR1 | ChemR23 | Chromosome 12:107,208,800-107,257,218 | 12 |
| SELPLG | CD162, PSGL1, CLA, 6-S | Chromosome 12:107,539,800-107,551,799 | 12 |
| ALKBH2 |  | Chromosome 12:108,010,379-108,015,660 | 12 |
| ALKBH2 |  | Chromosome 12:108,010,379-108,015,660 | 12 |
| FOXN4 |  | Chromosome 12:108,200,167-108,231,408 | 12 |
| FOXN4 |  | Chromosome 12:108,200,167-108,231,408 | 12 |
| RAD9B |  | Chromosome 12:109,424,388-109,454,274 | 12 |
| RAD9B |  | Chromosome 12:109,424,388-109,454,274 | 12 |


| ETV6 |  | Chromosome 12:11,694,055-11,939,603 | 12 |
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| ETV6 |  | Chromosome 12:11,694,055-11,939,603 | 12 |
| ETV6 |  | Chromosome 12:11,694,055-11,939,603 | 12 |
| ETV6 |  | Chromosome 12:11,694,055-11,939,603 | 12 |
| PTPN11 | SHP2 | Chromosome 12:111,340,919-111,432,100 | 12 |
| PTPN11 | SHP2 | Chromosome 12:111,340,919-111,432,100 | 12 |
| OAS1 |  | Chromosome 12:111,829,122-111,854,374 | 12 |
| DTX1 | DELTEX | Chromosome 12:111,980,045-112,020,216 | 12 |
| DTX1 | DELTEX | Chromosome 12:111,980,045-112,020,216 | 12 |
| HRK | harakiri | Chromosome 12:115,783,410-115,803,615 | 12 |
| HRK | harakiri | Chromosome 12:115,783,410-115,803,615 | 12 |
| NOS1 | NO synthase | Chromosome 12:116,135,362-116,283,965 | 12 |
| NOS1 | NO synthase | Chromosome 12:116,135,362-116,283,965 | 12 |
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| HSP B8 |  | Chromosome 12:118,100,978-118,116,934 | 12 |
| HSP B8 |  | Chromosome 12:118,100,978-118,116,934 | 12 |
| OASL |  | Chromosome 12:119,942,478-119,961,164 | 12 |
| OASL |  | Chromosome 12:119,942,478-119,961,164 | 12 |
| BCL2L14 |  | Chromosome 12:12,115,145-12,255,214 | 12 |
| BCL2L14 |  | Chromosome 12:12,115,145-12,255,214 | 12 |
| LRP6 |  | Chromosome 12:12,164,953-12,311,013 | 12 |
| DUSP16 |  | Chromosome 12:12,520,098-12,606,584 | 12 |
| DUSP16 |  | Chromosome 12:12,520,098-12,606,584 | 12 |
| DUSP16 |  | Chromosome 12:12,520,098-12,606,584 | 12 |
| DDX47 |  | Chromosome 12:12,770,130-12,874,182 | 12 |
| P2RX7 | P2X7, P2Z | Chromosome 12:120,055,061-120,108,259 | 12 |
| P2RX7 | P2X7, P2Z | Chromosome 12:120,055,061-120,108,259 | 12 |
| IL31 |  | Chromosome 12:121,222,530-121,224,699 | 12 |
| CLIP1 | RSN | Chromosome 12:121,321,934-121,473,069 | 12 |
| CLIP1 | RSN | Chromosome 12:121,321,934-121,473,069 | 12 |
| GPR109B | HM74 | Chromosome 12:121,765,256-121,767,297 | 12 |
| DDX55 |  | Chromosome 12:122,652,625-122,671,435 | 12 |
| GTF2H3 | TFIIH, TFIIK | Chromosome 12:122,684,333-122,711,573 | 12 |
| NCOR2 | SMRT | Chromosome 12:123,374,914-123,568,793 | 12 |
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| NCOR2 | SMRT | Chromosome 12:123,374,914-123,568,793 | 12 |
| SCARB1 | SR-BI | Chromosome 12:123,828,129-123,914,346 | 12 |
| SCARB1 |  | Chromosome 12:123,828,129-123,914,346 | 12 |
| DHX37 |  | Chromosome 12:123,997,325-124,039,620 | 12 |
| NLRP9P | NOD25 | Chromosome 12:128,063,805-128,067,640 | 12 |
| FZD10 | CD350 | Chromosome 12:129,212,957-129,216,238 | 12 |
| FZD10 | CD350 | Chromosome 12:129,212,957-129,216,238 | 12 |
| RAN |  | Chromosome 12:129,922,521-129,927,316 | 12 |
| P2RX2 | P2X2 | Chromosome 12:131,705,476-131,709,045 | 12 |
| ART4 | CD297 | Chromosome 12:14,873,512-14,887,680 | 12 |
| ART4 | CD297 | Chromosome 12:14,873,512-14,887,680 | 12 |
| ARHGDIB | Rho GD2 | Chromosome 12:14,986,217-15,005,870 | 12 |
| ARHGDIB | Rho GD2 | Chromosome 12:14,986,217-15,005,870 | 12 |
| PTPRO | PTPROt | Chromosome 12:15,366,754-15,641,602 | 12 |
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| PTPRO | PTPROt | Chromosome 12:15,366,754-15,641,602 | 12 |
| PIK3C2G |  | Chromosome 12:18,305,741-18,692,617 | 12 |
| PIK3C2G |  | Chromosome 12:18,305,741-18,692,617 | 12 |


| PIK3C2G |  | Chromosome 12:18,305,741-18,692,617 | 12 |
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| PIK3C2G |  | Chromosome 12:18,305,741-18,692,617 | 12 |
| FKBP4 | Calcineurin | Chromosome 12:2,774,414-2,783,385 | 12 |
| FKBP4 | Calcineurin | Chromosome 12:2,774,414-2,783,385 | 12 |
| KRAS |  | Chromosome 12:25,249,447-25,295,121 | 12 |
| KRAS |  | Chromosome 12:25,249,447-25,295,121 | 12 |
| ITPR2 |  | Chromosome 12:26,377,193-26,877,398 | 12 |
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| PTHLH | PTHrP | Chromosome 12:28,002,284-28,016,183 | 12 |
| PTHLH | PTHrP | Chromosome 12:28,002,284-28,016,183 | 12 |
| PTHLH | PTHrP | Chromosome 12:28,002,284-28,016,183 | 12 |
| ERGIC2 |  | Chromosome 12:29,381,556-29,425,410 | 12 |
| ERGIC2 |  | Chromosome 12:29,381,556-29,425,410 | 12 |
| PRMT8 | PRMT8 | Chromosome 12:3,470,686-3,573,400 | 12 |
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| DDX11 |  | Chromosome 12:31,118,061-31,148,992 | 12 |
| DDX11 |  | Chromosome 12:31,118,061-31,148,992 | 12 |
| CCND2 |  | Chromosome 12:4,253,199-4,284,777 | 12 |
| FGF6 | FGF6 | Chromosome 12:4,413,569-4,425,041 | 12 |
| DYRK4 |  | Chromosome 12:4,569,505-4,593,302 | 12 |
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| IRAK4 |  | Chromosome 12:42,439,047-42,468,166 | 12 |
| IRAK4 |  | Chromosome 12:42,439,047-42,468,166 | 12 |
| SFRS2IP | Caspase 11 | Chromosome 12:44,601,459-44,670,615 | 12 |
| SFRS2IP | Caspase 11 | Chromosome 12:44,601,459-44,670,615 | 12 |
| HDAC7A |  | Chromosome 12:46,462,772-46,499,924 | 12 |
| HDAC7A |  | Chromosome 12:46,462,772-46,499,924 | 12 |
| DDX23 |  | Chromosome 12:47,509,806-47,532,224 | 12 |
| DDX23 |  | Chromosome 12:47,509,806-47,532,224 | 12 |
| WNT1 |  | Chromosome 12:47,658,503-47,662,746 | 12 |
| DHH | desert Hh | Chromosome 12:47,769,471-47,774,869 | 12 |
| TUBA1A |  | Chromosome 12:47,864,847-47,869,153 | 12 |
| C1QL4 |  | Chromosome 12:48,012,467-48,017,238 | 12 |
| MCRS1 | Mcrs1 | Chromosome 12:48,238,352-48,248,178 | 12 |
| ATF1 |  | Chromosome 12:49,444,128-49,500,328 | 12 |
| ATF1 |  | Chromosome 12:49,444,128-49,500,328 | 12 |
| ATF1 |  | Chromosome 12:49,444,128-49,500,328 | 12 |
| VWF | vWf | Chromosome 12:5,928,301-6,104,097 | 12 |
| VWF | vWf | Chromosome 12:5,928,301-6,104,097 | 12 |
| VWF | vWf | Chromosome 12:5,928,301-6,104,097 | 12 |
| VWF | vWf | Chromosome 12:5,928,301-6,104,097 | 12 |
| VWF | vWf | Chromosome 12:5,928,301-6,104,097 | 12 |
| ELA1 |  | Chromosome 12:50,008,494-50,026,730 | 12 |
| KRT6A | Keratin 6 | Chromosome 12:51,167,231-51,173,289 | 12 |
| KRT8 | CARD2 | Chromosome 12:51,577,238-51,585,127 | 12 |
| KRT8 | CARD2 | Chromosome 12:51,577,238-51,585,127 | 12 |
| ITGB7 | LPAM (integrin alpha4/be | Chromosome 12:51,871,374-51,887,267 | 12 |
| RARG | RARgamma, NR1B3 | Chromosome 12:51,890,621-51,912,253 | 12 |


| SP1 |  | Chromosome 12:52,060,246-52,096,497 | 12 |
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| SP1 |  | Chromosome 12:52,060,246-52,096,497 | 12 |
| HOXC11 | HOX cluster | Chromosome 12:52,653,177-52,656,470 | 12 |
| NFE2 | NF-E2 p 45 | Chromosome 12:52,972,162-52,981,058 | 12 |
| NFE2 | NF-E2 p 45 | Chromosome 12:52,972,162-52,981,058 | 12 |
| ITGA5 | CD49e | Chromosome 12:53,075,312-53,099,317 | 12 |
| DCD | dermicidin | Chromosome 12:53,324,642-53,328,416 | 12 |
| DCD | dermicidin | Chromosome 12:53,324,642-53,328,416 | 12 |
| ITGA7 |  | Chromosome 12:54,364,619-54,387,949 | 12 |
| ITGA7 |  | Chromosome 12:54,364,619-54,387,949 | 12 |
| DGK alpha |  | Chromosome 12:54,611,213-54,634,074 | 12 |
| SILV | SIL | Chromosome 12:54,634,156-54,646,765 | 12 |
| IL23A |  | Chromosome 12:55,018,926-55,020,461 | 12 |
| NACA | NAC | Chromosome 12:55,392,484-55,407,248 | 12 |
| NACA | NAC | Chromosome 12:55,392,484-55,407,248 | 12 |
| NAB2 |  | Chromosome 12:55,769,157-55,775,526 | 12 |
| SHMT2 | serine hydroxymethyltran | Chromosome 12:55,909,819-55,914,981 | 12 |
| SHMT2 | serine hydroxymethyltran | Chromosome 12:55,909,819-55,914,981 | 12 |
| INHBC | inhibin | Chromosome 12:56,114,810-56,130,876 | 12 |
| INHBE | inhibin | Chromosome 12:56,135,363-56,138,058 | 12 |
| CDK4 |  | Chromosome 12:56,428,270-56,432,431 | 12 |
| CTDSP2 | NIF2 | Chromosome 12:56,499,977-56,527,014 | 12 |
| CD9 | MRP1 | Chromosome 12:6,179,134-6,217,688 | 12 |
| TNFRSF1A | CD120a | Chromosome 12:6,308,184-6,321,522 | 12 |
| TNFRSF1A | CD120a | Chromosome 12:6,308,184-6,321,522 | 12 |
| LTBR | LTbetaR | Chromosome 12:6,363,595-6,370,994 | 12 |
| GAPDH | GAPD | Chromosome 12:6,513,872-6,517,797 | 12 |
| GAPDH | GAPD | Chromosome 12:6,513,872-6,517,797 | 12 |
| CD4 | OKT4, Leu3a | Chromosome 12:6,768,912-6,800,237 | 12 |
| GNB3 | G-protein beta | Chromosome 12:6,819,636-6,826,819 | 12 |
| IRAK3 | IRAK M | Chromosome 12:64,869,270-64,928,684 | 12 |
| IRAK3 | IRAK M | Chromosome 12:64,869,270-64,928,684 | 12 |
| DYRK2 |  | Chromosome 12:66,329,021-66,340,410 | 12 |
| DYRK2 |  | Chromosome 12:66,329,021-66,340,410 | 12 |
| IFNG |  | Chromosome 12:66,834,816-66,839,790 | 12 |
| IFNG |  | Chromosome 12:66,834,816-66,839,790 | 12 |
| IFNG |  | Chromosome 12:66,834,816-66,839,790 | 12 |
| IFNG | Interferon gamma | Chromosome 12:66,834,816-66,839,790 | 12 |
| IL22 |  | Chromosome 12:66,928,292-66,933,651 | 12 |
| MDM2 |  | Chromosome 12:67,488,247-67,520,481 | 12 |
| MDM2 |  | Chromosome 12:67,488,247-67,520,481 | 12 |
| PTPRB |  | Chromosome 12:69,201,231-69,317,469 | 12 |
| PTPRB |  | Chromosome 12:69,201,231-69,317,469 | 12 |
| PTPRR |  | Chromosome 12:69,318,129-69,600,853 | 12 |
| PTPRR |  | Chromosome 12:69,318,129-69,600,853 | 12 |
| C1RL |  | Chromosome 12:7,138,291-7,153,069 | 12 |
| CLEC4C | BDCA2 | Chromosome 12:7,773,278-7,793,336 | 12 |
| CLEC4C | BDCA2 | Chromosome 12:7,773,278-7,793,336 | 12 |
| PAWR | PRKC1 | Chromosome 12:78,509,876-78,608,921 | 12 |
| PTPRQ |  | Chromosome 12:79,318,597-79,598,099 | 12 |
| PTPRQ |  | Chromosome 12:79,318,597-79,598,099 | 12 |
| PTPRQ |  | Chromosome 12:79,318,597-79,598,099 | 12 |
| FOXJ2 |  | Chromosome 12:8,076,626-8,099,385 | 12 |
| FOXJ2 |  | Chromosome 12:8,076,626-8,099,385 | 12 |
| CLEC4a | DCIR | Chromosome 12:8,167,493-8,182,470 | 12 |
| CLEC4D |  | Chromosome 12:8,557,403-8,566,229 | 12 |


| A2ML1 | alpha 2 microglobulin-like | Chromosome 12:8,866,484-8,920,646 | 12 |
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| KITLG | Stem cell factor=SCF | Chromosome 12:87,410,697-87,498,369 | 12 |
| KITLG | Stem cell factor=SCF | Chromosome 12:87,410,697-87,498,369 | 12 |
| DUSP6 |  | Chromosome 12:88,265,968-88,270,427 | 12 |
| DUSP6 |  | Chromosome 12:88,265,968-88,270,427 | 12 |
| KLRG1 |  | Chromosome 12:9,033,484-9,054,610 | 12 |
| DDX12 |  | Chromosome 12:9,460,894-9,492,092 | 12 |
| DDX12 |  | Chromosome 12:9,460,894-9,492,092 | 12 |
| DDX12 |  | Chromosome 12:9,460,894-9,492,092 | 12 |
| KLRB1 | CD161 | Chromosome 12:9,638,415-9,651,764 | 12 |
| KLRF1 |  | Chromosome 12:9,871,344-9,888,871 | 12 |
| DCN | Decorin | Chromosome 12:90,063,166-90,100,937 | 12 |
| DCN | Decorin | Chromosome 12:90,063,166-90,100,937 | 12 |
| BTG1 |  | Chromosome 12:91,061,030-91,063,751 | 12 |
| EEA1 |  | Chromosome 12:91,693,257-91,847,138 | 12 |
| EEA1 |  | Chromosome 12:91,693,257-91,847,138 | 12 |
| UBE2N | UBC13 | Chromosome 12:92,326,219-92,360,157 | 12 |
| SOCS2 |  | Chromosome 12:92,487,729-92,494,109 | 12 |
| SOCS2 |  | Chromosome 12:92,487,729-92,494,109 | 12 |
| CRADD | CED-3, RAIDD | Chromosome 12:92,595,282-92,768,663 | 12 |
| CRADD | CED-3, RAIDD | Chromosome 12:92,595,282-92,768,663 | 12 |
| CRADD | CED-3, RAIDD | Chromosome 12:92,595,282-92,768,663 | 12 |
| CRADD | CED-3, RAIDD | Chromosome 12:92,595,282-92,768,663 | 12 |
| PLXNC1 | CD232, Plexin C1 | Chromosome 12:93,066,630-93,223,356 | 12 |
| PLXNC1 | CD232, Plexin C1 | Chromosome 12:93,066,630-93,223,356 | 12 |
| PLXNC1 | CD232, Plexin C1 | Chromosome 12:93,066,630-93,223,356 | 12 |
| NR2C1 | NR2C1 | Chromosome 12:93,939,802-93,991,487 | 12 |
| LTA4H |  | Chromosome 12:94,918,742-94,953,496 | 12 |
| LTA4H |  | Chromosome 12:94,918,742-94,953,496 | 12 |
| TMPO | thymopoietin | Chromosome 12:97,433,527-97,468,250 | 12 |
| TMPO | thymopoietin | Chromosome 12:97,433,527-97,468,250 | 12 |
| APAF1 | CED-4 | Chromosome 12:97,563,209-97,653,342 | 12 |
| APAF1 | CED-4 | Chromosome 12:97,563,209-97,653,342 | 12 |
| APAF1 | CED-4 | Chromosome 12:97,563,209-97,653,342 | 12 |
| NR1H4 | NR1H4 | Chromosome 12:99,391,810-99,481,774 | 12 |
| NR1H4 | NR1H4 | Chromosome 12:99,391,810-99,481,774 | 12 |
| NR1H4 | NR1H4 | Chromosome 12:99,391,810-99,481,774 | 12 |
| NR1H4 | NR1H4 | Chromosome 12:99,391,810-99,481,774 | 12 |
| ITGBL1 |  | Chromosome 13:100,902,857-101,169,146 | 13 |
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| ITGBL1 |  | Chromosome 13:100,902,857-101,169,146 | 13 |
| ITGBL1 |  | Chromosome 13:100,902,857-101,169,146 | 13 |
| FGF14 | FGF14 | Chromosome 13:101,169,308-101,852,156 | 13 |
| FGF14 | FGF14 | Chromosome 13:101,169,308-101,852,156 | 13 |
| FGF14 | FGF14 | Chromosome 13:101,169,308-101,852,156 | 13 |
| FGF14 | FGF14 | Chromosome 13:101,169,308-101,852,156 | 13 |
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| FGF14 | FGF14 | Chromosome 13:101,169,308-101,852,156 | 13 |
| FGF14 | FGF14 | Chromosome 13:101,169,308-101,852,156 | 13 |
| ERCC5 |  | Chromosome 13:102,295,195-102,326,346 | 13 |
| LIG4 |  | Chromosome 13:107,657,791-107,668,717 | 13 |
| TNFSF13B | CD257, BAFF, BLYS | Chromosome 13:107,719,978-107,758,826 | 13 |
| COL4A1 |  | Chromosome 13:109,599,311-109,757,505 | 13 |
| COL4A1 |  | Chromosome 13:109,599,311-109,757,505 | 13 |
| COL4A2 |  | Chromosome 13:109,757,632-109,963,375 | 13 |
| LAMP1 | CD107a | Chromosome 13:112,999,557-113,025,746 | 13 |


| IL17D |  | Chromosome 13:20,175,479-20,195,237 | 13 |
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| SAP18 |  | Chromosome 13:20,612,650-20,621,221 | 13 |
| SAP18 |  | Chromosome 13:20,612,650-20,621,221 | 13 |
| FGF9 | FGF9 | Chromosome 13:21,143,170-21,176,637 | 13 |
| FGF9 | FGF9 | Chromosome 13:21,143,170-21,176,637 | 13 |
| TNFRSF19 |  | Chromosome 13:23,042,723-23,148,232 | 13 |
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| TNFRSF19 |  | Chromosome 13:23,042,723-23,148,232 | 13 |
| CDK8 |  | Chromosome 13:25,726,276-25,877,375 | 13 |
| CDK8 |  | Chromosome 13:25,726,276-25,877,375 | 13 |
| CDK8 |  | Chromosome 13:25,726,276-25,877,375 | 13 |
| WASF3 | Wiskott Aldrich | Chromosome 13:26,029,840-26,161,085 | 13 |
| GTF3A |  | Chromosome 13:26,896,681-26,907,823 | 13 |
| GTF3A |  | Chromosome 13:26,896,681-26,907,823 | 13 |
| CDX2 |  | Chromosome 13:27,434,273-27,441,317 | 13 |
| FLT3 | CD135 | Chromosome 13:27,475,411-27,572,729 | 13 |
| FLT3 | CD135 | Chromosome 13:27,475,411-27,572,729 | 13 |
| FLT1 | CD308, VEGFR1 | Chromosome 13:27,773,790-27,967,232 | 13 |
| FLT1 | CD308, VEGFR1 | Chromosome 13:27,773,790-27,967,232 | 13 |
| FLT1 | CD308, VEGFR1 | Chromosome 13:27,773,790-27,967,232 | 13 |
| HMGB1 |  | Chromosome 13:29,930,884-30,089,729 | 13 |
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| HMGB1 |  | Chromosome 13:29,930,884-30,089,729 | 13 |
| ALOX5AP |  | Chromosome 13:30,207,645-30,236,556 | 13 |
| BRCA2 |  | Chromosome 13:31,787,617-31,871,809 | 13 |
| CCNA1 |  | Chromosome 13:35,904,495-35,915,008 | 13 |
| RFXAP |  | Chromosome 13:36,291,339-36,301,740 | 13 |
| FOXO1A |  | Chromosome 13:40,027,801-40,138,734 | 13 |
| F0X01A |  | Chromosome 13:40,027,801-40,138,734 | 13 |
| TNFSF11 | CD254, TRANCE, OPGL, | Chromosome 13:42,034,872-42,080,148 | 13 |
| TNFSF11 | CD254, TRANCE, OPGL, | Chromosome 13:42,034,872-42,080,148 | 13 |
| GTF2F2 |  | Chromosome 13:44,592,650-44,756,237 | 13 |
| GTF2F2 |  | Chromosome 13:44,592,650-44,756,237 | 13 |
| GTF2F2 |  | Chromosome 13:44,592,650-44,756,237 | 13 |
| TPT1 |  | Chromosome 13:44,809,008-44,813,505 | 13 |
| HTR2A | 5-HT2A | Chromosome 13:46,305,514-46,368,179 | 13 |
| HTR2A | 5-HT2A | Chromosome 13:46,305,514-46,368,179 | 13 |
| P2RY5 |  | Chromosome 13:47,883,170-47,887,947 | 13 |
| P2RY5 |  | Chromosome 13:47,883,170-47,887,947 | 13 |
| CYSLTR2 |  | Chromosome 13:48,178,692-48,181,499 | 13 |
| LECT1 | chondromodulin 1 | Chromosome 13:52,175,400-52,211,948 | 13 |
| ABCC4 | ABCC4, MRP4 | Chromosome 13:94,470,084-94,751,688 | 13 |
| ABCC4 | ABCC4, MRP4 | Chromosome 13:94,470,084-94,751,688 | 13 |
| ABCC4 | ABCC4, MRP4 | Chromosome 13:94,470,084-94,751,688 | 13 |
| ABCC4 | ABCC4, MRP4 | Chromosome 13:94,470,084-94,751,688 | 13 |
| EBI2 |  | Chromosome 13:98,744,790-98,757,708 | 13 |
| EBI2 |  | Chromosome 13:98,744,790-98,757,708 | 13 |
| HSP90AA1 | HSP90 | Chromosome 14:101,617,139-101,675,776 | 14 |
| RAGE |  | Chromosome 14:101,762,375-101,841,284 | 14 |
| TRAF3 |  | Chromosome 14:102,313,569-102,442,381 | 14 |
| TRAF3 |  | Chromosome 14:102,313,569-102,442,381 | 14 |
| BAG5 |  | Chromosome 14:103,092,642-103,098,907 | 14 |
| XRCC3 |  | Chromosome 14:103,233,707-103,251,549 | 14 |
| SIVA |  | Chromosome 14:104,290,529-104,297,036 | 14 |
| GPR132 | G2A | Chromosome 14:104,586,782-104,602,799 | 14 |


| IGHD1-20 |  | Chromosome 14:105,428,094-105,428,110 | 14 |
| :---: | :---: | :---: | :---: |
| IGHD1-20 |  | Chromosome 14:105,428,094-105,428,110 | 14 |
| CCNB1IP1 |  | Chromosome 14:19,849,367-19,871,297 | 14 |
| CCNB1IP1 |  | Chromosome 14:19,849,367-19,871,297 | 14 |
| RNASE6 |  | Chromosome 14:20,319,050-20,320,464 | 14 |
| RNASE7 | ribonuclease | Chromosome 14:20,580,251-20,582,226 | 14 |
| TRA@ | TCR | Chromosome 14:21,159,897-22,090,915 | 14 |
| TRA@ | TCR | Chromosome 14:21,159,897-22,090,915 | 14 |
| TRA@ | TCR | Chromosome 14:21,159,897-22,090,915 | 14 |
| TRA@ | TCR | Chromosome 14:21,159,897-22,090,915 | 14 |
| TRA@ | TCR | Chromosome 14:21,159,897-22,090,915 | 14 |
| TRA@ | TCR | Chromosome 14:21,159,897-22,090,915 | 14 |
| TRA@ | TCR | Chromosome 14:21,159,897-22,090,915 | 14 |
| TRA@ | TCR | Chromosome 14:21,159,897-22,090,915 | 14 |
| TRA@ | TCR | Chromosome 14:21,159,897-22,090,915 | 14 |
| MMP14 |  | Chromosome 14:22,375,633-22,386,643 | 14 |
| MMP14 |  | Chromosome 14:22,375,633-22,386,643 | 14 |
| PRMT5 | PRMT5 | Chromosome 14:22,459,573-22,468,501 | 14 |
| CEBPE |  | Chromosome 14:22,656,355-22,658,665 | 14 |
| BCL2L2 |  | Chromosome 14:22,845,866-22,850,798 | 14 |
| IL25 | IL17E | Chromosome 14:22,911,858-22,915,452 | 14 |
| PSME1 |  | Chromosome 14:23,661,207-23,678,016 | 14 |
| CHMP4A | CHMP4A | Chromosome 14:23,748,627-23,755,020 | 14 |
| NFATC4 |  | Chromosome 14:23,907,094-23,918,650 | 14 |
| CTSG | carthepsin G | Chromosome 14:24,112,564-24,115,306 | 14 |
| GZMB | Granzyme B | Chromosome 14:24,170,000-24,173,313 | 14 |
| GZMB | Granzyme B | Chromosome 14:24,170,000-24,173,313 | 14 |
| FOXG1C |  | Chromosome 14:28,304,801-28,308,621 | 14 |
| FOXG1C |  | Chromosome 14:28,304,801-28,308,621 | 14 |
| PRKD1 |  | Chromosome 14:29,115,436-29,466,651 | 14 |
| PRKD1 |  | Chromosome 14:29,115,436-29,466,651 | 14 |
| PRKD1 |  | Chromosome 14:29,115,436-29,466,651 | 14 |
| PRKD1 |  | Chromosome 14:29,115,436-29,466,651 | 14 |
| PRKD1 |  | Chromosome 14:29,115,436-29,466,651 | 14 |
| CFL2 |  | Chromosome 14:34,249,398-34,253,649 | 14 |
| CFL2 |  | Chromosome 14:34,249,398-34,253,649 | 14 |
| NFKBIA | IkB alpha | Chromosome 14:34,940,468-34,943,703 | 14 |
| NFKBIA | IkB alpha | Chromosome 14:34,940,468-34,943,703 | 14 |
| FOXA1 |  | Chromosome 14:37,128,940-37,134,240 | 14 |
| FOXA1 |  | Chromosome 14:37,128,940-37,134,240 | 14 |
| SSTR1 | sst1 receptor | Chromosome 14:37,746,955-37,752,019 | 14 |
| SSTR1 | sst1 receptor | Chromosome 14:37,746,955-37,752,019 | 14 |
| FKBP3 | Calcineurin | Chromosome 14:44,654,859-44,674,272 | 14 |
| PPIL5 |  | Chromosome 14:49,135,165-49,151,140 | 14 |
| MAP4K5 |  | Chromosome 14:49,954,993-50,069,126 | 14 |
| MAP4K5 |  | Chromosome 14:49,954,993-50,069,126 | 14 |
| TXNDC |  | Chromosome 14:50,776,686-50,792,512 | 14 |
| PTGDR |  | Chromosome 14:51,804,181-51,813,192 | 14 |
| PTGER2 | EP2 | Chromosome 14:51,850,863-51,865,074 | 14 |
| PSMC6 |  | Chromosome 14:52,243,668-52,264,466 | 14 |
| BMP4 | BMP4 | Chromosome 14:53,486,207-53,493,362 | 14 |
| BMP4 | BMP4 | Chromosome 14:53,486,207-53,493,362 | 14 |
| CDKN3 |  | Chromosome 14:53,933,423-53,956,682 | 14 |
| SOCS4 |  | Chromosome 14:54,563,594-54,585,960 | 14 |
| PELI2 | Pellino2 | Chromosome 14:55,654,846-55,837,784 | 14 |
| PELI2 | Pellino2 | Chromosome 14:55,654,846-55,837,784 | 14 |


| PELI2 | Pellino2 | Chromosome 14:55,654,846-55,837,784 | 14 |
| :---: | :---: | :---: | :---: |
| PELI2 | Pellino2 | Chromosome 14:55,654,846-55,837,784 | 14 |
| PRKCH | PKCń | Chromosome 14:60,858,186-61,087,451 | 14 |
| PRKCH | PKCń | Chromosome 14:60,858,186-61,087,451 | 14 |
| PRKCH | PKCń | Chromosome 14:60,858,186-61,087,451 | 14 |
| PRKCH | PKCŕ | Chromosome 14:60,858,186-61,087,451 | 14 |
| GPHB5 | GPHB5 | Chromosome 14:62,849,395-62,854,316 | 14 |
| GPHB5 | GPHB5 | Chromosome 14:62,849,395-62,854,316 | 14 |
| ESR2 | Estrogen receptor beta, N | Chromosome 14:63,621,388-63,875,070 | 14 |
| ESR2 | Estrogen receptor beta, N | Chromosome 14:63,621,388-63,875,070 | 14 |
| ESR2 | Estrogen receptor beta, N | Chromosome 14:63,621,388-63,875,070 | 14 |
| GPX2 |  | Chromosome 14:64,475,625-64,479,284 | 14 |
| ACTN1 | alpha actinin 1 | Chromosome 14:68,410,793-68,515,747 | 14 |
| ACTN1 | alpha actinin 1 | Chromosome 14:68,410,793-68,515,747 | 14 |
| MAP3K9 |  | Chromosome 14:70,264,605-70,345,641 | 14 |
| MAP3K9 |  | Chromosome 14:70,264,605-70,345,641 | 14 |
| FOS | C-Fos | Chromosome 14:74,815,284-74,818,685 | 14 |
| BTAF |  | Chromosome 14:75,058,537-75,083,086 | 14 |
| BTAF |  | Chromosome 14:75,058,537-75,083,086 | 14 |
| TGFB3 |  | Chromosome 14:75,494,195-75,517,242 | 14 |
| TGFB3 |  | Chromosome 14:75,494,195-75,517,242 | 14 |
| ESRRB | NR3B2 | Chromosome 14:75,907,479-76,036,961 | 14 |
| ESRRB | NR3B2 | Chromosome 14:75,907,479-76,036,961 | 14 |
| ESRRB | NR3B2 | Chromosome 14:75,907,479-76,036,961 | 14 |
| ALKBH1 |  | Chromosome 14:77,208,502-77,244,109 | 14 |
| ALKBH1 |  | Chromosome 14:77,208,502-77,244,109 | 14 |
| TSHR | thyroid stimulating hormo | Chromosome 14:80,491,528-80,682,399 | 14 |
| TSHR | thyroid stimulating hormo | Chromosome 14:80,491,528-80,682,399 | 14 |
| GTF2A1 | TFIIA | Chromosome 14:80,716,147-80,757,328 | 14 |
| GALC | Lactosylceramide | Chromosome 14:87,469,111-87,529,660 | 14 |
| GALC | Lactosylceramide | Chromosome 14:87,469,111-87,529,660 | 14 |
| PTPN21 | PTPD1 | Chromosome 14:88,003,867-88,090,876 | 14 |
| PTPN21 | PTPD1 | Chromosome 14:88,003,867-88,090,876 | 14 |
| CHES1 | FOXN3 | Chromosome 14:88,692,274-88,953,127 | 14 |
| CHES1 | FOXN3 | Chromosome 14:88,692,274-88,953,127 | 14 |
| CHES1 | FOXN3 | Chromosome 14:88,692,274-88,953,127 | 14 |
| CHES1 | FOXN3 | Chromosome 14:88,692,274-88,953,127 | 14 |
| GPR68 | OGR1, I-2 | Chromosome 14:90,768,629-90,789,977 | 14 |
| GPR68 | OGR1, I-2 | Chromosome 14:90,768,629-90,789,977 | 14 |
| LGMN | AEP | Chromosome 14:92,239,907-92,284,765 | 14 |
| LGMN | AEP | Chromosome 14:92,239,907-92,284,765 | 14 |
| MOAP1 |  | Chromosome 14:92,718,294-92,721,002 | 14 |
| DDX24 |  | Chromosome 14:93,587,019-93,617,311 | 14 |
| DDX24 |  | Chromosome 14:93,587,019-93,617,311 | 14 |
| SERPINA2 | Serpin a3g | Chromosome 14:93,900,404-93,914,178 | 14 |
| SERPINA2 | Serpin a3g | Chromosome 14:93,900,404-93,914,178 | 14 |
| BDKRB2 | Bradykinin receptor | Chromosome 14:95,740,950-95,780,542 | 14 |
| BDKRB2 | Bradykinin receptor | Chromosome 14:95,740,950-95,780,542 | 14 |
| CYP46A1 |  | Chromosome 14:99,220,407-99,263,391 | 14 |
| CYP46A1 |  | Chromosome 14:99,220,407-99,263,391 | 14 |
| YY1 |  | Chromosome 14:99,774,855-99,814,557 | 14 |
| YY1 |  | Chromosome 14:99,774,855-99,814,557 | 14 |
| SNRPN |  | Chromosome 15:22,619,887-23,215,702 | 15 |
| SNRPN |  | Chromosome 15:22,619,887-23,215,702 | 15 |
| SNRPN |  | Chromosome 15:22,619,887-23,215,702 | 15 |
| SNRPN |  | Chromosome 15:22,619,887-23,215,702 | 15 |


| SNRPN |  | Chromosome 15:22,619,887-23,215,702 | 15 |
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| AVEN |  | Chromosome 15:31,945,720-32,118,595 | 15 |
| AVEN |  | Chromosome 15:31,945,720-32,118,595 | 15 |
| AVEN |  | Chromosome 15:31,945,720-32,118,595 | 15 |
| ACTC1 | F-actin | Chromosome 15:32,869,723-32,875,181 | 15 |
| ACTC1 | F-actin | Chromosome 15:32,869,723-32,875,181 | 15 |
| RASGRP1 | rasGRP | Chromosome 15:36,567,590-36,644,224 | 15 |
| THBS1 | thrombospondin | Chromosome 15:37,660,572-37,676,960 | 15 |
| THBS1 | thrombospondin | Chromosome 15:37,660,572-37,676,960 | 15 |
| PLCB2 | phospholipase beta 2 | Chromosome 15:38,366,448-38,387,330 | 15 |
| RAD51 |  | Chromosome 15:38,774,661-38,811,646 | 15 |
| DLL4 | Delta4 | Chromosome 15:39,008,839-39,018,529 | 15 |
| DLL4 | Delta4 | Chromosome 15:39,008,839-39,018,529 | 15 |
| NUSAP1 |  | Chromosome 15:39,412,361-39,460,538 | 15 |
| MGA |  | Chromosome 15:39,739,902-39,849,433 | 15 |
| MGA |  | Chromosome 15:39,739,902-39,849,433 | 15 |
| HISPPD2A | IPS1 | Chromosome 15:41,612,949-41,769,525 | 15 |
| HISPPD2A | IPS1 | Chromosome 15:41,612,949-41,769,525 | 15 |
| PDIA3 | ERp57 | Chromosome 15:41,825,882-41,852,096 | 15 |
| B2M | Beta-2 microglobulin | Chromosome 15:42,790,977-42,797,649 | 15 |
| B2M | Beta-2 microglobulin | Chromosome 15:42,790,977-42,797,649 | 15 |
| FGF7 | KGF | Chromosome 15:47,502,751-47,566,815 | 15 |
| FGF7 | KGF | Chromosome 15:47,502,751-47,566,815 | 15 |
| HDC |  | Chromosome 15:48,321,436-48,345,218 | 15 |
| CYP19A1 |  | Chromosome 15:49,288,961-49,418,086 | 15 |
| CYP19A1 |  | Chromosome 15:49,288,961-49,418,086 | 15 |
| MAPK6 |  | Chromosome 15:50,098,739-50,145,754 | 15 |
| BCL2L10 |  | Chromosome 15:50,189,114-50,192,264 | 15 |
| TCF12 | HEB/SCBP | Chromosome 15:54,998,125-55,368,008 | 15 |
| TCF12 | HEB/SCBP | Chromosome 15:54,998,125-55,368,008 | 15 |
| TCF12 | HEB/SCBP | Chromosome 15:54,998,125-55,368,008 | 15 |
| TCF12 | HEB/SCBP | Chromosome 15:54,998,125-55,368,008 | 15 |
| TCF12 | HEB/SCBP | Chromosome 15:54,998,125-55,368,008 | 15 |
| ADAM10 | CDw156c | Chromosome 15:56,675,802-56,829,469 | 15 |
| ADAM10 | CDw156c | Chromosome 15:56,675,802-56,829,469 | 15 |
| CCNB2 | cyclin B2 | Chromosome 15:57,184,612-57,204,536 | 15 |
| GTF2A2 |  | Chromosome 15:57,718,358-57,736,991 | 15 |
| FOXB1 |  | Chromosome 15:58,084,427-58,085,434 | 15 |
| FOXB1 |  | Chromosome 15:58,084,427-58,085,434 | 15 |
| ANXA2 | Annexin-2 | Chromosome 15:58,426,642-58,477,477 | 15 |
| RORA | RORa, NR1F1 | Chromosome 15:58,576,755-59,308,794 | 15 |
| RORA | RORa, NR1F1 | Chromosome 15:58,576,755-59,308,794 | 15 |
| RORA | RORa, NR1F1 | Chromosome 15:58,576,755-59,308,794 | 15 |
| RORA | RORa, NR1F1 | Chromosome 15:58,576,755-59,308,794 | 15 |
| RORA | RORa, NR1F1 | Chromosome 15:58,576,755-59,308,794 | 15 |
| RORA | RORa, NR1F1 | Chromosome 15:58,576,755-59,308,794 | 15 |
| RORA | RORa, NR1F1 | Chromosome 15:58,576,755-59,308,794 | 15 |
| DAPK2 |  | Chromosome 15:61,986,288-62,125,574 | 15 |
| DAPK2 |  | Chromosome 15:61,986,288-62,125,574 | 15 |
| DAPK2 |  | Chromosome 15:61,986,288-62,125,574 | 15 |
| PPIB |  | Chromosome 15:62,235,067-62,242,407 | 15 |
| ANKDD1A |  | Chromosome 15:62,995,046-63,038,086 | 15 |
| ANKDD1A |  | Chromosome 15:62,995,046-63,038,086 | 15 |
| PDCD7 |  | Chromosome 15:63,196,770-63,213,227 | 15 |
| MAP2K1 | MEK1 | Chromosome 15:64,466,674-64,570,936 | 15 |
| MAP2K1 | MEK1 | Chromosome 15:64,466,674-64,570,936 | 15 |


| MAP2K1 | MEK1 | Chromosome 15:64,466,674-64,570,936 | 15 |
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| SMAD3 |  | Chromosome 15:65,145,249-65,274,587 | 15 |
| SMAD3 |  | Chromosome 15:65,145,249-65,274,587 | 15 |
| MAP2K5 |  | Chromosome 15:65,622,075-65,886,506 | 15 |
| MAP2K5 |  | Chromosome 15:65,622,075-65,886,506 | 15 |
| MAP2K5 |  | Chromosome 15:65,622,075-65,886,506 | 15 |
| ITGA11 |  | Chromosome 15:66,381,096-66,511,546 | 15 |
| ITGA11 |  | Chromosome 15:66,381,096-66,511,546 | 15 |
| ITGA11 |  | Chromosome 15:66,381,096-66,511,546 | 15 |
| RPLP1 | P1 | Chromosome 15:67,532,177-67,534,939 | 15 |
| TLE3 |  | Chromosome 15:68,127,597-68,177,310 | 15 |
| TLE3 |  | Chromosome 15:68,127,597-68,177,310 | 15 |
| NR2E3 | NR2E3 | Chromosome 15:69,889,948-69,897,654 | 15 |
| NR2E3 | NR2E3 | Chromosome 15:69,889,948-69,897,654 | 15 |
| HEXA | Hexasaminidase | Chromosome 15:70,364,122-70,455,868 | 15 |
| NPTN | SDFR1 | Chromosome 15:71,639,410-71,712,806 | 15 |
| CD276 | B7-H3 | Chromosome 15:71,763,675-71,793,912 | 15 |
| CYP11A1 |  | Chromosome 15:72,417,157-72,447,134 | 15 |
| CYP11A1 |  | Chromosome 15:72,417,157-72,447,134 | 15 |
| CYP1A1 |  | Chromosome 15:72,798,943-72,804,930 | 15 |
| CYP1A1 |  | Chromosome 15:72,798,943-72,804,930 | 15 |
| PTPN9 |  | Chromosome 15:73,546,515-73,658,680 | 15 |
| NRG4 | NRG4 | Chromosome 15:74,020,333-74,091,842 | 15 |
| NRG4 | NRG4 | Chromosome 15:74,020,333-74,091,842 | 15 |
| PSTPIP1 |  | Chromosome 15:75,074,609-75,116,727 | 15 |
| PSTPIP1 |  | Chromosome 15:75,074,609-75,116,727 | 15 |
| CTSH | Cathepsin H | Chromosome 15:77,001,162-77,024,475 | 15 |
| CTSH | Cathepsin H | Chromosome 15:77,001,162-77,024,475 | 15 |
| BCL2A1 | Bfl-1, BCL2-A1, BCL2a1b | Chromosome 15:78,040,290-78,050,698 | 15 |
| IL16 |  | Chromosome 15:79,262,148-79,392,157 | 15 |
| IL16 |  | Chromosome 15:79,262,148-79,392,157 | 15 |
| IL16 |  | Chromosome 15:79,262,148-79,392,157 | 15 |
| AKAP13 | HA-3 | Chromosome 15:83,578,821-84,093,590 | 15 |
| AKAP13 | HA-3 | Chromosome 15:83,578,821-84,093,590 | 15 |
| AKAP13 | HA-3 | Chromosome 15:83,578,821-84,093,590 | 15 |
| AKAP13 | HA-3 | Chromosome 15:83,578,821-84,093,590 | 15 |
| AKAP13 | HA-3 | Chromosome 15:83,578,821-84,093,590 | 15 |
| AKAP13 | HA-3 | Chromosome 15:83,578,821-84,093,590 | 15 |
| ISG20 |  | Chromosome 15:86,983,039-87,000,684 | 15 |
| ISG20 |  | Chromosome 15:86,983,039-87,000,684 | 15 |
| ANPEP | CD13 | Chromosome 15:88,129,130-88,159,072 | 15 |
| ANPEP | CD13 | Chromosome 15:88,129,130-88,159,072 | 15 |
| BLM | Bloom syndrome | Chromosome 15:89,061,606-89,159,688 | 15 |
| BLM | Bloom syndrome | Chromosome 15:89,061,606-89,159,688 | 15 |
| BLM | Bloom syndrome | Chromosome 15:89,061,606-89,159,688 | 15 |
| FURIN |  | Chromosome 15:89,212,889-89,227,691 | 15 |
| NR2F2 | NR2F2 | Chromosome 15:94,674,950-94,683,048 | 15 |
| NR2F2 | NR2F2 | Chromosome 15:94,674,950-94,683,048 | 15 |
| IGF1R | CD221 | Chromosome 15:97,010,288-97,319,034 | 15 |
| IGF1R | CD221 | Chromosome 15:97,010,288-97,319,034 | 15 |
| IGF1R | CD221 | Chromosome 15:97,010,288-97,319,034 | 15 |
| IGF1R | CD221 | Chromosome 15:97,010,288-97,319,034 | 15 |
| MAPK8IP3 |  | Chromosome 16:1,696,222-1,760,319 | 16 |
| NOXO1 |  | Chromosome 16:1,968,919-1,971,441 | 16 |
| CIITA | MHC2TA=CIITA, C2ta | Chromosome 16:10,867,648-10,926,341 | 16 |
| CIITA | MHC2TA=CIITA, C2ta | Chromosome 16:10,867,648-10,926,341 | 16 |


| SOCS1 |  | Chromosome 16:11,255,775-11,257,540 | 16 |
| :---: | :---: | :---: | :---: |
| SOCS1 |  | Chromosome 16:11,255,775-11,257,540 | 16 |
| LITAF | lipopolysaccharide induce | Chromosome 16:11,549,357-11,588,823 | 16 |
| LITAF | lipopolysaccharide induce | Chromosome 16:11,549,357-11,588,823 | 16 |
| TNFRSF17 | CD269, BCMA | Chromosome 16:11,966,465-11,969,426 | 16 |
| TNFRSF17 | CD269, BCMA | Chromosome 16:11,966,465-11,969,426 | 16 |
| MYH11 |  | Chromosome 16:15,704,493-15,858,388 | 16 |
| MYH11 |  | Chromosome 16:15,704,493-15,858,388 | 16 |
| MYH11 |  | Chromosome 16:15,704,493-15,858,388 | 16 |
| ABCC1 | MRP | Chromosome 16:15,950,935-16,143,774 | 16 |
| HBA2 |  | Chromosome 16:162,875-163,708 | 16 |
| TRAF7 |  | Chromosome 16:2,145,800-2,168,131 | 16 |
| IGSF6 |  | Chromosome 16:21,559,426-21,571,473 | 16 |
| IGSF6 |  | Chromosome 16:21,559,426-21,571,473 | 16 |
| PLK1 | Plk1 | Chromosome 16:23,597,692-23,609,189 | 16 |
| PLK1 | Plk1 | Chromosome 16:23,597,692-23,609,189 | 16 |
| PRKCB1 | PKC beta | Chromosome 16:23,754,823-24,139,358 | 16 |
| PRKCB1 | PKC beta | Chromosome 16:23,754,823-24,139,358 | 16 |
| PRKCB1 | PKC beta | Chromosome 16:23,754,823-24,139,358 | 16 |
| PRKCB1 | PKC beta | Chromosome 16:23,754,823-24,139,358 | 16 |
| PRKCB1 | PKC beta | Chromosome 16:23,754,823-24,139,358 | 16 |
| PRKCB1 | PKC beta | Chromosome 16:23,754,823-24,139,358 | 16 |
| IL4R | CD124 | Chromosome 16:27,232,752-27,283,600 | 16 |
| IL4R | CD124 | Chromosome 16:27,232,752-27,283,600 | 16 |
| GTF3C1 |  | Chromosome 16:27,379,436-27,468,775 | 16 |
| AXIN1 |  | Chromosome 16:277,441-342,465 | 16 |
| IL27 |  | Chromosome 16:28,418,184-28,425,656 | 16 |
| IL27 |  | Chromosome 16:28,418,184-28,425,656 | 16 |
| SPN | CD43 | Chromosome 16:29,581,801-29,589,688 | 16 |
| MVP | lung resistance protein | Chromosome 16:29,731,591-29,766,842 | 16 |
| TNFRSF12A | CD266, TWEAKR | Chromosome 16:3,010,343-3,012,385 | 16 |
| TNFRSF12A | CD266, TWEAKR | Chromosome 16:3,010,343-3,012,385 | 16 |
| MEFV | Pyrin | Chromosome 16:3,232,029-3,246,628 | 16 |
| MEFV | Pyrin | Chromosome 16:3,232,029-3,246,628 | 16 |
| NLRC3 | NOD3 | Chromosome 16:3,531,826-3,567,290 | 16 |
| NLRC3 | NOD3 | Chromosome 16:3,531,826-3,567,290 | 16 |
| DNASE1 |  | Chromosome 16:3,630,847-3,654,064 | 16 |
| CREBBP | CBP | Chromosome 16:3,716,568-3,870,723 | 16 |
| CREBBP | CBP | Chromosome 16:3,716,568-3,870,723 | 16 |
| CREBBP | CBP | Chromosome 16:3,716,568-3,870,723 | 16 |
| SEPHS2 | SPS2 | Chromosome 16:30,362,453-30,364,725 | 16 |
| ITGAL | LFA1, CD11a | Chromosome 16:30,391,551-30,442,007 | 16 |
| CTF1 | CTF1 | Chromosome 16:30,815,429-30,822,382 | 16 |
| ITGAM | Mac-1, CD11B | Chromosome 16:31,178,789-31,251,714 | 16 |
| ITGAM | Mac-1, CD11B | Chromosome 16:31,178,789-31,251,714 | 16 |
| ITGAX | CD11c, CR4 | Chromosome 16:31,274,010-31,301,819 | 16 |
| HMOX2 |  | Chromosome 16:4,466,426-4,500,349 | 16 |
| ITFG1 | CDA08, TIP | Chromosome 16:45,746,798-46,052,519 | 16 |
| ITFG1 | CDA08, TIP | Chromosome 16:45,746,798-46,052,519 | 16 |
| ITFG1 | CDA08, TIP | Chromosome 16:45,746,798-46,052,519 | 16 |
| ABCC11 | MRP8 | Chromosome 16:46,758,323-46,838,806 | 16 |
| ABCC11 | MRP8 | Chromosome 16:46,758,323-46,838,806 | 16 |
| NOD2 | BLAU, CARD15, CD, PS | Chromosome 16:49,288,551-49,324,488 | 16 |
| NOD2 | BLAU, CARD15, CD, PS | Chromosome 16:49,288,551-49,324,488 | 16 |
| AKTIP | FTS | Chromosome 16:52,082,693-52,094,671 | 16 |
| AKTIP | FTS | Chromosome 16:52,082,693-52,094,671 | 16 |


| MMP2 |  | Chromosome 16:54,070,589-54,098,104 | 16 |
| :---: | :---: | :---: | :---: |
| MMP2 |  | Chromosome 16:54,070,589-54,098,104 | 16 |
| CES1 | Carboxylesterase 3 | Chromosome 16:54,394,264-54,424,576 | 16 |
| MT3 |  | Chromosome 16:55,180,768-55,182,501 | 16 |
| MT3 |  | Chromosome 16:55,180,768-55,182,501 | 16 |
| NLRC5 | NOD27 | Chromosome 16:55,581,018-55,673,941 | 16 |
| NLRC5 | NOD27 | Chromosome 16:55,581,018-55,673,941 | 16 |
| NLRC5 | NOD27 | Chromosome 16:55,581,018-55,673,941 | 16 |
| CCL22 |  | Chromosome 16:55,950,219-55,957,602 | 16 |
| CCL17 |  | Chromosome 16:55,996,180-56,007,475 | 16 |
| MMP15 |  | Chromosome 16:56,616,783-56,638,306 | 16 |
| MMP15 |  | Chromosome 16:56,616,783-56,638,306 | 16 |
| CDH5 | CD144, VE-cadherin | Chromosome 16:64,958,064-64,996,190 | 16 |
| CDH5 | CD144, VE-cadherin | Chromosome 16:64,958,064-64,996,190 | 16 |
| CKLF |  | Chromosome 16:65,143,967-65,170,463 | 16 |
| CBFB |  | Chromosome 16:65,620,551-65,692,462 | 16 |
| CBFB |  | Chromosome 16:65,620,551-65,692,462 | 16 |
| TRADD |  | Chromosome 16:65,745,605-65,751,306 | 16 |
| NOL3 | MYC | Chromosome 16:65,765,371-65,767,127 | 16 |
| PSMB10 | proteasome subunit | Chromosome 16:66,525,908-66,528,254 | 16 |
| PSMB10 | proteasome subunit | Chromosome 16:66,525,908-66,528,254 | 16 |
| NFATC3 |  | Chromosome 16:66,676,845-66,818,338 | 16 |
| PRMT7 | PRMT7 | Chromosome 16:66,902,446-66,948,663 | 16 |
| MPG |  | Chromosome 16:67,018-75,845 | 16 |
| CDH1 | CD324, E-cadherin | Chromosome 16:67,328,696-67,426,945 | 16 |
| NFAT5 |  | Chromosome 16:68,156,498-68,296,054 | 16 |
| NFAT5 |  | Chromosome 16:68,156,498-68,296,054 | 16 |
| DDX19B |  | Chromosome 16:68,890,573-68,925,232 | 16 |
| DHX38 |  | Chromosome 16:70,685,116-70,704,312 | 16 |
| DHX38 |  | Chromosome 16:70,685,116-70,704,312 | 16 |
| ATBF1 |  | Chromosome 16:71,378,456-71,639,775 | 16 |
| ATBF1 |  | Chromosome 16:71,378,456-71,639,775 | 16 |
| ATBF1 |  | Chromosome 16:71,378,456-71,639,775 | 16 |
| ATBF1 |  | Chromosome 16:71,378,456-71,639,775 | 16 |
| ATBF1 |  | Chromosome 16:71,378,456-71,639,775 | 16 |
| ATBF1 |  | Chromosome 16:71,378,456-71,639,775 | 16 |
| CFDP1 |  | Chromosome 16:73,885,109-74,024,888 | 16 |
| CFDP1 |  | Chromosome 16:73,885,109-74,024,888 | 16 |
| CFDP1 |  | Chromosome 16:73,885,109-74,024,888 | 16 |
| CLEC3A |  | Chromosome 16:76,613,992-76,623,499 | 16 |
| MAF | cMaf | Chromosome 16:78,185,732-78,192,112 | 16 |
| MAF | cMaf | Chromosome 16:78,185,732-78,192,112 | 16 |
| GINS2 |  | Chromosome 16:84,268,781-84,280,089 | 16 |
| IRF8 | ICSBP | Chromosome 16:84,490,275-84,513,713 | 16 |
| IRF8 | ICSBP | Chromosome 16:84,490,275-84,513,713 | 16 |
| FOXF1 |  | Chromosome 16:85,101,659-85,105,548 | 16 |
| FOXF1 |  | Chromosome 16:85,101,659-85,105,548 | 16 |
| SLC7A5 |  | Chromosome 16:86,421,130-86,460,615 | 16 |
| SLC7A5 |  | Chromosome 16:86,421,130-86,460,615 | 16 |
| IL17C |  | Chromosome 16:87,232,502-87,234,383 | 16 |
| MC1R | MC1 | Chromosome 16:88,512,527-88,529,713 | 16 |
| SCARF1 |  | Chromosome 17:1,483,902-1,495,792 | 17 |
| SCARF1 |  | Chromosome 17:1,483,902-1,495,792 | 17 |
| SMG6 |  | Chromosome 17:1,909,888-2,220,160 | 17 |
| SMG6 |  | Chromosome 17:1,909,888-2,220,160 | 17 |
| SMG6 |  | Chromosome 17:1,909,888-2,220,160 | 17 |


| SMG6 |  | Chromosome 17:1,909,888-2,220,160 | 17 |
| :---: | :---: | :---: | :---: |
| MYH2 | myosin heavy chain 2a | Chromosome 17:10,365,192-10,393,704 | 17 |
| MYH2 | myosin heavy chain 2a | Chromosome 17:10,365,192-10,393,704 | 17 |
| MAP2K4 | MKK4 | Chromosome 17:11,864,860-11,987,865 | 17 |
| IL6STP |  | Chromosome 17:15,616,046-15,629,130 | 17 |
| IL6STP |  | Chromosome 17:15,616,046-15,629,130 | 17 |
| NCOR1 | NCoR | Chromosome 17:15,875,983-16,059,570 | 17 |
| NCOR1 | NCoR | Chromosome 17:15,875,983-16,059,570 | 17 |
| TRPV2 |  | Chromosome 17:16,259,613-16,281,042 | 17 |
| TNFRSF13B | CD267, TAC1 | Chromosome 17:16,783,123-16,816,127 | 17 |
| SREBF1 | SREBP1a,b,c | Chromosome 17:17,655,794-17,681,050 | 17 |
| DRG2 | DRG2 | Chromosome 17:17,932,008-17,952,017 | 17 |
| DRG2 | DRG2 | Chromosome 17:17,932,008-17,952,017 | 17 |
| ALKBH5 |  | Chromosome 17:18,028,014-18,053,993 | 17 |
| FOXO3B |  | Chromosome 17:18,516,347-18,516,964 | 17 |
| MAPK7 |  | Chromosome 17:19,221,659-19,227,445 | 17 |
| PAFAH1B1 | PAF, platelet activating fa | Chromosome 17:2,443,686-2,535,638 | 17 |
| PAFAH1B1 | PAF, platelet activating fa | Chromosome 17:2,443,686-2,535,638 | 17 |
| PAFAH1B1 | PAF, platelet activating fa | Chromosome 17:2,443,686-2,535,638 | 17 |
| MAP2K3 |  | Chromosome 17:21,128,581-21,159,118 | 17 |
| LGALS9 | Galectin 9 | Chromosome 17:22,980,951-23,000,711 | 17 |
| NOS2A | NO synthase, iNOS | Chromosome 17:23,107,919-23,151,682 | 17 |
| NOS2A | NO synthase, iNOS | Chromosome 17:23,107,919-23,151,682 | 17 |
| VTN |  | Chromosome 17:23,718,425-23,721,844 | 17 |
| FOXN1 |  | Chromosome 17:23,875,086-23,889,302 | 17 |
| TIAF1 | TGF antiapoptotic factor | Chromosome 17:24,424,663-24,531,556 | 17 |
| TIAF1 | TGF antiapoptotic factor | Chromosome 17:24,424,663-24,531,556 | 17 |
| TIAF1 | TGF antiapoptotic factor | Chromosome 17:24,424,663-24,531,556 | 17 |
| TIAF1 | TGF antiapoptotic factor | Chromosome 17:24,424,663-24,531,556 | 17 |
| SLC6A4 | 5-HTT | Chromosome 17:25,549,032-25,586,831 | 17 |
| SLC6A4 | 5-HTT | Chromosome 17:25,549,032-25,586,831 | 17 |
| CRLF3 |  | Chromosome 17:26,133,828-26,175,826 | 17 |
| CRLF3 |  | Chromosome 17:26,133,828-26,175,826 | 17 |
| CCL2 | MCP1 | Chromosome 17:29,606,409-29,608,335 | 17 |
| CCL13 | MCP4 | Chromosome 17:29,707,584-29,709,742 | 17 |
| CCL1 |  | Chromosome 17:29,711,512-29,714,365 | 17 |
| TRPV1 | VR1 | Chromosome 17:3,415,491-3,459,454 | 17 |
| TRPV1 | VR1 | Chromosome 17:3,415,491-3,459,454 | 17 |
| CTNS |  | Chromosome 17:3,486,522-3,511,585 | 17 |
| ITGAE | HML-1, CD103 | Chromosome 17:3,564,671-3,660,578 | 17 |
| P2RX1 | P2X1 | Chromosome 17:3,746,634-3,766,709 | 17 |
| MMP28 |  | Chromosome 17:31,116,989-31,146,753 | 17 |
| CCL5 | RANTES | Chromosome 17:31,222,611-31,231,490 | 17 |
| LOC441792 | NO synthase | Chromosome 17:31,811,186-31,816,297 | 17 |
| ACACA | BCL2A1-ACC1 | Chromosome 17:32,516,040-32,841,015 | 17 |
| ACACA | BCL2A1-ACC1 | Chromosome 17:32,516,040-32,841,015 | 17 |
| ACACA | BCL2A1-ACC1 | Chromosome 17:32,516,040-32,841,015 | 17 |
| DUSP14 |  | Chromosome 17:32,924,064-32,947,709 | 17 |
| DDX52 |  | Chromosome 17:33,046,526-33,077,600 | 17 |
| ERBB2 | CD340 | Chromosome 17:35,097,919-35,138,441 | 17 |
| ERBB2 | CD340 | Chromosome 17:35,097,919-35,138,441 | 17 |
| CSF3 |  | Chromosome 17:35,425,214-35,427,592 | 17 |
| THRA | NR1A1 | Chromosome 17:35,472,589-35,503,646 | 17 |
| NR1D1 | NR1D1 | Chromosome 17:35,502,567-35,510,499 | 17 |
| NR1D1 | NR1D1 | Chromosome 17:35,502,567-35,510,499 | 17 |
| RARA | NR1B1 | Chromosome 17:35,718,972-35,767,420 | 17 |


| TOP2A |  | Chromosome 17:35,798,321-35,827,695 | 17 |
| :---: | :---: | :---: | :---: |
| CCR7 | CD197 | Chromosome 17:35,963,547-35,975,250 | 17 |
| CCR7 | CD197 | Chromosome 17:35,963,547-35,975,250 | 17 |
| KRT34 | HA-4 | Chromosome 17:36,787,447-36,792,181 | 17 |
| KRT35 | HA-5 | Chromosome 17:36,886,467-36,891,194 | 17 |
| KRT35 | HA-5 | Chromosome 17:36,886,467-36,891,194 | 17 |
| FKBP10 |  | Chromosome 17:37,222,727-37,232,995 | 17 |
| FKBP10 |  | Chromosome 17:37,222,727-37,232,995 | 17 |
| LGP2 |  | Chromosome 17:37,506,952-37,518,277 | 17 |
| LGP2 |  | Chromosome 17:37,506,952-37,518,277 | 17 |
| STAT5B |  | Chromosome 17:37,604,721-37,681,950 | 17 |
| STAT5B |  | Chromosome 17:37,604,721-37,681,950 | 17 |
| STAT5B |  | Chromosome 17:37,604,721-37,681,950 | 17 |
| CCR10 |  | Chromosome 17:38,084,961-38,087,371 | 17 |
| PSME3 |  | Chromosome 17:38,238,949-38,249,303 | 17 |
| DHX8 |  | Chromosome 17:38,916,860-38,957,206 | 17 |
| DHX8 |  | Chromosome 17:38,916,860-38,957,206 | 17 |
| DHX8 |  | Chromosome 17:38,916,860-38,957,206 | 17 |
| DUSP3 |  | Chromosome 17:39,199,015-39,211,872 | 17 |
| DUSP3 |  | Chromosome 17:39,199,015-39,211,872 | 17 |
| DUSP3 |  | Chromosome 17:39,199,015-39,211,872 | 17 |
| HDAC5 |  | Chromosome 17:39,509,647-39,556,540 | 17 |
| HDAC5 |  | Chromosome 17:39,509,647-39,556,540 | 17 |
| SLC4A1 | CD233 | Chromosome 17:39,682,566-39,700,993 | 17 |
| ITGA2B | CD41, HPA1 | Chromosome 17:39,805,076-39,822,399 | 17 |
| ALOX15 |  | Chromosome 17:4,480,963-4,491,709 | 17 |
| ARRB2 | Beta arrestin 2 | Chromosome 17:4,560,533-4,571,544 | 17 |
| PSMB6 |  | Chromosome 17:4,646,397-4,648,756 | 17 |
| MINK |  | Chromosome 17:4,683,351-4,742,135 | 17 |
| GP1BA | HPA5a,b, CD42b | Chromosome 17:4,776,372-4,779,067 | 17 |
| PFN1 | Profilin | Chromosome 17:4,789,692-4,793,067 | 17 |
| C1QL1 |  | Chromosome 17:40,392,587-40,401,170 | 17 |
| C1QL1 |  | Chromosome 17:40,392,587-40,401,170 | 17 |
| MAP3K14 | NIK | Chromosome 17:40,696,278-40,750,148 | 17 |
| MAP3K14 | NIK | Chromosome 17:40,696,278-40,750,148 | 17 |
| MAP3K14 | NIK | Chromosome 17:40,696,278-40,750,148 | 17 |
| CRF1R | CRHR1 | Chromosome 17:41,217,449-41,268,973 | 17 |
| CRF1R | CRHR1 | Chromosome 17:41,217,449-41,268,973 | 17 |
| CRF1R | CRHR1 | Chromosome 17:41,217,449-41,268,973 | 17 |
| WNT3 |  | Chromosome 17:42,196,855-42,251,081 | 17 |
| WNT3 |  | Chromosome 17:42,196,855-42,251,081 | 17 |
| WNT3 |  | Chromosome 17:42,196,855-42,251,081 | 17 |
| ITGB3 | CD60, HPA2, Mac-2, GPI | Chromosome 17:42,686,207-42,745,076 | 17 |
| ITGB3 | CD60, HPA2, Mac-2, GPI | Chromosome 17:42,686,207-42,745,076 | 17 |
| ITGB3 | CD60, HPA2, Mac-2, GPI | Chromosome 17:42,686,207-42,745,076 | 17 |
| TBX21 | T-bet | Chromosome 17:43,165,609-43,178,484 | 17 |
| TBX21 | T-bet | Chromosome 17:43,165,609-43,178,484 | 17 |
| TBX21 | T-bet | Chromosome 17:43,165,609-43,178,484 | 17 |
| NFE2L1 |  | Chromosome 17:43,480,720-43,493,841 | 17 |
| SKAP1 | SKAP55 | Chromosome 17:43,565,804-43,862,551 | 17 |
| SKAP1 | SKAP55 | Chromosome 17:43,565,804-43,862,551 | 17 |
| SKAP1 | SKAP55 | Chromosome 17:43,565,804-43,862,551 | 17 |
| HOXB4 | HOX cluster | Chromosome 17:44,007,868-44,010,742 | 17 |
| NGFR | CD271 | Chromosome 17:44,927,654-44,947,360 | 17 |
| DLX3 | Delta-like 2 | Chromosome 17:45,422,368-45,427,587 | 17 |
| COL1A1 | collagen alpha 1 | Chromosome 17:45,616,456-45,633,992 | 17 |


| COL1A1 | collagen alpha 1 | Chromosome 17:45,616,456-45,633,992 | 17 |
| :---: | :---: | :---: | :---: |
| NME1 | NM23H1 | Chromosome 17:46,585,919-46,604,103 | 17 |
| C1QBP |  | Chromosome 17:5,276,823-5,283,195 | 17 |
| DHX33 |  | Chromosome 17:5,284,956-5,312,905 | 17 |
| NLRP1 | NALP1, CARD7, DEFCAF | Chromosome 17:5,343,472-5,428,553 | 17 |
| HLF | HIf | Chromosome 17:50,697,370-50,755,886 | 17 |
| HLF | HIf | Chromosome 17:50,697,370-50,755,886 | 17 |
| TRIM25 | ZNF147 | Chromosome 17:52,320,269-52,346,408 | 17 |
| TRIM25 | ZNF147 | Chromosome 17:52,320,269-52,346,408 | 17 |
| EPX | eosinophil peroxidase | Chromosome 17:53,625,088-53,636,783 | 17 |
| MPO |  | Chromosome 17:53,702,201-53,713,295 | 17 |
| MPO |  | Chromosome 17:53,702,201-53,713,295 | 17 |
| DHX40 |  | Chromosome 17:54,997,668-55,040,484 | 17 |
| DHX40 |  | Chromosome 17:54,997,668-55,040,484 | 17 |
| MRC2 | Endo180, CD280, manno | Chromosome 17:58,058,494-58,124,629 | 17 |
| MRC2 | Endo180, CD280, manno | Chromosome 17:58,058,494-58,124,629 | 17 |
| MRC2 | Endo180, CD280, manno | Chromosome 17:58,058,494-58,124,629 | 17 |
| ACE | CD143 | Chromosome 17:58,908,166-58,952,935 | 17 |
| DDX42 |  | Chromosome 17:59,205,299-59,250,409 | 17 |
| GH1 | GH1 | Chromosome 17:59,348,294-59,349,930 | 17 |
| PECAM1 | CD31 | Chromosome 17:59,754,142-59,817,723 | 17 |
| DDX5 |  | Chromosome 17:59,926,200-59,932,872 | 17 |
| ALOX12 |  | Chromosome 17:6,840,108-6,856,220 | 17 |
| CLECSF10A | CD301, MGL1, CLEC10A | Chromosome 17:6,918,580-6,924,324 | 17 |
| GNA13 |  | Chromosome 17:60,437,295-60,483,216 | 17 |
| GNA13 |  | Chromosome 17:60,437,295-60,483,216 | 17 |
| AXIN2 |  | Chromosome 17:60,955,143-60,988,227 | 17 |
| AXIN2 |  | Chromosome 17:60,955,143-60,988,227 | 17 |
| PRKCA | PKC alpha | Chromosome 17:61,729,388-62,237,324 | 17 |
| PRKCA | PKC alpha | Chromosome 17:61,729,388-62,237,324 | 17 |
| PRKCA | PKC alpha | Chromosome 17:61,729,388-62,237,324 | 17 |
| PRKCA | PKC alpha | Chromosome 17:61,729,388-62,237,324 | 17 |
| PRKCA | PKC alpha | Chromosome 17:61,729,388-62,237,324 | 17 |
| PRKCA | PKC alpha | Chromosome 17:61,729,388-62,237,324 | 17 |
| PRKCA | PKC alpha | Chromosome 17:61,729,388-62,237,324 | 17 |
| MAP2K6 |  | Chromosome 17:64,922,433-65,051,067 | 17 |
| MAP2K6 |  | Chromosome 17:64,922,433-65,051,067 | 17 |
| SSTR2 | sst2 receptor | Chromosome 17:68,672,755-68,679,689 | 17 |
| SSTR2 | sst2 receptor | Chromosome 17:68,672,755-68,679,689 | 17 |
| SSTR2 | sst2 receptor | Chromosome 17:68,672,755-68,679,689 | 17 |
| CD300A | IGSF12, MAIR1, IRC1, IR | Chromosome 17:69,974,117-69,992,528 | 17 |
| CD300A | IGSF12, MAIR1, IRC1, IR | Chromosome 17:69,974,117-69,992,528 | 17 |
| GPS2 |  | Chromosome 17:7,156,702-7,173,362 | 17 |
| FGF11 | FGF11 | Chromosome 17:7,283,413-7,288,980 | 17 |
| TNFRSF13C | CD268, BAFFR | Chromosome 17:7,392,932-7,405,649 | 17 |
| TNFSF12-13 |  | Chromosome 17:7,393,099-7,405,649 | 17 |
| ALOX15B |  | Chromosome 17:7,883,083-7,893,177 | 17 |
| CD300C | CMRF35A1-6, -H, LIR | Chromosome 17:70,048,842-70,053,877 | 17 |
| ENSG0000020 | MAIR2, CD300d | Chromosome 17:70,087,099-70,100,017 | 17 |
| CD300LF | IREM1 | Chromosome 17:70,202,047-70,220,712 | 17 |
| SLC25A19 | Mup1 | Chromosome 17:70,780,669-70,797,109 | 17 |
| SLC25A19 | Mup1 | Chromosome 17:70,780,669-70,797,109 | 17 |
| ITGB4 | CD104 | Chromosome 17:71,229,111-71,265,494 | 17 |
| FOXJ1 |  | Chromosome 17:71,644,009-71,648,966 | 17 |
| FOXJ1 |  | Chromosome 17:71,644,009-71,648,966 | 17 |
| SPHK1 | sphingosine kinase | Chromosome 17:71,892,297-71,895,536 | 17 |


| SPHK1 | sphingosine kinase | Chromosome 17:71,892,297-71,895,536 | 17 |
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| PTDSR |  | Chromosome 17:72,220,514-72,234,158 | 17 |
| BIRC5 | SURVIVIN | Chromosome 17:73,721,872-73,733,311 | 17 |
| BIRC5 | SURVIVIN | Chromosome 17:73,721,872-73,733,311 | 17 |
| SOCS3 |  | Chromosome 17:73,864,454-73,867,753 | 17 |
| SOCS3 |  | Chromosome 17:73,864,454-73,867,753 | 17 |
| PSCD1 | Cytohesin-1 | Chromosome 17:74,181,727-74,289,971 | 17 |
| PSCD1 | Cytohesin-1 | Chromosome 17:74,181,727-74,289,971 | 17 |
| TIMP2 |  | Chromosome 17:74,360,654-74,433,067 | 17 |
| EIF4A3 |  | Chromosome 17:75,723,612-75,735,533 | 17 |
| EIF4A3 |  | Chromosome 17:75,723,612-75,735,533 | 17 |
| CHMP6 | CHMP6 | Chromosome 17:76,580,274-76,588,528 | 17 |
| AATK |  | Chromosome 17:76,705,160-76,754,467 | 17 |
| ACTG1 | F-actin | Chromosome 17:77,091,594-77,094,422 | 17 |
| ARHGDIA | Rho GD1 | Chromosome 17:77,418,886-77,422,527 | 17 |
| CD7 | gp40, gp41 | Chromosome 17:77,866,035-77,868,769 | 17 |
| SECTM1 |  | Chromosome 17:77,872,189-77,884,930 | 17 |
| FOXK2 | ILF1 | Chromosome 17:78,070,883-78,153,743 | 17 |
| FOXK2 | ILF1 | Chromosome 17:78,070,883-78,153,743 | 17 |
| PIK3R5 |  | Chromosome 17:8,722,953-8,756,559 | 17 |
| PIK3R5 |  | Chromosome 17:8,722,953-8,756,559 | 17 |
| PIK3R5 |  | Chromosome 17:8,722,953-8,756,559 | 17 |
| GNAL | G protein alpha | Chromosome 18:11,679,263-11,871,922 | 18 |
| GNAL | G protein alpha | Chromosome 18:11,679,263-11,871,922 | 18 |
| GNAL | G protein alpha | Chromosome 18:11,679,263-11,871,922 | 18 |
| GNAL | G protein alpha | Chromosome 18:11,679,263-11,871,922 | 18 |
| GNAL | G protein alpha | Chromosome 18:11,679,263-11,871,922 | 18 |
| PTPN2 |  | Chromosome 18:12,775,480-12,874,334 | 18 |
| PTPN2 |  | Chromosome 18:12,775,480-12,874,334 | 18 |
| RNMT |  | Chromosome 18:13,716,680-13,754,554 | 18 |
| RNMT |  | Chromosome 18:13,716,680-13,754,554 | 18 |
| MC5R | melanocortin receptor 5 | Chromosome 18:13,815,543-13,816,861 | 18 |
| ROCK1 |  | Chromosome 18:16,787,533-16,944,869 | 18 |
| ROCK1 |  | Chromosome 18:16,787,533-16,944,869 | 18 |
| GATA6 |  | Chromosome 18:18,003,414-18,036,225 | 18 |
| LAMA3 |  | Chromosome 18:19,523,560-19,789,028 | 18 |
| LAMA3 |  | Chromosome 18:19,523,560-19,789,028 | 18 |
| LAMA3 |  | Chromosome 18:19,523,560-19,789,028 | 18 |
| HRH4 |  | Chromosome 18:20,294,591-20,313,919 | 18 |
| HRH4 |  | Chromosome 18:20,294,591-20,313,919 | 18 |
| THOC1 |  | Chromosome 18:204,522-258,049 | 18 |
| CDH2 | CD325, N-cadherin | Chromosome 18:23,784,933-24,011,189 | 18 |
| CDH2 | CD325, N-cadherin | Chromosome 18:23,784,933-24,011,189 | 18 |
| CDH2 | CD325, N-cadherin | Chromosome 18:23,784,933-24,011,189 | 18 |
| CDH2 | CD325, N-cadherin | Chromosome 18:23,784,933-24,011,189 | 18 |
| CDH2 | CD325, N-cadherin | Chromosome 18:23,784,933-24,011,189 | 18 |
| TGIF1 |  | Chromosome 18:3,402,072-3,448,409 | 18 |
| TGIF1 |  | Chromosome 18:3,402,072-3,448,409 | 18 |
| TGIF1 |  | Chromosome 18:3,402,072-3,448,409 | 18 |
| COLEC12 | SRCL-1, SCARA4 | Chromosome 18:309,356-490,685 | 18 |
| COLEC12 | SRCL-1, SCARA4 | Chromosome 18:309,356-490,685 | 18 |
| PIK3C3 |  | Chromosome 18:37,789,197-37,915,446 | 18 |
| PIK3C3 |  | Chromosome 18:37,789,197-37,915,446 | 18 |
| PIK3C3 |  | Chromosome 18:37,789,197-37,915,446 | 18 |
| SLC14A1 | Kidd antigen | Chromosome 18:41,558,155-41,585,297 | 18 |
| SLC14A1 | Kidd antigen | Chromosome 18:41,558,155-41,585,297 | 18 |


| CD33L3 |  | Chromosome 18:41,659,543-41,678,045 | 18 |
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| SMAD7 |  | Chromosome 18:44,700,221-44,731,079 | 18 |
| SMAD7 |  | Chromosome 18:44,700,221-44,731,079 | 18 |
| MAPK4 |  | Chromosome 18:46,340,482-46,512,194 | 18 |
| MAPK4 |  | Chromosome 18:46,340,482-46,512,194 | 18 |
| MAPK4 |  | Chromosome 18:46,340,482-46,512,194 | 18 |
| TCF4 | ITF2 | Chromosome 18:51,045,967-51,406,858 | 18 |
| TCF4 | ITF2 | Chromosome 18:51,045,967-51,406,858 | 18 |
| TCF4 | ITF2 | Chromosome 18:51,045,967-51,406,858 | 18 |
| TCF4 | ITF2 | Chromosome 18:51,045,967-51,406,858 | 18 |
| MALT1 |  | Chromosome 18:54,489,598-54,568,350 | 18 |
| MALT1 |  | Chromosome 18:54,489,598-54,568,350 | 18 |
| RAX |  | Chromosome 18:55,085,251-55,091,605 | 18 |
| RAX |  | Chromosome 18:55,085,251-55,091,605 | 18 |
| LMAN1 | MBL1 | Chromosome 18:55,148,088-55,177,463 | 18 |
| MC4R | melanocortin receptor 4 | Chromosome 18:56,189,564-56,190,562 | 18 |
| MC4R | melanocortin receptor 4 | Chromosome 18:56,189,564-56,190,562 | 18 |
| TNFRSF11A | CD265, OPG, RANK | Chromosome 18:58,143,500-58,205,872 | 18 |
| TNFRSF11A | CD265, OPG, RANK | Chromosome 18:58,143,500-58,205,872 | 18 |
| BCL2 |  | Chromosome 18:58,941,559-59,137,593 | 18 |
| BCL2 |  | Chromosome 18:58,941,559-59,137,593 | 18 |
| BCL2 |  | Chromosome 18:58,941,559-59,137,593 | 18 |
| SERPINB2 |  | Chromosome 18:59,705,922-59,722,100 | 18 |
| HMSD |  | Chromosome 18:59,767,574-59,779,093 | 18 |
| LAMA1 | laminin | Chromosome 18:6,931,885-7,107,813 | 18 |
| LAMA1 | laminin | Chromosome 18:6,931,885-7,107,813 | 18 |
| LAMA1 | laminin | Chromosome 18:6,931,885-7,107,813 | 18 |
| TYMS | Thymidylate synthase, TS | Chromosome 18:647,619-663,492 | 18 |
| DNAM1 | CD226 | Chromosome 18:65,681,172-65,775,140 | 18 |
| DNAM1 | CD226 | Chromosome 18:65,681,172-65,775,140 | 18 |
| DNAM1 | CD226 | Chromosome 18:65,681,172-65,775,140 | 18 |
| SOCS6 |  | Chromosome 18:66,107,243-66,145,329 | 18 |
| SOCS6 |  | Chromosome 18:66,107,243-66,145,329 | 18 |
| PTPRM |  | Chromosome 18:7,557,817-8,396,854 | 18 |
| PTPRM |  | Chromosome 18:7,557,817-8,396,854 | 18 |
| PTPRM |  | Chromosome 18:7,557,817-8,396,854 | 18 |
| PTPRM |  | Chromosome 18:7,557,817-8,396,854 | 18 |
| PTPRM |  | Chromosome 18:7,557,817-8,396,854 | 18 |
| PTPRM |  | Chromosome 18:7,557,817-8,396,854 | 18 |
| PTPRM |  | Chromosome 18:7,557,817-8,396,854 | 18 |
| PTPRM |  | Chromosome 18:7,557,817-8,396,854 | 18 |
| PTPRM |  | Chromosome 18:7,557,817-8,396,854 | 18 |
| YES1 |  | Chromosome 18:711,592-802,547 | 18 |
| YES1 |  | Chromosome 18:711,592-802,547 | 18 |
| MBP |  | Chromosome 18:72,819,777-72,973,762 | 18 |
| MBP |  | Chromosome 18:72,819,777-72,973,762 | 18 |
| NFATC1 |  | Chromosome 18:75,256,760-75,390,311 | 18 |
| NFATC1 |  | Chromosome 18:75,256,760-75,390,311 | 18 |
| RALBP1 | RIP1 | Chromosome 18:9,465,007-9,528,106 | 18 |
| HMHA1 | HA-1 | Chromosome 19:1,018,174-1,037,627 | 19 |
| TCF3 | E12 | Chromosome 19:1,560,293-1,603,328 | 19 |
| TCF3 | E12 | Chromosome 19:1,560,293-1,603,328 | 19 |
| TCF3 | E12 | Chromosome 19:1,560,293-1,603,328 | 19 |
| P2RY11 | P2Y11 | Chromosome 19:10,083,197-10,087,065 | 19 |
| P2RY11 | P2Y11 | Chromosome 19:10,083,197-10,087,065 | 19 |
| ICAM4 | CD242 | Chromosome 19:10,258,650-10,260,198 | 19 |


| TYK2 |  | Chromosome 19:10,322,205-10,352,211 | 19 |
| :---: | :---: | :---: | :---: |
| KEAP1 |  | Chromosome 19:10,457,796-10,475,243 | 19 |
| DNM2 | Dynamin2 | Chromosome 19:10,673,106-10,803,579 | 19 |
| CARM1 | PRMT4 | Chromosome 19:10,843,253-10,894,448 | 19 |
| SMARCA4 | SWI/SNF | Chromosome 19:10,932,606-11,033,953 | 19 |
| LDLR | LDL receptor | Chromosome 19:11,061,132-11,105,490 | 19 |
| RAB3D | Rab3d | Chromosome 19:11,296,093-11,311,321 | 19 |
| EPOR |  | Chromosome 19:11,348,883-11,356,019 | 19 |
| ECSIT |  | Chromosome 19:11,477,744-11,500,972 | 19 |
| JUNB |  | Chromosome 19:12,763,286-12,765,129 | 19 |
| CALR | Calreticulin | Chromosome 19:12,910,423-12,916,303 | 19 |
| RAD23A | CARD1 | Chromosome 19:12,917,654-12,925,455 | 19 |
| RFX1 | RFX1 | Chromosome 19:13,933,352-13,978,097 | 19 |
| RFX1 | RFX1 | Chromosome 19:13,933,352-13,978,097 | 19 |
| PRKACA | PKA alpha | Chromosome 19:14,063,500-14,089,559 | 19 |
| CD97 |  | Chromosome 19:14,353,213-14,380,535 | 19 |
| PTGER1 | EP1 | Chromosome 19:14,444,278-14,447,174 | 19 |
| EMR3 | CD313 | Chromosome 19:14,570,918-14,646,810 | 19 |
| EMR2 | CD312 | Chromosome 19:14,704,205-14,750,353 | 19 |
| CASP14 |  | Chromosome 19:15,024,015-15,027,900 | 19 |
| CASP14 |  | Chromosome 19:15,024,015-15,027,900 | 19 |
| NOTCH3 |  | Chromosome 19:15,131,444-15,172,792 | 19 |
| PGLYRP2 | PGRP-L | Chromosome 19:15,440,463-15,451,312 | 19 |
| CYP4F8 |  | Chromosome 19:15,587,421-15,601,445 | 19 |
| CYP4F3 |  | Chromosome 19:15,612,707-15,634,634 | 19 |
| CYP4F2 |  | Chromosome 19:15,849,834-15,869,885 | 19 |
| CYP4F11 |  | Chromosome 19:15,884,181-15,906,326 | 19 |
| HSH2D |  | Chromosome 19:16,105,838-16,130,381 | 19 |
| HSH2D |  | Chromosome 19:16,105,838-16,130,381 | 19 |
| HSH2D |  | Chromosome 19:16,105,838-16,130,381 | 19 |
| KLF2 | KLF2 | Chromosome 19:16,296,648-16,299,345 | 19 |
| CHERP | calcium homooestasis red | Chromosome 19:16,489,705-16,514,248 | 19 |
| NR2F6 | NR2F6 | Chromosome 19:17,203,694-17,217,151 | 19 |
| NR2F6 | NR2F6 | Chromosome 19:17,203,694-17,217,151 | 19 |
| BST2 | CD317 | Chromosome 19:17,374,755-17,377,457 | 19 |
| BST2 | CD317 | Chromosome 19:17,374,755-17,377,457 | 19 |
| B3GNT3 |  | Chromosome 19:17,766,658-17,785,385 | 19 |
| B3GNT3 |  | Chromosome 19:17,766,658-17,785,385 | 19 |
| JAK3 |  | Chromosome 19:17,788,322-17,819,800 | 19 |
| IL12RB1 | CD212 | Chromosome 19:18,031,371-18,058,702 | 19 |
| PIK3R2 |  | Chromosome 19:18,125,016-18,142,343 | 19 |
| GDF15 | CA19-9, MIC1 | Chromosome 19:18,357,968-18,360,987 | 19 |
| GDF15 | CA19-9, MIC1 | Chromosome 19:18,357,968-18,360,987 | 19 |
| FKBP8 | Calcineurin | Chromosome 19:18,503,568-18,515,383 | 19 |
| DDX49 |  | Chromosome 19:18,891,494-18,900,436 | 19 |
| RFXANK |  | Chromosome 19:19,164,008-19,173,678 | 19 |
| GADD45B |  | Chromosome 19:2,427,135-2,429,257 | 19 |
| GADD45B |  | Chromosome 19:2,427,135-2,429,257 | 19 |
| TBXAR2 | thromboxane A2 receptor | Chromosome 19:3,545,504-3,557,658 | 19 |
| EEF2 |  | Chromosome 19:3,927,054-3,936,461 | 19 |
| PIAS4 |  | Chromosome 19:3,958,748-3,990,383 | 19 |
| CCNE1 |  | Chromosome 19:34,994,741-35,007,059 | 19 |
| CCNE1 |  | Chromosome 19:34,994,741-35,007,059 | 19 |
| PDCD5 |  | Chromosome 19:37,763,944-37,770,171 | 19 |
| CEBPA |  | Chromosome 19:38,482,776-38,485,160 | 19 |
| CEBPA |  | Chromosome 19:38,482,776-38,485,160 | 19 |


| CEBPA |  | Chromosome 19:38,482,776-38,485,160 | 19 |
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| MAP2K2 | MEK3 | Chromosome 19:4,041,319-4,075,126 | 19 |
| EBI3 | IL35 | Chromosome 19:4,180,495-4,188,525 | 19 |
| EBI3 | IL35 | Chromosome 19:4,180,495-4,188,525 | 19 |
| C19orf10 |  | Chromosome 19:4,608,557-4,621,415 | 19 |
| C19orf10 |  | Chromosome 19:4,608,557-4,621,415 | 19 |
| TICAM1 | TRIF | Chromosome 19:4,766,944-4,782,716 | 19 |
| TICAM1 | TRIF | Chromosome 19:4,766,944-4,782,716 | 19 |
| USF2 | USF2a,b | Chromosome 19:40,451,721-40,462,558 | 19 |
| USF2 | USF2a,b | Chromosome 19:40,451,721-40,462,558 | 19 |
| ZBTB32 | ROG | Chromosome 19:40,895,670-40,899,780 | 19 |
| TA-NFKBH |  | Chromosome 19:41,070,983-41,085,025 | 19 |
| TA-NFKBH |  | Chromosome 19:41,070,983-41,085,025 | 19 |
| TA-NFKBH |  | Chromosome 19:41,070,983-41,085,025 | 19 |
| ALKBH6 |  | Chromosome 19:41,191,863-41,196,981 | 19 |
| MAP4K1 |  | Chromosome 19:43,770,121-43,800,471 | 19 |
| MAP4K1 |  | Chromosome 19:43,770,121-43,800,471 | 19 |
| LGALS4 | galectin 4 | Chromosome 19:43,984,155-43,995,422 | 19 |
| NFKBIB | IkB beta | Chromosome 19:44,082,455-44,091,374 | 19 |
| NFKBIB | IkB beta | Chromosome 19:44,082,455-44,091,374 | 19 |
| NFKBIB | IkB beta | Chromosome 19:44,082,455-44,091,374 | 19 |
| IL28B |  | Chromosome 19:44,426,033-44,427,609 | 19 |
| IL28A |  | Chromosome 19:44,450,997-44,452,572 | 19 |
| ZFP36 | TTP | Chromosome 19:44,589,293-44,591,885 | 19 |
| LGALS13 | galectin13 | Chromosome 19:44,785,004-44,789,954 | 19 |
| LGALS14 |  | Chromosome 19:44,886,786-44,891,928 | 19 |
| CLC | galectin 10 | Chromosome 19:44,913,735-44,920,508 | 19 |
| MADCAM1 |  | Chromosome 19:447,490-456,342 | 19 |
| MADCAM1 |  | Chromosome 19:447,490-456,342 | 19 |
| MAP3K10 |  | Chromosome 19:45,389,491-45,413,314 | 19 |
| MAP3K10 |  | Chromosome 19:45,389,491-45,413,314 | 19 |
| MAP3K10 |  | Chromosome 19:45,389,491-45,413,314 | 19 |
| BLVRB |  | Chromosome 19:45,645,541-45,663,516 | 19 |
| BLVRB |  | Chromosome 19:45,645,541-45,663,516 | 19 |
| CYP2A6 |  | Chromosome 19:46,041,284-46,226,008 | 19 |
| CYP2A7 |  | Chromosome 19:46,041,286-46,226,008 | 19 |
| CYP2A7 |  | Chromosome 19:46,041,286-46,226,008 | 19 |
| CYP2S1 |  | Chromosome 19:46,390,955-46,405,284 | 19 |
| TGFB1 |  | Chromosome 19:46,528,254-46,551,656 | 19 |
| TGFB1 |  | Chromosome 19:46,528,254-46,551,656 | 19 |
| CEACAM5 | CD66e | Chromosome 19:46,904,377-46,925,686 | 19 |
| CEACAM5 | CD66e | Chromosome 19:46,904,377-46,925,686 | 19 |
| CEACAM3 | CD66d | Chromosome 19:46,992,381-47,007,431 | 19 |
| CEACAM3 | CD66d | Chromosome 19:46,992,381-47,007,431 | 19 |
| DEDD2 |  | Chromosome 19:47,394,592-47,416,115 | 19 |
| DEDD2 |  | Chromosome 19:47,394,592-47,416,115 | 19 |
| KIR2DL5B |  | Chromosome 19:47,577,500-47,579,250 | 19 |
| CEACAM1 | CD66a | Chromosome 19:47,703,298-47,724,479 | 19 |
| CEACAM8 | CD66b | Chromosome 19:47,776,235-47,790,890 | 19 |
| CEACAM8 | CD66b | Chromosome 19:47,776,235-47,790,890 | 19 |
| PSG1 | CD66f | Chromosome 19:48,063,198-48,075,711 | 19 |
| PSG1 | CD66f | Chromosome 19:48,063,198-48,075,711 | 19 |
| CD177 | CD177, PRV1 | Chromosome 19:48,549,651-48,559,368 | 19 |
| CD177 | CD177, PRV1 | Chromosome 19:48,549,651-48,559,368 | 19 |
| XRCC1 |  | Chromosome 19:48,739,032-48,771,998 | 19 |
| XRCC1 |  | Chromosome 19:48,739,032-48,771,998 | 19 |


| PLAUR | CD87 | Chromosome 19:48,842,088-48,866,539 | 19 |
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| IRGC | IGTP, TGTP | Chromosome 19:48,912,078-48,916,013 | 19 |
| IRGC | IGTP, TGTP | Chromosome 19:48,912,078-48,916,013 | 19 |
| PVR | CD155 | Chromosome 19:49,839,066-49,858,690 | 19 |
| PVR | CD155 | Chromosome 19:49,839,066-49,858,690 | 19 |
| BCL3 |  | Chromosome 19:49,943,820-49,955,140 | 19 |
| BCL3 |  | Chromosome 19:49,943,820-49,955,140 | 19 |
| PTPRS |  | Chromosome 19:5,157,379-5,237,399 | 19 |
| PTPRS |  | Chromosome 19:5,157,379-5,237,399 | 19 |
| FUT3 | CD174 | Chromosome 19:5,793,902-5,802,482 | 19 |
| FUT3 | CD174 | Chromosome 19:5,793,902-5,802,482 | 19 |
| RFX2 | RFX2 | Chromosome 19:5,944,175-6,061,554 | 19 |
| RFX2 | RFX2 | Chromosome 19:5,944,175-6,061,554 | 19 |
| RELB |  | Chromosome 19:50,196,539-50,233,292 | 19 |
| ERCC2 | XPD | Chromosome 19:50,546,686-50,565,669 | 19 |
| ERCC2 | XPD | Chromosome 19:50,546,686-50,565,669 | 19 |
| ERCC1 |  | Chromosome 19:50,604,712-50,619,017 | 19 |
| GPR4 |  | Chromosome 19:50,784,865-50,797,294 | 19 |
| FOXA3 |  | Chromosome 19:51,059,358-51,068,895 | 19 |
| FOXA3 |  | Chromosome 19:51,059,358-51,068,895 | 19 |
| FOXA3 |  | Chromosome 19:51,059,358-51,068,895 | 19 |
| PGLYRP1 | PGRP-Ia | Chromosome 19:51,214,255-51,218,163 | 19 |
| PKD2 |  | Chromosome 19:51,869,413-51,911,597 | 19 |
| KIR2DL4 |  | Chromosome 19:52,356,000-52.367.000 | 19 |
| BBC3 | PUMA | Chromosome 19:52,415,921-52,427,863 | 19 |
| C5R1 | CD88 | Chromosome 19:52,504,971-52,517,173 | 19 |
| DHX34 |  | Chromosome 19:52,544,386-52,577,795 | 19 |
| LIG1 |  | Chromosome 19:53,310,515-53,365,372 | 19 |
| CARD8 | TUCAN, CARDINAL | Chromosome 19:53,403,325-53,450,955 | 19 |
| EMP3 | Emp3 | Chromosome 19:53,520,441-53,525,623 | 19 |
| SPHK2 | sphingosine kinase | Chromosome 19:53,814,360-53,825,474 | 19 |
| FUT1 | CD173 | Chromosome 19:53,943,080-53,950,459 | 19 |
| FGF21 | FGF21 | Chromosome 19:53,950,628-53,953,395 | 19 |
| FGF21 | FGF21 | Chromosome 19:53,950,628-53,953,395 | 19 |
| BAX |  | Chromosome 19:54,149,929-54,156,867 | 19 |
| LHB | LHB | Chromosome 19:54,211,049-54,212,159 | 19 |
| CGB | CGB | Chromosome 19:54,217,939-54,244,212 | 19 |
| SNRP70 |  | Chromosome 19:54,280,277-54,303,682 | 19 |
| CD37 |  | Chromosome 19:54,530,240-54,535,675 | 19 |
| CD37 |  | Chromosome 19:54,530,240-54,535,675 | 19 |
| CD37 |  | Chromosome 19:54,530,240-54,535,675 | 19 |
| FCGRT | FCRn | Chromosome 19:54,708,304-54,721,402 | 19 |
| FCGRT | FCRn | Chromosome 19:54,708,304-54,721,402 | 19 |
| NOSIP | NO synthase | Chromosome 19:54,750,780-54,775,626 | 19 |
| IL4I1 |  | Chromosome 19:55,084,723-55,124,598 | 19 |
| NR1H2 | LXRbeta | Chromosome 19:55,571,515-55,578,051 | 19 |
| NR1H2 | LXRbeta | Chromosome 19:55,571,515-55,578,051 | 19 |
| SIGLEC9 | CDw329 | Chromosome 19:56,319,977-56,325,379 | 19 |
| SIGLEC9 | CDw329 | Chromosome 19:56,319,977-56,325,379 | 19 |
| SIGLEC10 | CDw330 | Chromosome 19:56,605,087-56,612,869 | 19 |
| SIGLEC6 | CDw327 | Chromosome 19:56,714,795-56,726,922 | 19 |
| SIGLEC5 | CD170 | Chromosome 19:56,806,996-56,831,696 | 19 |
| FPR1 |  | Chromosome 19:56,940,839-56,946,962 | 19 |
| FPRL1 | Annexin-1R, f-MLP | Chromosome 19:56,955,995-56,965,591 | 19 |
| BIRC8 |  | Chromosome 19:58,484,666-58,486,687 | 19 |
| BIRC8 |  | Chromosome 19:58,484,666-58,486,687 | 19 |


| NLRP12 | NALP12, RNO2, PYPAF7 | Chromosome 19:58,988,650-59,019,409 | 19 |
| :---: | :---: | :---: | :---: |
| NLRP12 | NALP12, RNO2, PYPAF7 | Chromosome 19:58,988,650-59,019,409 | 19 |
| PRKCG | PKC gamma, Cgamma | Chromosome 19:59,077,279-59,102,713 | 19 |
| PRKCG | PKC gamma, Cgamma | Chromosome 19:59,077,279-59,102,713 | 19 |
| OSCAR |  | Chromosome 19:59,289,745-59,297,806 | 19 |
| OSCAR |  | Chromosome 19:59,289,745-59,297,806 | 19 |
| LILRB3 | CD85a, PIRB (mouse) | Chromosome 19:59,412,549-59,418,709 | 19 |
| LILRA3 |  | Chromosome 19:59,491,666-59,496,077 | 19 |
| LAIR1 | CD305 | Chromosome 19:59,557,047-59,568,533 | 19 |
| TTYH1 |  | Chromosome 19:59,618,417-59,639,882 | 19 |
| KIR3DX1 | KIR3DL0 | Chromosome 19:59,738,595-59,748,862 | 19 |
| LILRP2 | ILT10 | Chromosome 19:59,911,791-59,916,501 | 19 |
| GTF2F1 | TFIIF | Chromosome 19:6,235,811-6,344,184 | 19 |
| ALKBH7 |  | Chromosome 19:6,323,444-6,326,040 | 19 |
| TNFSF9 | 4-1BBL | Chromosome 19:6,482,037-6,486,933 | 19 |
| TNFSF7 | CD70 | Chromosome 19:6,536,850-6,542,163 | 19 |
| C3 | C3bBb, C3bBb3b | Chromosome 19:6,628,878-6,671,660 | 19 |
| C3 | C3bBb, C3bBb3b | Chromosome 19:6,628,878-6,671,660 | 19 |
| VAV1 |  | Chromosome 19:6,723,722-6,808,371 | 19 |
| EMR1 | CD311 | Chromosome 19:6,838,577-6,891,464 | 19 |
| EMR1 | CD311 | Chromosome 19:6,838,577-6,891,464 | 19 |
| FCAR | CD89 | Chromosome 19:60,077,361-60,095,055 | 19 |
| NLRP2 | NALP2, LRR,PYPAF2, N | Chromosome 19:60,168,465-60,204,318 | 19 |
| PTPRH |  | Chromosome 19:60,384,428-60,412,654 | 19 |
| NLRP9 | NALP9, NOD6 | Chromosome 19:60,911,610-60,941,580 | 19 |
| NLRP9 | NALP9, NOD6 | Chromosome 19:60,911,610-60,941,580 | 19 |
| NLRP13 | NALP13, NOD14 | Chromosome 19:61,099,123-61,135,489 | 19 |
| CHMP2A | CHMP2A | Chromosome 19:63,754,745-63,758,298 | 19 |
| INSR | CD220, Insulin receptor | Chromosome 19:7,067,049-7,245,045 | 19 |
| INSR | CD220, Insulin receptor | Chromosome 19:7,067,049-7,245,045 | 19 |
| INSR | CD220, Insulin receptor | Chromosome 19:7,067,049-7,245,045 | 19 |
| FCER2 | CD23 | Chromosome 19:7,659,662-7,673,032 | 19 |
| FCER2 | CD23 | Chromosome 19:7,659,662-7,673,032 | 19 |
| CLEC4M | CD299, DC-SIGN-R, L-SI | Chromosome 19:7,734,081-7,740,491 | 19 |
| MAP2K7 |  | Chromosome 19:7,874,728-7,885,363 | 19 |
| AZU1 | HBP, CAP37 | Chromosome 19:776,097-783,017 | 19 |
| CCL25 |  | Chromosome 19:8,023,934-8,033,547 | 19 |
| MYO1F | IF | Chromosome 19:8,491,689-8,548,330 | 19 |
| CFD |  | Chromosome 19:810,665-814,624 | 19 |
| PIN1 |  | Chromosome 19:9,806,999-9,821,358 | 19 |
| COL5A3 |  | Chromosome 19:9,931,237-9,982,147 | 19 |
| PXDN |  | Chromosome 2:1,614,666-1,727,298 | 2 |
| KLF11 | TIEG | Chromosome 2:10,101,133-10,112,414 | 2 |
| MAP4K4 |  | Chromosome 2:101,680,920-101,877,584 | 2 |
| IL1R2 | CD121b | Chromosome 2:101,974,738-102,011,317 | 2 |
| IL1R1 | CD121a | Chromosome 2:102,125,678-102,162,766 | 2 |
| IL1RL2 |  | Chromosome 2:102,169,865-102,222,243 | 2 |
| IL1RL1 | ST2 | Chromosome 2:102,294,394-102,334,929 | 2 |
| IL18R1 | CDw218a | Chromosome 2:102,345,529-102,381,650 | 2 |
| IL18RAP | CDw218b | Chromosome 2:102,401,686-102,435,457 | 2 |
| SLC9A2 | NHE-2 | Chromosome 2:102,602,598-102,694,241 | 2 |
| SULT1C2 |  | Chromosome 2:108,360,853-108,370,702 | 2 |
| SULT1C2 |  | Chromosome 2:108,360,853-108,370,702 | 2 |
| RANBP2 |  | Chromosome 2:108,702,369-108,767,683 | 2 |
| EDAR |  | Chromosome 2:108,877,361-108,972,260 | 2 |
| ROCK2 | ROCKalpha | Chromosome 2:11,239,229-11,402,162 | 2 |


| ROCK2 | ROCKalpha | Chromosome 2:11,239,229-11,402,162 | 2 |
| :---: | :---: | :---: | :---: |
| ROCK2 | ROCKalpha | Chromosome 2:11,239,229-11,402,162 | 2 |
| BCL2L11 | Bim, BIM | Chromosome 2:111,597,781-111,641,058 | 2 |
| BCL2L11 | Bim, BIM | Chromosome 2:111,597,781-111,641,058 | 2 |
| MERTK | Mer Receptor Tyrosine kil | Chromosome 2:112,372,662-112,503,416 | 2 |
| MERTK | Mer Receptor Tyrosine kil | Chromosome 2:112,372,662-112,503,416 | 2 |
| IL1A |  | Chromosome 2:113,247,963-113,259,442 | 2 |
| IL1B |  | Chromosome 2:113,303,808-113,310,827 | 2 |
| IL1B |  | Chromosome 2:113,303,808-113,310,827 | 2 |
| IL1B |  | Chromosome 2:113,303,808-113,310,827 | 2 |
| IL1F7 |  | Chromosome 2:113,387,017-113,392,930 | 2 |
| IL1F7 |  | Chromosome 2:113,387,017-113,392,930 | 2 |
| IL1F7 |  | Chromosome 2:113,387,017-113,392,930 | 2 |
| IL1F9 |  | Chromosome 2:113,452,077-113,459,698 | 2 |
| IL1RN | CD25, IL1RA | Chromosome 2:113,591,941-113,608,064 | 2 |
| IL1RN | CD25, IL1RA | Chromosome 2:113,591,941-113,608,064 | 2 |
| DDX18 |  | Chromosome 2:118,288,725-118,306,425 | 2 |
| MARCO | SCARA2 | Chromosome 2:119,416,215-119,468,706 | 2 |
| MARCO | SCARA2 | Chromosome 2:119,416,215-119,468,706 | 2 |
| C1QL2 |  | Chromosome 2:119,630,289-119,632,941 | 2 |
| C1QL2 |  | Chromosome 2:119,630,289-119,632,941 | 2 |
| TMEM37 |  | Chromosome 2:119,905,950-119,911,486 | 2 |
| TMEM37 |  | Chromosome 2:119,905,950-119,911,486 | 2 |
| PTPN4 |  | Chromosome 2:120,233,677-120,451,507 | 2 |
| PTPN4 |  | Chromosome 2:120,233,677-120,451,507 | 2 |
| PTPN4 |  | Chromosome 2:120,233,677-120,451,507 | 2 |
| INHBB | Inhibin B | Chromosome 2:120,819,469-120,825,444 | 2 |
| INHBB | Inhibin B | Chromosome 2:120,819,469-120,825,444 | 2 |
| GLI2 | Gli2 | Chromosome 2:121,266,327-121,466,321 | 2 |
| GLI2 | Gli2 | Chromosome 2:121,266,327-121,466,321 | 2 |
| GLI2 | Gli2 | Chromosome 2:121,266,327-121,466,321 | 2 |
| GLI2 | Gli2 | Chromosome 2:121,266,327-121,466,321 | 2 |
| CLASP1 | clAP1 | Chromosome 2:121,811,825-122,123,522 | 2 |
| CLASP1 | clAP1 | Chromosome 2:121,811,825-122,123,522 | 2 |
| CLASP1 | CIAP1 | Chromosome 2:121,811,825-122,123,522 | 2 |
| CLASP1 | CIAP1 | Chromosome 2:121,811,825-122,123,522 | 2 |
| CLASP1 | cIAP1 | Chromosome 2:121,811,825-122,123,522 | 2 |
| GYPC | CD236c, CD236d, CD23¢ | Chromosome 2:127,130,154-127,170,716 | 2 |
| GYPC | CD236c, CD236d, CD23 | Chromosome 2:127,130,154-127,170,716 | 2 |
| GYPC | CD236c, CD236d, CD236 | Chromosome 2:127,130,154-127,170,716 | 2 |
| MAP3K2 |  | Chromosome 2:127,778,609-127,817,240 | 2 |
| MAP3K2 |  | Chromosome 2:127,778,609-127,817,240 | 2 |
| PROC | activated protein C | Chromosome 2:127,892,486-127,903,288 | 2 |
| PROC | activated protein C | Chromosome 2:127,892,486-127,903,288 | 2 |
| PTPN18 |  | Chromosome 2:130,830,088-130,848,614 | 2 |
| PTPN18 |  | Chromosome 2:130,830,088-130,848,614 | 2 |
| MCM6 |  | Chromosome 2:136,313,666-136,350,481 | 2 |
| MCM6 |  | Chromosome 2:136,313,666-136,350,481 | 2 |
| CXCR4 | CD184 | Chromosome 2:136,705,639-136,709,450 | 2 |
| CXCR4 | CD184 | Chromosome 2:136,705,639-136,709,450 | 2 |
| HNMT | HMT | Chromosome 2:138,438,278-138,490,404 | 2 |
| HNMT | HMT | Chromosome 2:138,438,278-138,490,404 | 2 |
| DDX1 |  | Chromosome 2:15,648,753-15,688,676 | 2 |
| DDX1 |  | Chromosome 2:15,648,753-15,688,676 | 2 |
| MYCN | n-myc | Chromosome 2:15,998,134-16,004,580 | 2 |
| MYCN | n-myc | Chromosome 2:15,998,134-16,004,580 | 2 |


| NMI |  | Chromosome 2:151,835,231-151,854,620 | 2 |
| :---: | :---: | :---: | :---: |
| NMI |  | Chromosome 2:151,835,231-151,854,620 | 2 |
| PRPF40A | FNBP3 | Chromosome 2:153,216,334-153,283,014 | 2 |
| PRPF40A | FNBP3 | Chromosome 2:153,216,334-153,283,014 | 2 |
| NR4A2 | NR4A2, NURR1 | Chromosome 2:156,889,194-156,897,474 | 2 |
| PSCDBP | CYTIP | Chromosome 2:157,979,377-158,008,850 | 2 |
| PSCDBP | CYTIP | Chromosome 2:157,979,377-158,008,850 | 2 |
| MARCH7 | Axotrophin | Chromosome 2:160,277,256-160,333,330 | 2 |
| MARCH7 | Axotrophin | Chromosome 2:160,277,256-160,333,330 | 2 |
| MARCH7 | Axotrophin | Chromosome 2:160,277,256-160,333,330 | 2 |
| LY75 | CD205, DEC-205 | Chromosome 2:160,368,118-160,469,493 | 2 |
| PLA2R1 | phospholipase A3 | Chromosome 2:160,505,506-160,627,367 | 2 |
| ITGB6 |  | Chromosome 2:160,664,438-160,765,009 | 2 |
| TANK | NFKB activator | Chromosome 2:161,701,712-161,800,928 | 2 |
| TANK | NFKB activator | Chromosome 2:161,701,712-161,800,928 | 2 |
| DPP4 | CD26 | Chromosome 2:162,557,001-162,639,298 | 2 |
| DPP4 | CD26 | Chromosome 2:162,557,001-162,639,298 | 2 |
| IFIH1 | MDA5, Ifit1 | Chromosome 2:162,831,835-162,883,285 | 2 |
| PPIG |  | Chromosome 2:170,149,096-170,202,500 | 2 |
| PPIG |  | Chromosome 2:170,149,096-170,202,500 | 2 |
| HAT1 |  | Chromosome 2:172,487,204-172,556,846 | 2 |
| HAT1 |  | Chromosome 2:172,487,204-172,556,846 | 2 |
| ITGA6 | CD49f | Chromosome 2:173,000,616-173,079,256 | 2 |
| PDK1 |  | Chromosome 2:173,129,025-173,172,108 | 2 |
| SP3 | chondromodulin transcrip | Chromosome 2:174,481,504-174,538,676 | 2 |
| SP3 | chondromodulin transcrip | Chromosome 2:174,481,504-174,538,676 | 2 |
| CIR |  | Chromosome 2:174,921,124-174,968,689 | 2 |
| CIR |  | Chromosome 2:174,921,124-174,968,689 | 2 |
| WIPF1 | WASPIP | Chromosome 2:175,132,548-175,255,873 | 2 |
| WIPF1 | WASPIP | Chromosome 2:175,132,548-175,255,873 | 2 |
| HOXD10 | HOX cluster | Chromosome 2:176,689,738-176,692,916 | 2 |
| HOXD10 | HOX cluster | Chromosome 2:176,689,738-176,692,916 | 2 |
| AGPS | ADAP, alkylglycerone phc | Chromosome 2:177,965,731-178,112,411 | 2 |
| AGPS | ADAP, alkylglycerone phq | Chromosome 2:177,965,731-178,112,411 | 2 |
| AGPS | ADAP, alkylglycerone phq | Chromosome 2:177,965,731-178,112,411 | 2 |
| ITGA4 | VLA4, CD49d | Chromosome 2:182,029,864-182,110,719 | 2 |
| ITGA4 | VLA4, CD49d | Chromosome 2:182,029,864-182,110,719 | 2 |
| FRZB | sFRP3 | Chromosome 2:183,406,982-183,439,743 | 2 |
| DUSP19 |  | Chromosome 2:183,651,732-183,673,616 | 2 |
| DUSP19 |  | Chromosome 2:183,651,732-183,673,616 | 2 |
| ITGAV | CD51 | Chromosome 2:187,163,045-187,253,873 | 2 |
| ITGAV | CD51 | Chromosome 2:187,163,045-187,253,873 | 2 |
| ITGAV | CD51 | Chromosome 2:187,163,045-187,253,873 | 2 |
| COL3A1 |  | Chromosome 2:189,547,344-189,585,717 | 2 |
| COL3A1 |  | Chromosome 2:189,547,344-189,585,717 | 2 |
| COL3A1 |  | Chromosome 2:189,547,344-189,585,717 | 2 |
| SLC40A1 | ferroportin | Chromosome 2:190,133,561-190,153,858 | 2 |
| SLC40A1 | ferroportin | Chromosome 2:190,133,561-190,153,858 | 2 |
| STAT1 |  | Chromosome 2:191,542,121-191,587,181 | 2 |
| STAT1 |  | Chromosome 2:191,542,121-191,587,181 | 2 |
| STAT4 |  | Chromosome 2:191,602,551-191,724,539 | 2 |
| GTF3C3 |  | Chromosome 2:197,336,917-197,372,670 | 2 |
| GTF3C3 |  | Chromosome 2:197,336,917-197,372,670 | 2 |
| HSPD1 | HSP65 | Chromosome 2:198,059,553-198,073,243 | 2 |
| SDC1 | Syndecan, CD138 | Chromosome 2:20,264,039-20,288,675 | 2 |
| PPIL3 |  | Chromosome 2:201,443,924-201,462,244 | 2 |


| PPIL3 |  | Chromosome 2:201,443,924-201,462,244 | 2 |
| :---: | :---: | :---: | :---: |
| CASP10 |  | Chromosome 2:201,756,100-201,802,372 | 2 |
| CFLAR | cFLIP, vFLIP | Chromosome 2:201,806,396-201,854,521 | 2 |
| BMPR2 |  | Chromosome 2:202,949,916-203,140,719 | 2 |
| BMPR2 |  | Chromosome 2:202,949,916-203,140,719 | 2 |
| CYP20A1 |  | Chromosome 2:203,811,658-203,878,579 | 2 |
| CYP20A1 |  | Chromosome 2:203,811,658-203,878,579 | 2 |
| CD28 |  | Chromosome 2:204,279,443-204,310,801 | 2 |
| CD28 |  | Chromosome 2:204,279,443-204,310,801 | 2 |
| CTLA4 | CD152 | Chromosome 2:204,440,754-204,446,928 | 2 |
| CTLA4 | CD152 | Chromosome 2:204,440,754-204,446,928 | 2 |
| CTLA4 | CD152 | Chromosome 2:204,440,754-204,446,928 | 2 |
| KLF7 |  | Chromosome 2:207,653,323-207,738,859 | 2 |
| KLF7 |  | Chromosome 2:207,653,323-207,738,859 | 2 |
| KLF7 |  | Chromosome 2:207,653,323-207,738,859 | 2 |
| CREB1 |  | Chromosome 2:208,102,931-208,171,818 | 2 |
| CPS1 |  | Chromosome 2:211,050,678-211,252,076 | 2 |
| CPS1 |  | Chromosome 2:211,050,678-211,252,076 | 2 |
| CPS1 |  | Chromosome 2:211,050,678-211,252,076 | 2 |
| CPS1 |  | Chromosome 2:211,050,678-211,252,076 | 2 |
| FN1 | fibronectin | Chromosome 2:215,933,409-216,009,041 | 2 |
| FN1 | fibronectin | Chromosome 2:215,933,409-216,009,041 | 2 |
| XRCC5 |  | Chromosome 2:216,680,435-216,779,248 | 2 |
| XRCC5 |  | Chromosome 2:216,680,435-216,779,248 | 2 |
| XRCC5 |  | Chromosome 2:216,680,435-216,779,248 | 2 |
| SMARCAL1 |  | Chromosome 2:216,985,441-217,056,021 | 2 |
| SMARCAL1 |  | Chromosome 2:216,985,441-217,056,021 | 2 |
| SMARCAL1 |  | Chromosome 2:216,985,441-217,056,021 | 2 |
| IGFBP2 |  | Chromosome 2:217,206,372-217,237,404 | 2 |
| IGFBP2 |  | Chromosome 2:217,206,372-217,237,404 | 2 |
| IL8RB | CD128b, CXCR2 | Chromosome 2:218,698,991-218,710,220 | 2 |
| SCL11A1 | NRAMP1 | Chromosome 2:218,955,161-218,968,994 | 2 |
| SCL11A1 | NRAMP1 | Chromosome 2:218,955,161-218,968,994 | 2 |
| CYP27A1 |  | Chromosome 2:219,354,745-219,388,259 | 2 |
| CYP27A1 |  | Chromosome 2:219,354,745-219,388,259 | 2 |
| CYP27A1 |  | Chromosome 2:219,354,745-219,388,259 | 2 |
| IHH | indian Hh | Chromosome 2:219,628,173-219,633,433 | 2 |
| IHH | indian Hh | Chromosome 2:219,628,173-219,633,433 | 2 |
| IHH | indian Hh | Chromosome 2:219,628,173-219,633,433 | 2 |
| TUBA1 |  | Chromosome 2:219,822,677-219,826,882 | 2 |
| INHA | Inhibin A | Chromosome 2:220,145,161-220,148,679 | 2 |
| COL4A4 |  | Chromosome 2:227,578,168-227,737,519 | 2 |
| COL4A4 |  | Chromosome 2:227,578,168-227,737,519 | 2 |
| COL4A3 |  | Chromosome 2:227,737,525-227,887,751 | 2 |
| HRB | HIV binding protein | Chromosome 2:228,045,286-228,130,548 | 2 |
| CCL20 |  | Chromosome 2:228,386,814-228,390,494 | 2 |
| CCL20 |  | Chromosome 2:228,386,814-228,390,494 | 2 |
| SP110 |  | Chromosome 2:230,741,896-230,792,932 | 2 |
| SP110 |  | Chromosome 2:230,741,896-230,792,932 | 2 |
| SP110 |  | Chromosome 2:230,741,896-230,792,932 | 2 |
| HTR2B | 5-HT2B | Chromosome 2:231,681,199-231,698,068 | 2 |
| HTR2B | 5-HT2B | Chromosome 2:231,681,199-231,698,068 | 2 |
| INPP5D | SHIP1 | Chromosome 2:233,633,433-233,781,288 | 2 |
| INPP5D | SHIP1 | Chromosome 2:233,633,433-233,781,288 | 2 |
| UGT1A1 | UGT1 | Chromosome 2:234,191,030-234,346,695 | 2 |
| UGT1A1 | UGT1 | Chromosome 2:234,191,030-234,346,695 | 2 |


| UGT1A1 | UGT1 | Chromosome 2:234,191,030-234,346,695 | 2 |
| :---: | :---: | :---: | :---: |
| UGT1A1 | UGT1 | Chromosome 2:234,191,030-234,346,695 | 2 |
| CXCR7 | CMKOR1 | Chromosome 2:237,143,182-237,155,730 | 2 |
| CXCR7 | CMKOR1 | Chromosome 2:237,143,182-237,155,730 | 2 |
| COL6A3 |  | Chromosome 2:237,897,401-237,987,559 | 2 |
| COL6A3 |  | Chromosome 2:237,897,401-237,987,559 | 2 |
| TRAF3IP1 |  | Chromosome 2:238,893,821-238,972,536 | 2 |
| TRAF3IP1 |  | Chromosome 2:238,893,821-238,972,536 | 2 |
| TRAF3IP1 |  | Chromosome 2:238,893,821-238,972,536 | 2 |
| ASB1 |  | Chromosome 2:239,000,365-239,025,630 | 2 |
| ASB1 |  | Chromosome 2:239,000,365-239,025,630 | 2 |
| HDAC4 |  | Chromosome 2:239,635,319-239,987,580 | 2 |
| HDAC4 |  | Chromosome 2:239,635,319-239,987,580 | 2 |
| HDAC4 |  | Chromosome 2:239,635,319-239,987,580 | 2 |
| HDAC4 |  | Chromosome 2:239,635,319-239,987,580 | 2 |
| FKBP1B | Calcineurin | Chromosome 2:24,126,075-24,140,055 | 2 |
| FKBP1B | Calcineurin | Chromosome 2:24,126,075-24,140,055 | 2 |
| DUSP28 |  | Chromosome 2:241,148,144-241,152,104 | 2 |
| DUSP28 |  | Chromosome 2:241,148,144-241,152,104 | 2 |
| SEPT2 |  | Chromosome 2:241,903,396-241,942,115 | 2 |
| SEPT2 |  | Chromosome 2:241,903,396-241,942,115 | 2 |
| BOK |  | Chromosome 2:242,146,865-242,162,226 | 2 |
| PDCD1 | CD279, PD1 | Chromosome 2:242,440,711-242,449,731 | 2 |
| POMC | POMC | Chromosome 2:25,237,226-25,245,063 | 2 |
| POMC | POMC | Chromosome 2:25,237,226-25,245,063 | 2 |
| UCN | urocortin | Chromosome 2:27,383,769-27,384,634 | 2 |
| ALK | CD246 | Chromosome 2:29,269,144-29,997,936 | 2 |
| ALK | CD246 | Chromosome 2:29,269,144-29,997,936 | 2 |
| ALK | CD246 | Chromosome 2:29,269,144-29,997,936 | 2 |
| ALK | CD246 | Chromosome 2:29,269,144-29,997,936 | 2 |
| ALK | CD246 | Chromosome 2:29,269,144-29,997,936 | 2 |
| ALK | CD246 | Chromosome 2:29,269,144-29,997,936 | 2 |
| XDH | Xanthine oxidoreductase | Chromosome 2:31,410,691-31,491,117 | 2 |
| XDH | Xanthine oxidoreductase | Chromosome 2:31,410,691-31,491,117 | 2 |
| XDH | Xanthine oxidoreductase | Chromosome 2:31,410,691-31,491,117 | 2 |
| XDH | Xanthine oxidoreductase | Chromosome 2:31,410,691-31,491,117 | 2 |
| NLRC4 | CARD12, IPAF, CLAN | Chromosome 2:32,303,022-32,344,427 | 2 |
| NLRC4 | CARD12, IPAF, CLAN | Chromosome 2:32,303,022-32,344,427 | 2 |
| BIRC6 |  | Chromosome 2:32,435,234-32,697,470 | 2 |
| BIRC6 |  | Chromosome 2:32,435,234-32,697,470 | 2 |
| LTBP1 |  | Chromosome 2:33,025,896-33,478,080 | 2 |
| LTBP1 |  | Chromosome 2:33,025,896-33,478,080 | 2 |
| LTBP1 |  | Chromosome 2:33,025,896-33,478,080 | 2 |
| LTBP1 |  | Chromosome 2:33,025,896-33,478,080 | 2 |
| LTBP1 |  | Chromosome 2:33,025,896-33,478,080 | 2 |
| LTBP1 |  | Chromosome 2:33,025,896-33,478,080 | 2 |
| LTBP1 |  | Chromosome 2:33,025,896-33,478,080 | 2 |
| LTBP1 |  | Chromosome 2:33,025,896-33,478,080 | 2 |
| LTBP1 |  | Chromosome 2:33,025,896-33,478,080 | 2 |
| RASGRP3 |  | Chromosome 2:33,514,920-33,643,162 | 2 |
| PKD3 |  | Chromosome 2:37,331,149-37,398,541 | 2 |
| PKD3 |  | Chromosome 2:37,331,149-37,398,541 | 2 |
| PKD3 |  | Chromosome 2:37,331,149-37,398,541 | 2 |
| CYP1B1 | Cytochromes | Chromosome 2:38,148,154-38,156,796 | 2 |
| CYP1B1 | Cytochromes | Chromosome 2:38,148,154-38,156,796 | 2 |
| DHX57 |  | Chromosome 2:38,878,375-38,956,525 | 2 |


| DHX57 |  | Chromosome 2:38,878,375-38,956,525 | 2 |
| :---: | :---: | :---: | :---: |
| DHX57 |  | Chromosome 2:38,878,375-38,956,525 | 2 |
| SOS1 |  | Chromosome 2:39,066,469-39,201,067 | 2 |
| SOS1 |  | Chromosome 2:39,066,469-39,201,067 | 2 |
| MAP4K3 |  | Chromosome 2:39,329,911-39,517,946 | 2 |
| MAP4K3 |  | Chromosome 2:39,329,911-39,517,946 | 2 |
| MAP4K3 |  | Chromosome 2:39,329,911-39,517,946 | 2 |
| PLEKHH2 | MAX1 | Chromosome 2:43,717,916-43,848,630 | 2 |
| PRKCD | PKC epsilon | Chromosome 2:45,732,547-46,268,633 | 2 |
| PRKCD | PKC epsilon | Chromosome 2:45,732,547-46,268,633 | 2 |
| PRKCD | PKC epsilon | Chromosome 2:45,732,547-46,268,633 | 2 |
| PRKCD | PKC epsilon | Chromosome 2:45,732,547-46,268,633 | 2 |
| PRKCD | PKC epsilon | Chromosome 2:45,732,547-46,268,633 | 2 |
| PIGF |  | Chromosome 2:46,661,580-46,697,708 | 2 |
| PIGF |  | Chromosome 2:46,661,580-46,697,708 | 2 |
| SOCS5 |  | Chromosome 2:46,779,595-46,843,431 | 2 |
| FOXN2 |  | Chromosome 2:48,395,374-48,459,938 | 2 |
| FOXN2 |  | Chromosome 2:48,395,374-48,459,938 | 2 |
| LHCGR | LH-R | Chromosome 2:48,767,471-48,836,321 | 2 |
| FSHR | follitropin receptor | Chromosome 2:49,043,156-49,235,134 | 2 |
| FSHR | follitropin receptor | Chromosome 2:49,043,156-49,235,134 | 2 |
| FSHR | follitropin receptor | Chromosome 2:49,043,156-49,235,134 | 2 |
| FSHR | follitropin receptor | Chromosome 2:49,043,156-49,235,134 | 2 |
| VRK2 |  | Chromosome 2:58,127,224-58,240,510 | 2 |
| VRK2 |  | Chromosome 2:58,127,224-58,240,510 | 2 |
| RSAD2 | CIG5 | Chromosome 2:6,935,247-6,955,821 | 2 |
| RSAD2 | CIG5 | Chromosome 2:6,935,247-6,955,821 | 2 |
| RSAD2 | CIG5 | Chromosome 2:6,935,247-6,955,821 | 2 |
| REL | cREL | Chromosome 2:60,962,254-61,003,682 | 2 |
| PELI1 | Pellino1 | Chromosome 2:64,173,499-64,225,062 | 2 |
| PELI1 | Pellino1 | Chromosome 2:64,173,499-64,225,062 | 2 |
| PPP3R1 |  | Chromosome 2:68,203,572-68,341,866 | 2 |
| PPP3R1 |  | Chromosome 2:68,203,572-68,341,866 | 2 |
| PPP3R1 |  | Chromosome 2:68,203,572-68,341,866 | 2 |
| MXD1 | MAD | Chromosome 2:69,995,707-70,023,581 | 2 |
| MXD1 | MAD | Chromosome 2:69,995,707-70,023,581 | 2 |
| MXD1 | MAD | Chromosome 2:69,995,707-70,023,581 | 2 |
| TGFA | TGF alpha | Chromosome 2:70,527,924-70,634,438 | 2 |
| TGFA | TGF alpha | Chromosome 2:70,527,924-70,634,438 | 2 |
| TGFA | TGF alpha | Chromosome 2:70,527,924-70,634,438 | 2 |
| CD207 | CD207, langerin | Chromosome 2:70,910,855-70,916,461 | 2 |
| CYP26B1 |  | Chromosome 2:72,209,875-72,228,471 | 2 |
| DUSP11 |  | Chromosome 2:73,842,837-73,860,756 | 2 |
| DQX1 |  | Chromosome 2:74,598,766-74,606,826 | 2 |
| DQX1 |  | Chromosome 2:74,598,766-74,606,826 | 2 |
| DOK1 |  | Chromosome 2:74,634,795-74,638,181 | 2 |
| SFTPB | SP-B | Chromosome 2:85,737,951-85,748,823 | 2 |
| SFTPB | SP-B | Chromosome 2:85,737,951-85,748,823 | 2 |
| ST3GAL5 | Lactosylceramide | Chromosome 2:85,919,782-85,969,648 | 2 |
| ST3GAL5 | Lactosylceramide | Chromosome 2:85,919,782-85,969,648 | 2 |
| JMJD1A | H3K9me | Chromosome 2:86,521,954-86,573,350 | 2 |
| JMJD1A | H3K9me | Chromosome 2:86,521,954-86,573,350 | 2 |
| JMJD1A | H3K9me | Chromosome 2:86,521,954-86,573,350 | 2 |
| CD8A |  | Chromosome 2:86,865,239-86,871,638 | 2 |
| CD8B1 | CD8B1 | Chromosome 2:86,895,971-86,942,549 | 2 |
| IGKV1-12 |  | Chromosome 2:89,120,836-89,121,310 | 2 |


| IGKV1-12 |  | Chromosome 2:89,120,836-89,121,310 | 2 |
| :---: | :---: | :---: | :---: |
| ITGB1BP1 |  | Chromosome 2:9,463,264-9,481,127 | 2 |
| YWHAQ |  | Chromosome 2:9,641,552-9,688,629 | 2 |
| DUSP2 | PAC-1 | Chromosome 2:96,172,638-96,174,906 | 2 |
| NCAPH | BRRN1 | Chromosome 2:96,365,211-96,405,001 | 2 |
| ZAP70 |  | Chromosome 2:97,696,461-97,722,755 | 2 |
| ZAP70 |  | Chromosome 2:97,696,461-97,722,755 | 2 |
| LYG2 | lysozyme like | Chromosome 2:99,225,141-99,238,002 | 2 |
| LYG2 | lysozyme like | Chromosome 2:99,225,141-99,238,002 | 2 |
| PSMF1 |  | Chromosome 20:1,041,939-1,097,022 | 20 |
| PSMF1 |  | Chromosome 20:1,041,939-1,097,022 | 20 |
| FKBP1A | Calcineurin | Chromosome 20:1,297,622-1,321,806 | 20 |
| SIRPB2 | CD172g | Chromosome 20:1,399,386-1,420,233 | 20 |
| SIRPB1 | CD172b | Chromosome 20:1,491,568-1,548,655 | 20 |
| SIRPB1 | CD172b | Chromosome 20:1,491,568-1,548,655 | 20 |
| SIRPA | SIRPalpha1 | Chromosome 20:1,822,813-1,868,543 | 20 |
| SIRPA | SIRPalpha1 | Chromosome 20:1,822,813-1,868,543 | 20 |
| JAG1 | CD339, Jagged-1 | Chromosome 20:10,566,334-10,602,636 | 20 |
| JAG1 | CD339, Jagged-1 | Chromosome 20:10,566,334-10,602,636 | 20 |
| DFB128 |  | Chromosome 20:116,527-118,264 | 20 |
| DFB129 |  | Chromosome 20:155,899-158,527 | 20 |
| TGM3 |  | Chromosome 20:2,224,647-2,269,725 | 20 |
| TGM3 |  | Chromosome 20:2,224,647-2,269,725 | 20 |
| PTPRA |  | Chromosome 20:2,769,366-2,967,320 | 20 |
| PTPRA |  | Chromosome 20:2,769,366-2,967,320 | 20 |
| PTPRA |  | Chromosome 20:2,769,366-2,967,320 | 20 |
| FOXA2 |  | Chromosome 20:22,509,643-22,514,102 | 20 |
| THBD | CD141, thrombomodulin | Chromosome 20:22,974,270-22,978,301 | 20 |
| THBD | CD141, thrombomodulin | Chromosome 20:22,974,270-22,978,301 | 20 |
| C1QR | CD93 | Chromosome 20:23,007,995-23,014,977 | 20 |
| CST1 | SN | Chromosome 20:23,676,190-23,679,574 | 20 |
| CST1 | SN | Chromosome 20:23,676,190-23,679,574 | 20 |
| DFB115 |  | Chromosome 20:29,309,128-29,311,096 | 20 |
| DFB122 |  | Chromosome 20:29,466,712-29,480,644 | 20 |
| HM13 | H13 | Chromosome 20:29,565,892-29,621,031 | 20 |
| BCL2L1 | BCL-XL, Bcl-x (L) | Chromosome 20:29,715,916-29,774,366 | 20 |
| DUSP15 |  | Chromosome 20:29,899,102-29,922,211 | 20 |
| PDRG1 | p53 | Chromosome 20:29,996,419-30,003,556 | 20 |
| SIGLEC1 | CD169 | Chromosome 20:3,615,617-3,635,775 | 20 |
| SIGLEC1 | CD169 | Chromosome 20:3,615,617-3,635,775 | 20 |
| SIGLEC1 | CD169 | Chromosome 20:3,615,617-3,635,775 | 20 |
| CDC25B |  | Chromosome 20:3,724,386-3,734,762 | 20 |
| HCK |  | Chromosome 20:30,103,715-30,153,320 | 20 |
| HCK |  | Chromosome 20:30,103,715-30,153,320 | 20 |
| CBFA2T2 |  | Chromosome 20:31,541,589-31,701,503 | 20 |
| CBFA2T2 |  | Chromosome 20:31,541,589-31,701,503 | 20 |
| CBFA2T2 |  | Chromosome 20:31,541,589-31,701,503 | 20 |
| CHMP4B | CHMP4B | Chromosome 20:31,862,780-31,905,831 | 20 |
| ITCH |  | Chromosome 20:32,414,702-32,562,859 | 20 |
| ITCH |  | Chromosome 20:32,414,702-32,562,859 | 20 |
| PROCR | EPCR, CD201 | Chromosome 20:33,212,131-33,228,828 | 20 |
| PROCR | EPCR, CD201 | Chromosome 20:33,212,131-33,228,828 | 20 |
| PROCR | EPCR, CD201 | Chromosome 20:33,212,131-33,228,828 | 20 |
| MMP24 |  | Chromosome 20:33,278,095-33,328,218 | 20 |
| SAMHD1 | Mg11 | Chromosome 20:34,953,761-35,013,590 | 20 |
| SRC |  | Chromosome 20:35,406,502-35,467,239 | 20 |


| SRC |  | Chromosome 20:35,406,502-35,467,239 | 20 |
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| BPI |  | Chromosome 20:36,365,991-36,399,321 | 20 |
| LBP |  | Chromosome 20:36,408,299-36,439,067 | 20 |
| PPP1R16B |  | Chromosome 20:36,867,762-36,985,081 | 20 |
| PPP1R16B |  | Chromosome 20:36,867,762-36,985,081 | 20 |
| DHX35 |  | Chromosome 20:37,024,409-37,101,778 | 20 |
| PLCG1 |  | Chromosome 20:39,199,291-39,237,775 | 20 |
| PRNP | CD230 | Chromosome 20:4,614,996-4,630,236 | 20 |
| PRNP | CD230 | Chromosome 20:4,614,996-4,630,236 | 20 |
| PTPRT |  | Chromosome 20:40,134,806-41,252,024 | 20 |
| PTPRT |  | Chromosome 20:40,134,806-41,252,024 | 20 |
| PTPRT |  | Chromosome 20:40,134,806-41,252,024 | 20 |
| PTPRT |  | Chromosome 20:40,134,806-41,252,024 | 20 |
| PTPRT |  | Chromosome 20:40,134,806-41,252,024 | 20 |
| PTPRT |  | Chromosome 20:40,134,806-41,252,024 | 20 |
| PTPRT |  | Chromosome 20:40,134,806-41,252,024 | 20 |
| PTPRT |  | Chromosome 20:40,134,806-41,252,024 | 20 |
| PTPRT |  | Chromosome 20:40,134,806-41,252,024 | 20 |
| PTPRT |  | Chromosome 20:40,134,806-41,252,024 | 20 |
| PTPRT |  | Chromosome 20:40,134,806-41,252,024 | 20 |
| PTPRT |  | Chromosome 20:40,134,806-41,252,024 | 20 |
| PPIAL |  | Chromosome 20:41,292,801-41,293,515 | 20 |
| HNF4A | HNF4 alpha 1,2,3,4, NR2, | Chromosome 20:42,417,855-42,493,444 | 20 |
| HNF4A | HNF4 alpha 1,2,3,4, NR2, | Chromosome 20:42,417,855-42,493,444 | 20 |
| ADA | adenosine deaminase | Chromosome 20:42,681,577-42,713,797 | 20 |
| ADA | adenosine deaminase | Chromosome 20:42,681,577-42,713,797 | 20 |
| STK4 | MST1/STK4 | Chromosome 20:43,028,529-43,142,014 | 20 |
| STK4 | MST1/STK4 | Chromosome 20:43,028,529-43,142,014 | 20 |
| STK4 | MST1/STK4 | Chromosome 20:43,028,529-43,142,014 | 20 |
| WFDC12 |  | Chromosome 20:43,185,480-43,186,520 | 20 |
| SEMG1 |  | Chromosome 20:43,269,052-43,271,827 | 20 |
| SDC4 | Syndecan | Chromosome 20:43,387,342-43,410,478 | 20 |
| SDC4 | Syndecan | Chromosome 20:43,387,342-43,410,478 | 20 |
| ACOT8 |  | Chromosome 20:43,903,768-43,919,442 | 20 |
| CTSA | Cathepsin A | Chromosome 20:43,952,190-43,960,866 | 20 |
| MMP9 |  | Chromosome 20:44,070,954-44,078,607 | 20 |
| MMP9 |  | Chromosome 20:44,070,954-44,078,607 | 20 |
| TNFRSF5 | CD40, 41 | Chromosome 20:44,180,313-44,366,257 | 20 |
| TNFRSF5 | CD40, 41 | Chromosome 20:44,180,313-44,366,257 | 20 |
| TNFRSF5 | CD40, 41 | Chromosome 20:44,180,313-44,366,257 | 20 |
| PTGIS |  | Chromosome 20:47,553,818-47,618,114 | 20 |
| PTGIS |  | Chromosome 20:47,553,818-47,618,114 | 20 |
| PTGIS |  | Chromosome 20:47,553,818-47,618,114 | 20 |
| SLC9A8 | NHE-8 | Chromosome 20:47,862,657-47,942,179 | 20 |
| SLC9A8 | NHE-8 | Chromosome 20:47,862,657-47,942,179 | 20 |
| UBE2V1 | UEV1A | Chromosome 20:48,131,068-48,203,678 | 20 |
| UBE2V1 | UEV1A | Chromosome 20:48,131,068-48,203,678 | 20 |
| PTPN1 |  | Chromosome 20:48,560,294-48,634,706 | 20 |
| PTPN1 |  | Chromosome 20:48,560,294-48,634,706 | 20 |
| PTPN1 |  | Chromosome 20:48,560,294-48,634,706 | 20 |
| NFATC2 |  | Chromosome 20:49,441,083-49,592,665 | 20 |
| NFATC2 |  | Chromosome 20:49,441,083-49,592,665 | 20 |
| NFATC2 |  | Chromosome 20:49,441,083-49,592,665 | 20 |
| MC3R | melanocortin receptor 3 | Chromosome 20:54,257,195-54,258,278 | 20 |
| MC3R | melanocortin receptor 3 | Chromosome 20:54,257,195-54,258,278 | 20 |
| TFAP2C |  | Chromosome 20:54,637,765-54,647,746 | 20 |


| BMP7 | BMP7 | Chromosome 20:55,177,211-55,275,091 | 20 |
| :---: | :---: | :---: | :---: |
| BMP7 | BMP7 | Chromosome 20:55,177,211-55,275,091 | 20 |
| BMP7 | BMP7 | Chromosome 20:55,177,211-55,275,091 | 20 |
| RAE1 |  | Chromosome 20:55,359,535-55,387,618 | 20 |
| CTSZ | Cathepsin Z | Chromosome 20:56,990,597-57,015,697 | 20 |
| CTSZ | Cathepsin Z | Chromosome 20:56,990,597-57,015,697 | 20 |
| EDN3 | endothelin 3 | Chromosome 20:57,308,877-57,334,442 | 20 |
| EDN3 | endothelin 3 | Chromosome 20:57,308,877-57,334,442 | 20 |
| BMP2 | BMP2 | Chromosome 20:6,696,311-6,708,927 | 20 |
| BMP2 | BMP2 | Chromosome 20:6,696,311-6,708,927 | 20 |
| HRH3 |  | Chromosome 20:60,223,421-60,228,718 | 20 |
| LAMA5 |  | Chromosome 20:60,317,510-60,375,763 | 20 |
| GATA5 |  | Chromosome 20:60,471,948-60,484,421 | 20 |
| DIDO1 | DATF1, Dio1 | Chromosome 20:60,979,535-61,039,743 | 20 |
| BIRC7 | LIVIN | Chromosome 20:61,337,680-61,342,299 | 20 |
| EEF1A2 | EF-1 alpha | Chromosome 20:61,589,810-61,600,949 | 20 |
| TNFRSF6B | DcR3 | Chromosome 20:61,759,607-61,800,495 | 20 |
| TNFRSF6B | DcR3 | Chromosome 20:61,759,607-61,800,495 | 20 |
| OPRL1 | ORL1 | Chromosome 20:62,181,932-62,215,047 | 20 |
| BAGE4 |  | Chromosome 21:10,042,713-10,120,798 | 21 |
| BAGE4 |  | Chromosome 21:10,042,713-10,120,798 | 21 |
| BTG3 |  | Chromosome 21:17,887,842-17,907,136 | 21 |
| BTG3 |  | Chromosome 21:17,887,842-17,907,136 | 21 |
| JAM2 | CD322 | Chromosome 21:25,933,460-26,009,106 | 21 |
| JAM2 | CD322 | Chromosome 21:25,933,460-26,009,106 | 21 |
| GRIK1 |  | Chromosome 21:29,831,125-30,234,153 | 21 |
| GRIK1 |  | Chromosome 21:29,831,125-30,234,153 | 21 |
| GRIK1 |  | Chromosome 21:29,831,125-30,234,153 | 21 |
| GRIK1 |  | Chromosome 21:29,831,125-30,234,153 | 21 |
| GRIK1 |  | Chromosome 21:29,831,125-30,234,153 | 21 |
| GRIK1 |  | Chromosome 21:29,831,125-30,234,153 | 21 |
| SOD1 | Superoxide dismutase, IP | Chromosome 21:31,953,806-31,963,115 | 21 |
| SOD1 | Superoxide dismutase, IP | Chromosome 21:31,953,806-31,963,115 | 21 |
| OLIG2 | RACK17 | Chromosome 21:33,320,023-33,323,374 | 21 |
| IFNAR2 |  | Chromosome 21:33,524,076-33,559,839 | 21 |
| IFNAR2 |  | Chromosome 21:33,524,076-33,559,839 | 21 |
| IFNAR1 |  | Chromosome 21:33,619,079-33,654,038 | 21 |
| IFNGR2 |  | Chromosome 21:33,697,072-33,731,698 | 21 |
| RUNX1 | RUNX | Chromosome 21:35,081,968-35,343,511 | 21 |
| RUNX1 | RUNX | Chromosome 21:35,081,968-35,343,511 | 21 |
| RUNX1 | RUNX | Chromosome 21:35,081,968-35,343,511 | 21 |
| RUNX1 | RUNX | Chromosome 21:35,081,968-35,343,511 | 21 |
| RUNX1 | RUNX | Chromosome 21:35,081,968-35,343,511 | 21 |
| RUNX1 | RUNX | Chromosome 21:35,081,968-35,343,511 | 21 |
| CHAF1B |  | Chromosome 21:36,679,559-36,710,995 | 21 |
| CHAF1B |  | Chromosome 21:36,679,559-36,710,995 | 21 |
| DYRK1A |  | Chromosome 21:37,661,729-37,809,347 | 21 |
| DYRK1A |  | Chromosome 21:37,661,729-37,809,347 | 21 |
| DYRK1A |  | Chromosome 21:37,661,729-37,809,347 | 21 |
| ERG |  | Chromosome 21:38,675,671-38,955,488 | 21 |
| ERG |  | Chromosome 21:38,675,671-38,955,488 | 21 |
| ERG |  | Chromosome 21:38,675,671-38,955,488 | 21 |
| ERG |  | Chromosome 21:38,675,671-38,955,488 | 21 |
| IGSF5 |  | Chromosome 21:40,039,204-40,095,893 | 21 |
| IGSF5 |  | Chromosome 21:40,039,204-40,095,893 | 21 |
| DSCAM |  | Chromosome 21:40,306,213-41,140,909 | 21 |


| DSCAM |  | Chromosome 21:40,306,213-41,140,909 | 21 |
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| DSCAM |  | Chromosome 21:40,306,213-41,140,909 | 21 |
| DSCAM |  | Chromosome 21:40,306,213-41,140,909 | 21 |
| DSCAM |  | Chromosome 21:40,306,213-41,140,909 | 21 |
| DSCAM |  | Chromosome 21:40,306,213-41,140,909 | 21 |
| DSCAM |  | Chromosome 21:40,306,213-41,140,909 | 21 |
| DSCAM |  | Chromosome 21:40,306,213-41,140,909 | 21 |
| DSCAM |  | Chromosome 21:40,306,213-41,140,909 | 21 |
| DSCAM |  | Chromosome 21:40,306,213-41,140,909 | 21 |
| MX2 |  | Chromosome 21:41,655,820-41,702,739 | 21 |
| MX2 |  | Chromosome 21:41,655,820-41,702,739 | 21 |
| MX1 |  | Chromosome 21:41,720,024-41,753,008 | 21 |
| TFF3 | Tff3 | Chromosome 21:42,599,751-42,608,775 | 21 |
| PDXK |  | Chromosome 21:43,963,406-44,006,608 | 21 |
| PDXK |  | Chromosome 21:43,963,406-44,006,608 | 21 |
| PDXK |  | Chromosome 21:43,963,406-44,006,608 | 21 |
| ICOSLG | CD275 | Chromosome 21:44,467,313-44,485,262 | 21 |
| AIRE |  | Chromosome 21:44,530,191-44,542,530 | 21 |
| ITGB2 | CD18, Mac-1, CR3 | Chromosome 21:45,130,296-45,173,181 | 21 |
| SLC19A1 | RFC | Chromosome 21:45,737,914-45,786,779 | 21 |
| COL6A1 |  | Chromosome 21:46,226,091-46,249,391 | 21 |
| COL6A1 |  | Chromosome 21:46,226,091-46,249,391 | 21 |
| COL6A2 |  | Chromosome 21:46,342,470-46,377,190 | 21 |
| PRMT2 | PRMT2 | Chromosome 21:46,879,507-46,909,291 | 21 |
| IL17RA | CDw217, IL17R | Chromosome 22:15,945,849-15,971,405 | 22 |
| BCL2L13 |  | Chromosome 22:16,501,485-16,593,383 | 22 |
| BID |  | Chromosome 22:16,591,460-16,631,812 | 22 |
| BID |  | Chromosome 22:16,591,460-16,631,812 | 22 |
| GGT2 |  | Chromosome 22:17,141,172-17,159,474 | 22 |
| DGCR2 |  | Chromosome 22:17,403,798-17,489,962 | 22 |
| DGCR2 |  | Chromosome 22:17,403,798-17,489,962 | 22 |
| GP1BB | CD42c, HPA3 | Chromosome 22:18,091,066-18,092,297 | 22 |
| GP1BB | CD42c, HPA3 | Chromosome 22:18,091,066-18,092,297 | 22 |
| SCARF2 |  | Chromosome 22:19,108,875-19,122,146 | 22 |
| CRKL | crkl | Chromosome 22:19,601,714-19,638,034 | 22 |
| CRKL | crkl | Chromosome 22:19,601,714-19,638,034 | 22 |
| P2RX6 | P2X6 | Chromosome 22:19,699,449-19,713,119 | 22 |
| SDF2L1 | stromal cell derived factor | Chromosome 22:20,326,542-20,328,588 | 22 |
| MAPK1 | p42 MAPK, Erk | Chromosome 22:20,443,946-20,551,970 | 22 |
| VPREB1 | CD179a | Chromosome 22:20,929,200-20,929,926 | 22 |
| PRAME |  | Chromosome 22:21,220,123-21,231,768 | 22 |
| BCR | B cell receptor | Chromosome 22:21,852,552-21,990,224 | 22 |
| BCR | B cell receptor | Chromosome 22:21,852,552-21,990,224 | 22 |
| IGLL1 | Ig kappa light chain, CD1 | Chromosome 22:22,245,312-22,252,495 | 22 |
| IGLL1 | Ig kappa light chain, CD1 | Chromosome 22:22,245,312-22,252,495 | 22 |
| MIF |  | Chromosome 22:22,369,647-22,567,417 | 22 |
| MIF |  | Chromosome 22:22,369,647-22,567,417 | 22 |
| GSST1 |  | Chromosome 22:22,706,141-22,714,271 | 22 |
| CABIN1 |  | Chromosome 22:22,737,765-22,904,596 | 22 |
| CABIN1 |  | Chromosome 22:22,737,765-22,904,596 | 22 |
| ADORA2 | Adenosin receptor 2 | Chromosome 22:23,153,537-23,168,325 | 22 |
| GGT1 | CD224 | Chromosome 22:23,309,718-23,354,972 | 22 |
| MN1 |  | Chromosome 22:26,474,265-26,527,486 | 22 |
| MN1 |  | Chromosome 22:26,474,265-26,527,486 | 22 |
| XBP1 | XBP1 | Chromosome 22:27,520,548-27,526,560 | 22 |
| LIF |  | Chromosome 22:28,966,441-28,972,748 | 22 |


| LIF |  | Chromosome 22:28,966,441-28,972,748 | 22 |
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| TCN2 | transcobalamin 2 | Chromosome 22:29,333,161-29,353,047 | 22 |
| DUSP18 |  | Chromosome 22:29,388,039-29,393,872 | 22 |
| DRG1 | DRG1 | Chromosome 22:30,125,539-30,160,438 | 22 |
| DRG1 | DRG1 | Chromosome 22:30,125,539-30,160,438 | 22 |
| TIMP3 |  | Chromosome 22:31,526,802-31,589,028 | 22 |
| TIMP3 |  | Chromosome 22:31,526,802-31,589,028 | 22 |
| HMOX1 | Heme oxygenease 1 | Chromosome 22:34,107,057-34,120,194 | 22 |
| APOL6 |  | Chromosome 22:34,374,370-34,394,402 | 22 |
| APOL6 |  | Chromosome 22:34,374,370-34,394,402 | 22 |
| MYH9 |  | Chromosome 22:35,007,272-35,113,958 | 22 |
| CACNG2 |  | Chromosome 22:35,290,050-35,428,849 | 22 |
| NCF3 | neutrophil cytosolic factor | Chromosome 22:35,586,976-35,604,005 | 22 |
| NCF3 | neutrophil cytosolic factor | Chromosome 22:35,586,976-35,604,005 | 22 |
| IL2RB | CD122 | Chromosome 22:35,851,824-35,875,908 | 22 |
| C1QTNF6 |  | Chromosome 22:35,906,152-35,914,276 | 22 |
| C1QTNF6 |  | Chromosome 22:35,906,152-35,914,276 | 22 |
| RAC2 | Rac2 | Chromosome 22:35,951,238-35,970,251 | 22 |
| CARD10 | CARD10 | Chromosome 22:36,216,346-36,245,193 | 22 |
| CARD10 | CARD10 | Chromosome 22:36,216,346-36,245,193 | 22 |
| PLA2G6 | Phospholipase | Chromosome 22:36,837,448-36,907,763 | 22 |
| PLA2G6 | Phospholipase | Chromosome 22:36,837,448-36,907,763 | 22 |
| DDX17 |  | Chromosome 22:37,209,389-37,232,262 | 22 |
| DDX17 |  | Chromosome 22:37,209,389-37,232,262 | 22 |
| UNC84B |  | Chromosome 22:37,460,681-37,481,928 | 22 |
| UNC84B |  | Chromosome 22:37,460,681-37,481,928 | 22 |
| PDGFB |  | Chromosome 22:37,949,310-37,971,006 | 22 |
| MAP3K7IP1 | TAB1 | Chromosome 22:38,125,692-38,163,078 | 22 |
| ATF4 |  | Chromosome 22:38,246,515-38,248,637 | 22 |
| GRAP2 | Gads | Chromosome 22:38,627,032-38,698,204 | 22 |
| GRAP2 | Gads | Chromosome 22:38,627,032-38,698,204 | 22 |
| EP300 |  | Chromosome 22:39,817,736-39,906,024 | 22 |
| CENPM | PANE1 | Chromosome 22:40,664,687-40,673,094 | 22 |
| CENPM | PANE1 | Chromosome 22:40,664,687-40,673,094 | 22 |
| CYP2D6 |  | Chromosome 22:40,852,445-40,856,827 | 22 |
| TCF20 |  | Chromosome 22:40,885,963-40,941,389 | 22 |
| NFAM1 |  | Chromosome 22:41,108,917-41,158,340 | 22 |
| NFAM1 |  | Chromosome 22:41,108,917-41,158,340 | 22 |
| POLDIP3 | S6K1 | Chromosome 22:41,309,671-41,340,906 | 22 |
| A4GALT | CD77, Lactosylceramide | Chromosome 22:41,418,071-41,446,820 | 22 |
| BIK |  | Chromosome 22:41,836,701-41,855,662 | 22 |
| BIK |  | Chromosome 22:41,836,701-41,855,662 | 22 |
| PPARA | PPARalpha, NR1C1 | Chromosome 22:44,925,163-45,018,317 | 22 |
| DIP | DIP | Chromosome 22:45,394,963-45,454,352 | 22 |
| DIP | DIP | Chromosome 22:45,394,963-45,454,352 | 22 |
| CERK | ceramide kinase | Chromosome 22:45,458,971-45,512,816 | 22 |
| ECGF1 |  | Chromosome 22:49,311,047-49,315,321 | 22 |
| MAPK8IP2 |  | Chromosome 22:49,385,997-49,396,843 | 22 |
| SHANK3 |  | Chromosome 22:49,459,936-49,518,507 | 22 |
| IRAK2 |  | Chromosome 3:10,181,563-10,260,427 | 3 |
| NFKBIZ |  | Chromosome 3:103,029,547-103,062,556 | 3 |
| NFKBIZ |  | Chromosome 3:103,029,547-103,062,556 | 3 |
| ALCAM | CD166 | Chromosome 3:106,568,403-106,778,434 | 3 |
| ALCAM | CD166 | Chromosome 3:106,568,403-106,778,434 | 3 |
| ALCAM | CD166 | Chromosome 3:106,568,403-106,778,434 | 3 |
| CBLB |  | Chromosome 3:106,859,799-107,070,577 | 3 |


| CBLB |  | Chromosome 3:106,859,799-107,070,577 | 3 |
| :---: | :---: | :---: | :---: |
| CBLB |  | Chromosome 3:106,859,799-107,070,577 | 3 |
| CD47 | IAP | Chromosome 3:109,244,631-109,292,625 | 3 |
| CD47 | IAP | Chromosome 3:109,244,631-109,292,625 | 3 |
| CD47 | IAP | Chromosome 3:109,244,631-109,292,625 | 3 |
| HRH1 |  | Chromosome 3:11,269,400-11,279,415 | 3 |
| HRH1 |  | Chromosome 3:11,269,400-11,279,415 | 3 |
| TRAT1 | TRIM | Chromosome 3:110,024,321-110,056,542 | 3 |
| PVRL3 | CDw113 | Chromosome 3:112,273,555-112,395,063 | 3 |
| PVRL3 | CDw113 | Chromosome 3:112,273,555-112,395,063 | 3 |
| CD96 |  | Chromosome 3:112,743,546-112,853,906 | 3 |
| CD96 |  | Chromosome 3:112,743,546-112,853,906 | 3 |
| CD96 |  | Chromosome 3:112,743,546-112,853,906 | 3 |
| CD200 |  | Chromosome 3:113,522,943-113,564,349 | 3 |
| CD200 |  | Chromosome 3:113,522,943-113,564,349 | 3 |
| BTLA | CD272=BTLA | Chromosome 3:113,667,463-113,701,066 | 3 |
| CD200R2 |  | Chromosome 3:114,017,246-114,047,487 | 3 |
| CD200R2 |  | Chromosome 3:114,017,246-114,047,487 | 3 |
| CD200R1 |  | Chromosome 3:114,122,746-114,176,650 | 3 |
| TIMP4 |  | Chromosome 3:12,169,568-12,175,851 | 3 |
| TIMP4 |  | Chromosome 3:12,169,568-12,175,851 | 3 |
| TIMP4 |  | Chromosome 3:12,169,568-12,175,851 | 3 |
| PPARG | PPARgamma, NR1C3 | Chromosome 3:12,304,359-12,450,843 | 3 |
| PPARG | PPARgamma, NR1C3 | Chromosome 3:12,304,359-12,450,843 | 3 |
| PPARG | PPARgamma, NR1C3 | Chromosome 3:12,304,359-12,450,843 | 3 |
| IGSF11 |  | Chromosome 3:120,102,167-120,347,588 | 3 |
| IGSF11 |  | Chromosome 3:120,102,167-120,347,588 | 3 |
| IGSF11 |  | Chromosome 3:120,102,167-120,347,588 | 3 |
| CD80 | B7-1 | Chromosome 3:120,725,832-120,761,139 | 3 |
| CD80 | B7-1 | Chromosome 3:120,725,832-120,761,139 | 3 |
| PLA1A | Phospholipase | Chromosome 3:120,792,984-120,831,342 | 3 |
| NR112 | NR112 | Chromosome 3:120,982,021-121,020,022 | 3 |
| GSK3B |  | Chromosome 3:121,028,233-121,295,954 | 3 |
| GTF2E1 | TFIIE | Chromosome 3:121,937,926-121,984,605 | 3 |
| CD86 | B7-2 | Chromosome 3:123,256,911-123,322,673 | 3 |
| CD86 | B7-2 | Chromosome 3:123,256,911-123,322,673 | 3 |
| CD86 | B7-2 | Chromosome 3:123,256,911-123,322,673 | 3 |
| MYLK | myosin light chain kinase | Chromosome 3:124,813,833-125,085,839 | 3 |
| MYLK | myosin light chain kinase | Chromosome 3:124,813,833-125,085,839 | 3 |
| ITGB5 |  | Chromosome 3:125,964,485-126,088,842 | 3 |
| ITGB5 |  | Chromosome 3:125,964,485-126,088,842 | 3 |
| ITGB5 |  | Chromosome 3:125,964,485-126,088,842 | 3 |
| PLXNA1 | Plexin-A1 | Chromosome 3:128,190,192-128,238,922 | 3 |
| PLXNA1 | Plexin-A1 | Chromosome 3:128,190,192-128,238,922 | 3 |
| MCM2 |  | Chromosome 3:128,799,943-128,823,969 | 3 |
| MCM2 |  | Chromosome 3:128,799,943-128,823,969 | 3 |
| SEC61A1 |  | Chromosome 3:129,253,902-129,273,216 | 3 |
| SEC61A1 |  | Chromosome 3:129,253,902-129,273,216 | 3 |
| GATA2 |  | Chromosome 3:129,680,960-129,694,718 | 3 |
| GP9 | CD42a | Chromosome 3:130,262,300-130,263,941 | 3 |
| PLXND1 | Plexin-D1 | Chromosome 3:130,756,708-130,808,351 | 3 |
| PLXND1 | Plexin-D1 | Chromosome 3:130,756,708-130,808,351 | 3 |
| TRH | TRH | Chromosome 3:131,176,253-131,179,470 | 3 |
| TRH | TRH | Chromosome 3:131,176,253-131,179,470 | 3 |
| TRH | TRH | Chromosome 3:131,176,253-131,179,470 | 3 |
| PIK3R4 |  | Chromosome 3:131,880,468-131,948,340) | 3 |


| PIK3R4 |  | Chromosome 3:131,880,468-131,948,340) | 3 |
| :---: | :---: | :---: | :---: |
| CCRL1 |  | Chromosome 3:133,798,784-133,804,072 | 3 |
| EPHB1 |  | Chromosome 3:135,996,950-136,461,999 | 3 |
| EPHB1 |  | Chromosome 3:135,996,950-136,461,999 | 3 |
| EPHB1 |  | Chromosome 3:135,996,950-136,461,999 | 3 |
| EPHB1 |  | Chromosome 3:135,996,950-136,461,999 | 3 |
| EPHB1 |  | Chromosome 3:135,996,950-136,461,999 | 3 |
| EPHB1 |  | Chromosome 3:135,996,950-136,461,999 | 3 |
| EPHB1 |  | Chromosome 3:135,996,950-136,461,999 | 3 |
| IL20RB |  | Chromosome 3:138,159,397-138,212,610 | 3 |
| IL20RB |  | Chromosome 3:138,159,397-138,212,610 | 3 |
| PIK3CB |  | Chromosome 3:139,856,921-139,960,875 | 3 |
| PIK3CB |  | Chromosome 3:139,856,921-139,960,875 | 3 |
| XPC |  | Chromosome 3:14,161,648-14,195,143 | 3 |
| NR2C2 | NR2C2 | Chromosome 3:14,964,240-15,065,784 | 3 |
| NR2C2 | NR2C2 | Chromosome 3:14,964,240-15,065,784 | 3 |
| FOXL2 |  | Chromosome 3:140,145,756-140,148,491 | 3 |
| FOXL2 |  | Chromosome 3:140,145,756-140,148,491 | 3 |
| RNF7 |  | Chromosome 3:142,939,741-142,947,933 | 3 |
| RNF7 |  | Chromosome 3:142,939,741-142,947,933 | 3 |
| ATP1B3 | CD298 | Chromosome 3:143,078,160-143,128,072 | 3 |
| SLC9A9 | NHE-9 | Chromosome 3:144,466,754-145,049,979 | 3 |
| SLC9A9 | NHE-9 | Chromosome 3:144,466,754-145,049,979 | 3 |
| SLC9A9 | NHE-9 | Chromosome 3:144,466,754-145,049,979 | 3 |
| SLC9A9 | NHE-9 | Chromosome 3:144,466,754-145,049,979 | 3 |
| SLC9A9 | NHE-9 | Chromosome 3:144,466,754-145,049,979 | 3 |
| SLC9A9 | NHE-9 | Chromosome 3:144,466,754-145,049,979 | 3 |
| SLC9A9 | NHE-9 | Chromosome 3:144,466,754-145,049,979 | 3 |
| SLC9A9 | NHE-9 | Chromosome 3:144,466,754-145,049,979 | 3 |
| SLC9A9 | NHE-9 | Chromosome 3:144,466,754-145,049,979 | 3 |
| AGTR1 | Angiotensin receptor | Chromosome 3:149,898,355-149,943,478 | 3 |
| AGTR1 | Angiotensin receptor | Chromosome 3:149,898,355-149,943,478 | 3 |
| AGTR1 | Angiotensin receptor | Chromosome 3:149,898,355-149,943,478 | 3 |
| HLTF |  | Chromosome 3:150,230,604-150,287,007 | 3 |
| HLTF |  | Chromosome 3:150,230,604-150,287,007 | 3 |
| HPS3 | Hermansky-Pudlak syndr | Chromosome 3:150,330,061-150,373,995 | 3 |
| P2RY14 | P2Y14 | Chromosome 3:152,412,595-152,478,847 | 3 |
| P2RY14 | P2Y14 | Chromosome 3:152,412,595-152,478,847 | 3 |
| IGSF10 |  | Chromosome 3:152,637,167-152,659,187 | 3 |
| P2RY1 | P2Y1 | Chromosome 3:154,035,426-154,038,535 | 3 |
| P2RY1 | P2Y1 | Chromosome 3:154,035,426-154,038,535 | 3 |
| DHX36 |  | Chromosome 3:155,476,152-155,524,971 | 3 |
| DHX36 |  | Chromosome 3:155,476,152-155,524,971 | 3 |
| MME | CD10 | Chromosome 3:156,280,153-156,384,186 | 3 |
| MME | CD10 | Chromosome 3:156,280,153-156,384,186 | 3 |
| PTX3 |  | Chromosome 3:158,637,301-158,644,071 | 3 |
| PLCL2 |  | Chromosome 3:16,949,586-17,107,089 | 3 |
| PLCL2 |  | Chromosome 3:16,949,586-17,107,089 | 3 |
| IL12A |  | Chromosome 3:161,189,323-161,196,500 | 3 |
| IL12A |  | Chromosome 3:161,189,323-161,196,500 | 3 |
| IL12A |  | Chromosome 3:161,189,323-161,196,500 | 3 |
| TRIM59 | MRF1 | Chromosome 3:161,427,938-161,650,320 | 3 |
| TRIM59 | MRF1 | Chromosome 3:161,427,938-161,650,320 | 3 |
| TRIM59 | MRF1 | Chromosome 3:161,427,938-161,650,320 | 3 |
| B3GALNT1 |  | Chromosome 3:162,284,365-162,305,854 | 3 |
| PDCD10 |  | Chromosome 3:168,884,388-168,935,345 | 3 |


| PDCD10 |  | Chromosome 3:168,884,388-168,935,345 | 3 |
| :---: | :---: | :---: | :---: |
| EVI1 |  | Chromosome 3:170,285,244-170,347,054 | 3 |
| EVI1 |  | Chromosome 3:170,285,244-170,347,054 | 3 |
| TERC |  | Chromosome 3:170,965,092-170,965,542 | 3 |
| TERC |  | Chromosome 3:170,965,092-170,965,542 | 3 |
| PRKCI | PKCӨ | Chromosome 3:171,422,919-171,506,459 | 3 |
| PRKCI | PKCO | Chromosome 3:171,422,919-171,506,459 | 3 |
| TNFSF10 | CD253, TRAIL | Chromosome 3:173,706,158-173,723,963 | 3 |
| TNFSF10 | CD253, TRAIL | Chromosome 3:173,706,158-173,723,963 | 3 |
| TBL1XR1 | TBLR1 | Chromosome 3:178,221,867-178,397,734 | 3 |
| TBL1XR1 | TBLR1 | Chromosome 3:178,221,867-178,397,734 | 3 |
| TBL1XR1 | TBLR1 | Chromosome 3:178,221,867-178,397,734 | 3 |
| PIK3CA |  | Chromosome 3:180,349,005-180,435,194 | 3 |
| PIK3CA |  | Chromosome 3:180,349,005-180,435,194 | 3 |
| LAMP3 | CD208, DC-LAMP | Chromosome 3:184,322,697-184,363,317 | 3 |
| LAMP3 | CD208, DC-LAMP | Chromosome 3:184,322,697-184,363,317 | 3 |
| THPO | Thrombopoietin | Chromosome 3:185,572,467-185,578,626 | 3 |
| THPO | Thrombopoietin | Chromosome 3:185,572,467-185,578,626 | 3 |
| MAP3K13 |  | Chromosome 3:186,563,664-186,683,322 | 3 |
| KNG1 | Bradykinin | Chromosome 3:187,917,814-187,944,437 | 3 |
| KNG1 | Bradykinin | Chromosome 3:187,917,814-187,944,437 | 3 |
| ST6GAL1 | CD75s | Chromosome 3:188,131,210-188,279,035 | 3 |
| ST6GAL1 | CD75s | Chromosome 3:188,131,210-188,279,035 | 3 |
| ST6GAL1 | CD75s | Chromosome 3:188,131,210-188,279,035 | 3 |
| ST6GAL1 | CD75s | Chromosome 3:188,131,210-188,279,035 | 3 |
| MASP1 | MASP3 | Chromosome 3:188,418,632-188,492,446 | 3 |
| IFRG28 |  | Chromosome 3:188,568,862-188,572,066 | 3 |
| IFRG28 |  | Chromosome 3:188,568,862-188,572,066 | 3 |
| SST | SST | Chromosome 3:188,869,388-188,870,895 | 3 |
| SST | SST | Chromosome 3:188,869,388-188,870,895 | 3 |
| SST | SST | Chromosome 3:188,869,388-188,870,895 | 3 |
| TP73L | p63 | Chromosome 3:190,831,910-191,097,759 | 3 |
| TP73L | p63 | Chromosome 3:190,831,910-191,097,759 | 3 |
| TP73L | p63 | Chromosome 3:190,831,910-191,097,759 | 3 |
| IL1RAP |  | Chromosome 3:191,714,585-191,858,537 | 3 |
| IL1RAP |  | Chromosome 3:191,714,585-191,858,537 | 3 |
| IL1RAP |  | Chromosome 3:191,714,585-191,858,537 | 3 |
| FGF12 | FGF12 | Chromosome 3:193,342,413-193,928,066 | 3 |
| FGF12 | FGF12 | Chromosome 3:193,342,413-193,928,066 | 3 |
| FGF12 | FGF12 | Chromosome 3:193,342,413-193,928,066 | 3 |
| FGF12 | FGF12 | Chromosome 3:193,342,413-193,928,066 | 3 |
| FGF12 | FGF12 | Chromosome 3:193,342,413-193,928,066 | 3 |
| FGF12 | FGF12 | Chromosome 3:193,342,413-193,928,066 | 3 |
| FGF12 | FGF12 | Chromosome 3:193,342,413-193,928,066 | 3 |
| FGF12 | FGF12 | Chromosome 3:193,342,413-193,928,066 | 3 |
| FGF12 | FGF12 | Chromosome 3:193,342,413-193,928,066 | 3 |
| FGF12 | FGF12 | Chromosome 3:193,342,413-193,928,066 | 3 |
| FGF12 | FGF12 | Chromosome 3:193,342,413-193,928,066 | 3 |
| GP5 | CD42d, HPA4 | Chromosome 3:195,595,348-195,601,523 | 3 |
| TFRC | transferrin receptor | Chromosome 3:197,260,553-197,293,343 | 3 |
| PAK2 |  | Chromosome 3:197,951,312-198,043,756 | 3 |
| PAK2 |  | Chromosome 3:197,951,312-198,043,756 | 3 |
| MFI2 | CD228 | Chromosome 3:198,214,553-198,241,043 | 3 |
| MFI2 | CD228 | Chromosome 3:198,214,553-198,241,043 | 3 |
| CHLI |  | Chromosome 3:213,650-426,098 | 3 |
| CHLI |  | Chromosome 3:213,650-426,098 | 3 |


| CHLI |  | Chromosome 3:213,650-426,098 | 3 |
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| NKIRAS1 |  | Chromosome 3:23,908,576-23,933,541 | 3 |
| NR1D2 |  | Chromosome 3:23,961,810-23,996,241 | 3 |
| THRB | NR1A2 | Chromosome 3:24,134,709-24,511,317 | 3 |
| THRB | NR1A2 | Chromosome 3:24,134,709-24,511,317 | 3 |
| THRB | NR1A2 | Chromosome 3:24,134,709-24,511,317 | 3 |
| THRB | NR1A2 | Chromosome 3:24,134,709-24,511,317 | 3 |
| THRB | NR1A2 | Chromosome 3:24,134,709-24,511,317 | 3 |
| RARB | NR1B2 | Chromosome 3:25,190,893-25,614,424 | 3 |
| RARB | NR1B2 | Chromosome 3:25,190,893-25,614,424 | 3 |
| RARB | NR1B2 | Chromosome 3:25,190,893-25,614,424 | 3 |
| RARB | NR1B2 | Chromosome 3:25,190,893-25,614,424 | 3 |
| RARB | NR1B2 | Chromosome 3:25,190,893-25,614,424 | 3 |
| EOMES | eomesodermin | Chromosome 3:27,732,872-27,738,807 | 3 |
| IL5RA | CD125 | Chromosome 3:3,086,421-3,127,031 | 3 |
| IL5RA | CD125 | Chromosome 3:3,086,421-3,127,031 | 3 |
| TGFBR2 |  | Chromosome 3:30,622,998-30,710,638 | 3 |
| TGFBR2 |  | Chromosome 3:30,622,998-30,710,638 | 3 |
| STT3B |  | Chromosome 3:31,549,495-31,652,560 | 3 |
| CMTM8 |  | Chromosome 3:32,255,175-32,386,817 | 3 |
| CMTM8 |  | Chromosome 3:32,255,175-32,386,817 | 3 |
| CMTM8 |  | Chromosome 3:32,255,175-32,386,817 | 3 |
| CMTM7 |  | Chromosome 3:32,408,167-32,471,337 | 3 |
| CMTM7 |  | Chromosome 3:32,408,167-32,471,337 | 3 |
| CMTM6 |  | Chromosome 3:32,497,808-32,519,869 | 3 |
| CMTM6 |  | Chromosome 3:32,497,808-32,519,869 | 3 |
| CCR4 | CD194 | Chromosome 3:32,968,070-32,972,840 | 3 |
| CLASP2 | c1AP2 | Chromosome 3:33,512,741-33,734,852 | 3 |
| PDCD6IP |  | Chromosome 3:33,814,561-33,886,198 | 3 |
| PDCD6IP |  | Chromosome 3:33,814,561-33,886,198 | 3 |
| ITGA9 |  | Chromosome 3:37,468,817-37,836,285 | 3 |
| ITGA9 |  | Chromosome 3:37,468,817-37,836,285 | 3 |
| ITGA9 |  | Chromosome 3:37,468,817-37,836,285 | 3 |
| ITGA9 |  | Chromosome 3:37,468,817-37,836,285 | 3 |
| ITGA9 |  | Chromosome 3:37,468,817-37,836,285 | 3 |
| ITGA9 |  | Chromosome 3:37,468,817-37,836,285 | 3 |
| ITGA9 |  | Chromosome 3:37,468,817-37,836,285 | 3 |
| CTDSPL | NIF1 | Chromosome 3:37,878,129-38,000,964 | 3 |
| CTDSPL | NIF1 | Chromosome 3:37,878,129-38,000,964 | 3 |
| MyD88 |  | Chromosome 3:38,155,009-38,159,517 | 3 |
| MyD88 |  | Chromosome 3:38,155,009-38,159,517 | 3 |
| CX3CR1 |  | Chromosome 3:39,279,989-39,298,190 | 3 |
| CX3CR1 |  | Chromosome 3:39,279,989-39,298,190 | 3 |
| CX3CR1 |  | Chromosome 3:39,279,989-39,298,190 | 3 |
| CCR8 | CD198 | Chromosome 3:39,346,219-39,351,077 | 3 |
| ITPR1 |  | Chromosome 3:4,510,136-4,864,081 | 3 |
| ITPR1 |  | Chromosome 3:4,510,136-4,864,081 | 3 |
| ITPR1 |  | Chromosome 3:4,510,136-4,864,081 | 3 |
| ITPR1 |  | Chromosome 3:4,510,136-4,864,081 | 3 |
| ITPR1 |  | Chromosome 3:4,510,136-4,864,081 | 3 |
| ITPR1 |  | Chromosome 3:4,510,136-4,864,081 | 3 |
| CTNNB1 | beta-catenin | Chromosome 3:41,216,004-41,256,938 | 3 |
| CTNNB1 | beta-catenin | Chromosome 3:41,216,004-41,256,938 | 3 |
| NKTR |  | Chromosome 3:42,617,151-42,665,237 | 3 |
| CCBP2 | D6 | Chromosome 3:42,825,980-42,883,779 | 3 |
| CYP8B1 |  | Chromosome 3:42,888,688-42,892,637 | 3 |


| CDCP1 | CD318 | Chromosome 3:45,098,773-45,162,918 | 3 |
| :---: | :---: | :---: | :---: |
| CDCP1 | CD318 | Chromosome 3:45,098,773-45,162,918 | 3 |
| CDCP1 | CD318 | Chromosome 3:45,098,773-45,162,918 | 3 |
| TMEM158 | RIS1 | Chromosome 3:45,240,962-45,242,758 | 3 |
| CCR9 | CD198, 199 | Chromosome 3:45,903,023-45,919,671 | 3 |
| XCR1 |  | Chromosome 3:46,037,295-46,043,983 | 3 |
| CCR1 | CD191 | Chromosome 3:46,218,204-46,224,836 | 3 |
| CCR3 | CD193 | Chromosome 3:46,227,186-46,283,166 | 3 |
| CCRL2 |  | Chromosome 3:46,423,725-46,426,018 | 3 |
| LTF | Lactoferrin | Chromosome 3:46,452,500-46,481,657 | 3 |
| PTHR1 | PTHR1 | Chromosome 3:46,894,240-46,926,585 | 3 |
| PTHR1 | PTHR1 | Chromosome 3:46,894,240-46,926,585 | 3 |
| PTPN23 |  | Chromosome 3:47,397,528-47,429,935 | 3 |
| DHX30 |  | Chromosome 3:47,819,625-47,866,687 | 3 |
| PLXNB1 | Plexin-B | Chromosome 3:48,420,266-48,446,464 | 3 |
| GPX1 | GSH peroxidase, cGPx | Chromosome 3:49,369,613-49,370,795 | 3 |
| RHOA | RHOA | Chromosome 3:49,371,582-49,424,530 | 3 |
| DAG1 | DAG | Chromosome 3:49,482,595-49,548,048 | 3 |
| DAG1 | DAG | Chromosome 3:49,482,595-49,548,048 | 3 |
| MST1 | MST1/STK4 | Chromosome 3:49,696,391-49,701,099 | 3 |
| MST1R | CDw136 | Chromosome 3:49,899,439-49,916,074 | 3 |
| MST1R | CDw136 | Chromosome 3:49,899,439-49,916,074 | 3 |
| MST1R | CDw136 | Chromosome 3:49,899,439-49,916,074 | 3 |
| DUSP7 |  | Chromosome 3:52,059,799-52,065,329 | 3 |
| DUSP7 |  | Chromosome 3:52,059,799-52,065,329 | 3 |
| TLR9 | CD289 | Chromosome 3:52,230,138-52,248,223 | 3 |
| TLR9 | CD289 | Chromosome 3:52,230,138-52,248,223 | 3 |
| STAB1 | FEEL-1 | Chromosome 3:52,504,396-52,533,551 | 3 |
| STAB1 | FEEL-1 | Chromosome 3:52,504,396-52,533,551 | 3 |
| PRKCD | PKC delta | Chromosome 3:53,170,263-53,201,773 | 3 |
| IL17RB |  | Chromosome 3:53,855,612-53,874,867 | 3 |
| IL17RB |  | Chromosome 3:53,855,612-53,874,867 | 3 |
| WNT5A |  | Chromosome 3:55,474,783-55,496,371 | 3 |
| ERC2 |  | Chromosome 3:55,517,376-56,477,431 | 3 |
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| ERC2 |  | Chromosome 3:55,517,376-56,477,431 | 3 |
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| ERC2 |  | Chromosome 3:55,517,376-56,477,431 | 3 |
| ERC2 |  | Chromosome 3:55,517,376-56,477,431 | 3 |
| ERC2 |  | Chromosome 3:55,517,376-56,477,431 | 3 |
| ERC2 |  | Chromosome 3:55,517,376-56,477,431 | 3 |
| ERC2 |  | Chromosome 3:55,517,376-56,477,431 | 3 |
| IL17RD |  | Chromosome 3:57,103,316-57,179,374 | 3 |
| IL17RD |  | Chromosome 3:57,103,316-57,179,374 | 3 |
| IL17RD |  | Chromosome 3:57,103,316-57,179,374 | 3 |
| PTPRG |  | Chromosome 3:61,522,285-62,255,613 | 3 |
| PTPRG |  | Chromosome 3:61,522,285-62,255,613 | 3 |
| PTPRG |  | Chromosome 3:61,522,285-62,255,613 | 3 |
| PTPRG |  | Chromosome 3:61,522,285-62,255,613 | 3 |
| PTPRG |  | Chromosome 3:61,522,285-62,255,613 | 3 |
| PTPRG |  | Chromosome 3:61,522,285-62,255,613 | 3 |
| PTPRG |  | Chromosome 3:61,522,285-62,255,613 | 3 |
| PTPRG |  | Chromosome 3:61,522,285-62,255,613 | 3 |
| ARL6IP5 | JWA | Chromosome 3:69,216,780-69,237,929 | 3 |
| MITF |  | Chromosome 3:69,871,323-70,100,177 | 3 |


| MITF |  | Chromosome 3:69,871,323-70,100,177 | 3 |
| :---: | :---: | :---: | :---: |
| MITF |  | Chromosome 3:69,871,323-70,100,177 | 3 |
| MITF |  | Chromosome 3:69,871,323-70,100,177 | 3 |
| FOXP1 |  | Chromosome 3:71,087,426-71,715,830 | 3 |
| FOXP1 |  | Chromosome 3:71,087,426-71,715,830 | 3 |
| FOXP1 |  | Chromosome 3:71,087,426-71,715,830 | 3 |
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| FOXP1 |  | Chromosome 3:71,087,426-71,715,830 | 3 |
| FOXP1 |  | Chromosome 3:71,087,426-71,715,830 | 3 |
| FOXP1 |  | Chromosome 3:71,087,426-71,715,830 | 3 |
| FOXP1 |  | Chromosome 3:71,087,426-71,715,830 | 3 |
| FOXP1 |  | Chromosome 3:71,087,426-71,715,830 | 3 |
| FOXP1 |  | Chromosome 3:71,087,426-71,715,830 | 3 |
| FOXP1 |  | Chromosome 3:71,087,426-71,715,830 | 3 |
| CADM2 |  | Chromosome 3:85,858,322-86,200,641 | 3 |
| CADM2 |  | Chromosome 3:85,858,322-86,200,641 | 3 |
| CADM2 |  | Chromosome 3:85,858,322-86,200,641 | 3 |
| CADM2 |  | Chromosome 3:85,858,322-86,200,641 | 3 |
| CADM2 |  | Chromosome 3:85,858,322-86,200,641 | 3 |
| CHMP2B | CHMP2B | Chromosome 3:87,359,140-87,387,339 | 3 |
| CHMP2B | CHMP2B | Chromosome 3:87,359,140-87,387,339 | 3 |
| CHMP2B | CHMP2B | Chromosome 3:87,359,140-87,387,339 | 3 |
| IL17RE |  | Chromosome 3:9,919,150-9,933,086 | 3 |
| IL17RC |  | Chromosome 3:9,933,782-9,950,314 | 3 |
| Protein S |  | Chromosome 3:95,074,647-95,175,412 | 3 |
| FGFR3 | CD333 | Chromosome 4:1,764,832-1,780,396 | 4 |
| H2AFZ |  | Chromosome 4:101,088,265-101,090,535 | 4 |
| PPP3CA |  | Chromosome 4:102,163,610-102,487,376 | 4 |
| PPP3CA |  | Chromosome 4:102,163,610-102,487,376 | 4 |
| PPP3CA |  | Chromosome 4:102,163,610-102,487,376 | 4 |
| PPP3CA |  | Chromosome 4:102,163,610-102,487,376 | 4 |
| PPP3CA |  | Chromosome 4:102,163,610-102,487,376 | 4 |
| PPP3CA |  | Chromosome 4:102,163,610-102,487,376 | 4 |
| BANK1 | B cell scaffold protein | Chromosome 4:102,930,919-103,214,918 | 4 |
| BANK1 | B cell scaffold protein | Chromosome 4:102,930,919-103,214,918 | 4 |
| BANK1 | B cell scaffold protein | Chromosome 4:102,930,919-103,214,918 | 4 |
| NFKB1 | p105, p50 | Chromosome 4:103,641,518-103,757,507 | 4 |
| NFKB1 | p105, p50 | Chromosome 4:103,641,518-103,757,507 | 4 |
| SCYE1 | SCYE1 | Chromosome 4:107,456,302-107,489,097 | 4 |
| SCYE1 | SCYE1 | Chromosome 4:107,456,302-107,489,097 | 4 |
| CYP2U1 |  | Chromosome 4:109,072,166-109,094,062 | 4 |
| CYP2U1 |  | Chromosome 4:109,072,166-109,094,062 | 4 |
| HS3ST1 |  | Chromosome 4:11,009,086-11,040,487 | 4 |
| HS3ST1 |  | Chromosome 4:11,009,086-11,040,487 | 4 |
| CFI |  | Chromosome 4:110,881,301-110,942,590 | 4 |
| EGF1 |  | Chromosome 4:111,053,499-111,152,868 | 4 |
| EGF1 |  | Chromosome 4:111,053,499-111,152,868 | 4 |
| EGF1 |  | Chromosome 4:111,053,499-111,152,868 | 4 |
| EGF1 |  | Chromosome 4:111,053,499-111,152,868 | 4 |
| ENPEP | CD249 | Chromosome 4:111,616,697-111,702,872 | 4 |
| ENPEP | CD249 | Chromosome 4:111,616,697-111,702,872 | 4 |
| ANK2 |  | Chromosome 4:114,190,319-114,524,337 | 4 |
| ANK2 |  | Chromosome 4:114,190,319-114,524,337 | 4 |
| ANK2 |  | Chromosome 4:114,190,319-114,524,337 | 4 |
| ANK2 |  | Chromosome 4:114,190,319-114,524,337 | 4 |
| CAMK2D | CaMKII | Chromosome 4:114,593,021-114,902,177 | 4 |


| CAMK2D | CaMKII | Chromosome 4:114,593,021-114,902,177 | 4 |
| :---: | :---: | :---: | :---: |
| CAMK2D | CaMKII | Chromosome 4:114,593,021-114,902,177 | 4 |
| CAMK2D | CaMKII | Chromosome 4:114,593,021-114,902,177 | 4 |
| CAMK2D | CaMKII | Chromosome 4:114,593,021-114,902,177 | 4 |
| MAD2LI |  | Chromosome 4:121,200,029-121,207,411 | 4 |
| MAD2LI |  | Chromosome 4:121,200,029-121,207,411 | 4 |
| ANXA5 |  | Chromosome 4:122,808,598-122,837,626 | 4 |
| CCNA2 |  | Chromosome 4:122,957,975-122,964,505 | 4 |
| IL2 |  | Chromosome 4:123,592,075-123,597,339 | 4 |
| IL21 |  | Chromosome 4:123,753,221-123,761,662 | 4 |
| FGF2 | FGF2 | Chromosome 4:123,967,313-124,038,840 | 4 |
| FGF2 | FGF2 | Chromosome 4:123,967,313-124,038,840 | 4 |
| PLK4 | Sak | Chromosome 4:129,021,551-129,039,377 | 4 |
| C1QTNF7 |  | Chromosome 4:14,950,658-15,056,887 | 4 |
| C1QTNF7 |  | Chromosome 4:14,950,658-15,056,887 | 4 |
| CCRN4L |  | Chromosome 4:140,156,393-140,186,543 | 4 |
| IL15 |  | Chromosome 4:142,777,204-142,874,062 | 4 |
| IL15 |  | Chromosome 4:142,777,204-142,874,062 | 4 |
| SMARCA5 |  | Chromosome 4:144,654,066-144,694,017 | 4 |
| SMARCA5 |  | Chromosome 4:144,654,066-144,694,017 | 4 |
| GYPB | MNSs antigen, CD235b | Chromosome 4:145,136,707-145,159,946 | 4 |
| GYPA | MNSs antigen, CD235a | Chromosome 4:145,249,906-145,281,294 | 4 |
| SMAD1 |  | Chromosome 4:146,622,401-146,699,778 | 4 |
| GTF2F2L |  | Chromosome 4:148,646,691-148,647,812 | 4 |
| GTF2F2L |  | Chromosome 4:148,646,691-148,647,812 | 4 |
| GTF2F2L |  | Chromosome 4:148,646,691-148,647,812 | 4 |
| NR3C2 | NR3C2 | Chromosome 4:149,219,370-149,582,973 | 4 |
| NR3C2 | NR3C2 | Chromosome 4:149,219,370-149,582,973 | 4 |
| NR3C2 | NR3C2 | Chromosome 4:149,219,370-149,582,973 | 4 |
| NR3C2 | NR3C2 | Chromosome 4:149,219,370-149,582,973 | 4 |
| NR3C2 | NR3C2 | Chromosome 4:149,219,370-149,582,973 | 4 |
| BST1 | CD157 | Chromosome 4:15,313,738-15,343,508 | 4 |
| BST1 | CD157 | Chromosome 4:15,313,738-15,343,508 | 4 |
| CD38 |  | Chromosome 4:15,388,999-15,460,167 | 4 |
| PROM1 | CD133, SCA | Chromosome 4:15,578,955-15,686,664 | 4 |
| PROM1 | CD133, SCA | Chromosome 4:15,578,955-15,686,664 | 4 |
| FBXW7 |  | Chromosome 4:153,461,860-153,675,622 | 4 |
| FBXW7 |  | Chromosome 4:153,461,860-153,675,622 | 4 |
| FBXW7 |  | Chromosome 4:153,461,860-153,675,622 | 4 |
| FBXW7 |  | Chromosome 4:153,461,860-153,675,622 | 4 |
| TLR2 | CD282 | Chromosome 4:154,824,891-154,846,693 | 4 |
| TLR2 | CD282 | Chromosome 4:154,824,891-154,846,693 | 4 |
| FGB | fibrinogen | Chromosome 4:155,703,596-155,711,688 | 4 |
| NPY2R |  | Chromosome 4:156,349,231-156,357,678 | 4 |
| PPID |  | Chromosome 4:159,849,729-159,864,002 | 4 |
| PPID |  | Chromosome 4:159,849,729-159,864,002 | 4 |
| CPE |  | Chromosome 4:166,519,538-166,638,926 | 4 |
| CPE |  | Chromosome 4:166,519,538-166,638,926 | 4 |
| HPGD |  | Chromosome 4:175,647,955-175,680,213 | 4 |
| HPGD |  | Chromosome 4:175,647,955-175,680,213 | 4 |
| HPGD |  | Chromosome 4:175,647,955-175,680,213 | 4 |
| HPGD |  | Chromosome 4:175,647,955-175,680,213 | 4 |
| VEGFC |  | Chromosome 4:177,841,685-177,950,889 | 4 |
| IRF2 |  | Chromosome 4:185,545,909-185,632,697 | 4 |
| IRF2 |  | Chromosome 4:185,545,909-185,632,697 | 4 |
| IRF2 |  | Chromosome 4:185,545,909-185,632,697 | 4 |


| TLR3 | CD283 | Chromosome 4:187,227,303-187,243,246 | 4 |
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| CYP4V2 |  | Chromosome 4:187,349,668-187,371,606 | 4 |
| CYP4V2 |  | Chromosome 4:187,349,668-187,371,606 | 4 |
| RNF4 |  | Chromosome 4:2,440,605-2,487,382 | 4 |
| ADD1 | alpha adducin | Chromosome 4:2,815,382-2,901,587 | 4 |
| DHX15 |  | Chromosome 4:24,138,185-24,195,282 | 4 |
| DHX15 |  | Chromosome 4:24,138,185-24,195,282 | 4 |
| SOD3 | IP01 | Chromosome 4:24,405,153-24,411,562 | 4 |
| SOD3 | IP01 | Chromosome 4:24,405,153-24,411,562 | 4 |
| Pl4K2B | PI4K2B | Chromosome 4:24,844,751-24,889,811 | 4 |
| RBPSUH | RBP-Jk | Chromosome 4:25,930,430-26,045,851 | 4 |
| RBPSUH | RBP-Jk | Chromosome 4:25,930,430-26,045,851 | 4 |
| RBPSUH | RBP-Jk | Chromosome 4:25,930,430-26,045,851 | 4 |
| RBPSUH | RBP-Jk | Chromosome 4:25,930,430-26,045,851 | 4 |
| TLR10 | CD290 | Chromosome 4:38,450,255-38,460,984 | 4 |
| TLR10 | CD290 | Chromosome 4:38,450,255-38,460,984 | 4 |
| TLR6 | CD286 | Chromosome 4:38,504,618-38,507,555 | 4 |
| RFC1 | RFC | Chromosome 4:38,965,471-39,044,390 | 4 |
| RFC1 | RFC | Chromosome 4:38,965,471-39,044,390 | 4 |
| RFC1 | RFC | Chromosome 4:38,965,471-39,044,390 | 4 |
| RHOH | RhoH | Chromosome 4:39,874,965-39,922,663 | 4 |
| RHOH | RhoH | Chromosome 4:39,874,965-39,922,663 | 4 |
| TXK | Txk, Rlk, thioredoxin | Chromosome 4:47,762,988-47,831,030 | 4 |
| TXK | Txk, Rlk, thioredoxin | Chromosome 4:47,762,988-47,831,030 | 4 |
| TXK | Txk, Rlk, thioredoxin | Chromosome 4:47,762,988-47,831,030 | 4 |
| CYTL1 | cytokine-like 1 | Chromosome 4:5,067,214-5,072,100 | 4 |
| CYTL1 | cytokine-like 1 | Chromosome 4:5,067,214-5,072,100 | 4 |
| PDGFRA | CD140a | Chromosome 4:54,790,204-54,859,171 | 4 |
| PDGFRA | CD140a | Chromosome 4:54,790,204-54,859,171 | 4 |
| KIT | FLT3-ITD, CD117, c-KIT | Chromosome 4:55,218,842-55,301,638 | 4 |
| KIT | FLT3-ITD, CD117, c-KIT | Chromosome 4:55,218,842-55,301,638 | 4 |
| KIT | FLT3-ITD, CD117, c-KIT | Chromosome 4:55,218,842-55,301,638 | 4 |
| KDR | CD309, VEGFR2 | Chromosome 4:55,639,401-55,686,519 | 4 |
| BRDG1 |  | Chromosome 4:68,107,041-68,155,206 | 4 |
| BRDG1 |  | Chromosome 4:68,107,041-68,155,206 | 4 |
| GNRHR | GNRH1-R | Chromosome 4:68,285,688-68,304,399 | 4 |
| TMPRSS11D | HAT | Chromosome 4:68,369,189-68,432,311 | 4 |
| TMPRSS11D | HAT | Chromosome 4:68,369,189-68,432,311 | 4 |
| UGT2B17 | UGT2 | Chromosome 4:69,085,497-69,116,840 | 4 |
| UGT2B7 |  | Chromosome 4:69,996,782-70,013,293 | 4 |
| UGT2B28 |  | Chromosome 4:70,180,783-70,323,496 | 4 |
| UGT2B28 |  | Chromosome 4:70,180,783-70,323,496 | 4 |
| HTN3 | histatin 3 | Chromosome 4:70,928,761-70,936,836 | 4 |
| HTN3 | histatin 3 | Chromosome 4:70,928,761-70,936,836 | 4 |
| IGJ |  | Chromosome 4:71,740,548-71,751,128 | 4 |
| IL8 | CXCL8 | Chromosome 4:74,825,139-74,828,297 | 4 |
| CXCL6 |  | Chromosome 4:74,921,277-74,923,341 | 4 |
| PF4 | CXCL4 | Chromosome 4:75,065,660-75,066,541 | 4 |
| CXCL3 |  | Chromosome 4:75,121,170-75,123,354 | 4 |
| EREG | EREG | Chromosome 4:75,449,724-75,473,341 | 4 |
| AREG | AREG | Chromosome 4:75,529,717-75,539,590 | 4 |
| AREG | AREG | Chromosome 4:75,529,717-75,539,590 | 4 |
| BTC | BTC betacellulin | Chromosome 4:75,889,001-75,938,853 | 4 |
| BTC | BTC betacellulin | Chromosome 4:75,889,001-75,938,853 | 4 |
| CXCL9 | Mig | Chromosome 4:77,141,523-77,147,665 | 4 |
| SCARB2 |  | Chromosome 4:77,298,918-77,354,059 | 4 |


| SCARB2 |  | Chromosome 4:77,298,918-77,354,059 | 4 |
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| CXCL13 |  | Chromosome 4:78,651,931-78,752,010 | 4 |
| CXCL13 |  | Chromosome 4:78,651,931-78,752,010 | 4 |
| CXCL13 |  | Chromosome 4:78,651,931-78,752,010 | 4 |
| FGF5 | FGF5 | Chromosome 4:81,406,766-81,431,195 | 4 |
| FGF5 | FGF5 | Chromosome 4:81,406,766-81,431,195 | 4 |
| BMP3 | BMP3 | Chromosome 4:82,171,143-82,193,749 | 4 |
| BMP3 | BMP3 | Chromosome 4:82,171,143-82,193,749 | 4 |
| HPA |  | Chromosome 4:84,432,639-84,475,330 | 4 |
| MAPK10 |  | Chromosome 4:87,156,656-87,593,307 | 4 |
| MAPK10 |  | Chromosome 4:87,156,656-87,593,307 | 4 |
| MAPK10 |  | Chromosome 4:87,156,656-87,593,307 | 4 |
| MAPK10 |  | Chromosome 4:87,156,656-87,593,307 | 4 |
| MAPK10 |  | Chromosome 4:87,156,656-87,593,307 | 4 |
| PTPN13 |  | Chromosome 4:87,734,909-87,955,326 | 4 |
| PTPN13 |  | Chromosome 4:87,734,909-87,955,326 | 4 |
| PTPN13 |  | Chromosome 4:87,734,909-87,955,326 | 4 |
| SPP1 | Osteopontin, Eta1 | Chromosome 4:89,115,826-89,123,592 | 4 |
| SPP1 | Osteopontin, Eta1 | Chromosome 4:89,115,826-89,123,592 | 4 |
| ABCG2 | CD338, BCRP, MRX, MX | Chromosome 4:89,230,440-89,299,035 | 4 |
| HERC6 | FLJ20637 | Chromosome 4:89,518,915-89,583,272 | 4 |
| HERC6 | FLJ20637 | Chromosome 4:89,518,915-89,583,272 | 4 |
| HERC6 | FLJ20637 | Chromosome 4:89,518,915-89,583,272 | 4 |
| DFB131 |  | Chromosome 4:9,055,358-9,061,338 | 4 |
| PGDS |  | Chromosome 4:95,438,730-95,483,050 | 4 |
| BMPR1B | CD293 | Chromosome 4:95,898,151-96,295,099 | 4 |
| BMPR1B | CD293 | Chromosome 4:95,898,151-96,295,099 | 4 |
| BMPR1B | CD293 | Chromosome 4:95,898,151-96,295,099 | 4 |
| BMPR1B | CD293 | Chromosome 4:95,898,151-96,295,099 | 4 |
| BMPR1B | CD293 | Chromosome 4:95,898,151-96,295,099 | 4 |
| IBD5 |  | Chromosome 5 | 5 |
| IBD5 |  | Chromosome 5 | 5 |
| TERT |  | Chromosome 5:1,306,282-1,348,162 | 5 |
| TERT |  | Chromosome 5:1,306,282-1,348,162 | 5 |
| DAP |  | Chromosome 5:10,732,343-10,814,344 | 5 |
| DAP |  | Chromosome 5:10,732,343-10,814,344 | 5 |
| TSLP | Thymic stromal lymphopo | Chromosome 5:110,433,677-110,441,623 | 5 |
| TSLP | Thymic stromal lymphopo | Chromosome 5:110,433,677-110,441,623 | 5 |
| CAMK4 | CaMKIV | Chromosome 5:110,587,968-110,858,483 | 5 |
| CAMK4 | CaMKIV | Chromosome 5:110,587,968-110,858,483 | 5 |
| CAMK4 | CaMKIV | Chromosome 5:110,587,968-110,858,483 | 5 |
| CAMK4 | CaMKIV | Chromosome 5:110,587,968-110,858,483 | 5 |
| APC |  | Chromosome 5:112,101,483-112,209,835 | 5 |
| APC |  | Chromosome 5:112,101,483-112,209,835 | 5 |
| TCAM2 | TIRP | Chromosome 5:114,942,247-114,989,610 | 5 |
| TCAM2 | TIRP | Chromosome 5:114,942,247-114,989,610 | 5 |
| CDO1 |  | Chromosome 5:115,168,329-115,180,304 | 5 |
| PPIC |  | Chromosome 5:122,386,977-122,400,324 | 5 |
| PPIC |  | Chromosome 5:122,386,977-122,400,324 | 5 |
| IL3 |  | Chromosome 5:131,424,121-131,426,796 | 5 |
| SLC22A4 | OCTN1 | Chromosome 5:131,658,035-131,707,798 | 5 |
| SLC22A4 | OCTN1 | Chromosome 5:131,658,035-131,707,798 | 5 |
| SLC22A4 | OCTN1 | Chromosome 5:131,658,035-131,707,798 | 5 |
| SLC22A5 | OCTN2 | Chromosome 5:131,733,343-131,759,205 | 5 |
| IRF1 |  | Chromosome 5:131,845,200-131,854,389 | 5 |
| RAD50 | rad50 | Chromosome 5:131,920,529-132,007,651 | 5 |


| LEAP2 | liver expressed antimicrob | Chromosome 5:132,235,913-132,238,637 | 5 |
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| LEAP2 | liver expressed antimicrob | Chromosome 5:132,235,913-132,238,637 | 5 |
| HSPA4 |  | Chromosome 5:132,415,561-132,468,608 | 5 |
| DDX46 |  | Chromosome 5:134,122,360-134,194,710 | 5 |
| DDX46 |  | Chromosome 5:134,122,360-134,194,710 | 5 |
| CXCL14 |  | Chromosome 5:134,934,274-134,942,868 | 5 |
| IL9 |  | Chromosome 5:135,255,834-135,259,415 | 5 |
| IL9 |  | Chromosome 5:135,255,834-135,259,415 | 5 |
| IL9 |  | Chromosome 5:135,255,834-135,259,415 | 5 |
| BRD8 | SMAP | Chromosome 5:137,503,358-137,542,257 | 5 |
| BRD8 | SMAP | Chromosome 5:137,503,358-137,542,257 | 5 |
| BRD8 | SMAP | Chromosome 5:137,503,358-137,542,257 | 5 |
| EGR1 |  | Chromosome 5:137,829,080-137,832,903 | 5 |
| EGR1 |  | Chromosome 5:137,829,080-137,832,903 | 5 |
| NRG2 | NRG2 | Chromosome 5:139,207,444-139,403,063 | 5 |
| NRG2 | NRG2 | Chromosome 5:139,207,444-139,403,063 | 5 |
| NRG2 | NRG2 | Chromosome 5:139,207,444-139,403,063 | 5 |
| HBEGF | HBEGF | Chromosome 5:139,692,612-139,706,359 | 5 |
| HBEGF | HBEGF | Chromosome 5:139,692,612-139,706,359 | 5 |
| CD14 |  | Chromosome 5:139,991,501-139,993,439 | 5 |
| NDUFA2 | B8/HY | Chromosome 5:140,005,142-140,007,424 | 5 |
| PCDHB5 |  | Chromosome 5:140,494,984-140,497,888 | 5 |
| PCDHB16 |  | Chromosome 5:140,541,164-140,545,980 | 5 |
| HDAC3 |  | Chromosome 5:140,980,627-140,996,596 | 5 |
| FGF1 |  | Chromosome 5:141,951,927-142,046,134 | 5 |
| FGF1 |  | Chromosome 5:141,951,927-142,046,134 | 5 |
| NR3C1 | GR | Chromosome 5:142,637,689-142,795,270 | 5 |
| NR3C1 | GR | Chromosome 5:142,637,689-142,795,270 | 5 |
| HMHB1 | HB-1 | Chromosome 5:143,171,919-143,180,477 | 5 |
| HMHB1 | HB-1 | Chromosome 5:143,171,919-143,180,477 | 5 |
| IL17B |  | Chromosome 5:148,734,023-148,739,031 | 5 |
| IL17B |  | Chromosome 5:148,734,023-148,739,031 | 5 |
| CSNK1A1 | casein kinase 1 | Chromosome 5:148,855,038-148,911,200 | 5 |
| CSF1R | CD115 | Chromosome 5:149,413,051-149,473,128 | 5 |
| CSF1R | CD115 | Chromosome 5:149,413,051-149,473,128 | 5 |
| PDGFRB | CD140b | Chromosome 5:149,473,595-149,515,615 | 5 |
| PDGFRB | CD140b | Chromosome 5:149,473,595-149,515,615 | 5 |
| CD74 | p41 | Chromosome 5:149,761,393-149,772,685 | 5 |
| IRGM | LRG47 | Chromosome 5:150,207,879-150,260,488 | 5 |
| GPX3 |  | Chromosome 5:150,380,112-150,388,747 | 5 |
| HAVCR2 | TIM-3 | Chromosome 5:156,445,421-156,468,716 | 5 |
| ITK | Itk | Chromosome 5:156,540,432-156,614,687 | 5 |
| ITK | Itk | Chromosome 5:156,540,432-156,614,687 | 5 |
| EBF |  | Chromosome 5:158,058,006-158,459,347 | 5 |
| EBF |  | Chromosome 5:158,058,006-158,459,347 | 5 |
| EBF |  | Chromosome 5:158,058,006-158,459,347 | 5 |
| EBF |  | Chromosome 5:158,058,006-158,459,347 | 5 |
| EBF |  | Chromosome 5:158,058,006-158,459,347 | 5 |
| EBF |  | Chromosome 5:158,058,006-158,459,347 | 5 |
| IL12B |  | Chromosome 5:158,674,369-158,690,059 | 5 |
| IL12B |  | Chromosome 5:158,674,369-158,690,059 | 5 |
| IL12B |  | Chromosome 5:158,674,369-158,690,059 | 5 |
| C1QTNF2 |  | Chromosome 5:159,707,339-159,730,207 | 5 |
| C1QTNF2 |  | Chromosome 5:159,707,339-159,730,207 | 5 |
| CCNG1 |  | Chromosome 5:162,797,155-162,804,600 | 5 |
| HMMR | CD168 | Chromosome 5:162,820,241-162,851,525 | 5 |


| DOCK2 | dedicator of cytokinesis 2 | Chromosome 5:168,996,871-169,442,959 | 5 |
| :---: | :---: | :---: | :---: |
| DOCK2 | dedicator of cytokinesis 2 | Chromosome 5:168,996,871-169,442,959 | 5 |
| DOCK2 | dedicator of cytokinesis 2 | Chromosome 5:168,996,871-169,442,959 | 5 |
| DOCK2 | dedicator of cytokinesis 2 | Chromosome 5:168,996,871-169,442,959 | 5 |
| DOCK2 | dedicator of cytokinesis 2 | Chromosome 5:168,996,871-169,442,959 | 5 |
| DOCK2 | dedicator of cytokinesis 2 | Chromosome 5:168,996,871-169,442,959 | 5 |
| DOCK2 | dedicator of cytokinesis 2 | Chromosome 5:168,996,871-169,442,959 | 5 |
| DOCK2 | dedicator of cytokinesis 2 | Chromosome 5:168,996,871-169,442,959 | 5 |
| FLJ46831 | FOXI1 | Chromosome 5:169,465,495-169,469,305 | 5 |
| LCP2 | SLP76 | Chromosome 5:169,607,666-169,657,400 | 5 |
| LPC2 | SLP-76 | Chromosome 5:169,607,666-169,657,400 | 5 |
| DUSP1 | MKP1 | Chromosome 5:172,127,707-172,130,809 | 5 |
| BNIP1 |  | Chromosome 5:172,504,130-172,523,989 | 5 |
| BNIP1 |  | Chromosome 5:172,504,130-172,523,989 | 5 |
| BNIP1 |  | Chromosome 5:172,504,130-172,523,989 | 5 |
| HRH2 |  | Chromosome 5:175,017,637-175,045,847 | 5 |
| HRH2 |  | Chromosome 5:175,017,637-175,045,847 | 5 |
| FGFR4 | CD334 | Chromosome 5:176,446,493-176,457,733 | 5 |
| DDX41 |  | Chromosome 5:176,871,184-176,876,573 | 5 |
| DDX41 |  | Chromosome 5:176,871,184-176,876,573 | 5 |
| CANX |  | Chromosome 5:179,058,536-179,091,248 | 5 |
| MAPK9 |  | Chromosome 5:179,595,388-179,640,218 | 5 |
| FLT4 | CD310, VEGFR3 | Chromosome 5:179,945,812-180,009,172 | 5 |
| FLT4 | CD310, VEGFR3 | Chromosome 5:179,945,812-180,009,172 | 5 |
| FLT4 | CD310, VEGFR3 | Chromosome 5:179,945,812-180,009,172 | 5 |
| FOXO1B |  | Chromosome 5:180,458,383-180,460,484 | 5 |
| FOXO1B |  | Chromosome 5:180,458,383-180,460,484 | 5 |
| PDCD6 |  | Chromosome 5:324,739-488,225 | 5 |
| PDCD6 |  | Chromosome 5:324,739-488,225 | 5 |
| PDCD6 |  | Chromosome 5:324,739-488,225 | 5 |
| C1QTNF3 |  | Chromosome 5:34,022,040-34,160,396 | 5 |
| C1QTNF3 |  | Chromosome 5:34,022,040-34,160,396 | 5 |
| C1QTNF3 |  | Chromosome 5:34,022,040-34,160,396 | 5 |
| PRLR | PRLR | Chromosome 5:35,084,621-35,266,334 | 5 |
| PRLR | PRLR | Chromosome 5:35,084,621-35,266,334 | 5 |
| PRLR | PRLR | Chromosome 5:35,084,621-35,266,334 | 5 |
| PRLR | PRLR | Chromosome 5:35,084,621-35,266,334 | 5 |
| IL7R | CD127 | Chromosome 5:35,892,748-35,915,462 | 5 |
| IL7R | CD127 | Chromosome 5:35,892,748-35,915,462 | 5 |
| SKP2 |  | Chromosome 5:36,187,946-36,219,904 | 5 |
| SKP2 |  | Chromosome 5:36,187,946-36,219,904 | 5 |
| SKP2 |  | Chromosome 5:36,187,946-36,219,904 | 5 |
| LIFR | CD118 | Chromosome 5:38,510,822-38,631,253 | 5 |
| LIFR | CD118 | Chromosome 5:38,510,822-38,631,253 | 5 |
| LIFR | CD118 | Chromosome 5:38,510,822-38,631,253 | 5 |
| FYB | SLAP-130, ADAP | Chromosome 5:39,141,114-39,255,432 | 5 |
| FYB | SLAP-130, ADAP | Chromosome 5:39,141,114-39,255,432 | 5 |
| FYB | SLAP-130, ADAP | Chromosome 5:39,141,114-39,255,432 | 5 |
| C9 |  | Chromosome 5:39,320,061-39,400,412 | 5 |
| PTGER4 | EP4 | Chromosome 5:40,715,789-40,729,594 | 5 |
| PTGER4 | EP4 | Chromosome 5:40,715,789-40,729,594 | 5 |
| CARD6 |  | Chromosome 5:40,877,043-40,896,025 | 5 |
| C7 |  | Chromosome 5:40,945,356-41,018,798 | 5 |
| C6 |  | Chromosome 5:41,178,093-41,297,297 | 5 |
| C6 |  | Chromosome 5:41,178,093-41,297,297 | 5 |
| GHR | growth hormone receptor | Chromosome 5:42,459,783-42,757,736 | 5 |


| GHR | growth hormone receptor | Chromosome 5:42,459,783-42,757,736 | 5 |
| :---: | :---: | :---: | :---: |
| GHR | growth hormone receptor | Chromosome 5:42,459,783-42,757,736 | 5 |
| GHR | growth hormone receptor | Chromosome 5:42,459,783-42,757,736 | 5 |
| CCL28 |  | Chromosome 5:43,229,915-43,448,250 | 5 |
| CCL28 |  | Chromosome 5:43,229,915-43,448,250 | 5 |
| CCL28 |  | Chromosome 5:43,229,915-43,448,250 | 5 |
| FGF10 | FGF10 | Chromosome 5:44,340,854-44,424,541 | 5 |
| FGF10 | FGF10 | Chromosome 5:44,340,854-44,424,541 | 5 |
| ITGA1 | VLA1-3, CD49a | Chromosome 5:52,119,531-52,285,242 | 5 |
| ITGA1 | VLA1-3, CD49a | Chromosome 5:52,119,531-52,285,242 | 5 |
| ITGA1 | VLA1-3, CD49a | Chromosome 5:52,119,531-52,285,242 | 5 |
| ITGA2 | CD49b | Chromosome 5:52,321,014-52,423,947 | 5 |
| ITGA2 | CD49b | Chromosome 5:52,321,014-52,423,947 | 5 |
| GZMK |  | Chromosome 5:54,355,838-54,366,155 | 5 |
| GZMA | Granzyme A | Chromosome 5:54,434,230-54,441,837 | 5 |
| DHX29 |  | Chromosome 5:54,587,830-54,639,278 | 5 |
| DDX4 |  | Chromosome 5:55,069,609-55,148,362 | 5 |
| DDX4 |  | Chromosome 5:55,069,609-55,148,362 | 5 |
| DDX4 |  | Chromosome 5:55,069,609-55,148,362 | 5 |
| IL31RA |  | Chromosome 5:55,183,091-55,248,922 | 5 |
| IL6ST | CD130, gp130 | Chromosome 5:55,266,680-55,326,529 | 5 |
| MAP3K1 | MEKK1 | Chromosome 5:56,146,022-56,227,736 | 5 |
| MAP3K1 | MEKK1 | Chromosome 5:56,146,022-56,227,736 | 5 |
| PDE4D | Phosphodiesterases | Chromosome 5:58,302,468-59,320,301 | 5 |
| PDE4D | Phosphodiesterases | Chromosome 5:58,302,468-59,320,301 | 5 |
| PDE4D | Phosphodiesterases | Chromosome 5:58,302,468-59,320,301 | 5 |
| PDE4D | Phosphodiesterases | Chromosome 5:58,302,468-59,320,301 | 5 |
| PDE4D | Phosphodiesterases | Chromosome 5:58,302,468-59,320,301 | 5 |
| PDE4D | Phosphodiesterases | Chromosome 5:58,302,468-59,320,301 | 5 |
| PDE4D | Phosphodiesterases | Chromosome 5:58,302,468-59,320,301 | 5 |
| PDE4D | Phosphodiesterases | Chromosome 5:58,302,468-59,320,301 | 5 |
| PDE4D | Phosphodiesterases | Chromosome 5:58,302,468-59,320,301 | 5 |
| PDE4D | Phosphodiesterases | Chromosome 5:58,302,468-59,320,301 | 5 |
| PDE4D | Phosphodiesterases | Chromosome 5:58,302,468-59,320,301 | 5 |
| PDE4D | Phosphodiesterases | Chromosome 5:58,302,468-59,320,301 | 5 |
| PDE4D | Phosphodiesterases | Chromosome 5:58,302,468-59,320,301 | 5 |
| PDE4D | Phosphodiesterases | Chromosome 5:58,302,468-59,320,301 | 5 |
| LY64 | CD180 | Chromosome 5:66,513,872-66,528,368 | 5 |
| LY64 | CD180 | Chromosome 5:66,513,872-66,528,368 | 5 |
| LY64 | CD180 | Chromosome 5:66,513,872-66,528,368 | 5 |
| PIK3R1 |  | Chromosome 5:67,547,360-67,633,405 | 5 |
| PIK3R1 |  | Chromosome 5:67,547,360-67,633,405 | 5 |
| CCNB1 |  | Chromosome 5:68,498,593-68,509,828 | 5 |
| CCNB1 |  | Chromosome 5:68,498,593-68,509,828 | 5 |
| CCNB1 |  | Chromosome 5:68,498,593-68,509,828 | 5 |
| TAF9 | TAFII32 | Chromosome 5:68,682,567-68,701,596 | 5 |
| TAF9 | TAFII32 | Chromosome 5:68,682,567-68,701,596 | 5 |
| MTRR |  | Chromosome 5:7,922,217-7,954,237 | 5 |
| MTRR |  | Chromosome 5:7,922,217-7,954,237 | 5 |
| FOXD1 |  | Chromosome 5:72,777,839-72,780,108 | 5 |
| FOXD1 |  | Chromosome 5:72,777,839-72,780,108 | 5 |
| ENC1 |  | Chromosome 5:73,958,990-73,973,005 | 5 |
| ENC1 |  | Chromosome 5:73,958,990-73,973,005 | 5 |
| ENC1 |  | Chromosome 5:73,958,990-73,973,005 | 5 |
| HMGCR | HMG-CoA reductase | Chromosome 5:74,668,790-74,693,685 | 5 |
| HMGCR | HMG-CoA reductase | Chromosome 5:74,668,790-74,693,685 | 5 |


| COL4A3BP |  | Chromosome 5:74,702,684-74,843,719 | 5 |
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| F2RL2 | PAR3 | Chromosome 5:75,947,063-75,954,996 | 5 |
| F2RL2 | PAR3 | Chromosome 5:75,947,063-75,954,996 | 5 |
| F2R | PAR1 | Chromosome 5:76,047,542-76,067,054 | 5 |
| F2RL1 | PAR2 | Chromosome 5:76,150,610-76,166,896 | 5 |
| LHFPL2 |  | Chromosome 5:77,816,794-77,841,979 | 5 |
| LHFPL2 |  | Chromosome 5:77,816,794-77,841,979 | 5 |
| DHFR | ok | Chromosome 5:79,957,801-79,986,556) | 5 |
| DHFR | ok | Chromosome 5:79,957,801-79,986,556) | 5 |
| CSPG2 | versican | Chromosome 5:82,803,339-82,912,737 | 5 |
| CSPG2 | versican | Chromosome 5:82,803,339-82,912,737 | 5 |
| CSPG2 | versican | Chromosome 5:82,803,339-82,912,737 | 5 |
| RASA1 |  | Chromosome 5:86,599,461-86,723,489 | 5 |
| RASA1 |  | Chromosome 5:86,599,461-86,723,489 | 5 |
| RASA1 |  | Chromosome 5:86,599,461-86,723,489 | 5 |
| NR2F1 | NR2F1 | Chromosome 5:92,944,799-92,956,077 | 5 |
| ARTS1 | ERAP1 | Chromosome 5:96,122,277-96,169,559 | 5 |
| ARTS1 | ERAP1 | Chromosome 5:96,122,277-96,169,559 | 5 |
| ARTS1 | ERAP1 | Chromosome 5:96,122,277-96,169,559 | 5 |
| IBD3 |  | Chromosome 6 | 6 |
| IBD3 |  | Chromosome 6 | 6 |
| FOXQ1 |  | Chromosome 6:1,257,675-1,259,983 | 6 |
| FOXQ1 |  | Chromosome 6:1,257,675-1,259,983 | 6 |
| FOXF2 |  | Chromosome 6:1,335,068-1,340,831 | 6 |
| FOXC1 |  | Chromosome 6:1,555,206-1,559,131 | 6 |
| GMDS |  | Chromosome 6:1,569,040-2,190,845 | 6 |
| GMDS |  | Chromosome 6:1,569,040-2,190,845 | 6 |
| GMDS |  | Chromosome 6:1,569,040-2,190,845 | 6 |
| GMDS |  | Chromosome 6:1,569,040-2,190,845 | 6 |
| GCNT2 |  | Chromosome 6:10,636,575-10,737,587 | 6 |
| GCNT2 |  | Chromosome 6:10,636,575-10,737,587 | 6 |
| SIM1 |  | Chromosome 6:100,939,606-101,019,494 | 6 |
| SIM1 |  | Chromosome 6:100,939,606-101,019,494 | 6 |
| GRIK2 |  | Chromosome 6:101,953,385-102,624,651 | 6 |
| HACE1 |  | Chromosome 6:105,282,661-105,414,867 | 6 |
| HACE1 |  | Chromosome 6:105,282,661-105,414,867 | 6 |
| PRDM1 | Blimp1 | Chromosome 6:106,640,888-106,664,507 | 6 |
| PRDM1 | Blimp1 | Chromosome 6:106,640,888-106,664,507 | 6 |
| ATG5 | Atg5 | Chromosome 6:106,739,044-106,880,388 | 6 |
| ATG5 | Atg5 | Chromosome 6:106,739,044-106,880,388 | 6 |
| SOBP |  | Chromosome 6:107,918,010-108,089,195 | 6 |
| SOBP |  | Chromosome 6:107,918,010-108,089,195 | 6 |
| NR2D1 | NR2D1 | Chromosome 6:108,593,955-108,616,706 | 6 |
| NR2D1 | NR2D1 | Chromosome 6:108,593,955-108,616,706 | 6 |
| NR2D1 | NR2D1 | Chromosome 6:108,593,955-108,616,706 | 6 |
| FOXO3A |  | Chromosome 6:108,987,719-109,108,661 | 6 |
| FOXO3A |  | Chromosome 6:108,987,719-109,108,661 | 6 |
| CD164 |  | Chromosome 6:109,794,412-109,810,353 | 6 |
| PPIL6 |  | Chromosome 6:109,820,624-109,868,524 | 6 |
| NEDD9 | Cas-L, HEF1 | Chromosome 6:11,291,517-11,490,535 | 6 |
| NEDD9 | Cas-L, HEF1 | Chromosome 6:11,291,517-11,490,535 | 6 |
| NEDD9 | Cas-L, HEF1 | Chromosome 6:11,291,517-11,490,535 | 6 |
| NEDD9 | Cas-L, HEF1 | Chromosome 6:11,291,517-11,490,535 | 6 |
| NEDD9 | Cas-L, HEF1 | Chromosome 6:11,291,517-11,490,535 | 6 |
| WASF1 | Wiskott Aldrich | Chromosome 6:110,527,715-110,607,900 | 6 |
| WASF1 | Wiskott Aldrich | Chromosome 6:110,527,715-110,607,900 | 6 |


| BXDC1 |  | Chromosome 6:111,409,984-111,453,487 | 6 |
| :---: | :---: | :---: | :---: |
| FYN | FynT | Chromosome 6:112,088,228-112,301,348 | 6 |
| FYN | FynT | Chromosome 6:112,088,228-112,301,348 | 6 |
| FYN | FynT | Chromosome 6:112,088,228-112,301,348 | 6 |
| FYN | FynT | Chromosome 6:112,088,228-112,301,348 | 6 |
| LAMA4 |  | Chromosome 6:112,536,654-112,682,605 | 6 |
| LAMA4 |  | Chromosome 6:112,536,654-112,682,605 | 6 |
| LAMA4 |  | Chromosome 6:112,536,654-112,682,605 | 6 |
| HDAC2 |  | Chromosome 6:114,368,571-114,399,029 | 6 |
| HDAC2 |  | Chromosome 6:114,368,571-114,399,029 | 6 |
| FRK | FynT | Chromosome 6:116,369,386-116,488,614 | 6 |
| FRK | FynT | Chromosome 6:116,369,386-116,488,614 | 6 |
| RFXDC1 | RFXDC1 | Chromosome 6:117,305,068-117,360,008 | 6 |
| C6orf204 |  | Chromosome 6:118,892,932-119,137,924 | 6 |
| C6orf204 |  | Chromosome 6:118,892,932-119,137,924 | 6 |
| C6orf204 |  | Chromosome 6:118,892,932-119,137,924 | 6 |
| C6orf204 |  | Chromosome 6:118,892,932-119,137,924 | 6 |
| MAN1A1 |  | Chromosome 6:119,540,965-119,712,625 | 6 |
| MAN1A1 |  | Chromosome 6:119,540,965-119,712,625 | 6 |
| EDN1 |  | Chromosome 6:12,398,582-12,405,413 | 6 |
| EDN1 |  | Chromosome 6:12,398,582-12,405,413 | 6 |
| PHACTR1 |  | Chromosome 6:12,825,819-13,396,624 | 6 |
| NKAIN2 |  | Chromosome 6:124,166,768-125,188,502 | 6 |
| NCOA7 |  | Chromosome 6:126,144,000-126,293,950 | 6 |
| C6orf174,KIAA0408 |  | Chromosome 6:127,813,023-127,879,540 | 6 |
| PTPRK |  | Chromosome 6:128,331,625-128,883,453 | 6 |
| PTPRK |  | Chromosome 6:128,331,625-128,883,453 | 6 |
| PTPRK |  | Chromosome 6:128,331,625-128,883,453 | 6 |
| PTPRK |  | Chromosome 6:128,331,625-128,883,453 | 6 |
| PTPRK |  | Chromosome 6:128,331,625-128,883,453 | 6 |
| PTPRK |  | Chromosome 6:128,331,625-128,883,453 | 6 |
| LAMA2 |  | Chromosome 6:129,246,035-129,879,407 | 6 |
| LAMA2 |  | Chromosome 6:129,246,035-129,879,407 | 6 |
| LAMA2 |  | Chromosome 6:129,246,035-129,879,407 | 6 |
| LAMA2 |  | Chromosome 6:129,246,035-129,879,407 | 6 |
| LAMA2 |  | Chromosome 6:129,246,035-129,879,407 | 6 |
| LAMA2 |  | Chromosome 6:129,246,035-129,879,407 | 6 |
| LAMA2 |  | Chromosome 6:129,246,035-129,879,407 | 6 |
| EPB41L2 |  | Chromosome 6:131,202,180-131,426,017 | 6 |
| EPB41L2 |  | Chromosome 6:131,202,180-131,426,017 | 6 |
| AKAP7 |  | Chromosome 6:131,508,154-131,646,366 | 6 |
| AKAP7 |  | Chromosome 6:131,508,154-131,646,366 | 6 |
| ARG1 | Arginase 1 | Chromosome 6:131,935,977-131,947,165 | 6 |
| ARG1 | Arginase 1 | Chromosome 6:131,935,977-131,947,165 | 6 |
| ARG1 | Arginase 1 | Chromosome 6:131,935,977-131,947,165 | 6 |
| ENPP3 | CD203c | Chromosome 6:132,000,135-132,110,243 | 6 |
| ENPP3 | CD203c | Chromosome 6:132,000,135-132,110,243 | 6 |
| CTGF |  | Chromosome 6:132,310,199-132,314,206 | 6 |
| CTGF |  | Chromosome 6:132,310,199-132,314,206 | 6 |
| MOXD1 |  | Chromosome 6:132,658,887-132,764,357 | 6 |
| MOXD1 |  | Chromosome 6:132,658,887-132,764,357 | 6 |
| VNN1 |  | Chromosome 6:133,044,422-133,076,881 | 6 |
| VNN1 |  | Chromosome 6:133,044,422-133,076,881 | 6 |
| SGK |  | Chromosome 6:134,532,081-134,680,889 | 6 |
| SGK |  | Chromosome 6:134,532,081-134,680,889 | 6 |
| SGK |  | Chromosome 6:134,532,081-134,680,889 | 6 |


| MAP7 |  | Chromosome 6:136,705,565-136,913,485 | 6 |
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| MAP3K5 |  | Chromosome 6:136,919,878-137,155,349 | 6 |
| MAP3K5 |  | Chromosome 6:136,919,878-137,155,349 | 6 |
| MAP3K5 |  | Chromosome 6:136,919,878-137,155,349 | 6 |
| MAP3K5 |  | Chromosome 6:136,919,878-137,155,349 | 6 |
| IL20RA |  | Chromosome 6:137,362,801-137,407,991 | 6 |
| IL20RA |  | Chromosome 6:137,362,801-137,407,991 | 6 |
| IL22RA2 |  | Chromosome 6:137,506,650-137,536,478 | 6 |
| TNFAIP3 |  | Chromosome 6:138,230,274-138,246,142) | 6 |
| TNFAIP3 |  | Chromosome 6:138,230,274-138,246,142) | 6 |
| C6orf91 |  | Chromosome 6:139,158,950-139,266,900 | 6 |
| C6orf91 |  | Chromosome 6:139,158,950-139,266,900 | 6 |
| HECA |  | Chromosome 6:139,497,942-139,543,639 | 6 |
| HECA |  | Chromosome 6:139,497,942-139,543,639 | 6 |
| CITED2 | p300 | Chromosome 6:139,735,089-139,737,478 | 6 |
| CD83 |  | Chromosome 6:14,225,715-14,245,128 | 6 |
| CD83 |  | Chromosome 6:14,225,715-14,245,128 | 6 |
| HIVEP2 | Schnurri 2 | Chromosome 6:143,114,297-143,308,031 | 6 |
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| HIVEP2 | Schnurri 2 | Chromosome 6:143,114,297-143,308,031 | 6 |
| HIVEP2 | Schnurri 2 | Chromosome 6:143,114,297-143,308,031 | 6 |
| ADAT2 |  | Chromosome 6:143,788,765-143,813,517 | 6 |
| ADAT2 |  | Chromosome 6:143,788,765-143,813,517 | 6 |
| PHACTR2 |  | Chromosome 6:143,971,010-144,194,014 | 6 |
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| PHACTR2 |  | Chromosome 6:143,971,010-144,194,014 | 6 |
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| STX11 | Syntaxin 11 | Chromosome 6:144,513,356-144,551,200 | 6 |
| STX11 | Syntaxin 11 | Chromosome 6:144,513,356-144,551,200 | 6 |
| UTRN |  | Chromosome 6:144,654,566-145,215,863 | 6 |
| STXBP5 |  | Chromosome 6:147,566,565-147,748,588 | 6 |
| STXBP5 |  | Chromosome 6:147,566,565-147,748,588 | 6 |
| STXBP5 |  | Chromosome 6:147,566,565-147,748,588 | 6 |
| FLJ43763 |  | Chromosome 6:148,313-151,392 | 6 |
| FLJ43763 |  | Chromosome 6:148,313-151,392 | 6 |
| MAP3K7IP2 | TAB2 | Chromosome 6:149,680,756-149,774,442 | 6 |
| MAP3K7IP2 | TAB2 | Chromosome 6:149,680,756-149,774,442 | 6 |
| MAP3K7IP2 | TAB2 | Chromosome 6:149,680,756-149,774,442 | 6 |
| MAP3K7IP2 | TAB2 | Chromosome 6:149,680,756-149,774,442 | 6 |
| MAP3K7IP2 | TAB2 | Chromosome 6:149,680,756-149,774,442 | 6 |
| MAP3K7IP2 | TAB2 | Chromosome 6:149,680,756-149,774,442 | 6 |
| PPIL4 |  | Chromosome 6:149,867,324-149,908,864 | 6 |
| PCMT1 |  | Chromosome 6:150,112,273-150,174,249 | 6 |
| RAET1E | ULBP4 | Chromosome 6:150,251,294-150,253,863 | 6 |
| ULBP2 |  | Chromosome 6:150,304,829-150,312,064 | 6 |
| AKAP12 |  | Chromosome 6:151,603,202-151,719,602 | 6 |
| AKAP12 |  | Chromosome 6:151,603,202-151,719,602 | 6 |
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| ESR1 | Estrogen receptor alpha, | Chromosome 6:152,170,379-152,466,099 | 6 |
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| SYNE1 |  | Chromosome 6:152,484,515-153,000,227 | 6 |
| SYNE1 |  | Chromosome 6:152,484,515-153,000,227 | 6 |
| VIP |  | Chromosome 6:153,113,626-153,122,593 | 6 |


| OPMR1 | MOR | Chromosome 6:154,402,136-154,609,693 | 6 |
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| TFB1M |  | Chromosome 6:155,620,488-155,686,932 | 6 |
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| NOX3 |  | Chromosome 6:155,758,194-155,818,729 | 6 |
| NOX3 |  | Chromosome 6:155,758,194-155,818,729 | 6 |
| ARID1B |  | Chromosome 6:157,140,756-157,572,094 | 6 |
| ARID1B |  | Chromosome 6:157,140,756-157,572,094 | 6 |
| SNX9 |  | Chromosome 6:158,164,282-158,286,097 | 6 |
| SNX9 |  | Chromosome 6:158,164,282-158,286,097 | 6 |
| GTF2H5 |  | Chromosome 6:158,509,372-158,535,008 | 6 |
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| GTF2H5 |  | Chromosome 6:158,509,372-158,535,008 | 6 |
| ATXN1 |  | Chromosome 6:16,407,322-16,869,700 | 6 |
| SOD2 | IP01 | Chromosome 6:160,020,138-160,034,343 | 6 |
| WTAP |  | Chromosome 6:160,066,607-160,097,341 | 6 |
| IGF2R | CD222 | Chromosome 6:160,310,121-160,447,573 | 6 |
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| IGF2R | CD222 | Chromosome 6:160,310,121-160,447,573 | 6 |
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| SLC22A1 |  | Chromosome 6:160,462,853-160,499,740 | 6 |
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| PARK2 |  | Chromosome 6:161,688,442-163,068,793 | 6 |
| PACRG |  | Chromosome 6:163,068,154-163,656,514 | 6 |
| PDE10A |  | Chromosome 6:165,660,766-165,995,578 | 6 |
| RPS6KA2 |  | Chromosome 6:166,742,844-167,195,761 | 6 |
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| CCR6 | CD196 | Chromosome 6:167,332,660-167,473,174 | 6 |
| C6orf123 |  | Chromosome 6:167,928,066-167,940,388 | 6 |
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| WRNIP1 |  | Chromosome 6:2,710,665-2,731,926 | 6 |
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| CDKAL1 |  | Chromosome 6:20,642,667-21,340,614 | 6 |
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| PRL | Prolactin | Chromosome 6:22,395,459-22,405,709 | 6 |
| HDGFL1 |  | Chromosome 6:22,677,657-22,679,871 | 6 |
| HDGFL1 |  | Chromosome 6:22,677,657-22,679,871 | 6 |
| DUSP22 |  | Chromosome 6:237,053-296,355 | 6 |
| DUSP22 |  | Chromosome 6:237,053-296,355 | 6 |
| TTRAP |  | Chromosome 6:24,758,184-24,775,240 | 6 |
| TTRAP |  | Chromosome 6:24,758,184-24,775,240 | 6 |
| LRRC16A |  | Chromosome 6:25,387,285-25,728,737 | 6 |
| HFE |  | Chromosome 6:26,195,427-26,205,038 | 6 |
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| BTN3A1 | CD277 | Chromosome 6:26,510,460-26,523,445 | 6 |
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| PRSS16 | thymus serine protease | Chromosome 6:27,323,487-27,332,327 | 6 |
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| ZNF192 |  | Chromosome 6:28,217,695-28,233,215 | 6 |
| ZNF193 |  | Chromosome 6:28,301,046-28,309,239 | 6 |
| FAT10 |  | Chromosome 6:29,631,368-29,778,041 | 6 |
| UBD | Ubiquitin | Chromosome 6:29,631,368-29,778,041 | 6 |
| MOG |  | Chromosome 6:29,732,755-29,748,128 | 6 |
| HLA-F |  | Chromosome 6:29,798,531-29,803,052 | 6 |
| HLA-F |  | Chromosome 6:29,799,096-29,803,052 | 6 |
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| HLA-F |  | Chromosome 6:29,799,096-29,803,052 | 6 |
| HLA-F |  | Chromosome 6:29,799,096-29,803,052 | 6 |
| RPL23AP1 |  | Chromosome 6:29,802,425-29,802,895 | 6 |
| HLA-A |  | Chromosome 6:29,902,723-30,021,633 | 6 |
| RIPK1 |  | Chromosome 6:3,009,212-3,060,420 | 6 |
| HLA-A | M6S204 | Chromosome 6:30,018,310-30,021,633 | 6 |
| HLA-A | M6S213 | Chromosome 6:30,018,310-30,021,633 | 6 |
| HLA-A |  | Chromosome 6:30,018,310-30,021,633 | 6 |
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| HLA-A |  | Chromosome 6:30,018,310-30,021,633 | 6 |
| HCG4P4 |  | Chromosome 6:30,030,962-30,031,390 | 6 |
| ETF1P1 |  | Chromosome 6:30,107,469-30,109,633 | 6 |
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| TRIM10 |  | Chromosome 6:30,227,701-30,236,690 | 6 |
| RPP21 |  | Chromosome 6:30,420,877-30,422,649 | 6 |
| RANP1 |  | Chromosome 6:30,561,651-30,562,700 | 6 |
| HLA-E |  | Chromosome 6:30,565,250-30,569,077 | 6 |
| HLA-E |  | Chromosome 6:30,565,250-30,569,077 | 6 |
| GNL1 |  | Chromosome 6:30,621,633-30,632,987 | 6 |
| MDC1 |  | Chromosome 6:30,775,563-30,793,645 | 6 |
| hcg-2038200 |  | Chromosome 6:30,888,622-30,906,415 | $\underline{6}$ |
| GTF2H4 |  | Chromosome 6:30,983,956-30,989,859 | 6 |
| C6orf205 |  | Chromosome 6:31,059,474-31,065,654 | 6 |
| HLABC-CA |  | Chromosome 6:31,344,505-31,432,935 | 6 |
| HLA-C | MICB-CA | Chromosome 6:31,344,505-31,432,935 | 6 |
| HLA-C |  | Chromosome 6:31,344,505-31,432,935 | 6 |
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| HLA-C |  | Chromosome 6:31,344,505-31,432,935 | 6 |
| HLA-B |  | Chromosome 6:31,429,628-31,432,914 | 6 |
| HLA-B |  | Chromosome 6:31,429,628-31,432,914 | 6 |
| HLA-B |  | Chromosome 6:31,429,628-31,432,914 | 6 |
| MICA |  | Chromosome 6:31,475,540-31,491,069 | 6 |
| HCP5 |  | Chromosome 6:31,538,938-31,541,565 | 6 |
| MCCD1 | LOC401250 | Chromosome 6:31,604,718-31,605,987 | 6 |
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| LTA | TNFb | Chromosome 6:31,648,042-31,650,080 | 6 |
| LTA | TNFb | Chromosome 6:31,648,042-31,650,080 | 6 |
| LTA |  | Chromosome 6:31,648,072-31,650,077 | 6 |
| TNF |  | Chromosome 6:31,651,329-31,654,091 | 6 |
| AIF1 |  | Chromosome 6:31,690,984-31,692,781 | 6 |
| BAT2, BAT2 GT, BAT2CA |  | Chromosome 6:31,696,429-31,713,533 | 6 |
| LY6G5C |  | Chromosome 6:31,752,440-31,759,796 | 6 |
| HCG22 |  | Chromosome 6:31129963-31135632 | 6 |


| C2 |  | Chromosome 6:32,003,473-32,021,428 | 6 |
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| TNXB |  | Chromosome 6:32,084,175-32,185,131 | 6 |
| FKBPL |  | Chromosome 6:32,204,462-32,206,045 | 6 |
| NOTCH4 |  | Chromosome 6:32,266,521-32,299,822 | 6 |
| GPSM3 |  | Chromosome 6:32,266,521-32,299,822 | 6 |
| NOTCH4 |  | Chromosome 6:32,270,598-32,299,822 | 6 |
| C6orf10 | TNFa/b | Chromosome 6:32,368,453-32,460,310 | 6 |
| C6orf10 |  | Chromosome 6:32,368,464-32,447,662 | 6 |
| HLA-DRA |  | Chromosome 6:32,515,597-32,520,943 | 6 |
| HLA-DRA | DRA_CA, HLA-DRB1 | Chromosome 6:32,515,625-32,520,801 | 6 |
| HLA-DRB1 |  | Chromosome 6:32,654,524-32,665,603 | 6 |
| HLA-DQA1 |  | Chromosome 6:32,713,112-32,719,407 | 6 |
| HLA-DPB1 |  | Chromosome 6:32,735,222-32,754,296 | 6 |
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| HLA-DQB1 |  | Chromosome 6:32,735,225-32,742,572 | 6 |
| HLA-DQB2 |  | Chromosome 6:32,831,445-32,839,446 | 6 |
| HLA-DOB |  | Chromosome 6:32,888,518-32,892,803 | 6 |
| TAP2 |  | Chromosome 6:32,897,588-32,914,525 | 6 |
| BRD2 |  | Chromosome 6:33,044,415-33,057,075 | 6 |
| HLA-DPB2 |  | Chromosome 6:33,188,206-33,204,868 | 6 |
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| COL11A2 |  | Chromosome 6:33,238,447-33,268,223 | 6 |
| SLC39A7 |  | Chromosome 6:33,276,631-33,280,192 | 6 |
| VPS52 |  | Chromosome 6:33,326,027-33,347,640 | 6 |
| TAPBP | Tapasin | Chromosome 6:33,375,449-33,390,142 | 6 |
| BAK1 |  | Chromosome 6:33,648,307-33,655,997 | 6 |
| ITPR3 |  | Chromosome 6:33,696,500-33,772,329 | 6 |
| PACSIN1 |  | Chromosome 6:34,541,883-34,610,984 | 6 |
| PACSIN1 |  | Chromosome 6:34,541,883-34,610,984 | 6 |
| PPARD | PPARbeta, NR1C2 | Chromosome 6:35,418,313-35,503,933 | 6 |
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| FKBP5 | Calcineurin, Lymphoid eff | Chromosome 6:35,649,345-35,804,338 | 6 |
| SRPK1 | SFRS | Chromosome 6:35,908,789-35,996,942 | 6 |
| SRPK1 | SFRS | Chromosome 6:35,908,789-35,996,942 | 6 |
| MAPK14 | p38MAPK | Chromosome 6:36,103,551-36,186,513 | 6 |
| MAPK13 |  | Chromosome 6:36,129,769-36,215,820 | 6 |
| MAPK13 |  | Chromosome 6:36,129,769-36,215,820 | 6 |
| STK38 |  | Chromosome 6:36,569,647-36,623,234 | 6 |
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| CDKN1A |  | Chromosome 6:36,754,413-36,763,094 | 6 |
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| PPIL1 |  | Chromosome 6:36,930,581-36,950,778 | 6 |
| PPIL1 |  | Chromosome 6:36,930,581-36,950,778 | 6 |
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| ZFAND3 |  | Chromosome 6:37,895,285-38,230,375 | 6 |
| BTBD9 |  | Chromosome 6:38,250,711-38,673,848 | 6 |
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| DNAH8 |  | Chromosome 6:38,792,313-39,106,545 | 6 |
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| DNAH8 |  | Chromosome 6:38,792,313-39,106,545 | 6 |
| DNAH8 |  | Chromosome 6:38,792,313-39,106,545 | 6 |
| DAAM2 |  | Chromosome 6:39,868,120-39,980,622 | 6 |


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| TREML3 |  | Chromosome 6:41,284,270-41,298,360 | 6 |
| NCR2 | CD336, NKp44 | Chromosome 6:41,411,505-41,426,603 | 6 |
| FOXP4 |  | Chromosome 6:41,622,142-41,678,100 | 6 |
| FOXP4 |  | Chromosome 6:41,622,142-41,678,100 | 6 |
| FOXP4 |  | Chromosome 6:41,622,142-41,678,100 | 6 |
| CCND3 |  | Chromosome 6:42,010,649-42,124,404 | 6 |
| CCND3 |  | Chromosome 6:42,010,649-42,124,404 | 6 |
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| TRERF1 |  | Chromosome 6:42,300,647-42,527,767 | 6 |
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| RUNX2 |  | Chromosome 6:45,404,032-45,626,797 | 6 |
| CYP39A1 |  | Chromosome 6:46,625,404-46,728,482 | 6 |
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| CYP39A1 |  | Chromosome 6:46,625,404-46,728,482 | 6 |
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| TNFRSF21 |  | Chromosome 6:47,307,227-47,385,639 | 6 |
| CD2AP |  | Chromosome 6:47,553,899-47,702,620 | 6 |
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| RHAG | CD241 | Chromosome 6:49,680,830-49,712,511 | 6 |
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| DFB110 |  | Chromosome 6:50,084,810-50,097,607 | 6 |
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| IL17F |  | Chromosome 6:52,209,438-52,217,257 | 6 |
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| EFHC1 |  | Chromosome 6:52,392,953-52,468,540 | 6 |
| TRAM2 |  | Chromosome 6:52,470,159-52,549,821 | 6 |
| GSTA1 | Gluthation-S-transferase | Chromosome 6:52,764,183-52,776,616 | 6 |
| GCLC |  | Chromosome 6:53,470,098-53,517,790 | 6 |
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| GFRAL |  | Chromosome 6:55,300,226-55,375,250 | 6 |
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| BMP5 | BMP5 | Chromosome 6:55,726,402-55,848,334 | 6 |


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| ZNF451 |  | Chromosome 6:57,019,470-57,143,057 | 6 |
| BAG2 |  | Chromosome 6:57,145,083-57,157,694 | 6 |
| LY86 | MD1 | Chromosome 6:6,533,340-6,600,215 | 6 |
| LY86 | MD1 | Chromosome 6:6,533,340-6,600,215 | 6 |
| LY86 | MD1 | Chromosome 6:6,533,340-6,600,215 | 6 |
| KHDRBS2 |  | Chromosome 6:62,447,824-63,054,091 | 6 |
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| FKBP1AC |  | Chromosome 6:63,964,538-63,980,909 | 6 |
| PTP4A1 |  | Chromosome 6:64,339,879-64,351,448 | 6 |
| EYS |  | Chromosome 6:66,095,895-66,473,839 | 6 |
| BAI3 |  | Chromosome 6:69,401,980-70,156,124 | 6 |
| DSP |  | Chromosome 6:7,486,869-7,531,945 | 6 |
| DSP |  | Chromosome 6:7,486,869-7,531,945 | 6 |
| DSP |  | Chromosome 6:7,486,869-7,531,945 | 6 |
| BMP6 | BMP6 | Chromosome 6:7,672,009-7,826,752 | 6 |
| BMP6 | BMP6 | Chromosome 6:7,672,009-7,826,752 | 6 |
| KCNQ5 |  | Chromosome 6:73,388,241-73,965,295 | 6 |
| DDX43 |  | Chromosome 6:74,161,192-74,184,013 | 6 |
| DDX43 |  | Chromosome 6:74,161,192-74,184,013 | 6 |
| EEF1A1 | EF-1 alpha | Chromosome 6:74,282,194-74,288,344 | 6 |
| EEF1A1 | EF-1 alpha | Chromosome 6:74,282,194-74,288,344 | 6 |
| CD109 |  | Chromosome 6:74,462,548-74,591,509 | 6 |
| CD109 |  | Chromosome 6:74,462,548-74,591,509 | 6 |
| CD109 |  | Chromosome 6:74,462,548-74,591,509 | 6 |
| HTR1B | 5-HTR1B | Chromosome 6:78,228,641-78,229,900 | 6 |
| HTR1B | 5-HTR1B | Chromosome 6:78,228,641-78,229,900 | 6 |
| IRAK1BP1 |  | Chromosome 6:79,633,908-79,665,039 | 6 |
| IRAK1BP1 |  | Chromosome 6:79,633,908-79,665,039 | 6 |
| HMGN3 |  | Chromosome 6:79,967,681-80,001,174 | 6 |
| HMGN3 |  | Chromosome 6:79,967,681-80,001,174 | 6 |
| BCKDHB |  | Chromosome 6:80,873,083-81,112,706 | 6 |
| BCKDHB |  | Chromosome 6:80,873,083-81,112,706 | 6 |
| BCKDHB |  | Chromosome 6:80,873,083-81,112,706 | 6 |
| NT5E | CD73 | Chromosome 6:86,216,528-86,262,215 | 6 |
| NT5E | CD73 | Chromosome 6:86,216,528-86,262,215 | 6 |
| LOC643962 |  | Chromosome 6:87,597,028-87,709,921 | 6 |
| LOC643962 |  | Chromosome 6:87,597,028-87,709,921 | 6 |
| CGA | CGA | Chromosome 6:87,851,935-87,861,569 | 6 |
| OFCC1 |  | Chromosome 6:9,813,644-10,168,908 | 6 |
| ANKRD6 |  | Chromosome 6:90,199,616-90,400,123 | 6 |
| ANKRD6 |  | Chromosome 6:90,199,616-90,400,123 | 6 |
| ANKRD6 |  | Chromosome 6:90,199,616-90,400,123 | 6 |
| CASP8AP2 | FLASH | Chromosome 6:90,596,349-90,640,876 | 6 |
| MAP3K7 | TAK1 | Chromosome 6:91,280,013-91,353,628 | 6 |
| MAP3K7 | TAK1 | Chromosome 6:91,280,013-91,353,628 | 6 |
| EPHA7 |  | Chromosome 6:94,007,860-94,185,993 | 6 |
| EPHA7 |  | Chromosome 6:94,007,860-94,185,993 | 6 |
| KLHL32 |  | Chromosome 6:97,479,217-97,695,351 | 6 |
| KLHL32 |  | Chromosome 6:97,479,217-97,695,351 | 6 |
| KLHL32 |  | Chromosome 6:97,479,217-97,695,351 | 6 |


| EPO |  | Chromosome 7:100,156,359-100,159,259 | 7 |
| :---: | :---: | :---: | :---: |
| ACHE |  | Chromosome 7:100,325,551-100,331,651 | 7 |
| ACHE |  | Chromosome 7:100,325,551-100,331,651 | 7 |
| ACHE |  | Chromosome 7:100,325,551-100,331,651 | 7 |
| SERPINE1 | Plasminogen activator inh | Chromosome 7:100,557,172-100,569,026 | 7 |
| SERPINE1 | Plasminogen activator inh | Chromosome 7:100,557,172-100,569,026 | 7 |
| APS | SH2 adaptor protein | Chromosome 7:101,715,172-101,748,898 | 7 |
| ALKBH4 |  | Chromosome 7:101,883,690-101,892,293 | 7 |
| ALKBH4 |  | Chromosome 7:101,883,690-101,892,293 | 7 |
| RELN |  | Chromosome 7:102,899,473-103,417,199 | 7 |
| RELN |  | Chromosome 7:102,899,473-103,417,199 | 7 |
| RELN |  | Chromosome 7:102,899,473-103,417,199 | 7 |
| RELN |  | Chromosome 7:102,899,473-103,417,199 | 7 |
| RELN |  | Chromosome 7:102,899,473-103,417,199 | 7 |
| RELN |  | Chromosome 7:102,899,473-103,417,199 | 7 |
| SRPK2 | SFRS | Chromosome 7:104,544,059-104,816,577 | 7 |
| SRPK2 | SFRS | Chromosome 7:104,544,059-104,816,577 | 7 |
| SRPK2 | SFRS | Chromosome 7:104,544,059-104,816,577 | 7 |
| SRPK2 | SFRS | Chromosome 7:104,544,059-104,816,577 | 7 |
| SRPK2 | SFRS | Chromosome 7:104,544,059-104,816,577 | 7 |
| PBEF1 | visfatin | Chromosome 7:105,677,892-105,712,603 | 7 |
| PBEF1 | visfatin | Chromosome 7:105,677,892-105,712,603 | 7 |
| PBEF1 | visfatin | Chromosome 7:105,677,892-105,712,603 | 7 |
| PIK3CG |  | Chromosome 7:106,292,977-106,334,828 | 7 |
| PIK3CG |  | Chromosome 7:106,292,977-106,334,828 | 7 |
| LAMB1 |  | Chromosome 7:107,351,499-107,431,040 | 7 |
| LAMB1 |  | Chromosome 7:107,351,499-107,431,040 | 7 |
| LAMB4 |  | Chromosome 7:107,451,232-107,558,036 | 7 |
| IFRD1 |  | Chromosome 7:111,850,462-111,903,483 | 7 |
| IFRD1 |  | Chromosome 7:111,850,462-111,903,483 | 7 |
| FOXP2 |  | Chromosome 7:113,842,288-114,117,391 | 7 |
| FOXP2 |  | Chromosome 7:113,842,288-114,117,391 | 7 |
| FOXP2 |  | Chromosome 7:113,842,288-114,117,391 | 7 |
| FOXP2 |  | Chromosome 7:113,842,288-114,117,391 | 7 |
| CAV1 | Caveolin-1 | Chromosome 7:115,952,075-115,988,466 | 7 |
| CAV1 | Caveolin-1 | Chromosome 7:115,952,075-115,988,466 | 7 |
| CAV1 | Caveolin-1 | Chromosome 7:115,952,075-115,988,466 | 7 |
| MET |  | Chromosome 7:116,099,695-116,225,676 | 7 |
| MET |  | Chromosome 7:116,099,695-116,225,676 | 7 |
| MET |  | Chromosome 7:116,099,695-116,225,676 | 7 |
| PTPRZ1 |  | Chromosome 7:121,300,395-121,489,326 | 7 |
| PTPRZ1 |  | Chromosome 7:121,300,395-121,489,326 | 7 |
| PTPRZ1 |  | Chromosome 7:121,300,395-121,489,326 | 7 |
| PTPRZ1 |  | Chromosome 7:121,300,395-121,489,326 | 7 |
| POT1 |  | Chromosome 7:124,250,549-124,357,110 | 7 |
| POT1 |  | Chromosome 7:124,250,549-124,357,110 | 7 |
| POT1 |  | Chromosome 7:124,250,549-124,357,110 | 7 |
| LEP |  | Chromosome 7:127,668,567-127,684,917 | 7 |
| LEP |  | Chromosome 7:127,668,567-127,684,917 | 7 |
| IRF5 |  | Chromosome 7:128,365,230-128,377,325 | 7 |
| IRF5 |  | Chromosome 7:128,365,230-128,377,325 | 7 |
| SMO |  | Chromosome 7:128,615,949-128,640,622 | 7 |
| SMO |  | Chromosome 7:128,615,949-128,640,622 | 7 |
| NRF1 | NRF1 | Chromosome 7:129,038,791-129,184,158 | 7 |
| NRF1 | NRF1 | Chromosome 7:129,038,791-129,184,158 | 7 |
| NRF1 | NRF1 | Chromosome 7:129,038,791-129,184,158 | 7 |


| ETV1 |  | Chromosome 7:13,897,379-13,995,289 | 7 |
| :---: | :---: | :---: | :---: |
| ETV1 |  | Chromosome 7:13,897,379-13,995,289 | 7 |
| ETV1 |  | Chromosome 7:13,897,379-13,995,289 | 7 |
| ETV1 |  | Chromosome 7:13,897,379-13,995,289 | 7 |
| MKLN1 | Mkln1 | Chromosome 7:130,663,175-130,831,931 | 7 |
| MKLN1 | Mkln1 | Chromosome 7:130,663,175-130,831,931 | 7 |
| MKLN1 | Mkln1 | Chromosome 7:130,663,175-130,831,931 | 7 |
| MKLN1 | Mkln1 | Chromosome 7:130,663,175-130,831,931 | 7 |
| KLRG2 |  | Chromosome 7:138,786,805-138,818,998 | 7 |
| TBXAS1 |  | Chromosome 7:139,124,668-139,366,560 | 7 |
| TBXAS1 |  | Chromosome 7:139,124,668-139,366,560 | 7 |
| TBXAS1 |  | Chromosome 7:139,124,668-139,366,560 | 7 |
| TBXAS1 |  | Chromosome 7:139,124,668-139,366,560 | 7 |
| TBXAS1 |  | Chromosome 7:139,124,668-139,366,560 | 7 |
| TBXAS1 |  | Chromosome 7:139,124,668-139,366,560 | 7 |
| RAB19 | Rab19 | Chromosome 7:139,753,916-139,772,419 | 7 |
| CLEC5A | MDL1 | Chromosome 7:141,273,626-141,293,252 | 7 |
| CLEC5A | MDL1 | Chromosome 7:141,273,626-141,293,252 | 7 |
| TRB@ | TCR | Chromosome 7:141,645,314-142,221,097 | 7 |
| TRB@ | TCR | Chromosome 7:141,645,314-142,221,097 | 7 |
| TRB@ | TCR | Chromosome 7:141,645,314-142,221,097 | 7 |
| TRB@ | TCR | Chromosome 7:141,645,314-142,221,097 | 7 |
| TRB@ | TCR | Chromosome 7:141,645,314-142,221,097 | 7 |
| TRB@ | TCR | Chromosome 7:141,645,314-142,221,097 | 7 |
| TRB@ | TCR | Chromosome 7:141,645,314-142,221,097 | 7 |
| KEL | CD238 | Chromosome 7:142,348,323-142,369,625 | 7 |
| KEL | CD238 | Chromosome 7:142,348,323-142,369,625 | 7 |
| CASP2 |  | Chromosome 7:142,695,524-142,714,907 | 7 |
| EPHA1 | Ephrin R | Chromosome 7:142,798,327-142,816,107 | 7 |
| EZH2 |  | Chromosome 7:148,135,408-148,212,347 | 7 |
| EZH2 |  | Chromosome 7:148,135,408-148,212,347 | 7 |
| EZH2 |  | Chromosome 7:148,135,408-148,212,347 | 7 |
| RARRES2 | chemerin | Chromosome 7:149,666,351-149,669,696 | 7 |
| RARRES2 | chemerin | Chromosome 7:149,666,351-149,669,696 | 7 |
| NOS3 | NO synthase | Chromosome 7:150,319,080-150,342,609 | 7 |
| NOS3 | NO synthase | Chromosome 7:150,319,080-150,342,609 | 7 |
| CDK5 |  | Chromosome 7:150,381,832-150,385,929 | 7 |
| SHH | sonic Hh | Chromosome 7:155,288,319-155,297,728 | 7 |
| SHH | sonic Hh | Chromosome 7:155,288,319-155,297,728 | 7 |
| PTPRN2 | PTPRP | Chromosome 7:157,024,516-158,073,179 | 7 |
| PTPRN2 | PTPRP | Chromosome 7:157,024,516-158,073,179 | 7 |
| PTPRN2 | PTPRP | Chromosome 7:157,024,516-158,073,179 | 7 |
| PTPRN2 | PTPRP | Chromosome 7:157,024,516-158,073,179 | 7 |
| PTPRN2 | PTPRP | Chromosome 7:157,024,516-158,073,179 | 7 |
| PTPRN2 | PTPRP | Chromosome 7:157,024,516-158,073,179 | 7 |
| PTPRN2 | PTPRP | Chromosome 7:157,024,516-158,073,179 | 7 |
| HDAC9 |  | Chromosome 7:18,501,894-19,003,518 | 7 |
| HDAC9 |  | Chromosome 7:18,501,894-19,003,518 | 7 |
| HDAC9 |  | Chromosome 7:18,501,894-19,003,518 | 7 |
| HDAC9 |  | Chromosome 7:18,501,894-19,003,518 | 7 |
| HDAC9 |  | Chromosome 7:18,501,894-19,003,518 | 7 |
| HDAC9 |  | Chromosome 7:18,501,894-19,003,518 | 7 |
| HDAC9 |  | Chromosome 7:18,501,894-19,003,518 | 7 |
| HDAC9 |  | Chromosome 7:18,501,894-19,003,518 | 7 |
| HDAC9 |  | Chromosome 7:18,501,894-19,003,518 | 7 |
| HDAC9 |  | Chromosome 7:18,501,894-19,003,518 | 7 |


| TWIST1 | Twist1 | Chromosome 7:19,121,616-19,123,820 | 7 |
| :---: | :---: | :---: | :---: |
| TWIST1 | Twist1 | Chromosome 7:19,121,616-19,123,820 | 7 |
| CARD11 | CARMA1 | Chromosome 7:2,912,308-3,050,025 | 7 |
| CARD11 | CARMA1 | Chromosome 7:2,912,308-3,050,025 | 7 |
| ITGB8 |  | Chromosome 7:20,337,250-20,421,907 | 7 |
| ITGB8 |  | Chromosome 7:20,337,250-20,421,907 | 7 |
| ITGB8 |  | Chromosome 7:20,337,250-20,421,907 | 7 |
| IL6 |  | Chromosome 7:22,732,028-22,738,141 | 7 |
| IL6 |  | Chromosome 7:22,732,028-22,738,141 | 7 |
| GPNMB |  | Chromosome 7:23,252,841-23,281,254 | 7 |
| NPY |  | Chromosome 7:24,290,332-24,298,002 | 7 |
| NPY |  | Chromosome 7:24,290,332-24,298,002 | 7 |
| CYCS |  | Chromosome 7:25,124,800-25,131,480 | 7 |
| HOXA5 | HOX cluster | Chromosome 7:27,147,521-27,149,812 | 7 |
| HOXA5 | HOX cluster | Chromosome 7:27,147,521-27,149,812 | 7 |
| NOD1 | CARD4 | Chromosome 7:30,430,672-30,484,833 | 7 |
| CRHR2 |  | Chromosome 7:30,658,725-30,706,244 | 7 |
| CRF2R | CRF2R | Chromosome 7:30,658,725-30,706,244 | 7 |
| AQP1 | Aquaporin 1 | Chromosome 7:30,917,993-30,931,656 | 7 |
| AQP1 | Aquaporin 1 | Chromosome 7:30,917,993-30,931,656 | 7 |
| FKBP9 |  | Chromosome 7:32,963,577-33,013,067 | 7 |
| FKBP9 |  | Chromosome 7:32,963,577-33,013,067 | 7 |
| TRG@ | TCR | Chromosome 7:38,246,150-38,374,181 | 7 |
| TRG@ | TCR | Chromosome 7:38,246,150-38,374,181 | 7 |
| TRG@ | TCR | Chromosome 7:38,246,150-38,374,181 | 7 |
| TRG@ | TCR | Chromosome 7:38,246,150-38,374,181 | 7 |
| FOXK1 |  | Chromosome 7:4,688,456-4,777,600 | 7 |
| FOXK1 |  | Chromosome 7:4,688,456-4,777,600 | 7 |
| INHBA | Inhibin B, Activin A | Chromosome 7:41,695,126-41,709,231 | 7 |
| INHBA | Inhibin B, Activin A | Chromosome 7:41,695,126-41,709,231 | 7 |
| GLI3 | Gli3 | Chromosome 7:41,970,196-42,241,712 | 7 |
| GLI3 | Gli3 | Chromosome 7:41,970,196-42,241,712 | 7 |
| GLI3 | Gli3 | Chromosome 7:41,970,196-42,241,712 | 7 |
| GLI3 | Gli3 | Chromosome 7:41,970,196-42,241,712 | 7 |
| STK17A |  | Chromosome 7:43,589,251-43,632,247 | 7 |
| STK17A |  | Chromosome 7:43,589,251-43,632,247 | 7 |
| STK17A |  | Chromosome 7:43,589,251-43,632,247 | 7 |
| GCK |  | Chromosome 7:44,150,395-44,195,563 | 7 |
| GCK |  | Chromosome 7:44,150,395-44,195,563 | 7 |
| DDX56 |  | Chromosome 7:44,571,928-44,581,175 | 7 |
| DDX56 |  | Chromosome 7:44,571,928-44,581,175 | 7 |
| PPIA |  | Chromosome 7:44,802,777-44,809,240 | 7 |
| PPIA |  | Chromosome 7:44,802,777-44,809,240 | 7 |
| MYO1G | HA-2 | Chromosome 7:44,968,786-44,985,203 | 7 |
| ACTB | F-actin | Chromosome 7:5,533,312-5,536,747 | 7 |
| IKZF1 | Ikaros | Chromosome 7:50,314,924-50,438,053 | 7 |
| GRB10 | Grb10/Grb1R | Chromosome 7:50,625,259-50,828,652 | 7 |
| GRB10 | Grb10/Grb1R | Chromosome 7:50,625,259-50,828,652 | 7 |
| GRB10 | Grb10/Grb1R | Chromosome 7:50,625,259-50,828,652 | 7 |
| EGFR |  | Chromosome 7:55,054,219-55,242,525 | 7 |
| FKBP9L |  | Chromosome 7:55,716,261-55,748,439 | 7 |
| RAC1 | Rac1 | Chromosome 7:6,380,651-6,410,123 | 7 |
| RAC1 | Rac1 | Chromosome 7:6,380,651-6,410,123 | 7 |
| GUSB | glucuronidase beta - MPS | Chromosome 7:65,063,110-65,084,635 | 7 |
| GUSB | glucuronidase beta - MPS | Chromosome 7:65,063,110-65,084,635 | 7 |
| RPA3 |  | Chromosome 7:7,643,100-7,724,763 | 7 |


| RPA3 |  | Chromosome 7:7,643,100-7,724,763 | 7 |
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| FZD9 | CD349 | Chromosome 7:72,486,045-72,488,386 | 7 |
| CLDN3 |  | Chromosome 7:72,821,263-72,822,536 | 7 |
| LAT2 | NTA, LAB, TFII-1, WBSC | Chromosome 7:73,262,023-73,282,100 | 7 |
| LAT2 | NTA, LAB, TFII-1, WBSC | Chromosome 7:73,262,023-73,282,100 | 7 |
| CLIP2 | CYLN2 | Chromosome 7:73,341,741-73,458,201 | 7 |
| CLIP2 | CYLN2 | Chromosome 7:73,341,741-73,458,201 | 7 |
| GTF2IRD1 | TFII-1 | Chromosome 7:73,506,056-73,654,853 | 7 |
| GTF2IRD1 | TFII-1 | Chromosome 7:73,506,056-73,654,853 | 7 |
| GTF2I |  | Chromosome 7:73,709,966-73,812,958 | 7 |
| NCF1 | neutrophil cytosolic factor | Chromosome 7:73,826,245-73,841,595 | 7 |
| GTF2IRD2 |  | Chromosome 7:73,848,420-73,905,777 | 7 |
| PMS2L3 | PMS2L9 | Chromosome 7:74,975,005-74,995,389 | 7 |
| PMS2L3 | PMS2L9 | Chromosome 7:74,975,005-74,995,389 | 7 |
| CCL26 |  | Chromosome 7:75,236,778-75,257,150 | 7 |
| CCL26 |  | Chromosome 7:75,236,778-75,257,150 | 7 |
| HSPB1 | HSP27/28 | Chromosome 7:75,769,859-75,771,548 | 7 |
| FGL2 |  | Chromosome 7:76,662,535-76,667,080 | 7 |
| FGL2 |  | Chromosome 7:76,662,535-76,667,080 | 7 |
| PTPN12 |  | Chromosome 7:77,004,351-77,107,324 | 7 |
| PTPN12 |  | Chromosome 7:77,004,351-77,107,324 | 7 |
| GNAI1 | Gi | Chromosome 7:79,602,076-79,686,661 | 7 |
| GNAI1 | Gi | Chromosome 7:79,602,076-79,686,661 | 7 |
| CD36 | SCARB3 | Chromosome 7:80,069,459-80,144,262 | 7 |
| CD36 | SCARB3 | Chromosome 7:80,069,459-80,144,262 | 7 |
| HGF | hepatocyte growth factor | Chromosome 7:81,166,258-81,237,388 | 7 |
| HGF | hepatocyte growth factor | Chromosome 7:81,166,258-81,237,388 | 7 |
| SEMA3E |  | Chromosome 7:82,831,158-83,116,260 | 7 |
| SEMA3E |  | Chromosome 7:82,831,158-83,116,260 | 7 |
| SEMA3E |  | Chromosome 7:82,831,158-83,116,260 | 7 |
| SEMA3E |  | Chromosome 7:82,831,158-83,116,260 | 7 |
| SEMA3E |  | Chromosome 7:82,831,158-83,116,260 | 7 |
| SEMA3E |  | Chromosome 7:82,831,158-83,116,260 | 7 |
| SEMA3E |  | Chromosome 7:82,831,158-83,116,260 | 7 |
| SEMA3E |  | Chromosome 7:82,831,158-83,116,260 | 7 |
| ABCB1 | CD243, MDR1 | Chromosome 7:86,970,884-87,180,500 | 7 |
| ABCB1 | CD243, MDR1 | Chromosome 7:86,970,884-87,180,500 | 7 |
| ABCB1 | CD243, MDR1 | Chromosome 7:86,970,884-87,180,500 | 7 |
| ABCB1 | CD243, MDR1 | Chromosome 7:86,970,884-87,180,500 | 7 |
| CYP51A1 |  | Chromosome 7:91,579,402-91,601,946 | 7 |
| CYP51A1 |  | Chromosome 7:91,579,402-91,601,946 | 7 |
| CYP51A1 |  | Chromosome 7:91,579,402-91,601,946 | 7 |
| CDK6 |  | Chromosome 7:92,072,171-92,301,148 | 7 |
| CDK6 |  | Chromosome 7:92,072,171-92,301,148 | 7 |
| CDK6 |  | Chromosome 7:92,072,171-92,301,148 | 7 |
| COL1A2 | collagen alpha 2 | Chromosome 7:93,861,809-93,898,480 | 7 |
| COL1A2 | collagen alpha 2 | Chromosome 7:93,861,809-93,898,480 | 7 |
| COL1A2 | collagen alpha 2 | Chromosome 7:93,861,809-93,898,480 | 7 |
| DSS1 |  | Chromosome 7:96,156,015-96,177,139 | 7 |
| DSS1 |  | Chromosome 7:96,156,015-96,177,139 | 7 |
| Tac1 | Substance P | Chromosome 7:97,199,311-97,207,720 | 7 |
| Tac1 | Substance P | Chromosome 7:97,199,311-97,207,720 | 7 |
| BAIAP2L1 |  | Chromosome 7:97,760,007-97,868,316 | 7 |
| BAIAP2L1 |  | Chromosome 7:97,760,007-97,868,316 | 7 |
| CYP2W1 |  | Chromosome 7:989,361-995,802 | 7 |
| CYP2W1 |  | Chromosome 7:989,361-995,802 | 7 |


| CYP3A3 | CYP3A4 | Chromosome 7:99,083,437-99,219,744 | 7 |
| :---: | :---: | :---: | :---: |
| CYP3A3 | CYP3A4 | Chromosome 7:99,083,437-99,219,744 | 7 |
| CYP3A43 |  | Chromosome 7:99,263,572-99,302,109 | 7 |
| ZNF3 |  | Chromosome 7:99,499,406-99,517,299 | 7 |
| PILB | PILRbeta | Chromosome 7:99,771,673-99,803,388 | 7 |
| PILB | PILRbeta | Chromosome 7:99,771,673-99,803,388 | 7 |
| DEFA1A3 |  | Chromosome 8 | 8 |
| DEFA1A3 |  | Chromosome 8 | 8 |
| PINX1 |  | Chromosome 8:10,659,883-10,734,796 | 8 |
| PINX1 |  | Chromosome 8:10,659,883-10,734,796 | 8 |
| PINX1 |  | Chromosome 8:10,659,883-10,734,796 | 8 |
| YWHA2 |  | Chromosome 8:102,000,090-102,034,745 | 8 |
| YWHA2 |  | Chromosome 8:102,000,090-102,034,745 | 8 |
| KLF10 | TIEG1 | Chromosome 8:103,730,188-103,737,128 | 8 |
| KLF10 | TIEG1 | Chromosome 8:103,730,188-103,737,128 | 8 |
| BAALC |  | Chromosome 8:104,222,097-104,311,709 | 8 |
| BAALC |  | Chromosome 8:104,222,097-104,311,709 | 8 |
| TM7SF4 | DC-HIL | Chromosome 8:105,421,228-105,438,092 | 8 |
| TM7SF4 | DC-HIL | Chromosome 8:105,421,228-105,438,092 | 8 |
| ANGPT1 | Angiopoietin 1 | Chromosome 8:108,330,886-108,579,459 | 8 |
| ANGPT1 | Angiopoietin 1 | Chromosome 8:108,330,886-108,579,459 | 8 |
| ANGPT1 | Angiopoietin 1 | Chromosome 8:108,330,886-108,579,459 | 8 |
| ANGPT1 | Angiopoietin 1 | Chromosome 8:108,330,886-108,579,459 | 8 |
| ANGPT1 | Angiopoietin 1 | Chromosome 8:108,330,886-108,579,459 | 8 |
| BLK |  | Chromosome 8:11,388,919-11,459,522 | 8 |
| BLK |  | Chromosome 8:11,388,919-11,459,522 | 8 |
| BLK |  | Chromosome 8:11,388,919-11,459,522 | 8 |
| GATA4 |  | Chromosome 8:11,599,122-11,654,920 | 8 |
| DFB137 |  | Chromosome 8:11,868,871-11,869,517 | 8 |
| TRHR | TRH-R | Chromosome 8:110,168,900-110,200,989 | 8 |
| TRHR | TRH-R | Chromosome 8:110,168,900-110,200,989 | 8 |
| DFB130 |  | Chromosome 8:12,212,843-12,220,196 | 8 |
| TNFRSF11B |  | Chromosome 8:120,004,977-120,033,492 | 8 |
| TNFRSF11B |  | Chromosome 8:120,004,977-120,033,492 | 8 |
| NOV | CCN3 | Chromosome 8:120,497,882-120,505,776 | 8 |
| NOV | CCN3 | Chromosome 8:120,497,882-120,505,776 | 8 |
| COL14A1 | undulin | Chromosome 8:121,206,533-121,453,454 | 8 |
| COL14A1 | undulin | Chromosome 8:121,206,533-121,453,454 | 8 |
| COL14A1 | undulin | Chromosome 8:121,206,533-121,453,454 | 8 |
| MYC | c-myc | Chromosome 8:128,817,498-128,822,856 | 8 |
| MYC | c-myc | Chromosome 8:128,817,498-128,822,856 | 8 |
| CHRAC1 | nucleosome remodelling | Chromosome 8:141,590,586-141,596,434 | 8 |
| CHRAC1 | nucleosome remodelling | Chromosome 8:141,590,586-141,596,434 | 8 |
| CHRAC1 | nucleosome remodelling | Chromosome 8:141,590,586-141,596,434 | 8 |
| PTK2 | FAK | Chromosome 8:141,737,683-142,080,514 | 8 |
| PTK2 | FAK | Chromosome 8:141,737,683-142,080,514 | 8 |
| PTK2 | FAK | Chromosome 8:141,737,683-142,080,514 | 8 |
| PTP4A3 |  | Chromosome 8:142,501,189-142,510,802 | 8 |
| CYP11B1 | 11 beta hydroxylase | Chromosome 8:143,950,775-143,958,238 | 8 |
| LY6E | Ly6E/A | Chromosome 8:144,171,274-144,175,199 | 8 |
| MAPK15 |  | Chromosome 8:144,870,498-144,876,619 | 8 |
| MSR1 | CD204, SR-A, SCARA1 | Chromosome 8:16,009,761-16,094,595 | 8 |
| MSR1 | CD204, SR-A, SCARA1 | Chromosome 8:16,009,761-16,094,595 | 8 |
| MSR1 | CD204, SR-A, SCARA1 | Chromosome 8:16,009,761-16,094,595 | 8 |
| FGF20 | FGF20 | Chromosome 8:16,894,049-16,904,061 | 8 |
| FGF20 | FGF20 | Chromosome 8:16,894,049-16,904,061 | 8 |


| FGF20 | FGF20 | Chromosome 8:16,894,049-16,904,061 | 8 |
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| NAT2 |  | Chromosome 8:18,293,035-18,303,003 | 8 |
| NAT2 |  | Chromosome 8:18,293,035-18,303,003 | 8 |
| DOK2 |  | Chromosome 8:21,822,330-21,827,151 | 8 |
| DOK2 |  | Chromosome 8:21,822,330-21,827,151 | 8 |
| FGF17 | FGF17 | Chromosome 8:21,955,883-21,962,266 | 8 |
| FGF17 | FGF17 | Chromosome 8:21,955,883-21,962,266 | 8 |
| PPP3CC |  | Chromosome 8:22,354,541-22,454,583 | 8 |
| PPP3CC |  | Chromosome 8:22,354,541-22,454,583 | 8 |
| EGR3 |  | Chromosome 8:22,601,117-22,606,760 | 8 |
| EGR3 |  | Chromosome 8:22,601,117-22,606,760 | 8 |
| EGR3 |  | Chromosome 8:22,601,117-22,606,760 | 8 |
| TNFRSF10B | CD262, TRAILR2 | Chromosome 8:22,933,591-22,982,637 | 8 |
| TNFRSF10B | CD262, TRAILR2 | Chromosome 8:22,933,591-22,982,637 | 8 |
| TNFRSF10C | CD263, TRAILR3 | Chromosome 8:23,016,377-23,030,895 | 8 |
| TNFRSF10D | CD264, TRAILR4 | Chromosome 8:23,049,046-23,077,488 | 8 |
| TNFRSF10D | CD264, TRAILR4 | Chromosome 8:23,049,046-23,077,488 | 8 |
| TNFRSF10A | CD261, TRAILR1 | Chromosome 8:23,104,009-23,138,584 | 8 |
| TNFRSF10A | CD261, TRAILR1 | Chromosome 8:23,104,009-23,138,584 | 8 |
| CHMP7 | CHMP7 | Chromosome 8:23,157,114-23,175,452 | 8 |
| GNRH1 | GNRH1 | Chromosome 8:25,332,693-25,338,087 | 8 |
| EBF2 |  | Chromosome 8:25,757,490-25,958,292 | 8 |
| EBF2 |  | Chromosome 8:25,757,490-25,958,292 | 8 |
| EBF2 |  | Chromosome 8:25,757,490-25,958,292 | 8 |
| EBF2 |  | Chromosome 8:25,757,490-25,958,292 | 8 |
| BNIP3L |  | Chromosome 8:26,296,331-26,326,562 | 8 |
| PTK 2B | PYK2 | Chromosome 8:27,224,916-27,372,824 | 8 |
| PTK 2B | PYK2 | Chromosome 8:27,224,916-27,372,824 | 8 |
| PTK 2B | PYK2 | Chromosome 8:27,224,916-27,372,824 | 8 |
| CLU |  | Chromosome 8:27,510,351-27,528,288 | 8 |
| CLU |  | Chromosome 8:27,510,351-27,528,288 | 8 |
| SCARA3 | SCARA3 | Chromosome 8:27,547,304-27,590,211 | 8 |
| SCARA5 |  | Chromosome 8:27,783,655-27,906,117 | 8 |
| DUSP4 |  | Chromosome 8:29,249,530-29,264,104 | 8 |
| DUSP4 |  | Chromosome 8:29,249,530-29,264,104 | 8 |
| GTF2E2 |  | Chromosome 8:30,555,422-30,635,274 | 8 |
| GTF2E2 |  | Chromosome 8:30,555,422-30,635,274 | 8 |
| GTF2E2 |  | Chromosome 8:30,555,422-30,635,274 | 8 |
| DUSP26 |  | Chromosome 8:33,568,393-33,577,043 | 8 |
| DUSP26 |  | Chromosome 8:33,568,393-33,577,043 | 8 |
| BAG4 |  | Chromosome 8:38,153,263-38,189,966 | 8 |
| BAG4 |  | Chromosome 8:38,153,263-38,189,966 | 8 |
| FGFR1 |  | Chromosome 8:38,389,406-38,445,296 | 8 |
| FGFR1 |  | Chromosome 8:38,389,406-38,445,296 | 8 |
| FGFR1 |  | Chromosome 8:38,389,406-38,445,296 | 8 |
| TACC1 |  | Chromosome 8:38,734,008-38,829,703 | 8 |
| TACC1 |  | Chromosome 8:38,734,008-38,829,703 | 8 |
| INDO | IDO | Chromosome 8:39,890,485-39,905,120 | 8 |
| INDO | IDO | Chromosome 8:39,890,485-39,905,120 | 8 |
| ANK1 |  | Chromosome 8:41,629,901-41,873,437 | 8 |
| ANK1 |  | Chromosome 8:41,629,901-41,873,437 | 8 |
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| ANK1 |  | Chromosome 8:41,629,901-41,873,437 | 8 |
| IKBKB | IKK2 | Chromosome 8:42,247,986-42,309,130 | 8 |
| IKBKB | IKK2 | Chromosome 8:42,247,986-42,309,130 | 8 |
| PRKDC | DNA-PK | Chromosome 8:48,848,222-49,035,296 | 8 |


| OPRK1 | KOR | Chromosome 8:54,300,829-54,326,747 | 8 |
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| OPRK1 | KOR | Chromosome 8:54,300,829-54,326,747 | 8 |
| LYN | Lyn | Chromosome 8:56,954,926-57,086,493 | 8 |
| LYN | Lyn | Chromosome 8:56,954,926-57,086,493 | 8 |
| LYN | Lyn | Chromosome 8:56,954,926-57,086,493 | 8 |
| CYP7A1 |  | Chromosome 8:59,565,292-59,575,275 | 8 |
| CYP7A1 |  | Chromosome 8:59,565,292-59,575,275 | 8 |
| NSMAF |  | Chromosome 8:59,658,617-59,734,940 | 8 |
| ANGPT2 | Angiopoietin-2 | Chromosome 8:6,344,580-6,408,338 | 8 |
| ANGPT2 | Angiopoietin-2 | Chromosome 8:6,344,580-6,408,338 | 8 |
| CYP7B1 |  | Chromosome 8:65,671,246-65,873,902 | 8 |
| CYP7B1 |  | Chromosome 8:65,671,246-65,873,902 | 8 |
| CYP7B1 |  | Chromosome 8:65,671,246-65,873,902 | 8 |
| CRH |  | Chromosome 8:67,251,166-67,253,380 | 8 |
| TRAM1 |  | Chromosome 8:71,648,227-71,683,158 | 8 |
| LY96 | MD2 | Chromosome 8:75,066,141-75,103,859 | 8 |
| LY96 | MD2 | Chromosome 8:75,066,141-75,103,859 | 8 |
| HNF4G | NR2A2 | Chromosome 8:76,482,826-76,641,623 | 8 |
| HNF4G | NR2A2 | Chromosome 8:76,482,826-76,641,623 | 8 |
| HNF4G | NR2A2 | Chromosome 8:76,482,826-76,641,623 | 8 |
| PXMP3 |  | Chromosome 8:78,057,713-78,074,994 | 8 |
| PXMP3 |  | Chromosome 8:78,057,713-78,074,994 | 8 |
| IL7 |  | Chromosome 8:79,807,560-79,880,313 | 8 |
| IL7 |  | Chromosome 8:79,807,560-79,880,313 | 8 |
| IL7 |  | Chromosome 8:79,807,560-79,880,313 | 8 |
| CLDN23 |  | Chromosome 8:8,597,076-8,599,026 | 8 |
| CLDN23 |  | Chromosome 8:8,597,076-8,599,026 | 8 |
| PAG1 | PAG | Chromosome 8:82,042,600-82,186,858 | 8 |
| PAG1 | PAG | Chromosome 8:82,042,600-82,186,858 | 8 |
| FABP5 | FABP5 | Chromosome 8:82,355,326-82,359,563 | 8 |
| FABP5 | FABP5 | Chromosome 8:82,355,326-82,359,563 | 8 |
| FABP4 | FABP4 | Chromosome 8:82,553,481-82,558,023 | 8 |
| CHMP4C | CHMP4C | Chromosome 8:82,807,243-82,834,305 | 8 |
| CHMP4C | CHMP4C | Chromosome 8:82,807,243-82,834,305 | 8 |
| MMP16 |  | Chromosome 8:89,118,576-89,408,892 | 8 |
| MMP16 |  | Chromosome 8:89,118,576-89,408,892 | 8 |
| MMP16 |  | Chromosome 8:89,118,576-89,408,892 | 8 |
| MMP16 |  | Chromosome 8:89,118,576-89,408,892 | 8 |
| MMP16 |  | Chromosome 8:89,118,576-89,408,892 | 8 |
| RIPK2 | CARD3, RIP2, RICK | Chromosome 8:90,839,110-90,872,433 | 8 |
| RIPK2 | CARD3, RIP2, RICK | Chromosome 8:90,839,110-90,872,433 | 8 |
| NBS1 | NBS1 | Chromosome 8:91,014,740-91,066,075 | 8 |
| RUNX1T1 |  | Chromosome 8:93,040,328-93,176,619 | 8 |
| RUNX1T1 |  | Chromosome 8:93,040,328-93,176,619 | 8 |
| RUNX1T1 |  | Chromosome 8:93,040,328-93,176,619 | 8 |
| GEM | GEM | Chromosome 8:95,330,657-95,343,733 | 8 |
| GEM | GEM | Chromosome 8:95,330,657-95,343,733 | 8 |
| CCNE2 |  | Chromosome 8:95,961,628-95,976,660 | 8 |
| CCNE2 |  | Chromosome 8:95,961,628-95,976,660 | 8 |
| PTDSS1 |  | Chromosome 8:97,343,340-97,415,950 | 8 |
| PTDSS1 |  | Chromosome 8:97,343,340-97,415,950 | 8 |
| PTDSS1 |  | Chromosome 8:97,343,340-97,415,950 | 8 |
| PTDSS1 |  | Chromosome 8:97,343,340-97,415,950 | 8 |
| SDC2 | Syndecan | Chromosome 8:97,575,058-97,693,213 | 8 |
| SDC2 | Syndecan | Chromosome 8:97,575,058-97,693,213 | 8 |
| SDC2 | Syndecan | Chromosome 8:97,575,058-97,693,213 | 8 |


| POP1 |  | Chromosome 8:99,199,244-99,239,816 | 8 |
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| POP1 |  | Chromosome 8:99,199,244-99,239,816 | 8 |
| POP1 |  | Chromosome 8:99,199,244-99,239,816 | 8 |
| STK3 |  | Chromosome 8:99,536,041-99,907,085 | 8 |
| STK3 |  | Chromosome 8:99,536,041-99,907,085 | 8 |
| STK3 |  | Chromosome 8:99,536,041-99,907,085 | 8 |
| STK3 |  | Chromosome 8:99,536,041-99,907,085 | 8 |
| STK3 |  | Chromosome 8:99,536,041-99,907,085 | 8 |
| IL6RL1 |  | Chromosome 9 | 9 |
| IL6RL1 |  | Chromosome 9 | 9 |
| TGFBR1 |  | Chromosome 9:100,907,233-100,956,406 | 9 |
| TGFBR1 |  | Chromosome 9:100,907,233-100,956,406 | 9 |
| SEC61B |  | Chromosome 9:101,024,380-101,032,722 | 9 |
| NR4A3 |  | Chromosome 9:101,623,958-101,668,994 | 9 |
| NR4A3 |  | Chromosome 9:101,623,958-101,668,994 | 9 |
| PPP3R2 |  | Chromosome 9:103,393,718-103,397,104 | 9 |
| PPP3R2 |  | Chromosome 9:103,393,718-103,397,104 | 9 |
| SLC44A1 | CDW92 | Chromosome 9:107,046,724-107,241,273 | 9 |
| SLC44A1 | CDW92 | Chromosome 9:107,046,724-107,241,273 | 9 |
| TAL2 |  | Chromosome 9:107,464,599-107,465,214 | 9 |
| ACTL7B |  | Chromosome 9:110,656,692-110,659,068 | 9 |
| ACTL7B |  | Chromosome 9:110,656,692-110,659,068 | 9 |
| PTPN3 |  | Chromosome 9:111,177,800-111,300,407 | 9 |
| PTPN3 |  | Chromosome 9:111,177,800-111,300,407 | 9 |
| TXN | Txk, Rlk, thioredoxin | Chromosome 9:112,045,912-112,058,741 | 9 |
| TXN | Txk, Rlk, thioredoxin | Chromosome 9:112,045,912-112,058,741 | 9 |
| SVEP1/MUSK |  | Chromosome 9:112,431,057-112,431,557 | 9 |
| LTB4DH |  | Chromosome 9:113,364,678-113,401,917 | 9 |
| LTB4DH |  | Chromosome 9:113,364,678-113,401,917 | 9 |
| UGCG |  | Chromosome 9:113,698,867-113,737,470 | 9 |
| UGCG |  | Chromosome 9:113,698,867-113,737,470 | 9 |
| UGCG |  | Chromosome 9:113,698,867-113,737,470 | 9 |
| FKBP15 |  | Chromosome 9:114,967,620-115,024,010 | 9 |
| FKBP15 |  | Chromosome 9:114,967,620-115,024,010 | 9 |
| POLE3 |  | Chromosome 9:115,209,336-115,212,773 | 9 |
| POLE3 |  | Chromosome 9:115,209,336-115,212,773 | 9 |
| ORM1 | ORM3 | Chromosome 9:116,125,157-116,128,578 | 9 |
| ORM2 |  | Chromosome 9:116,131,890-116,135,357 | 9 |
| TNFSF15 |  | Chromosome 9:116,591,421-116,608,229 | 9 |
| TNFSF15 |  | Chromosome 9:116,591,421-116,608,229 | 9 |
| TNFSF8 | CD153 | Chromosome 9:116,704,945-116,732,591 | 9 |
| TNC | tenascin | Chromosome 9:116,822,634-116,920,260 | 9 |
| TNC | tenascin | Chromosome 9:116,822,634-116,920,260 | 9 |
| TLR4 | CD284 | Chromosome 9:119,506,405-119,519,589 | 9 |
| TLR4 | CD284 | Chromosome 9:119,506,405-119,519,589 | 9 |
| TRAF1 |  | Chromosome 9:122,704,492-122,730,868 | 9 |
| TRAF1 |  | Chromosome 9:122,704,492-122,730,868 | 9 |
| C5 |  | Chromosome 9:122,754,434-122,852,375 | 9 |
| PTGS1 | PG synthase | Chromosome 9:124,173,050-124,197,802 | 9 |
| PTGS1 | PG synthase | Chromosome 9:124,173,050-124,197,802 | 9 |
| PSMB7 |  | Chromosome 9:126,155,565-126,217,542 | 9 |
| PSMB7 |  | Chromosome 9:126,155,565-126,217,542 | 9 |
| NR5A1 |  | Chromosome 9:126,283,336-126,309,530 | 9 |
| NR6A1 | NR6A1 | Chromosome 9:126,319,380-126,573,410 | 9 |
| NR6A1 | NR6A1 | Chromosome 9:126,319,380-126,573,410 | 9 |
| HSPA5 | grp78 | Chromosome 9:127,036,953-127,043,430 | 9 |


| CDK9 | pTEFb | Chromosome 9:129,587,898-129,592,887 | 9 |
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| CDK9 | pTEFb | Chromosome 9:129,587,898-129,592,887 | 9 |
| ENG | CD105, TGF beta receptd | Chromosome 9:129,617,112-129,656,856 | 9 |
| LCN2 | LCN2 | Chromosome 9:129,951,171-129,956,333 | 9 |
| SET |  | Chromosome 9:130,485,844-130,498,488 | 9 |
| PPP2R4 | PP2A | Chromosome 9:130,913,050-130,951,046 | 9 |
| PTGES |  | Chromosome 9:131,540,433-131,555,165 | 9 |
| PTGES |  | Chromosome 9:131,540,433-131,555,165 | 9 |
| GPR107 |  | Chromosome 9:131,855,526-131,942,264 | 9 |
| GPR107 |  | Chromosome 9:131,855,526-131,942,264 | 9 |
| GPR107 |  | Chromosome 9:131,855,526-131,942,264 | 9 |
| LAMC3 |  | Chromosome 9:132,874,325-132,958,267 | 9 |
| LAMC3 |  | Chromosome 9:132,874,325-132,958,267 | 9 |
| RAPGEF1 | C3G | Chromosome 9:133,441,978-133,605,282 | 9 |
| RAPGEF1 | C3G | Chromosome 9:133,441,978-133,605,282 | 9 |
| DDX31 |  | Chromosome 9:134,458,205-134,535,609 | 9 |
| DDX31 |  | Chromosome 9:134,458,205-134,535,609 | 9 |
| GTF3C5 |  | Chromosome 9:134,895,897-134,923,709 | 9 |
| GTF3C5 |  | Chromosome 9:134,895,897-134,923,709 | 9 |
| ABO |  | Chromosome 9:135,120,384-135,140,451 | 9 |
| ADAMTS13 |  | Chromosome 9:135,276,941-135,314,329 | 9 |
| ADAMTS13 |  | Chromosome 9:135,276,941-135,314,329 | 9 |
| VAV2 |  | Chromosome 9:135,616,837-135,847,547 | 9 |
| VAV2 |  | Chromosome 9:135,616,837-135,847,547 | 9 |
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| VAV2 |  | Chromosome 9:135,616,837-135,847,547 | 9 |
| VAV2 |  | Chromosome 9:135,616,837-135,847,547 | 9 |
| COL5A1 |  | Chromosome 9:136,673,473-136,876,510 | 9 |
| FCN1 | ficolin | Chromosome 9:136,940,837-136,949,630 | 9 |
| NOTCH1 | Notch 1 | Chromosome 9:138,508,717-138,560,135 | 9 |
| TRAF2 |  | Chromosome 9:138,900,786-138,940,888 | 9 |
| NOXA1 |  | Chromosome 9:139,437,668-139,448,679 | 9 |
| NOXA1 |  | Chromosome 9:139,437,668-139,448,679 | 9 |
| LEDGF | p75 | Chromosome 9:15,454,064-15,501,017 | 9 |
| LEDGF | p75 | Chromosome 9:15,454,064-15,501,017 | 9 |
| SMARCA2 | SWI/SNF | Chromosome 9:2,005,342-2,183,624 | 9 |
| SMARCA2 | SWI/SNF | Chromosome 9:2,005,342-2,183,624 | 9 |
| SMARCA2 | SWI/SNF | Chromosome 9:2,005,342-2,183,624 | 9 |
| KIAA0020 | HA-8, PEN, CD162R | Chromosome 9:2,794,152-2,834,095 | 9 |
| KIAA0020 | HA-8, PEN, CD162R | Chromosome 9:2,794,152-2,834,095 | 9 |
| IFNB1 | IRG47 | Chromosome 9:21,067,104-21,067,962 | 9 |
| IFNB1 | IRG47 | Chromosome 9:21,067,104-21,067,962 | 9 |
| IFNW1 |  | Chromosome 9:21,130,213-21,132,144 | 9 |
| IFNA7 |  | Chromosome 9:21,191,234-21,229,990 | 9 |
| IFNA13 |  | Chromosome 9:21,357,423-21,358,961 | 9 |
| IFNE1 |  | Chromosome 9:21,470,838-21,472,312 | 9 |
| IFNE1 |  | Chromosome 9:21,470,838-21,472,312 | 9 |
| CDKN2A |  | Chromosome 9:21,957,751-21,984,490 | 9 |
| CDKN2A |  | Chromosome 9:21,957,751-21,984,490 | 9 |
| CDKN2B |  | Chromosome 9:21,992,902-21,999,312 | 9 |
| PLAA | PLA activatinp protein | Chromosome 9:26,894,081-26,937,461 | 9 |
| TEK | CD202b | Chromosome 9:27,099,236-27,220,173 | 9 |
| TEK | CD202b | Chromosome 9:27,099,236-27,220,173 | 9 |
| TEK | CD202b | Chromosome 9:27,099,236-27,220,173 | 9 |
| IFNK |  | Chromosome 9:27,514,302-27,516,496 | 9 |
| RFX3 | RFX3 | Chromosome 9:3,208,297-3,515,983 | 9 |


| RFX3 | RFX3 | Chromosome 9:3,208,297-3,515,983 | 9 |
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| RFX3 | RFX3 | Chromosome 9:3,208,297-3,515,983 | 9 |
| RFX3 | RFX3 | Chromosome 9:3,208,297-3,515,983 | 9 |
| RFX3 | RFX3 | Chromosome 9:3,208,297-3,515,983 | 9 |
| DDX58 |  | Chromosome 9:32,445,300-32,516,322 | 9 |
| DDX58 |  | Chromosome 9:32,445,300-32,516,322 | 9 |
| BAG1 |  | Chromosome 9:33,242,469-33,254,744 | 9 |
| CHMP5 | CHMP5 | Chromosome 9:33,254,167-33,271,525 | 9 |
| IL11RA |  | Chromosome 9:34,636,635-34,651,884 | 9 |
| IL11RA |  | Chromosome 9:34,636,635-34,651,884 | 9 |
| Cd72 |  | Chromosome 9:35,599,976-35,608,753 | 9 |
| TLN1 | Talin | Chromosome 9:35,687,334-35,722,369 | 9 |
| TLN1 | Talin | Chromosome 9:35,687,334-35,722,369 | 9 |
| MELK | Melk | Chromosome 9:36,562,873-36,667,679 | 9 |
| MELK | Melk | Chromosome 9:36,562,873-36,667,679 | 9 |
| SHB |  | Chromosome 9:37,909,131-38,059,249 | 9 |
| SHB |  | Chromosome 9:37,909,131-38,059,249 | 9 |
| JAK2 |  | Chromosome 9:4,975,245-5,118,183 | 9 |
| JAK2 |  | Chromosome 9:4,975,245-5,118,183 | 9 |
| CD274 | CD274, PD1 ligand 1, PD | Chromosome 9:5,440,525-5,460,547 | 9 |
| CD274 | CD274, PD1 ligand 1, PD | Chromosome 9:5,440,525-5,460,547 | 9 |
| PDCD1LG2 | CD273, PD1 ligand 2, PD | Chromosome 9:5,500,570-5,561,252 | 9 |
| IL33 |  | Chromosome 9:6,205,809-6,247,983 | 9 |
| IL33 |  | Chromosome 9:6,205,809-6,247,983 | 9 |
| PRKACG |  | Chromosome 9:70,817,241-70,818,849 | 9 |
| ANXA1 | Annexin-1 | Chromosome 9:74,956,493-74,975,129 | 9 |
| ANXA1 | Annexin-1 | Chromosome 9:74,956,493-74,975,129 | 9 |
| RORB | RORb, NR1F2 | Chromosome 9:76,302,072-76,491,937 | 9 |
| RORB | RORb, NR1F2 | Chromosome 9:76,302,072-76,491,937 | 9 |
| RORB | RORb, NR1F2 | Chromosome 9:76,302,072-76,491,937 | 9 |
| FOXB2 |  | Chromosome 9:78,824,391-78,825,689 | 9 |
| PTPRD |  | Chromosome 9:8,304,246-9,008,735 | 9 |
| PTPRD |  | Chromosome 9:8,304,246-9,008,735 | 9 |
| PTPRD |  | Chromosome 9:8,304,246-9,008,735 | 9 |
| PTPRD |  | Chromosome 9:8,304,246-9,008,735 | 9 |
| PTPRD |  | Chromosome 9:8,304,246-9,008,735 | 9 |
| PTPRD |  | Chromosome 9:8,304,246-9,008,735 | 9 |
| GAS1 |  | Chromosome 9:88,749,098-88,751,924 | 9 |
| DAPK1 |  | Chromosome 9:89,301,963-89,513,369 | 9 |
| DAPK1 |  | Chromosome 9:89,301,963-89,513,369 | 9 |
| DAPK1 |  | Chromosome 9:89,301,963-89,513,369 | 9 |
| DAPK1 |  | Chromosome 9:89,301,963-89,513,369 | 9 |
| CKS2 |  | Chromosome 9:91,115,925-91,121,438 | 9 |
| SEMA4D | CD100 | Chromosome 9:91,181,972-91,302,708 | 9 |
| SYK |  | Chromosome 9:92,603,890-92,700,652 | 9 |
| SYK |  | Chromosome 9:92,603,890-92,700,652 | 9 |
| NFIL3 | nuclear factor interleukin | Chromosome 9:93,211,148-93,225,965 | 9 |
| NFIL3 | nuclear factor interleukin | Chromosome 9:93,211,148-93,225,965 | 9 |
| PTCH1 | Ptc | Chromosome 9:97,245,083-97,318,923 | 9 |
| PTCH1 | Ptc | Chromosome 9:97,245,083-97,318,923 | 9 |
| PTCH1 | Ptc | Chromosome 9:97,245,083-97,318,923 | 9 |
| XPA |  | Chromosome 9:99,477,012-99,499,460 | 9 |
| FOXE1 |  | Chromosome 9:99,655,357-99,658,818 | 9 |
| ANP32B | acidic nucelar phosphopr | Chromosome 9:99,785,462-99,818,046 | 9 |
| BTK |  | Chromosome X:100,491,091-100,527,839 | X |
| BTK |  | Chromosome X:100,491,091-100,527,839 | X |


| IL1RAPL2 |  | Chromosome X:103,697,652-104,898,478 | X |
| :---: | :---: | :---: | :---: |
| IL1RAPL2 |  | Chromosome X:103,697,652-104,898,478 | X |
| IL1RAPL2 |  | Chromosome X:103,697,652-104,898,478 | X |
| IL1RAPL2 |  | Chromosome X:103,697,652-104,898,478 | X |
| IL1RAPL2 |  | Chromosome X:103,697,652-104,898,478 | X |
| IL1RAPL2 |  | Chromosome X:103,697,652-104,898,478 | X |
| IL1RAPL2 |  | Chromosome X:103,697,652-104,898,478 | X |
| IL1RAPL2 |  | Chromosome X:103,697,652-104,898,478 | X |
| IL1RAPL2 |  | Chromosome X:103,697,652-104,898,478 | X |
| IL1RAPL2 |  | Chromosome X:103,697,652-104,898,478 | X |
| IL1RAPL2 |  | Chromosome X:103,697,652-104,898,478 | X |
| IL1RAPL2 |  | Chromosome X:103,697,652-104,898,478 | X |
| IL1RAPL2 |  | Chromosome X:103,697,652-104,898,478 | X |
| GRAIL |  | Chromosome X:105,823,724-105,926,902 | X |
| GRAIL |  | Chromosome X:105,823,724-105,926,902 | X |
| GRAIL |  | Chromosome X:105,823,724-105,926,902 | X |
| TSC22D3 | DIP | Chromosome X:106,843,107-106,905,858 | X |
| TSC22D3 | DIP | Chromosome X:106,843,107-106,905,858 | X |
| NLRP3P | NOD13 | Chromosome X:107,103,687-107,105,431 | X |
| COL4A6 |  | Chromosome X:107,285,493-107,569,383 | X |
| COL4A6 |  | Chromosome X:107,285,493-107,569,383 | X |
| COL4A6 |  | Chromosome X:107,285,493-107,569,383 | X |
| COL4A6 |  | Chromosome X:107,285,493-107,569,383 | X |
| COL4A5 |  | Chromosome X:107,569,810-107,827,431 | X |
| COL4A5 |  | Chromosome X:107,569,810-107,827,431 | X |
| PAK3 |  | Chromosome X:110,226,244-110,350,816 | X |
| IL13RA2 | CD213a2 | Chromosome X:114,144,794-114,159,792 | X |
| AGTR2 | Angiotensin receptor | Chromosome X:115,216,003-115,220,253 | X |
| IL13RA1 | CD213a1 | Chromosome X:117,745,563-117,812,530 | X |
| IL13RA1 | CD213a1 | Chromosome X:117,745,563-117,812,530 | X |
| LAMP2 | CD107b | Chromosome X:119,446,367-119,487,189 | X |
| LAMP2 | CD107b | Chromosome X:119,446,367-119,487,189 | X |
| TLR7 | CD287 | Chromosome X:12,795,123-12,818,420 | X |
| TLR8 | CD288 | Chromosome X:12,834,679-12,851,209 | X |
| TMSB4X | thymosin beta 4 | Chromosome X:12,903,148-12,905,267 | X |
| BIRC4 | XIAP | Chromosome X:122,821,558-122,875,510 | X |
| BIRC4 | XIAP | Chromosome X:122,821,558-122,875,510 | X |
| SH2D1A | EAT2a, SAP | Chromosome X:123,307,875-123,334,686 | X |
| SH2D1A | EAT2a, SAP | Chromosome X:123,307,875-123,334,686 | X |
| SMARCA1 | ISWI | Chromosome X:128,408,159-128,485,158 | X |
| SMARCA1 | ISWI | Chromosome X:128,408,159-128,485,158 | X |
| SMARCA1 | ISWI | Chromosome X:128,408,159-128,485,158 | X |
| APLN | apelin | Chromosome X:128,607,006-128,616,595 | X |
| APLN | apelin | Chromosome X:128,607,006-128,616,595 | X |
| AIFM1 | PDCD8 | Chromosome X:129,091,018-129,127,489 | X |
| AIFM1 | PDCD8 | Chromosome X:129,091,018-129,127,489 | X |
| IGSF1 |  | Chromosome X:130,235,161-130,361,358 | X |
| IGSF1 |  | Chromosome X:130,235,161-130,361,358 | X |
| DDX26B |  | Chromosome X:134,482,215-134,544,100 | X |
| SLC9A6 | NHE-6 | Chromosome X:134,895,264-134,957,089 | X |
| SLC9A6 | NHE-6 | Chromosome X:134,895,264-134,957,089 | X |
| CD40LG | CD40L, CD154, TNFSF5 | Chromosome X:135,558,002-135,570,215 | X |
| CD40LG | CD40L, CD154, TNFSF5 | Chromosome X:135,558,002-135,570,215 | X |
| FGF13 | FGF13 | Chromosome X:137,541,401-137,894,912 | X |
| FGF13 | FGF13 | Chromosome X:137,541,401-137,894,912 | X |
| FGF13 | FGF13 | Chromosome X:137,541,401-137,894,912 | X |


| FGF13 | FGF13 | Chromosome X:137,541,401-137,894,912 | X |
| :---: | :---: | :---: | :---: |
| FGF13 | FGF13 | Chromosome X:137,541,401-137,894,912 | X |
| FGF13 | FGF13 | Chromosome X:137,541,401-137,894,912 | X |
| CD99L2 |  | Chromosome X:149,685,467-149,817,837 | X |
| CD99L2 |  | Chromosome X:149,685,467-149,817,837 | X |
| FIGF | VEGFD | Chromosome X:15,273,639-15,312,498 | X |
| FIGF | VEGFD | Chromosome X:15,273,639-15,312,498 | X |
| FIGF | VEGFD | Chromosome X:15,273,639-15,312,498 | X |
| BMX |  | Chromosome X:15,392,290-15,484,573 | X |
| ACE2 |  | Chromosome X:15,489,077-15,530,199 | X |
| BGN |  | Chromosome X:152,413,591-152,428,206 | X |
| BGN |  | Chromosome X:152,413,591-152,428,206 | X |
| DUSP9 |  | Chromosome X:152,561,182-152,569,975 | X |
| L1CAM | CD171 | Chromosome X:152,780,163-152,804,802 | X |
| IRAK1 | Pelle | Chromosome X:152,929,145-152,938,625 | X |
| G6PD | G6PDH | Chromosome X:153,412,800-153,428,981 | X |
| DKC1 | dyskeratosis congenita | Chromosome X:153,644,229-153,659,158 | X |
| SCML2 |  | Chromosome X:18,167,355-18,282,768 | X |
| SCML2 |  | Chromosome X:18,167,355-18,282,768 | X |
| MAP3K15 |  | Chromosome X:19,288,095-19,443,363 | X |
| MAP3K15 |  | Chromosome X:19,288,095-19,443,363 | X |
| MAP3K15 |  | Chromosome X:19,288,095-19,443,363 | X |
| SH3KBP1 |  | Chromosome X:19,462,014-19,815,640 | X |
| SH3KBP1 |  | Chromosome X:19,462,014-19,815,640 | X |
| SH3KBP1 |  | Chromosome X:19,462,014-19,815,640 | X |
| XG |  | Chromosome X:2,680,115-2,743,968 | X |
| DDX53 |  | Chromosome X:22,927,999-22,931,627 | X |
| DDX53 |  | Chromosome X:22,927,999-22,931,627 | X |
| DDX53 |  | Chromosome X:22,927,999-22,931,627 | X |
| PRDX4 |  | Chromosome X:23,592,300-23,614,437 | X |
| IL1RAPL1 |  | Chromosome X:28,515,437-29,884,761 | X |
| IL1RAPL1 |  | Chromosome X:28,515,437-29,884,761 | X |
| IL1RAPL1 |  | Chromosome X:28,515,437-29,884,761 | X |
| IL1RAPL1 |  | Chromosome X:28,515,437-29,884,761 | X |
| IL1RAPL1 |  | Chromosome X:28,515,437-29,884,761 | X |
| IL1RAPL1 |  | Chromosome X:28,515,437-29,884,761 | X |
| IL1RAPL1 |  | Chromosome X:28,515,437-29,884,761 | X |
| IL1RAPL1 |  | Chromosome X:28,515,437-29,884,761 | X |
| IL1RAPL1 |  | Chromosome X:28,515,437-29,884,761 | X |
| IL1RAPL1 |  | Chromosome X:28,515,437-29,884,761 | X |
| IL1RAPL1 |  | Chromosome X:28,515,437-29,884,761 | X |
| NR0B1 | NR0B1 | Chromosome X:30,232,244-30,237,636 | X |
| NR0B1 | NR0B1 | Chromosome X:30,232,244-30,237,636 | X |
| NR0B1 | NR0B1 | Chromosome X:30,232,244-30,237,636 | X |
| XK |  | Chromosome X:37,429,931-37,476,322 | X |
| CYBB | NOX2 | Chromosome X:37,524,208-37,557,658 | X |
| TSPAN7 | CD231 | Chromosome X:38,305,553-38,433,118 | X |
| TSPAN7 | CD231 | Chromosome X:38,305,553-38,433,118 | X |
| DDX3X |  | Chromosome X:41,077,595-41,108,669 | X |
| DDX3X |  | Chromosome X:41,077,595-41,108,669 | X |
| DUSP21 |  | Chromosome X:44,588,193-44,589,078 | X |
| DUSP21 |  | Chromosome X:44,588,193-44,589,078 | X |
| SLC9A7 | NHE-7 | Chromosome X:46,349,697-46,503,434 | X |
| SLC9A7 | NHE-7 | Chromosome X:46,349,697-46,503,434 | X |
| SLC9A7 | NHE-7 | Chromosome X:46,349,697-46,503,434 | X |
| TIMP1 |  | Chromosome X:47,326,634-47,331,134 | X |


| CFP | properdin, PFC | Chromosome X:47,368,557-47,374,648 | X |
| :---: | :---: | :---: | :---: |
| WAS | Wiskott Aldrich, WASP | Chromosome X:48,427,112-48,434,762 | X |
| SUV39H1 | Suv39h1 | Chromosome X:48,439,930-48,452,347 | X |
| GATA1 |  | Chromosome X:48,529,906-48,537,662 | X |
| HDAC6 |  | Chromosome X:48,545,170-48,568,336 | X |
| FOXP3 |  | Chromosome X:48,993,841-49,008,232 | X |
| CCNB3 |  | Chromosome X:49,856,156-50,111,653 | X |
| CCNB3 |  | Chromosome X:49,856,156-50,111,653 | X |
| TSPYL2 | CDA1 | Chromosome X:53,128,274-53,134,447 | X |
| TSPYL2 | CDA1 | Chromosome X:53,128,274-53,134,447 | X |
| JARID1C | SMCY homolog | Chromosome X:53,238,059-53,271,329 | X |
| JARID1C | SMCY homolog | Chromosome X:53,238,059-53,271,329 | X |
| ALAS2 | Alas2 | Chromosome X:55,052,213-55,074,136 | X |
| ALAS2 | Alas2 | Chromosome X:55,052,213-55,074,136 | X |
| NLRP2P | NOD24 | Chromosome X:57,719,936-57,723,438 | X |
| AR | Androgen receptor, NR3ర | Chromosome X:66,680,599-66,867,186 | X |
| AR | Androgen receptor, NR3O | Chromosome X:66,680,599-66,867,186 | X |
| EDA |  | Chromosome X:68,752,636-69,176,047 | X |
| EDA |  | Chromosome X:68,752,636-69,176,047 | X |
| EDA |  | Chromosome X:68,752,636-69,176,047 | X |
| P2RY4 | P2Y4 | Chromosome X:69,394,741-69,396,379 | X |
| P2RY4 | P2Y4 | Chromosome X:69,394,741-69,396,379 | X |
| MLLT7 | FOXO4 | Chromosome X:70,232,772-70,240,110 | X |
| IL2RG | CD132 | Chromosome X:70,243,979-70,248,188 | X |
| CXCR3 | CD183 | Chromosome X:70,752,491-70,755,092 | X |
| CXCR3 | CD183 | Chromosome X:70,752,491-70,755,092 | X |
| FLJ20105 |  | Chromosome X:71,341,232-71,375,602 | X |
| FLJ20105 |  | Chromosome X:71,341,232-71,375,602 | X |
| FLJ20105 |  | Chromosome X:71,341,232-71,375,602 | X |
| HDAC8 |  | Chromosome X:71,466,091-71,709,623 | X |
| HDAC8 |  | Chromosome X:71,466,091-71,709,623 | X |
| HDAC8 |  | Chromosome X:71,466,091-71,709,623 | X |
| FGF16 | FGF16 | Chromosome X:76,596,303-76,598,669 | X |
| FGF16 | FGF16 | Chromosome X:76,596,303-76,598,669 | X |
| CYSLTR1 |  | Chromosome X:77,413,617-77,469,743 | X |
| CYSLTR1 |  | Chromosome X:77,413,617-77,469,743 | X |
| TBL1X |  | Chromosome X:9,391,369-9,647,778 | X |
| TBL1X |  | Chromosome X:9,391,369-9,647,778 | X |
| TBL1X |  | Chromosome X:9,391,369-9,647,778 | X |
| TBL1X |  | Chromosome X:9,391,369-9,647,778 | X |
| DIAPH2 |  | Chromosome X:95,826,365-96,746,652 | X |
| DIAPH2 |  | Chromosome X:95,826,365-96,746,652 | X |
| DIAPH2 |  | Chromosome X:95,826,365-96,746,652 | X |
| DIAPH2 |  | Chromosome X:95,826,365-96,746,652 | X |
| DIAPH2 |  | Chromosome X:95,826,365-96,746,652 | X |
| DIAPH2 |  | Chromosome X:95,826,365-96,746,652 | X |
| DIAPH2 |  | Chromosome X:95,826,365-96,746,652 | X |
| DIAPH2 |  | Chromosome X:95,826,365-96,746,652 | X |
| DIAPH2 |  | Chromosome X:95,826,365-96,746,652 | X |
| DIAPH2 |  | Chromosome X:95,826,365-96,746,652 | X |
| DIAPH2 |  | Chromosome X:95,826,365-96,746,652 | X |
| NOX1 |  | Chromosome X:99,984,969-100,015,990 | X |
| NOX1 |  | Chromosome X:99,984,969-100,015,990 | X |
| USP9Y | A1/HY, DFFRY | Chromosome Y:13,322,554-13,482,162 | Y |
| DDX3Y | DRB1*1501/HY, DBY, DQ | Chromosome Y:13,525,413-13,541,784 | Y |
| DDX3Y | DRB1*1501/HY, DBY, DQ | Chromosome Y:13,525,413-13,541,784 | Y |


| UTY | B60/HY | Chromosome $\mathrm{Y}: 13,869,653-14,101,947$ | Y |
| :--- | :--- | :--- | :---: |
| UTY | B60/HY | Chromosome $\mathrm{Y}: 13,869,653-14,101,947$ | Y |
| TMSB4Y | A33/HY | Chromosome $\mathrm{Y}: 14,324,841-14,327,298$ | Y |
| CD24 |  | Chromosome $\mathrm{Y}: 19,611,898-19,614,093$ | Y |
| RPS4Y1 | B52/HY | Chromosome $\mathrm{Y}: 2,769,527-2,794,997$ | Y |
| RPS4Y1 | B52/HY | Chromosome $\mathrm{Y}: 2,769,527-2,794,997$ | Y |
| JARID1D | A2/HY, B7/HY, HLA-DQ5 | Chromosome $\mathrm{Y}: 20,326,689-20,366,212$ | Y |
| TBL1Y |  | Chromosome $\mathrm{Y}: 6,838,727-7,019,724$ | Y |
| TBL1Y |  | Chromosome $\mathrm{Y}: 6,838,727-7,019,724$ | Y |


| Gene Start Position | Gene End Position | Internal marker name | Marker database name |
| :---: | :---: | :---: | :---: |
| 1128751 | 1131952 | TNFRSF18 | new design |
| 1,260,521 | 1,274,623 | 0119C03 | D1S1394i |
| 1,557,337 | 1,623,109 | 0109H08 | D1S1252i |
| 1,971,769 | 2,106,694 | 0114H07 | D1S1708i |
| 10,439,166 | 10,455,200 | 0113C07 | D1S0305i |
| 100,957,885 | 100,977,189 | 0113E08 | D1S0467i |
| 101,475,032 | 101,479,662 | 0108A11 | D1S206 |
| 107,400,824 | 107,403,439 | 0119H04 | D1S1192i |
| 107,915,305 | 108,309,108 | 0111G07 | D1S0718i |
| 107,915,305 | 108,309,108 | 0108H03 | D1S1026i |
| 107,915,305 | 108,309,108 | 0117D07 | D1S1469i |
| 107,915,305 | 108,309,108 | 0104H04 | D1S1607i |
| 109,406,644 | 109,420,147 | 0101H03 | G09558 |
| 109,892,824 | 109,938,498 | 0117B09 | D1S0465i |
| 109,892,824 | 109,938,498 | 0109E10 | D1S1307i |
| 11,009,167 | 11,029,877 | 0114H04 | D1S0758i |
| 11,009,167 | 11,029,877 | 0116B01 | D1S1176i |
| 11,089,179 | 11,245,176 | 0116C01 | D1S0037i |
| 11,089,179 | 11,245,176 | 0105B06 | D1S0070i |
| 11,089,179 | 11,245,176 | 0113B06 | D1S0307i |
| 11,768,367 | 11,788,702 | 0108B09 | D1S0057i |
| 11,828,353 | 11,830,989 | 0109E01 | D1S2740 |
| 110,254,778 | 110,275,144 | 9902C03 | D1S1016i |
| 111,215,344 | 111,244,081 | 0117 A 09 | D1S0464i |
| 111,215,344 | 111,244,081 | 004 C 07 | D1S2809 |
| 111,571,804 | 111,587,585 | 0114 C 11 | D1S0836i |
| 111,827,493 | 111,908,107 | 0117G08 | D1S0458i |
| 111,827,493 | 111,908,107 | 0102C07 | D1S2789 |
| 111,886,363 | 112,060,836 | 0104 A 07 | D1S2837 |
| 113,045,251 | 113,051,579 | 0108H10 | D1S2756 |
| 114,092,981 | 114,215,904 | 0111B02 | D1S0280i |
| 114,092,981 | 114,215,904 | 0113E02 | D1S0471i |
| 114,911,701 | 114,925,788 | 0117C06 | D1S1220i |
| 115,048,613 | 115,102,147 | 0119C07 | G12424 |
| 115,373,938 | 115,378,464 | 0102A02 | D1S1693i |
| 115,630,060 | 115,682,380 | 0110E08 | D1S0034i |
| 116,858,680 | 116,915,184 | 0112F09 | D1S1147i |
| 116,918,554 | 117,011,898 | 0113B12 | D1S0455i |
| 117,098,530 | 117,113,374 | 335G10 | AL365325.4_109510 |
| 117,254,202 | 117,334,503 | 0106E06 | D1S1532i |
| 117,254,202 | 117,334,503 | 0109A03 | D1S252 |
| 117,487,732 | 117,555,079 | 0111G04 | D1S0475i |
| 117,487,732 | 117,555,079 | 0115G02 | D1S0644i |
| 119,851,356 | 119,859,200 | 0112C03 | D1S0895i |
| 12,046,021 | 12,126,851 | 0116F09 | D1S1559i |
| 12,149,647 | 12,191,872 | 264A06 | AL355998.4_40197 |
| 120,255,699 | 120,413,799 | 9902B01 | D1S1785i |
| 120728502 | 120737460 | FCGR1B | new design |
| 120728502 | 120737460 | FCGR1B2 | new design |
| 144,236,248 | 144,255,225 | T002F11 | D1S2344 |
| 144,407,155 | 144,426,971 | 0102G04 | D1S442 |
| 146,418,535 | 146,422,374 | 0110H02 | D1S1378i |
| 146,567,361 | 146,577,147 | 159F12 | AL022240.8_87561 |
| 148,167,168 | 148,175,396 | 0113B01 | D1S1407i |
| 148,813,658 | 148,818,760 | 0119C12 | D1S1299i |


| 148,969,175 | 149,005,057 | 0101B10 | D1S1006i |
| :---: | :---: | :---: | :---: |
| 149,035,311 | 149,047,436 | 0111A05 | D1S0479i |
| 149,035,311 | 149,047,436 | 0119D06 | D1S0483i |
| 149,275,670 | 149,286,700 | 0113G07 | D1S0513i |
| 149,531,037 | 149,566,815 | 0118F04 | D1S0139i |
| 15,655,811 | 15,690,482 | 0104E01 | D1S0157i |
| 15,655,811 | 15,690,482 | 0118A12 | D1S436 |
| 150,039,364 | 150,070,972 | 0108F12 | D1S0489i |
| 150,271,606 | 150,276,135 | 0119G12 | D1S1490i |
| 151,536,962 | 151,549,818 | 270 A11 | AL161636.7_109719 |
| 151,612,808 | 151,614,749 | 0119F05 | D1S1558i |
| 151,773,699 | 151,775,344 | 0111H06 | D1S0685i |
| 151,782,713 | 151,789,236 | 270G12 | AL157404.7_111158 |
| 151,900,905 | 151,910,148 | 0110D05 | D1S1454i |
| 151,900,905 | 151,910,148 | 0101C02 | Z67234 |
| 152,229,853 | 152,231,250 | 0116F05 | D1S0498i |
| 152,229,853 | 152,231,250 | 0109B03 | D1S2858 |
| 152644293 | 152708550 | IL6R1 | new design |
| 152644293 | 152708550 | IL6R2 | new design |
| 153,213,753 | 153,218,348 | 336 C 06 | AC027440.2_106016 |
| 153,366,560 | 153,374,010 | 0101F08 | D1S1677i |
| 153,424,924 | 153,429,330 | 0105D10 | D1S0493i |
| 154,700,143 | 154,737,244 | 690D09 | DISD22_0000791 |
| 155,042,659 | 155,053,270 | 478G05 | chr1.fa.O7frz. 178252111 |
| 155,042,659 | 155,053,270 | 099D11 | HUMUT2150 |
| 155,749,791 | 155,788,934 | 478C06 | chr1.fa.O7frz. 178941521 |
| 155,810,163 | 155,834,494 | 0119H03 | D1S1317i |
| 155,982,145 | 156,013,546 | 0115G06 | D1S0009i |
| 155,982,145 | 156,013,546 | 0116G01 | D1S0030i |
| 156,416,361 | 156,421,310 | 0108D08 | D1S0898i |
| 156,526,200 | 156,530,044 | 0117F06 | D1S1291i |
| 157,236,382 | 157,291,569 | 0112G02 | D1S0886i |
| 157,236,382 | 157,291,569 | 0105C04 | Z67479 |
| 157,408,023 | 157,442,914 | 0106F05 | D1S2635 |
| 157,526,128 | 157,544,638 | 0103A11 | D1S1544i |
| 157,948,703 | 157,951,003 | 0101E01 | HUMUT1234 |
| 158,017,346 | 158,018,957 | 0102D01 | D1S1636i |
| 158,163,453 | 158,182,010 | 0119F10 | D1S1617i |
| 158,327,754 | 158,335,103 | 0105B03 | D1S2707 |
| 158,603,481 | 158,609,262 | 0112D03 | D1S0899i |
| 158,721,444 | 158,759,676 | 0117E09 | D1S0505i |
| 158,915,160 | 158,948,265 | 0110H08 | D1S0053i |
| 159,032,552 | 159,064,669 | 0108D02 | D1S0163i |
| 159,231,625 | 159,275,404 | 336E06 | AL354714.7_79226 |
| 159,231,625 | 159,275,404 | 0110F03 | D1S1414i |
| 159,451,693 | 159,457,113 | 0119H10 | D1S1455i |
| 159,466,079 | 159,474,590 | 0110G05 | D1S1463i |
| 159741844 | 159755984 | FCGR2A | new design |
| 159,859,610 | 159,867,620 | 0114F08 | D1S0904i |
| 16,046,946 | 16,139,542 | 0111C09 | D1S0770i |
| 160,631,680 | 160,648,552 | 0107H03 | D1S0515i |
| 163,636,778 | 163,681,057 | 0107C03 | D1S2878 |
| 165,666,501 | 165,754,471 | 0109H06 | D1S0005i |
| 165,666,501 | 165,754,471 | $0115 \mathrm{G11}$ | D1S1554i |
| 165,957,832 | 166,026,684 | 0109C03 | D1S2750 |
| 166,776,626 | 166,779,859 | 0102H01 | D1S0187i |


| 166,812,335 | 166,817,939 | 0115C05 | D1S0109i |
| :---: | :---: | :---: | :---: |
| 167,750,028 | 167,822,450 | 0110A10 | D1S0095i |
| 167,824,661 | 167,866,031 | 0110D11 | D1S0164i |
| 167,824,661 | 167,866,031 | 0104F06 | D1S0168i |
| 167,926,432 | 167,947,463 | 0111B01 | D1S0249i |
| 17,507,277 | 17,563,086 | 0107H09 | D1S1144i |
| 170,894,777 | 170,902,637 | 0110G11 | D1S0194i |
| 170,894,777 | 170,902,637 | 0116D10 | D1S0257i |
| 171,275,723 | 171,286,679 | 0104C12 | D1S0032i |
| 171,275,723 | 171,286,679 | 0117C05 | D1S0271i |
| 171,419,493 | 171,443,094 | 0119F02 | D1S0235i |
| 171,419,493 | 171,443,094 | 0104G09 | D1S1477i |
| 171,713,028 | 171,724,569 | 0101F06 | D1S1070i |
| 172,139,562 | 172,153,139 | 479E02 | chr1.fa.O7frz. 196094961 |
| 173,303,617 | 173,383,825 | 0113 E 10 | D1S0159i |
| 173,558,558 | 173,979,529 | 160E02 | AL021919.4_42117 |
| 173,558,558 | 173,979,529 | 0116G09 | D1S0031i |
| 173,558,558 | 173,979,529 | 0105F06 | D1S0200i |
| 173,558,558 | 173,979,529 | 0111G01 | D1S0265i |
| 173,558,558 | 173,979,529 | 0109H05 | D1S1565 |
| 173,558,558 | 173,979,529 | 104D05 | HUMUT417 |
| 177,085,293 | 177,106,838 | 0112G09 | D1S1153i |
| 177,317,735 | 177,333,653 | 0116G07 | D1S0510i |
| 177,317,735 | 177,333,653 | 0113F11 | D1S0533i |
| 178,390,591 | 178,439,788 | 0113F05 | D1S0517i |
| 178,390,591 | 178,439,788 | 0106 C 11 | D1S2883 |
| 179,269,762 | 179,292,312 | 0115B10 | D1S0542i |
| 18,306,827 | 18,577,563 | 0116A06 | D1S0315i |
| 18,306,827 | 18,577,563 | 0112C12 | D1S1227i |
| 18,306,827 | 18,577,563 | 0102E10 | D1S2826 |
| 18,306,827 | 18,577,563 | 0118E02 | Z67335 |
| 181,075,127 | 181,123,510 | 0110H05 | D1S1464i |
| 181,075,127 | 181,123,510 | 0103B08 | D1S2623 |
| 181,422,022 | 181,480,662 | 0113H02 | D1S0682i |
| 181,422,022 | 181,480,662 | 0112 E 12 | D1S1236i |
| 181,791,320 | 181,826,634 | 0110H06 | D1S1488i |
| 184,532,034 | 184,550,317 | 270C08 | AC074116.5_2995 |
| 184,907,546 | 184,916,179 | 0113 D 07 | D1S0134i |
| 190,871,905 | 190,896,059 | 0101C07 | D1S0261i |
| 190,871,905 | 190,896,059 | 0112G08 | D1S1098i |
| 194,887,631 | 194,983,257 | 0116C10 | D1S0102i |
| 194,887,631 | 194,983,257 | 0118B06 | D1S0250i |
| 195,010,571 | 195,154,386 | 9902C05 | D1S0537i |
| 195,179,520 | 195,194,979 | 0104B03 | D1S1417i |
| 196,874,424 | 196,993,035 | 0101A08 | Z67221 |
| 198,263,353 | 198,413,175 | 0107C11 | D1S0540i |
| 198,859,647 | 198,905,749 | 0115D11 | D1S0687i |
| 198,859,647 | 198,905,749 | 0110F07 | D1S1514i |
| 2479150 | 2486613 | TNF142 | new design |
| 2479150 | 2486613 | TNFRSF14 | new design |
| 20,311,019 | 20,318,637 | 0119B05 | D1S1199i |
| 20,311,019 | 20,318,637 | 0117C02 | D1S1264i |
| 20,311,019 | 20,318,637 | 0105A10 | D1S2843 |
| 20,788,028 | 20,817,988 | 333C12 | AL365439.2_120643 |
| 20,788,028 | 20,817,988 | 079D08 | D1S1571 |
| 200,243,696 | 200,252,939 | 005G10 | D1S2615 |


| 200,382,764 | 200,397,332 | 0109F03 | D1S1727 |
| :---: | :---: | :---: | :---: |
| 200,403,802 | 200,425,104 | 0107D06 | D1S0482i |
| 201,326,405 | 201,403,156 | 0111F05 | D1S0528i |
| 201,326,405 | 201,403,156 | 0104D02 | D1S0693i |
| 201,326,405 | 201,403,156 | 0103D10 | D1S2683 |
| 201,576,375 | 201,587,240 | 0108C09 | D1S0543i |
| 202,000,957 | 202,012,123 | 0113A12 | HUMUT8081B |
| 202,308,866 | 202,363,494 | 005D12 | D1S2668 |
| 202,308,866 | 202,363,494 | 0118D09 | D1S2717 |
| 202,658,379 | 202,726,175 | 9902D07 | D1S1504i |
| 202,752,134 | 202,793,871 | 0105G03 | D1S0955i |
| 203,278,963 | 203,313,761 | 337H04 | AL359927.2_44777 |
| 203,278,963 | 203,313,761 | 0108D05 | D1S1579i |
| 203,833,330 | 203,868,623 | 0113F12 | D1S0523i |
| 204,710,414 | 204,736,846 | 0119B10 | D1S1245i |
| 204,875,504 | 204,924,381 | 0105F10 | D1S1678i |
| 204,924,912 | 204,974,251 | 0110H09 | D1S0094i |
| 205,007,570 | 205,012,462 | IL10G | new design |
| 205,038,838 | 205,082,949 | 0107C07 | D1S2772 |
| 205,105,322 | 205,109,191 | 0111G05 | D1S0548i |
| 205,137,411 | 205,144,107 | 0106C01 | D1S2735 |
| 205,328,810 | 205,339,961 | 006H01 | D1S2727 |
| 205,344,230 | 205,384,940 | 0114A09 | D1S1043i |
| 205,561,476 | 205,600,934 | 0117G09 | D1S0534i |
| 205,694,198 | 205,729,863 | 0115A01 | D1S1205i |
| 205,736,096 | 205,881,733 | 9902G06 | D1S1218i |
| 205,736,096 | 205,881,733 | 0117H06 | D1S1351i |
| 205,736,096 | 205,881,733 | 0109A04 | D1S2796 |
| 205,992,025 | 206,035,481 | 0105E03 | D1S2685 |
| 206,116,942 | 206,151,370 | 0102C01 | D1S0571i |
| 206,116,942 | 206,151,370 | 0103G08 | D1S2692 |
| 206,116,942 | 206,151,370 | 702B02 | DID22N_0002526 |
| 207,854,838 | 207,892,443 | 0115F02 | D1S0017i |
| 207,854,838 | 207,892,443 | 0110F10 | D1S0116i |
| 207,854,838 | 207,892,443 | 0102F11 | D1S471 |
| 208,025,659 | 208,046,102 | 0110C10 | D1S0097i |
| 209,566,580 | 209,614,911 | 0111A06 | D1S0578i |
| 209,566,580 | 209,614,911 | 0103A08 | D1S2810 |
| 210,805,374 | 210,860,742 | 0115H07 | D1S0456i |
| 210,805,374 | 210,860,742 | 0113E12 | D1S0556i |
| 212,597,474 | 212,791,265 | 0103G05 | D1S237 |
| 212,597,474 | 212,791,265 | 0119F01 | D1S419 |
| 212,843,155 | 212,904,537 | 0117D12 | D1S1134i |
| 214,743,211 | 215,377,720 | 0111H04 | D1S0478i |
| 214,743,211 | 215,377,720 | 0112G04 | D1S0944i |
| 214,743,211 | 215,377,720 | 0101C08 | D1S0967i |
| 214,743,211 | 215,377,720 | 0112H06 | D1S1047i |
| 214,743,211 | 215,377,720 | 0112B07 | D1S1053i |
| 214,743,211 | 215,377,720 | 0109D04 | D1S227 |
| 214,743,211 | 215,377,720 | 0118E09 | Z66862 |
| 216,586,200 | 216,684,584 | 0103C08 | D1S1390i |
| 219,119,366 | 219,125,022 | 0113A01 | D1S1593i |
| 219,119,366 | 219,125,022 | 0105D06 | D1S2641 |
| 219,941,389 | 219,982,141 | 0113F03 | D1S0557i |
| 219,941,389 | 219,982,141 | 0102C04 | D1S2894 |
| 219,941,389 | 219,982,141 | 0113H10 | HUMUT7354 |


| 22,021,324 | 22,136,377 | 0110E04 | D1S1431i |
| :---: | :---: | :---: | :---: |
| 22,318,177 | 22,342,197 | 0113F09 | D1S0170i |
| 22,835,705 | 22,838,762 | 333G05 | AC025929.3_119355 |
| 22,835,705 | 22,838,762 | 0106C12 | D1S0006i |
| 22,835,705 | 22,838,762 | 0108H06 | D1S1655i |
| 22,910,045 | 23,114,405 | 0101B02 | D1S2698 |
| 221,350,270 | 221,383,247 | 0110G01 | D1S1361i |
| 224,064,459 | 224,099,884 | 0101H06 | D1S0569i |
| 224,064,459 | 224,099,884 | T001F03 | D1S1230i |
| 224,615,015 | 224,662,414 | 0107G03 | D1S1644 |
| 224,615,015 | 224,662,414 | 0109C06 | Z66645 |
| 224,886,014 | 224,993,647 | 0114C01 | D1S0570i |
| 224,886,014 | 224,993,647 | 0115B12 | D1S1143i |
| 226,711,303 | 226,712,197 | 0106G10 | D1S1344i |
| 227,633,615 | 227,636,468 | 0114D06 | D1S1211i |
| 227,633,615 | 227,636,468 | 0119D12 | D1S1334i |
| 233,890,964 | 234,113,563 | 336E08 | AL390765.5_154842 |
| 233,890,964 | 234,113,563 | 0107H05 | D1S0103i |
| 233,890,964 | 234,113,563 | 0115H09 | D1S0509i |
| 233,890,964 | 234,113,563 | 0119A03 | D1S235 |
| 234,205,753 | 234,303,706 | 0114H05 | D1S0980i |
| 234,205,753 | 234,303,706 | 0114 D 11 | D1S1209i |
| 234,624,303 | 234,714,649 | 0111H05 | D1S0566i |
| 234,624,303 | 234,714,649 | 9902H04 | D1S1680 |
| 234,624,303 | 234,714,649 | 0109E04 | D1S2850 |
| 234,916,422 | 234,994,554 | 0115H01 | D1S1049i |
| 24,318,848 | 24,342,198 | 0119E08 | D1S1285i |
| 24,318,848 | 24,342,198 | 0115F11 | D1S1707i |
| 24,318,848 | 24,342,198 | 0103C07 | D1S1709i |
| 240,078,105 | 240,119,864 | 0102F02 | D1S0204i |
| 241,718,158 | 242,080,053 | 0115F07 | D1S0973i |
| 241,718,158 | 242,080,053 | 0112E11 | D1S1215i |
| 241,718,158 | 242,080,053 | 0109C12 | D1S1335i |
| 241,718,158 | 242,080,053 | 0104E04 | D1S1609 |
| 241,718,158 | 242,080,053 | T002B01 | D1S2811 |
| 245,647,974 | 245,679,033 | 0104B09 | D1S0507i |
| 245,647,974 | 245,679,033 | 0112F05 | D1S0976i |
| 25,098,596 | 25,164,062 | 0115D07 | D1S0052i |
| 25,098,596 | 25,164,062 | 0114E04 | D1S0760i |
| 25,471,568 | 25,529,523 | 0110B04 | D1S1424i |
| 25,561,327 | 25,629,270 | 0110C04 | D1S1428i |
| 26,158,845 | 26,197,235 | 0115H10 | D1S0613i |
| 26,158,845 | 26,197,235 | 0110F04 | D1S1432i |
| 26,516,998 | 26,519,601 | 0110A05 | D1S1438i |
| 27,089,567 | 27,099,549 | 0110E03 | D1S1403i |
| 27,110,566 | 27,113,047 | 002C08 | D1S455 |
| 27,297,893 | 27,366,059 | 0101D08 | D1S0325i |
| 27,297,893 | 27,366,059 | 689G11 | DISD22_0011961 |
| 27,811,162 | 27,834,375 | 0118E04 | D1S0278i |
| 27,811,162 | 27,834,375 | 002F08 | D1S2639 |
| 28,346,264 | 28,392,971 | 0106D12 | D1S1260i |
| 28,346,264 | 28,392,971 | 0108C11 | D1S1443i |
| 29,435,611 | 29,525,899 | 0101D01 | D1S0753i |
| 29,435,611 | 29,525,899 | 0108C06 | D1S1237i |
| 3,763,705 | 3,791,853 | 0116H01 | D1S0321i |
| 31,114,901 | 31,166,301 | 0101D10 | D1S513 |


| 31,114,901 | 31,166,301 | 095G12 | HUMUT2521 |
| :---: | :---: | :---: | :---: |
| 31,610,687 | 31,618,510 | 0111H08 | D1S0754i |
| 31,610,687 | 31,618,510 | 0102B09 | D1S1575i |
| 32,144,609 | 32,176,578 | 0101G03 | D1S0339i |
| 32,144,609 | 32,176,578 | 0113B11 | D1S0608i |
| 32,252,017 | 32,299,037 | 0114F03 | D1S0768i |
| 32,489,480 | 32,524,353 | 0101H07 | D1S0612i |
| 32,489,480 | 32,524,353 | 0105E06 | HUMUT7543 |
| 32,530,274 | 32,571,823 | 0218A07 | D2S1413i |
| 36,704,231 | 36,721,466 | 0210C02 | D2S0082i |
| 36,704,231 | 36,721,466 | 0218F01 | D2S0144i |
| 37,805,004 | 37,834,109 | T002C12 | D2S177 |
| 39,977,117 | 40,002,173 | 0210F02 | D2S0106i |
| 39,977,117 | 40,002,173 | 0203E12 | D2S2238 |
| 40,133,685 | 40,140,274 | 0216G09 | D2S0011i |
| 40,929,829 | 41,009,864 | 0219H10 | D2S1747i |
| 40,929,829 | 41,009,864 | 0220B07 | D2S1830i |
| 41,217,951 | 41,250,815 | 091G10 | HUMUT862 |
| 41,265,461 | 41,480,375 | 0203F12 | D2S1229i |
| 41,265,461 | 41,480,375 | 0213G06 | D2S1251i |
| 42,414,797 | 42,574,135 | 0216C06 | D2S0976i |
| 42,414,797 | 42,574,135 | 0205H06 | D2S2306 |
| 42,896,635 | 42,915,016 | 338B07 | AC013396.4_29951 |
| 43,055,363 | 43,083,247 | 0209F05 | D2S1577i |
| 43,055,363 | 43,083,247 | 0207F07 | D2S1580i |
| 43,576,062 | 43,592,722 | 0212B08 | D2S0968i |
| 43,597,213 | 43,601,461 | 0104F08 | D1S0068i |
| 43,769,134 | 43,861,924 | 0215D02 | D2S0447i |
| 43,769,134 | 43,861,924 | 0202C02 | D2S2294 |
| 44,171,579 | 44,175,499 | 555F11 | chr2.fa.O7frz. 45953438 |
| 45,038,623 | 45,049,479 | 0204G12 | D2S2174 |
| 46,278,399 | 46,371,054 | 0213H06 | D2S1252i |
| 47,037,305 | 47,057,672 | 0210D11 | D2S0428i |
| 47,037,305 | 47,057,672 | 716E12 | DIJ28_10008354 |
| 47,261,827 | 47,289,010 | 0207C11 | D2S1748i |
| 47,454,550 | 47,469,974 | 0212C08 | D2S0969i |
| 47,454,550 | 47,469,974 | 0205A07 | D2S1669i |
| 47,654,331 | 47,656,311 | 0209H05 | D2S1591i |
| 47,674,276 | 47,678,950 | 0212G08 | D2S0975i |
| 50,677,738 | 51,198,524 | 170A08 | AC007560.3_79097 |
| 50,677,738 | 51,198,524 | 0202C12 | D2S0009i |
| 50,677,738 | 51,198,524 | 0221B12 | D2S0056i |
| 50,677,738 | 51,198,524 | 0217F12 | D2S0100i |
| 50,677,738 | 51,198,524 | 0205G02 | D2S0978i |
| 50,677,738 | 51,198,524 | 0208A02 | D2S2316 |
| 56,732,527 | 56,817,845 | 0206H06 | D2S0079i |
| 56,732,527 | 56,817,845 | 0219B03 | D2S1406i |
| 56,732,527 | 56,817,845 | 0213E11 | D2S1414i |
| 56,883,583 | 56,953,596 | 0207G01 | D2S0021i |
| 57,093,065 | 57,156,482 | 0221A09 | D2S378 |
| 59,019,048 | 59,022,587 | 0210B02 | D2S0081i |
| 6,443,798 | 6,502,708 | 0102E07 | D1S1391i |
| 6,443,798 | 6,502,708 | 0105A07 | D1S1448i |
| 60,131,568 | 60,165,050 | 0116F10 | D1S0367i |
| 63,561,300 | 63,563,385 | 0104B01 | HUMUT6372 |
| 65,071,500 | 65,204,775 | 334G11 | AC025866.2_14272 |


| 65,071,500 | 65,204,775 | 266E11 | AL354878.8_58838 |
| :---: | :---: | :---: | :---: |
| 65,071,500 | 65,204,775 | 0101G02 | D1S0640i |
| 65,658,858 | 65,879,830 | 0114C05 | D1S0737i |
| 65,658,858 | 65,879,830 | 0104B05 | D1S1384i |
| 65,658,858 | 65,879,830 | T002E10 | D1S2866 |
| 66,030,781 | 66,612,850 | 0116H10 | D1S0390i |
| 66,030,781 | 66,612,850 | 0114C04 | D1S0713i |
| 66,030,781 | 66,612,850 | 0105E05 | D1S0716i |
| 66,030,781 | 66,612,850 | 0102B02 | D1S0721i |
| 66,030,781 | 66,612,850 | 0105H11 | D1S1484i |
| 67,404,671 | 67,498,250 | 0116E06 | D1S0407i |
| 67,404,671 | 67,498,250 | 0106E05 | D1S1158i |
| 67,545,635 | 67,635,171 | 0107D10 | D1S2806 |
| 7,902,494 | 7,923,513 | 0105G07 | D1S0016i |
| 7,902,494 | 7,923,513 | 0107A08 | D1S0027i |
| 71,090,624 | 71,286,079 | 159F03 | AL031429.11_5207 |
| 71,090,624 | 71,286,079 | 0118 D 07 | D1S0071i |
| 71,090,624 | 71,286,079 | 0118A05 | D1S0119i |
| 71,090,624 | 71,286,079 | 0117G05 | D1S0801i |
| 78,542,156 | 78,778,974 | 0111F10 | D1S0802i |
| 78,542,156 | 78,778,974 | 0107G09 | D1S0812i |
| 78,542,156 | 78,778,974 | T002H10 | D1S2876 |
| 78,888,104 | 78,902,351 | 0113H01 | D1S0408i |
| 78,888,104 | 78,902,351 | 0116G11 | D1S0412i |
| 79,128,037 | 79,279,105 | 0111F11 | D1S0818i |
| 79,128,037 | 79,279,105 | 0116F02 | D1S1196i |
| 84,316,329 | 84,476,769 | 0111H11 | D1S0824i |
| 84,316,329 | 84,476,769 | 0117E02 | D1S1301i |
| 85,504,519 | 85,516,359 | 0114 A 07 | D1S0428i |
| 85,504,519 | 85,516,359 | 0111G10 | D1S0803i |
| 85,556,756 | 85,703,415 | 0117H04 | D1S0035i |
| 85,556,756 | 85,703,415 | 0103D05 | D1S1676i |
| 89,091,203 | 89,129,889 | 0108 D 07 | D1S1380i |
| 89,244,948 | 89,261,132 | $0116 \mathrm{B12}$ | D1S0421i |
| 89,344,403 | 89,414,311 | 0109B07 | D1S0391i |
| 89,498,853 | 89,511,119 | 9902D01 | D1S2004i |
| 9,634,390 | 9,711,564 | 0104B07 | D1S1626i |
| 91,918,488 | 92,144,147 | 0119H02 | D1S0406i |
| 91,918,488 | 92,144,147 | 0112E10 | D1S1185i |
| 91,918,488 | 92,144,147 | 0106C07 | D1S1314i |
| 938,666 | 939,783 | 0105A06 | D1S1425i |
| 94,123,349 | 94,147,600 | 0105A03 | D1S0252i |
| 94,767,369 | 94,779,944 | 268F03 | AL390314.6_23296 |
| 101,899,841 | 101,979,366 | 486G01 | chr10.fa.O7frz. 109393646 |
| 101,899,841 | 101,979,366 | 1010A07 | D10S0301i |
| 103,103,810 | 103,307,068 | 1006F10 | D10S0600i |
| 103,103,810 | 103,307,068 | T001B11 | D10S0605i |
| 103,103,810 | 103,307,068 | 1010E04 | HUMUT7406 |
| 103,519,877 | 103,525,817 | 1004D10 | D10S0764i |
| 104,144,320 | 104,152,271 | 1008D06 | D10S0888i |
| 104,525,996 | 104,566,011 | $1003 \mathrm{B12}$ | D10S1692 |
| 104,580,278 | 104,587,280 | 1002B10 | D10S0155i |
| 11,087,290 | 11,418,680 | 353D02 | AC026887.4_12044 |
| 11,087,290 | 11,418,680 | 1002C02 | D10S0127i |
| 11,087,290 | 11,418,680 | 1008H04 | D10S0169i |
| 11,087,290 | 11,418,680 | 1011A02 | D10S0698i |


| 11,087,290 | 11,418,680 | 1007A03 | D10S0709i |
| :---: | :---: | :---: | :---: |
| 11,087,290 | 11,418,680 | 1001H05 | D10S0715i |
| 112,247,586 | 112,261,292 | 1009 H 06 | D10S0682i |
| 112,317,439 | 112,354,384 | 1009D11 | D10S0516i |
| 112,317,439 | 112,354,384 | 1005B04 | G08785 |
| 115,428,925 | 115,480,654 | 1011F10 | D10S0796i |
| 12,211,642 | 12,251,966 | 1004F04 | D10S0375i |
| 12,211,642 | 12,251,966 | 1001B11 | Z67393 |
| 121,400,872 | 121,427,321 | 212G07 | AF134471.1_33637 |
| 121,400,872 | 121,427,321 | 1002F08 | D10S0120i |
| 122,473,377 | 123,347,962 | 320B08 | AC009989.8_114829 |
| 122,473,377 | 123,347,962 | 591G08 | chr10.fa.O7frz. 13253093 |
| 122,473,377 | 123,347,962 | $578 \mathrm{H05}$ | chr10.fa.O7frz. 133333262 |
| 122,473,377 | 123,347,962 | 1008F09 | D10S0252i |
| 122,473,377 | 123,347,962 | 1008B05 | D10S0255i |
| 122,473,377 | 123,347,962 | 1010A12 | D10S0307i |
| 122,473,377 | 123,347,962 | 1003D07 | D10S0503i |
| 122,473,377 | 123,347,962 | 1006D08 | D10S0539i |
| 122,473,377 | 123,347,962 | 1006H10 | D10S0613i |
| 122,473,377 | 123,347,962 | 1010D12 | D10S0650i |
| 122,473,377 | 123,347,962 | 1007D06 | D10S0756i |
| 122,473,377 | 123,347,962 | 1004E10 | D10S0881i |
| 122,473,377 | 123,347,962 | 1011E05 | D10S0910i |
| 124,310,171 | 124,393,242 | 1009F08 | D10S0089 |
| 124,310,171 | 124,393,242 | 1003A09 | D10S0523i |
| 124,903,783 | 124,914,876 | 1009E10 | D10S0580i |
| 124,903,783 | 124,914,876 | 1007G01 | D10S0858i |
| 127,445,012 | 127,454,380 | 1006B10 | D10S0584i |
| 127,514,896 | 127,575,017 | 1003E10 | D10S0860i |
| 127,690,940 | 128,067,055 | 1003F02 | D10S0696i |
| 127,690,940 | 128,067,055 | 1011C09 | D10S0854i |
| 127,690,940 | 128,067,055 | 1007G08 | D10S0856i |
| 127,690,940 | 128,067,055 | 1007B08 | D10S0861i |
| 129,425,504 | 129,429,440 | T002F01 | D10S217 |
| 129,595,315 | 129,774,155 | 1001D09 | D10S0236i |
| 129,595,315 | 129,774,155 | 1001F08 | D10S0593i |
| 131,155,456 | 131,455,358 | 1006H07 | D10S0530i |
| 131,155,456 | 131,455,358 | 1006A08 | D10S0532i |
| 131,155,456 | 131,455,358 | 1007C09 | D10S0766i |
| 133,631,181 | 133,645,450 | 1003E12 | D10S0871i |
| 134,925,898 | 134,940,362 | 1003E03 | D10S0907i |
| 135,190,857 | 135,224,714 | 1007F12 | D10S0157i |
| 14,979,364 | 15,036,437 | 1006H01 | D10S0376i |
| 14,979,364 | 15,036,437 | 1002G08 | D10S0893i |
| 15,595,954 | 15,802,130 | 1010D05 | D10S0667i |
| 15,595,954 | 15,802,130 | 1010G12 | D10S0672i |
| 15,595,954 | 15,802,130 | 1003A05 | D10S0959 |
| 15,595,954 | 15,802,130 | 1003D06 | D10S1653 |
| 16,595,748 | 16,604,010 | 1002C12 | D10S1477 |
| 16,595,748 | 16,604,010 | 1003 H 05 | D10S674 |
| 17,311,283 | 17,319,598 | 1007A12 | HUMUT463 |
| 17,891,368 | 17,993,184 | 1009E12 | D10S0346i |
| 18,138,358 | 18,240,097 | 1002A09 | D10S0961i |
| 22,650,146 | 22,660,194 | 1003 A03 | D10S0938i |
| 26,767,138 | 26,896,738 | 1008G11 | D10S0179 |
| 26,767,138 | 26,896,738 | 1009C01 | D10S0280i |


| 26,767,138 | 26,896,738 | 1005H03 | G10204 |
| :---: | :---: | :---: | :---: |
| 30,762,872 | 30,790,768 | 1006F09 | D10S0570i |
| 30,762,872 | 30,790,768 | 1011G06 | D10S1674 |
| 31,647,430 | 31,858,748 | 212G02 | AC005877.3_160520 |
| 31,647,430 | 31,858,748 | 212D08 | AF225898.1_164426 |
| 31,647,430 | 31,858,748 | 146F11 | AF225898.1_167611 |
| 31,647,430 | 31,858,748 | 1004G06 | D10S0182i |
| 31,647,430 | 31,858,748 | 1011D07 | D10S208 |
| 31,647,430 | 31,858,748 | 1011F03 | D10S565 |
| 33,229,326 | 33,287,204 | 318C04 | AL133333.9_189168 |
| 33,229,326 | 33,287,204 | 1009C11 | D10S0668i |
| 33,506,426 | 33,665,196 | 1011D05 | D10S0035i |
| 33,506,426 | 33,665,196 | 1005E06 | D10S0071i |
| 35,455,807 | 35,541,892 | 1008G09 | D10S0201i |
| 35,455,807 | 35,541,892 | 1008A12 | D10S0420i |
| 35,455,807 | 35,541,892 | 1001B08 | HUMUT5962 |
| 44,185,611 | 44,200,548 | 1003D03 | D10S0927i |
| 45,189,635 | 45,261,571 | 483G05 | chr10.fa.O7frz.44957867 |
| 45,189,635 | 45,261,571 | 1008B11 | D10S0203i |
| 48,926,216 | 49,033,022 | 1004C01 | D10S0911i |
| 49,184,739 | 49,317,409 | 419G02 | chr10.fa.O7frz. 48816071 |
| 49,184,739 | 49,317,409 | 1008D07 | D10S0556i |
| 54,195,146 | 54,201,466 | 1005A12 | D10S0330i |
| 59,764,745 | 59,800,515 | 1002E01 | D10S0909i |
| 59,764,745 | 59,800,515 | 1005F03 | Z67552 |
| 6,034,340 | 6,060,156 | 1006C09 | D10S0559 |
| 6,092,658 | 6,144,294 | 1009A11 | D10S0177i |
| 6,509,111 | 6,662,269 | 1010B12 | D10S0063i |
| 6,509,111 | 6,662,269 | 1011E11 | D10S0096i |
| 6,509,111 | 6,662,269 | 1002C04 | D10S0951i |
| 61,458,165 | 61,819,494 | 1006F04 | D10S0460i |
| 61,458,165 | 61,819,494 | 1006E08 | D10S0549 |
| 61,458,165 | 61,819,494 | 1001E08 | D10S0892i |
| 61,458,165 | 61,819,494 | 1007F01 | D10S0919i |
| 61,458,165 | 61,819,494 | 1002E12 | D10S0971i |
| 62,205,690 | 62,224,616 | 1006D11 | D10S0627i |
| 64,241,762 | 64,246,133 | 1007B11 | D10S0087i |
| 64,241,762 | 64,246,133 | 1006D01 | D10S0351i |
| 70,331,040 | 70,376,609 | 1002E02 | D10S1678 |
| 70,517,834 | 70,534,573 | 578A01 | chr10.fa.O7frz. 73903894 |
| 72,027,110 | 72,032,521 | 1003G12 | D10S537 |
| 74,866,192 | 74,925,765 | 1010D11 | D10S0186i |
| 74,866,192 | 74,925,765 | 1007E01 | D10S0781i |
| 75,340,896 | 75,347,261 | 603A10 | chr10.fa.O7frz. 79381928 |
| 75,580,971 | 76,139,067 | 1005C12 | D10S0335i |
| 75,580,971 | 76,139,067 | 1005G12 | D10S0345i |
| 75,580,971 | 76,139,067 | 1009D08 | D10S0356i |
| 75,580,971 | 76,139,067 | 1006G03 | D10S0445i |
| 75,580,971 | 76,139,067 | 1010C09 | D10S0637i |
| 75,580,971 | 76,139,067 | 1006G12 | D10S0660i |
| 76,524,196 | 76,538,976 | 1005B12 | D10S0332i |
| 79,220,557 | 79,356,384 | 1001B04 | D10S0603i |
| 79,220,557 | 79,356,384 | 1009C06 | D10S0648i |
| 8,136,662 | 8,157,170 | 1009C03 | D10S0158i |
| 8,136,662 | 8,157,170 | 9905E11 | D10S0380i |
| 80,777,226 | 80,785,096 | 1001B03 | D10S0304i |


| 80,777,226 | 80,785,096 | 1002D08 | D10S201 |
| :---: | :---: | :---: | :---: |
| 81,687,476 | 81,698,841 | 9905F11 | D10S0459i |
| 83,624,786 | 84,736,913 | 1003F08 | D10S0039i |
| 83,624,786 | 84,736,913 | 1001F05 | D10S0072i |
| 83,624,786 | 84,736,913 | 1005B07 | D10S0097i |
| 83,624,786 | 84,736,913 | 1007B10 | D10S0109i |
| 83,624,786 | 84,736,913 | 1005D09 | D10S0218i |
| 83,624,786 | 84,736,913 | 1007H11 | D10S0239i |
| 83,624,786 | 84,736,913 | 1008C01 | D10S0467i |
| 83,624,786 | 84,736,913 | 1006F05 | D10S0475i |
| 83,624,786 | 84,736,913 | 1003C09 | D10S0487i |
| 83,624,786 | 84,736,913 | 1006E11 | D10S0629i |
| 83,624,786 | 84,736,913 | 1007E06 | D10S0731i |
| 83,624,786 | 84,736,913 | 1004E01 | D10S0750i |
| 83,624,786 | 84,736,913 | 1001E11 | D10S0960i |
| 83,624,786 | 84,736,913 | 1001E09 | D10S1786 |
| 88,506,376 | 88,674,925 | 1008C12 | D10S0585i |
| 89,612,850 | 89,721,667 | 1001C07 | D10S1765 |
| 89,612,850 | 89,721,667 | 1005D03 | Z67254 |
| 90,739,206 | 90,765,522 | 1001F09 | D10S1739 |
| 90,963,306 | 91,164,294 | 1008E02 | D10S0221i |
| 90,963,306 | 91,164,294 | 1006D06 | D10S0485i |
| 90,963,306 | 91,164,294 | 9906A02 | D10S0771i |
| 94,811,011 | 94,818,444 | 1010H09 | D10S0671i |
| 96,433,368 | 96,485,937 | 1005E04 | HUMUT925 |
| 96512371 | 96603007 | CYP2C191 | new design |
| 96512371 | 96603007 | CYP2C192 | new design |
| 96688418 | 96739137 | CYP2C9 | new design |
| 96,786,519 | 96,819,244 | 1006B06 | D10S0481i |
| 97,461,526 | 97,619,442 | 1003D10 | D10S0154i |
| 97,461,526 | 97,619,442 | 1001D06 | D10S0899i |
| 97,461,526 | 97,619,442 | 1011A08 | D10S0953i |
| 97,941,445 | 98,021,316 | 1005C10 | D10S0254i |
| 98,054,075 | 98,088,311 | 1001D03 | D10S0939i |
| 98,054,075 | 98,088,311 | 1002H01 | D10S0946i |
| 1,730,558 | 1,741,798 | 1108B01 | D11S0457i |
| 1,730,558 | 1,741,798 | 1111E09 | D11S0827i |
| 1,830,776 | 1,870,069 | 1101A06 | D11S0967i |
| 10,283,172 | 10,285,499 | 399D09 | AC018539.4_114043 |
| 100,414,313 | 100,506,465 | 1106G10 | D11S0557i |
| 100,414,313 | 100,506,465 | 1104H02 | D11S0935i |
| 101,693,404 | 101,713,675 | 1109G04 | D11S0870i |
| 101,896,449 | 101,906,688 | $1110 \mathrm{G10}$ | HUMUT1283 |
| 101,952,776 | 102,001,273 | 1110B12 | D11S4108 |
| 102,067,625 | 102,081,678 | 1105B12 | D11S0258i |
| 102,146,444 | 102,156,569 | 1105E12 | D11S0269i |
| 102,211,738 | 102,219,552 | 1104G07 | Z66956 |
| 102,318,934 | 102,331,672 | 1109H10 | D11S0391i |
| 102,318,934 | 102,331,672 | 1102G06 | D11S0599i |
| 104,261,876 | 104,274,607 | 1106H10 | D11S0558i |
| 104,318,804 | 104,345,373 | 1105D02 | D11S1886 |
| 104,370,180 | 104,384,909 | 1107A05 | D11S0774i |
| 104,513,879 | 104,515,663 | 1107F02 | D11S0764i |
| 104,513,879 | 104,515,663 | 1109B11 | HUMUT2064 |
| 106,878,664 | 106,941,637 | 1102C11 | D11S0559i |
| 106,878,664 | 106,941,637 | 1103C02 | D11S0944i |


| 107,598,769 | 107,745,036 | 1108F07 | D11S0615i |
| :---: | :---: | :---: | :---: |
| 107,598,769 | 107,745,036 | 1109G09 | D11S0619i |
| 107,598,769 | 107,745,036 | 1111 E 02 | D11S0622i |
| 107,598,769 | 107,745,036 | 9906E05 | D11S0848i |
| 108,041,014 | 108,316,866 | 1109C07 | D11S0404i |
| 108,041,014 | 108,316,866 | 1102A06 | D11S0409i |
| 108,041,014 | 108,316,866 | 1104A09 | D11S0952i |
| 109,605,376 | 109,672,647 | 1103G06 | D11S0285i |
| 109,605,376 | 109,672,647 | 037 E 05 | D11S927 |
| 110,728,190 | 110,755,627 | 1108A07 | D11S0278i |
| 110,728,190 | 110,755,627 | 1106C02 | D11S0312i |
| 111,519,186 | 111,540,050 | 1107B01 | D11S0018i |
| 111,519,186 | 111,540,050 | 1101D05 | D11S0894i |
| 112,337,368 | 112,653,781 | 1106B10 | D11S0541i |
| 112,337,368 | 112,653,781 | 1105G02 | D11S3179 |
| 112,337,368 | 112,653,781 | 1105B03 | Z67379 |
| 114,550,227 | 114,880,325 | 1108E09 | D11S0537i |
| 114,550,227 | 114,880,325 | 1109H05 | D11S0542i |
| 114,550,227 | 114,880,325 | T001D11 | D11S0568i |
| 114,550,227 | 114,880,325 | 1110C08 | D11S1885 |
| 114,550,227 | 114,880,325 | 684 E 12 | DISD22_0006557 |
| 114,550,227 | 114,880,325 | 1102D05 | Z67490 |
| 117,362,319 | 117,377,404 | 1104H05 | D11S1356 |
| 117,362,319 | 117,377,404 | 111D09 | U73649.1_16072 |
| 117,680,662 | 117,692,100 | 403B09 | AC068591.2_86017 |
| 117,710,475 | 117,718,669 | 403 A09 | AP001582.3_115137 |
| 118,125,623 | 118,167,082 | 1108G11 | D11S0300i |
| 118,125,623 | 118,167,082 | 1110B06 | D11S4104 |
| 118,259,777 | 118,272,181 | 704C08 | DID22N_0040945 |
| 118,259,777 | 118,272,181 | 745B10 | DIJ28_10013816 |
| 118,684,444 | 118,693,050 | 1109C12 | D11S0315i |
| 119,014,018 | 119,104,645 | 1101D10 | D11S0560i |
| 119,014,018 | 119,104,645 | 1107B04 | D11S0857i |
| 119,616,256 | 119,695,863 | 1108B06 | D11S0563i |
| 119,616,256 | 119,695,863 | 1111B03 | D11S0632i |
| 119,616,256 | 119,695,863 | 1103 HO 2 | Z67522 |
| 120,828,130 | 121,005,621 | 1107A01 | D11S0765i |
| 120,828,130 | 121,005,621 | 1107F05 | D11S0778i |
| 125,279,550 | 125,298,215 | 1106E03 | D11S0341i |
| 125,279,550 | 125,298,215 | 1108F01 | D11S0427i |
| 125,658,192 | 125,672,683 | 1106D11 | D11S0573i |
| 125,658,192 | 125,672,683 | 037H11 | D11S934 |
| 13,470,177 | 13,474,143 | 1102A12 | D11S926 |
| 133,290,395 | 133,327,321 | 1109A10 | D11S0418i |
| 133,290,395 | 133,327,321 | 1103E01 | D11S968 |
| 133,290,395 | 133,327,321 | 1101H11 | Z67178 |
| 133,444,030 | 133,526,861 | 1107C03 | D11S0779 |
| 133,444,030 | 133,526,861 | 1104G11 | D11S4125 |
| 133,753,608 | 133,787,022 | 1104B12 | D11S0676i |
| 14,856,131 | 14,870,327 | 1111E01 | D11S0104i |
| 17,067,861 | 17,147,864 | 1111C04 | D11S0661i |
| 17,067,861 | 17,147,864 | 1103F07 | D11S4160 |
| 18,090,596 | 18,094,695 | 1109E03 | D11S0687i |
| 18,223,365 | 18,226,758 | 1109G08 | D11S0663i |
| 18,223,365 | 18,226,758 | 1111G09 | D11S0834i |
| 18,300,719 | 18,345,153 | 1107F07 | D11S0840i |


| 18,682,435 | 18,704,353 | 1106D07 | D11S0451i |
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| 18,706,051 | 18,769,965 | 740D01 | DIJ28_10032405 |
| 2,106,918 | 2,125,616 | 1102D03 | D11S0973i |
| 2,355,096 | 2,375,225 | 1106 A07 | D11S0445i |
| 2,355,096 | 2,375,225 | 1105B01 | D11S1318 |
| 2,861,019 | 2,863,577 | 085D12 | HUMUT6724B |
| 20,365,679 | 20,487,404 | 1105C06 | D11S0094i |
| 20,365,679 | 20,487,404 | 1110B10 | D11S4190 |
| 22,646,230 | 22,791,123 | $1107 \mathrm{C08}$ | D11S0353i |
| 22,646,230 | 22,791,123 | 1106C04 | D11S0356i |
| 22,646,230 | 22,791,123 | $1101 \mathrm{E12}$ | D11S0363i |
| 268,570 | 275,304 | 1108H05 | D11S0006i |
| 268,570 | 275,304 | 1101A04 | D11S0978i |
| 27,633,016 | 27,699,872 | 1111D01 | D11S0077i |
| 27,633,016 | 27,699,872 | 1110C05 | D11S0144i |
| 3,622,937 | 3,642,222 | 1104D09 | D11S0838i |
| 33,681,132 | 33,714,600 | 1107C12 | D11S0166i |
| 33,681,132 | 33,714,600 | 1111A02 | D11S0460i |
| 34,417,054 | 34,450,183 | 1111C07 | D11S0739 |
| 35,116,993 | 35,210,525 | 1105A04 | D11S0012i |
| 35,116,993 | 35,210,525 | 1105C04 | D11S0020i |
| 35,116,993 | 35,210,525 | 1105H07 | D11S0139i |
| 36,467,299 | 36,488,398 | T003H08 | D11S4083 |
| 36,546,139 | 36,557,877 | 1101E02 | G10015 |
| 395716 | 407397 | SIGIRR1 | new design |
| 395716 | 407397 | SIGIRR2 | new design |
| 4,745,076 | 4,970,235 | 1105E06 | D11S0102i |
| 4,745,076 | 4,970,235 | 1108E08 | D11S0105i |
| 4,745,076 | 4,970,235 | 1108H06 | D11S0582i |
| 4,745,076 | 4,970,235 | 1103D12 | D11S4181 |
| 43,290,109 | 43,322,655 | 1110C11 | D11S0722i |
| 43,858,971 | 43,898,392 | 591F11 | chr11.fa.O7frz. 46072222 |
| 44,543,717 | 44,597,915 | 1109F02 | D11S0504i |
| 44,543,717 | 44,597,915 | 1109A08 | D11S0639 |
| 45,863,778 | 45,884,592 | 1107D05 | D11S0728i |
| 45,863,778 | 45,884,592 | 1102G01 | D11S0737i |
| 46,697,331 | 46,717,631 | 1110C07 | D11S0817i |
| 46,697,331 | 46,717,631 | 1102B06 | D11S0970i |
| 47,227,083 | 47,246,972 | 1106D04 | D11S0359i |
| 47,227,083 | 47,246,972 | 1102B01 | D11S0692i |
| 47,543,464 | 47,562,690 | 400B11 | AC067943.4_5205 |
| 47,543,464 | 47,562,690 | 1108A08 | D11S0366i |
| 47,567,792 | 47,580,516 | 1108F02 | D11S0492i |
| 47,958,689 | 48,146,246 | 1103 F 12 | D11S1350 |
| 47,958,689 | 48,146,246 | 1101A10 | D11S1784 |
| 5,203,270 | 5,212,454 | 1105A05 | D11S0069i |
| 5,203,270 | 5,212,454 | 1111C09 | D11S0819i |
| 5,667,495 | 5,688,669 | 487E06 | chr11.fa.O7frz. 5327809 |
| 5,667,495 | 5,688,669 | 1108 A 02 | D11S0447i |
| 5,667,495 | 5,688,669 | 1110D01 | D11S0453i |
| 56,757,630 | 56,761,489 | 1105A09 | D11S0182i |
| 56,862,525 | 56,894,125 | 1102A01 | D11S0694i |
| 56,910,832 | 56,914,706 | 1101E03 | Z67514 |
| 57,075,705 | 57,092,333 | T002G05 | D11S0635i |
| 57,121,603 | 57,138,902 | 1101E05 | D11S0598i |
| 57,121,603 | 57,138,902 | 1101B09 | D11S1777 |


| 59,580,677 | 59,595,164 | 1108C05 | D11S0872i |
| :---: | :---: | :---: | :---: |
| 59,953,638 | 59,971,841 | 1104D05 | D11S0690i |
| 6,581,540 | 6,588,677 | 399A08 | AC009796.4_61451 |
| 6,581,540 | 6,588,677 | 420 HO 3 | chr11.fa.O7frz.6316475 |
| 6,581,540 | 6,588,677 | 1109 A02 | D11S0577i |
| 60,374,983 | 60,380,020 | 1111 C 12 | D11S0927i |
| 60,626,543 | 60,651,900 | 1109F05 | D11S0506i |
| 61,039,361 | 61,104,874 | 1102A08 | D11S0075i |
| 61,039,361 | 61,104,874 | 1110F06 | D11S4076 |
| 61,316,726 | 61,321,286 | 1109H08 | D11S0685i |
| 62,316,219 | 62,329,529 | 1102G04 | D11S0785i |
| 62,380,094 | 62,412,929 | 1107F03 | D11S0703i |
| 63,030,132 | 63,040,815 | 1101E08 | D11S0702i |
| 63,758,646 | 63,762,835 | 1109G02 | D11S0640i |
| 63,764,989 | 63,768,262 | 1111D08 | D11S0787i |
| 63,829,616 | 63,840,786 | 1109E08 | D11S0644i |
| 64,250,959 | 64,269,504 | 1106F05 | D11S0405i |
| 64,458,519 | 64,459,936 | 563E04 | chr11.fa.O7frz.68864733 |
| 64,458,519 | 64,459,936 | 682B03 | DISO7_10017975 |
| 64,786,006 | 64,821,664 | 1108G05 | D11S0645i |
| 65,062,850 | 65,082,275 | 1101H05 | D11S0388i |
| 65,062,850 | 65,082,275 | 1102B05 | HUMUT2269 |
| 65,177,649 | 65,186,959 | 1105A10 | D11S0200i |
| 65,378,858 | 65,383,462 | 401H03 | AP001191.2_77147 |
| 65,378,858 | 65,383,462 | 1105D10 | D11S0204i |
| 65,838,534 | 65,841,091 | 1108G01 | D11S0393i |
| 65,990,974 | 66,001,382 | 1102F04 | D11S0887i |
| 66,070,967 | 66,087,373 | 1108B05 | D11S0406i |
| 66,087,511 | 66,092,623 | 1110F05 | D11S0818i |
| 66,580,897 | 66,596,060 | 1109D02 | D11S0389 |
| 66,580,897 | 66,596,060 | 1101G01 | Z67088 |
| 66,888,215 | 66,897,782 | 1109H09 | D11S0190i |
| 66,888,215 | 66,897,782 | 1103A04 | D11S0920i |
| 67,007,097 | 67,015,150 | 1102A02 | D11S1889 |
| 67,107,862 | 67,110,701 | 1103D10 | D11S4155 |
| 67,515,151 | 67,528,169 | 1101C11 | D11S0681i |
| 67,563,059 | 67,574,942 | 1108G07 | D11S0871i |
| 67,836,674 | 67,973,317 | 1111G01 | D11S0205i |
| 67,836,674 | 67,973,317 | 1107B11 | D11S0212i |
| 67,836,674 | 67,973,317 | 1104A01 | D11S1337 |
| 67,836,674 | 67,973,317 | 1110F11 | D11S4178 |
| 67,836,674 | 67,973,317 | 1104C06 | HUMUT5620 |
| 69,165,054 | 69,178,423 | 1111D06 | D11S0717i |
| 69,296,978 | 69,299,352 | 036B08 | D11S4136 |
| 69,726,917 | 69,731,144 | 1105H03 | HUMUT6360 |
| 7,016,373 | 7,049,333 | 1111D09 | D11S0823i |
| 7,016,373 | 7,049,333 | 1111F09 | D11S0828i |
| 7,937,547 | 7,941,780 | 1108C06 | D11S0883i |
| 7,937,547 | 7,941,780 | 726C10 | DIJ28_10040944 |
| 71,221,894 | 71,226,256 | 1106D12 | D11S0610i |
| 71,387,587 | 71,394,409 | 1102E10 | D11S0651i |
| 72,606,992 | 72,625,045 | 1108F06 | D11S0491i |
| 72,606,992 | 72,625,045 | 1107 A 03 | D11S0720i |
| 72,606,992 | 72,625,045 | 1107G02 | D11S0725i |
| 72,765,053 | 72,786,167 | 1101D03 | D11S0691i |
| 73,023,592 | 73,051,512 | 1104C05 | D11S916 |


| 73,981,277 | 74,031,413 | 1103 A08 | D11S0588i |
| :---: | :---: | :---: | :---: |
| 74,654,130 | 74,740,521 | 1111D04 | D11S0667i |
| 74,654,130 | 74,740,521 | 1102E06 | D11S0964i |
| 75,203,923 | 75,531,342 | 1108B09 | D11S0641i |
| 75,203,923 | 75,531,342 | 1102D06 | D11S0893i |
| 75,203,923 | 75,531,342 | 1108 F 12 | D11S0929i |
| 75,203,923 | 75,531,342 | 1104A11 | D11S1321 |
| 75,738,652 | 75,769,528 | 1104E06 | D11S0174i |
| 75,738,652 | 75,769,528 | 1109 A07 | D11S0515i |
| 76,710,708 | 76,862,581 | 401D09 | AP000486.4_115712 |
| 76,710,708 | 76,862,581 | 1103 A 07 | D11S4186 |
| 77,603,990 | 77,806,414 | 1109G10 | D11S0522i |
| 77,603,990 | 77,806,414 | 1102F03 | D11S0961i |
| 77,603,990 | 77,806,414 | 1103 B 10 | D11S937 |
| 822952 | 828835 | CD151 | new design |
| 86,334,369 | 86,344,081 | 1109A06 | D11S0221i |
| 86,334,369 | 86,344,081 | 1111C11 | D11S0903i |
| 88,550,268 | 88,668,474 | 1108H03 | D11S0254i |
| 88,550,268 | 88,668,474 | 1101G07 | D11S0260i |
| 88,550,268 | 88,668,474 | 1106G09 | D11S0529i |
| 88,699,160 | 88,864,301 | 1105E10 | D11S0223i |
| 93,790,114 | 93,866,688 | 1101A11 | D11S0226i |
| 93,790,114 | 93,866,688 | 1106H09 | D11S0532i |
| 10,015,281 | 10,029,461 | 1205C11 | D12S0288i |
| 10,015,281 | 10,029,461 | 1203D03 | D12S1674 |
| 10,113,421 | 10,142,872 | 1203F09 | D12S1690 |
| 10,202,167 | 10,216,004 | 1211 E02 | D12S1696 |
| 10,202,167 | 10,216,004 | 1203C08 | D12S77 |
| 10,351,684 | 10,359,983 | 1204B09 | D12S0390i |
| 10,456,181 | 10,464,461 | 1209H09 | D12S0218i |
| 10,633,039 | 10,643,431 | 1203F08 | D12S0095i |
| 10,742,955 | 10,767,171 | 1209C08 | D12S0140i |
| 101,313,806 | 101,398,471 | 1205C05 | D12S0036i |
| 101,313,806 | 101,398,471 | 1205F05 | D12S0043i |
| 101,875,594 | 101,878,421 | 1204A12 | D12S0533i |
| 101,875,594 | 101,878,421 | 9907E01 | D12S0929i |
| 102,505,181 | 102,684,635 | 357C05 | AC063946.10_124699 |
| 102,505,181 | 102,684,635 | 9906B10 | D12S0841i |
| 102,505,181 | 102,684,635 | $1207 \mathrm{B03}$ | D12S0848i |
| 102,505,181 | 102,684,635 | 9906H05 | D12S360 |
| 102,505,181 | 102,684,635 | 1211B02 | D12S865 |
| 102,848,290 | 102,865,833 | 099H02 | HUMUT835B |
| 103,034,988 | 103,056,170 | 565G02 | ehr12.fa.O7frz. 11199728 |
| 103,034,988 | 103,056,170 | 1210G04 | D12S338 |
| 103,204,857 | 103,268,192 | 1209H03 | D12S0467i |
| 105,501,163 | 105,680,711 | 1209B08 | D12S0093i |
| 105,501,163 | 105,680,711 | 1201C12 | D12S0574i |
| 105,501,163 | 105,680,711 | 1202G05 | D12S330 |
| 107,208,800 | 107,257,218 | 1202A05 | D12S1605 |
| 107,539,800 | 107,551,799 | 1204B04 | D12S84 |
| 108,010,379 | 108,015,660 | 223A08 | AC007637.9_40136 |
| 108,010,379 | 108,015,660 | 668F08 | ehr12.fa.O7frz. 117355013 |
| 108,200,167 | 108,231,408 | 1210H01 | D12S0004i |
| 108,200,167 | 108,231,408 | 1210C05 | D12S0070i |
| 109,424,388 | 109,454,274 | 1204A03 | D12S0030i |
| 109,424,388 | 109,454,274 | 1210H05 | D12S0174i |


| 11,694,055 | 11,939,603 | 1204B01 | D12S0024i |
| :---: | :---: | :---: | :---: |
| 11,694,055 | 11,939,603 | 1205C09 | D12S0172i |
| 11,694,055 | 11,939,603 | T003B10 | D12S89 |
| 11,694,055 | 11,939,603 | 1202E01 | D12S98 |
| 111,340,919 | 111,432,100 | 1207F10 | D12S0747i |
| 111,340,919 | 111,432,100 | 1208A06 | HUMUT5428 |
| 111,829,122 | 111,854,374 | 040C04 | D12S1340 |
| 111,980,045 | 112,020,216 | 1205G11 | D12S0309i |
| 111,980,045 | 112,020,216 | 1201F08 | D12S0483i |
| 115,783,410 | 115,803,615 | 1207E05 | D12S0781i |
| 115,783,410 | 115,803,615 | 1207D06 | D12S0882i |
| 116,135,362 | 116,283,965 | 1208H09 | D12S0177i |
| 116,135,362 | 116,283,965 | 9906H11 | D12S0749 |
| 116,135,362 | 116,283,965 | 9906A12 | D12S0761i |
| 118,100,978 | 118,116,934 | 1202A06 | D12S0756i |
| 118,100,978 | 118,116,934 | 658E07 | DISO7_10003646 |
| 119,942,478 | 119,961,164 | 1211G05 | D12S0076i |
| 119,942,478 | 119,961,164 | 1201B03 | D12S0573i |
| 12,115,145 | 12,255,214 | 1209E11 | D12S0050i |
| 12,115,145 | 12,255,214 | 1205C10 | D12S0217i |
| 12,164,953 | 12,311,013 | 1204H10 | D12S391 |
| 12,520,098 | 12,606,584 | 1208 E 12 | D12S0041i |
| 12,520,098 | 12,606,584 | 1203 A 07 | D12S0983i |
| 12,520,098 | 12,606,584 | T003F09 | D12S358 |
| 12,770,130 | 12,874,182 | 1204A02 | D12S0120i |
| 120,055,061 | 120,108,259 | 357F09 | AC069209.16_27083 |
| 120,055,061 | 120,108,259 | 1209D12 | D12S0011i |
| 121,222,530 | 121,224,699 | 1206G02 | D12S0378i |
| 121,321,934 | 121,473,069 | 9906H08 | D12S0382i |
| 121,321,934 | 121,473,069 | 9906B09 | D12S0550i |
| 121,765,256 | 121,767,297 | 9906G09 | D12S0767i |
| 122,652,625 | 122,671,435 | 1210A09 | D12S0307i |
| 122,684,333 | 122,711,573 | 1210F12 | D12S0606i |
| 123,374,914 | 123,568,793 | 357B10 | AC027706.2_46645 |
| 123,374,914 | 123,568,793 | 1209H07 | D12S0808i |
| 123,374,914 | 123,568,793 | T001A02 | D12S0811i |
| 123,374,914 | 123,568,793 | 1201A12 | D12S1611 |
| 123,374,914 | 123,568,793 | 040A07 | D12S1612 |
| 123,828,129 | 123,914,346 | 1207B12 | D12S0751i |
| 123,828,129 | 123,914,346 | 1207E08 | D12S0884i |
| 123,997,325 | 124,039,620 | 1210C12 | D12S0768i |
| 128,063,805 | 128,067,640 | 1210D07 | D12S0785i |
| 129,212,957 | 129,216,238 | 1206C10 | D12S0576i |
| 129,212,957 | 129,216,238 | 1201G12 | D12S0921i |
| 129,922,521 | 129,927,316 | 1204D08 | D12S0818i |
| 131,705,476 | 131,709,045 | 1204D01 | D12S357 |
| 14,873,512 | 14,887,680 | 1205A07 | D12S0089 |
| 14,873,512 | 14,887,680 | 1209F12 | D12S0130i |
| 14,986,217 | 15,005,870 | 1204C01 | D12S0105i |
| 14,986,217 | 15,005,870 | 1206H03 | D12S0399i |
| 15,366,754 | 15,641,602 | 1206B04 | D12S0404i |
| 15,366,754 | 15,641,602 | 1201 E 10 | D12S0948i |
| 15,366,754 | 15,641,602 | 1201E07 | D12S0974i |
| 15,366,754 | 15,641,602 | 1205E03 | G08975 |
| 18,305,741 | 18,692,617 | 1206D03 | D12S0392i |
| 18,305,741 | 18,692,617 | 1204H11 | D12S0629i |


| 18,305,741 | 18,692,617 | 1207H10 | D12S0849i |
| :---: | :---: | :---: | :---: |
| 18,305,741 | 18,692,617 | 1207B08 | D12S0855i |
| 18,305,741 | 18,692,617 | 1210G07 | D12S0859i |
| 2,774,414 | 2,783,385 | 1205C08 | D12S0129i |
| 2,774,414 | 2,783,385 | 1201E09 | D12S0132i |
| 25,249,447 | 25,295,121 | 1204C10 | D12S0103i |
| 25,249,447 | 25,295,121 | 1203H11 | D12S0922i |
| 26,377,193 | 26,877,398 | 1205E12 | D12S0325i |
| 26,377,193 | 26,877,398 | 1206E01 | D12S0354i |
| 26,377,193 | 26,877,398 | 1204A10 | D12S0523i |
| 26,377,193 | 26,877,398 | 1207D04 | D12S0648i |
| 26,377,193 | 26,877,398 | 9906C11 | D12S0658i |
| 26,377,193 | 26,877,398 | 1202C05 | D12S0667i |
| 26,377,193 | 26,877,398 | 1207B05 | D12S0671i |
| 26,377,193 | 26,877,398 | 1207C10 | D12S0679i |
| 28,002,284 | 28,016,183 | 221B11 | AC008011.11_31766 |
| 28,002,284 | 28,016,183 | 1203D06 | D12S0923i |
| 28,002,284 | 28,016,183 | 1211D01 | HUMUT7594 |
| 29,381,556 | 29,425,410 | 1205E06 | D12S0069i |
| 29,381,556 | 29,425,410 | 1208B03 | D12S0215i |
| 3,470,686 | 3,573,400 | 1205C07 | D12S0096i |
| 3,470,686 | 3,573,400 | 1201D03 | D12S1050 |
| 3,470,686 | 3,573,400 | 1201G01 | D12S1062 |
| 31,118,061 | 31,148,992 | 1201C09 | D12S0047i |
| 31,118,061 | 31,148,992 | 1204C07 | D12S0417i |
| 4,253,199 | 4,284,777 | 1203A10 | D12S1725 |
| 4,413,569 | 4,425,041 | 1205G09 | D12S0185i |
| 4,569,505 | 4,593,302 | 1205H07 | D12S0117i |
| 4,569,505 | 4,593,302 | 1201F07 | D12S0191i |
| 42,439,047 | 42,468,166 | 1206E10 | D12S0580i |
| 42,439,047 | 42,468,166 | 1205 H 01 | D12S1663 |
| 44,601,459 | 44,670,615 | 1209B03 | D12S0225i |
| 44,601,459 | 44,670,615 | 1209E05 | D12S0296i |
| 46,462,772 | 46,499,924 | 1205H06 | D12S0086i |
| 46,462,772 | 46,499,924 | 1207E06 | D12S0661i |
| 47,509,806 | 47,532,224 | 1202B05 | D12S0593i |
| 47,509,806 | 47,532,224 | 1209H01 | D12S0596i |
| 47,658,503 | 47,662,746 | 9906C09 | D12S0577i |
| 47,769,471 | 47,774,869 | 1208B05 | D12S0578i |
| 47,864,847 | 47,869,153 | 432G04 | chr12.fa.O7frz. 51899683 |
| 48,012,467 | 48,017,238 | 1203G08 | D12S1627 |
| 48,238,352 | 48,248,178 | 1206A01 | D12S0333i |
| 49,444,128 | 49,500,328 | 9906E06 | D12S1135i |
| 49,444,128 | 49,500,328 | 9906F06 | D12S1137i |
| 49,444,128 | 49,500,328 | 9906G07 | D12S1214i |
| 5,928,301 | 6,104,097 | 1210A05 | D12S0009i |
| 5,928,301 | 6,104,097 | 1205A10 | D12S0199i |
| 5,928,301 | 6,104,097 | 1203E01 | D12S0939i |
| 5,928,301 | 6,104,097 | 1203 A02 | D12S0966i |
| 5,928,301 | 6,104,097 | 1201H05 | D12S374 |
| 50,008,494 | 50,026,730 | 1202H07 | D12S0865i |
| 51,167,231 | 51,173,289 | 1210D01 | G08922 |
| 51,577,238 | 51,585,127 | 1211E09 | D12S0653i |
| 51,577,238 | 51,585,127 | 9906C10 | D12S0877i |
| 51,871,374 | 51,887,267 | 1208F05 | D12S0430i |
| 51,890,621 | 51,912,253 | 1201B11 | D12S0890i |


| 52,060,246 | 52,096,497 | 1201F06 | D12S0662i |
| :---: | :---: | :---: | :---: |
| 52,060,246 | 52,096,497 | 1202B09 | D12S1604 |
| 52,653,177 | 52,656,470 | 9906D06 | D12S1131i |
| 52,972,162 | 52,981,058 | 1207H01 | D12S0426i |
| 52,972,162 | 52,981,058 | 1206F10 | D12S0589i |
| 53,075,312 | 53,099,317 | 1209A02 | D12S0248i |
| 53,324,642 | 53,328,416 | 1208E08 | D12S0431i |
| 53,324,642 | 53,328,416 | 1201A03 | D12S0609i |
| 54,364,619 | 54,387,949 | 1201G05 | D12S0801i |
| 54,364,619 | 54,387,949 | 1201G02 | D12S0950i |
| 54,611,213 | 54,634,074 | 1202E02 | D12S0802i |
| 54,634,156 | 54,646,765 | 1203B02 | D12S0585i |
| 55,018,926 | 55,020,461 | 9906H09 | D12S0797i |
| 55,392,484 | 55,407,248 | 1206B06 | D12S0449i |
| 55,392,484 | 55,407,248 | 1206E11 | D12S0605i |
| 55,769,157 | 55,775,526 | 221D12 | AF067572.1_2832 |
| 55909819 | 55914981 | SHMT21 | new design |
| 55909819 | 55914981 | SHMT22 | new design |
| 56,114,810 | 56,130,876 | 1203D11 | D12S0982i |
| 56,135,363 | 56,138,058 | 1206F06 | D12S0455i |
| 56,428,270 | 56,432,431 | 1206C11 | D12S0601i |
| 56,499,977 | 56,527,014 | 685G06 | DISD22_0008338 |
| 6,179,134 | 6,217,688 | 1201F02 | D12S0385i |
| 6,308,184 | 6,321,522 | 1206B03 | D12S0389i |
| 6,308,184 | 6,321,522 | 1206D12 | D12S0635i |
| 6,363,595 | 6,370,994 | 1208A07 | D12S0917i |
| 6,513,872 | 6,517,797 | 1208B01 | D12S0831i |
| 6,513,872 | 6,517,797 | 1204A11 | D12S0969 |
| 6,768,912 | 6,800,237 | 1211C05 | D12S0170i |
| 6,819,636 | 6,826,819 | 1210B12 | D12S0824i |
| 64,869,270 | 64,928,684 | 322G08 | AC025603.1_117614 |
| 64,869,270 | 64,928,684 | 1208F08 | D12S0456i |
| 66,329,021 | 66,340,410 | 1207A03 | D12S0187i |
| 66,329,021 | 66,340,410 | 1204A04 | D12S335 |
| 66,834,816 | 66,839,790 | 224F06 | AC007458.13_25451 |
| 66,834,816 | 66,839,790 | 1205C04 | D12S0002i |
| 66,834,816 | 66,839,790 | 1205D08 | D12S0133i |
| 66,834,816 | 66,839,790 | IFNG | new design |
| 66,928,292 | 66,933,651 | 1206D06 | D12S0451i |
| 67,488,247 | 67,520,481 | 1209F05 | D12S0544i |
| 67,488,247 | 67,520,481 | 1207G01 | D12S0857i |
| 69,201,231 | 69,317,469 | 1202G03 | D12S1043 |
| 69,201,231 | 69,317,469 | 1203C06 | D12S1722 |
| 69,318,129 | 69,600,853 | 1202B12 | D12S0294i |
| 69,318,129 | 69,600,853 | 1206H01 | D12S0362i |
| 7,138,291 | 7,153,069 | 1209G04 | D12S0826i |
| 7,773,278 | 7,793,336 | 1211E11 | D12S0142i |
| 7,773,278 | 7,793,336 | 1208A10 | D12S0886i |
| 78,509,876 | 78,608,921 | 1209 H 08 | D12S0453i |
| 79,318,597 | 79,598,099 | 1209C10 | D12S0339 |
| 79,318,597 | 79,598,099 | 1210F10 | D12S0520i |
| 79,318,597 | 79,598,099 | 1202D01 | D12S0955i |
| 8,076,626 | 8,099,385 | 1205E08 | D12S0148i |
| 8,076,626 | 8,099,385 | 069E07 | D12S397 |
| 8,167,493 | 8,182,470 | 1207D08 | D12S0842i |
| 8,557,403 | 8,566,229 | 1206E12 | D12S0636i |


| 8,866,484 | 8,920,646 | 1210A07 | D12S0638i |
| :---: | :---: | :---: | :---: |
| 87,410,697 | 87,498,369 | 1210D08 | D12S0267i |
| 87,410,697 | 87,498,369 | 1209A03 | D12S0907i |
| 88,265,968 | 88,270,427 | 097 F 07 | HUMUT2007B |
| 88,265,968 | 88,270,427 | 1202F02 | Z67021 |
| 9,033,484 | 9,054,610 | T003E09 | D12S1695 |
| 9,460,894 | 9,492,092 | 1211 E 10 | D12S0023i |
| 9,460,894 | 9,492,092 | 1210G05 | D12S0158i |
| 9,460,894 | 9,492,092 | 1210A06 | D12S0195i |
| 9,638,415 | 9,651,764 | 1204G07 | HUMUT1833 |
| 9,871,344 | 9,888,871 | 1204E02 | Z67367 |
| 90,063,166 | 90,100,937 | 1205F04 | D12S0012i |
| 90,063,166 | 90,100,937 | 1205F03 | G09612 |
| 91,061,030 | 91,063,751 | 1201D11 | D12S0261i |
| 91,693,257 | 91,847,138 | 1211B11 | D12S0820i |
| 91,693,257 | 91,847,138 | 1207D10 | D12S0847i |
| 92,326,219 | 92,360,157 | 1209H05 | D12S0484i |
| 92,487,729 | 92,494,109 | 1211C10 | D12S0116i |
| 92,487,729 | 92,494,109 | 1209B11 | D12S0810i |
| 92,595,282 | 92,768,663 | 1204D10 | D12S0064i |
| 92,595,282 | 92,768,663 | 1209 E 12 | D12S0127i |
| 92,595,282 | 92,768,663 | 1206A09 | D12S0529 |
| 92,595,282 | 92,768,663 | T003F10 | D12S1346 |
| 93,066,630 | 93,223,356 | 453 H 02 | chr12.fa.O7frz. 100507297 |
| 93,066,630 | 93,223,356 | 1210G02 | D12S0464i |
| 93,066,630 | 93,223,356 | 1210C04 | D12S327 |
| 93,939,802 | 93,991,487 | 565D01 | chr12.fa.O7frz. 10164098 ¢ |
| 94,918,742 | 94,953,496 | 1207F08 | D12S0853i |
| 94,918,742 | 94,953,496 | 1211F04 | D12S309 |
| 97,433,527 | 97,468,250 | 1208G05 | D12S0255i |
| 97,433,527 | 97,468,250 | 1205B02 | D12S1706 |
| 97,563,209 | 97,653,342 | 220B04 | AC011248.8_84442 |
| 97,563,209 | 97,653,342 | 1203C05 | D12S0074i |
| 97,563,209 | 97,653,342 | 1208B09 | D12S0258i |
| 99,391,810 | 99,481,774 | 219F07 | AC010200.7_102437 |
| 99,391,810 | 99,481,774 | 224 A 12 | AC010200.7_92076 |
| 99,391,810 | 99,481,774 | 1205D06 | D12S0063i |
| 99,391,810 | 99,481,774 | 1208E02 | D12S0068i |
| 100,902,857 | 101,169,146 | 1304H06 | D13S0112i |
| 100,902,857 | 101,169,146 | 1303F09 | D13S0608i |
| 100,902,857 | 101,169,146 | 1305 F 12 | D13S0611i |
| 100,902,857 | 101,169,146 | 1303B08 | D13S0614i |
| 101,169,308 | 101,852,156 | 1302G09 | D13S0657i |
| 101,169,308 | 101,852,156 | 1301B09 | D13S0665i |
| 101,169,308 | 101,852,156 | 1301C04 | D13S0672i |
| 101,169,308 | 101,852,156 | 1308D04 | D13S1266 |
| 101,169,308 | 101,852,156 | 1304G02 | D13S1323 |
| 101,169,308 | 101,852,156 | 1302F04 | D13S174 |
| 101,169,308 | 101,852,156 | 1301G02 | G10095 |
| 102,295,195 | 102,326,346 | 1308A03 | D13S0024i |
| 107,657,791 | 107,668,717 | 1305B01 | D13S0342i |
| 107,719,978 | 107,758,826 | 1305H04 | D13S0419i |
| 109,599,311 | 109,757,505 | 1307E09 | D13S0634i |
| 109,599,311 | 109,757,505 | 1301B05 | Z66602 |
| 109,757,632 | 109,963,375 | 1307E03 | D13S0204i |
| 112,999,557 | 113,025,746 | 1306A08 | D13S0145i |


| 20,175,479 | 20,195,237 | $1301 \mathrm{B12}$ | D13S0638i |
| :---: | :---: | :---: | :---: |
| 20,612,650 | 20,621,221 | 1306B02 | D13S0585i |
| 20,612,650 | 20,621,221 | 1301F12 | HUMUT6291 |
| 21,143,170 | 21,176,637 | 1304F05 | D13S0077i |
| 21,143,170 | 21,176,637 | 1307H07 | D13S0129i |
| 23,042,723 | 23,148,232 | 1303H10 | D13S0522i |
| 23,042,723 | 23,148,232 | 1302A04 | D13S0692i |
| 23,042,723 | 23,148,232 | 040E10 | D13S292 |
| 25,726,276 | 25,877,375 | 1306F09 | D13S0173i |
| 25,726,276 | 25,877,375 | 1306G08 | D13S0358i |
| 25,726,276 | 25,877,375 | 1305F08 | D13S0507i |
| 26,029,840 | 26,161,085 | 1306B12 | D13S0366i |
| 26,896,681 | 26,907,823 | 1303D03 | D13S0372i |
| 26,896,681 | 26,907,823 | 1306B10 | D13S0475i |
| 27,434,273 | 27,441,317 | 1301B07 | D13S0079i |
| 27,475,411 | 27,572,729 | 1304C10 | D13S0242i |
| 27,475,411 | 27,572,729 | 1305F07 | D13S0482i |
| 27,773,790 | 27,967,232 | 1305A05 | D13S0420i |
| 27,773,790 | 27,967,232 | 1301G07 | D13S0655i |
| 27,773,790 | 27,967,232 | 1303H06 | D13S1242 |
| 29,930,884 | 30,089,729 | 1306E09 | D13S0095i |
| 29,930,884 | 30,089,729 | 1306G03 | D13S0126i |
| 29,930,884 | 30,089,729 | 1303B09 | D13S1246 |
| 29,930,884 | 30,089,729 | 1302A02 | D13S289 |
| 30,207,645 | 30,236,556 | 1304C01 | D13S1238 |
| 31,787,617 | 31,871,809 | 1306H07 | D13S0458i |
| 35,904,495 | 35,915,008 | 1305A09 | D13S0518i |
| 36,291,339 | 36,301,740 | 1302G03 | D13S0368i |
| 40,027,801 | 40,138,734 | 1304B06 | D13S0090i |
| 40,027,801 | 40,138,734 | 1301A08 | D13S1233 |
| 42,034,872 | 42,080,148 | 1304 E 10 | D13S0244i |
| 42,034,872 | 42,080,148 | 041B03 | D13S1297 |
| 44,592,650 | 44,756,237 | 565H08 | chr13.fa.O7frz. 44272759 |
| 44,592,650 | 44,756,237 | 1308B06 | D13S0229i |
| 44,592,650 | 44,756,237 | 1307B08 | D13S0455i |
| 44,809,008 | 44,813,505 | 1308H03 | D13S1312 |
| 46,305,514 | 46,368,179 | 325E05 | AL136958.7_37916 |
| 46,305,514 | 46,368,179 | 1307F09 | D13S0134i |
| 47,883,170 | 47,887,947 | 9907B02 | D13S0042i |
| 47,883,170 | 47,887,947 | 1304H10 | D13S0260i |
| 48,178,692 | 48,181,499 | 1304E01 | D13S1307 |
| 52,175,400 | 52,211,948 | 1303A10 | D13S0047i |
| 94,470,084 | 94,751,688 | 1304A12 | D13S0303i |
| 94,470,084 | 94,751,688 | 1303 HO 2 | D13S0506i |
| 94,470,084 | 94,751,688 | 1307D06 | D13S0635i |
| 94,470,084 | 94,751,688 | 1301F04 | HUMUT7403 |
| 98,744,790 | 98,757,708 | 1301D09 | D13S0582i |
| 98,744,790 | 98,757,708 | 1308B03 | D13S1271 |
| 101,617,139 | 101,675,776 | 1401H06 | D14S0309i |
| 101,762,375 | 101,841,284 | 1405 H 03 | D14S0453i |
| 102,313,569 | 102,442,381 | 1406B04 | D14S0349i |
| 102,313,569 | 102,442,381 | 1401F08 | D14S272 |
| 103,092,642 | 103,098,907 | 1405C03 | D14S0472i |
| 103,233,707 | 103,251,549 | 1402C11 | D14S0512i |
| 104,290,529 | 104,297,036 | 1405F06 | D14S0488i |
| 104,586,782 | 104,602,799 | 1405H10 | D14S0024i |


| 105,428,094 | 105,428,110 | 147E07 | AB019441.1_31065 |
| :---: | :---: | :---: | :---: |
| 105,428,094 | 105,428,110 | 1406F10 | D14S0475i |
| 19,849,367 | 19,871,297 | 1401G07 | D14S0500i |
| 19,849,367 | 19,871,297 | 732G03 | DIJ28_10035246 |
| 20,319,050 | 20,320,464 | 1406B07 | D14S0499i |
| 20,580,251 | 20,582,226 | 1403F01 | D14S1070 |
| 21,159,897 | 22,090,915 | 1404D01 | D14S0121i |
| 21,159,897 | 22,090,915 | 9907E08 | D14S0463i |
| 21,159,897 | 22,090,915 | 1405E05 | D14S0467i |
| 21,159,897 | 22,090,915 | 1405E03 | D14S0514i |
| 21,159,897 | 22,090,915 | 1402B07 | D14S0532i |
| 21,159,897 | 22,090,915 | 1402F02 | D14S0552i |
| 21,159,897 | 22,090,915 | 1401D04 | D14S1003 |
| 21,159,897 | 22,090,915 | 1401F11 | D14S283 |
| 21,159,897 | 22,090,915 | 1403B06 | G10057 |
| 22,375,633 | 22,386,643 | 1405B01 | D14S0528i |
| 22,375,633 | 22,386,643 | 1407E05 | HUMUT1079 |
| 22,459,573 | 22,468,501 | 1402H02 | Z67550 |
| 22,656,355 | 22,658,665 | 1401G11 | D14S990 |
| 22,845,866 | 22,850,798 | 1401B08 | D14S0468i |
| 22,911,858 | 22,915,452 | 9907F08 | D14S0470i |
| 23,661,207 | 23,678,016 | 1403B01 | D14S64 |
| 23,748,627 | 23,755,020 | 1401H02 | D14S0173i |
| 23,907,094 | 23,918,650 | 1405G11 | D14S0313i |
| 24,112,564 | 24,115,306 | 1404D04 | D14S0177i |
| 24170000 | 24173313 | GZMB1 | new design |
| 24170000 | 24173313 | GZMB2 | new design |
| 28,304,801 | 28,308,621 | 1402A12 | D14S1042 |
| 28,304,801 | 28,308,621 | 1401H05 | Z66624 |
| 29,115,436 | 29,466,651 | 9907D08 | D14S0440i |
| 29,115,436 | 29,466,651 | 1401C02 | D14S0448i |
| 29,115,436 | 29,466,651 | 1401F12 | D14S0451i |
| 29,115,436 | 29,466,651 | 1401C08 | D14S0516i |
| 29,115,436 | 29,466,651 | 1407C01 | D14S252 |
| 34,249,398 | 34,253,649 | 1406B06 | D14S0213i |
| 34,249,398 | 34,253,649 | 1402D02 | HUMUT7222 |
| 34,940,468 | 34,943,703 | 1406G02 | D14S0190i |
| 34,940,468 | 34,943,703 | 1406D09 | D14S0321i |
| 37,128,940 | 37,134,240 | 1406D12 | D14S0144i |
| 37,128,940 | 37,134,240 | 1406B05 | D14S0148i |
| 37,746,955 | 37,752,019 | 1407A10 | D14S0138i |
| 37,746,955 | 37,752,019 | 1406C06 | D14S0329 |
| 44,654,859 | 44,674,272 | 1405B12 | D14S0136i |
| 49,135,165 | 49,151,140 | 1402F09 | Z67256 |
| 49,954,993 | 50,069,126 | 1407B06 | D14S0158i |
| 49,954,993 | 50,069,126 | 1407C06 | D14S0162i |
| 50,776,686 | 50,792,512 | 9907G08 | D14S0490i |
| 51,804,181 | 51,813,192 | 1406 E 10 | D14S0323i |
| 51,850,863 | 51,865,074 | 1406F08 | D14S0160i |
| 52,243,668 | 52,264,466 | 1404A11 | D14S0331i |
| 53,486,207 | 53,493,362 | 1404A06 | D14S0217i |
| 53,486,207 | 53,493,362 | 1401G05 | Z67708 |
| 53,933,423 | 53,956,682 | 1405G10 | D14S0230i |
| 54,563,594 | 54,585,960 | 1401C10 | D14S0548i |
| 55,654,846 | 55,837,784 | 1401C01 | D14S0112i |
| 55,654,846 | 55,837,784 | 1407E10 | D14S0342i |


| 55,654,846 | 55,837,784 | 1403A03 | D14S1056 |
| :---: | :---: | :---: | :---: |
| 55,654,846 | 55,837,784 | 1403D02 | D14S1064 |
| 60,858,186 | 61,087,451 | 1404C03 | D14S0154i |
| 60,858,186 | 61,087,451 | 1404E03 | D14S0159i |
| 60,858,186 | 61,087,451 | 1404E11 | D14S0339i |
| 60,858,186 | 61,087,451 | 1405E04 | D14S0375i |
| 62,849,395 | 62,854,316 | 1404G07 | D14S0262i |
| 62,849,395 | 62,854,316 | T003B12 | D14S1012 |
| 63,621,388 | 63,875,070 | 1406F12 | D14S0503i |
| 63,621,388 | 63,875,070 | 1406A03 | D14S0506i |
| 63,621,388 | 63,875,070 | T003C12 | D14S63 |
| 64,475,625 | 64,479,284 | 1407E07 | D14S0257i |
| 68,410,793 | 68,515,747 | 1406E09 | D14S0360i |
| 68,410,793 | 68,515,747 | 1403B05 | Z66914 |
| 70,264,605 | 70,345,641 | 1407E09 | D14S0263i |
| 70,264,605 | 70,345,641 | 1407F02 | D14S1002 |
| 74,815,284 | 74,818,685 | 043 D 07 | D14S76 |
| 75,058,537 | 75,083,086 | 1402F08 | D14S0014i |
| 75,058,537 | 75,083,086 | 1405C11 | D14S0082i |
| 75,494,195 | 75,517,242 | 1403B09 | D14S0033i |
| 75,494,195 | 75,517,242 | 1407G05 | D14S0050i |
| 75,907,479 | 76,036,961 | 1406E06 | D14S0356i |
| 75,907,479 | 76,036,961 | 1407C11 | D14S0376i |
| 75,907,479 | 76,036,961 | 1401A09 | D14S0515i |
| 77,208,502 | 77,244,109 | 1405E06 | D14S594 |
| 77,208,502 | 77,244,109 | 1403C07 | HUMUT1235 |
| 80,491,528 | 80,682,399 | 1403F10 | D14S0071i |
| 80,491,528 | 80,682,399 | 1407G07 | D14S0275i |
| 80,716,147 | 80,757,328 | 1403A09 | D14S0031i |
| 87,469,111 | 87,529,660 | 1407D03 | D14S0045i |
| 87,469,111 | 87,529,660 | 1404H09 | D14S0301i |
| 88,003,867 | 88,090,876 | 1402H11 | D14S0272i |
| 88,003,867 | 88,090,876 | 1403C05 | Z67182 |
| 88,692,274 | 88,953,127 | 1403D09 | D14S0036i |
| 88,692,274 | 88,953,127 | 1402C02 | D14S0070i |
| 88,692,274 | 88,953,127 | 1401A07 | D14S0418i |
| 88,692,274 | 88,953,127 | 1401D12 | D14S0422i |
| 90,768,629 | 90,789,977 | 1406B03 | D14S0428i |
| 90,768,629 | 90,789,977 | 1406F03 | D14S0539 |
| 92,239,907 | 92,284,765 | 1406D05 | D14S0407i |
| 92,239,907 | 92,284,765 | 1405D06 | D14S0447i |
| 92,718,294 | 92,721,002 | 1405F07 | D14S0483i |
| 93,587,019 | 93,617,311 | 1401D01 | D14S0287i |
| 93,587,019 | 93,617,311 | 1401E03 | D14S0398i |
| 93,900,404 | 93,914,178 | 1404B10 | D14S0304i |
| 93,900,404 | 93,914,178 | 1407B05 | D14S0534i |
| 95,740,950 | 95,780,542 | 1402G03 | D14S0345i |
| 95,740,950 | 95,780,542 | 1401F04 | D14S0555i |
| 99,220,407 | 99,263,391 | 1406C05 | D14S0355i |
| 99,220,407 | 99,263,391 | 9907H08 | D14S0491i |
| 99,774,855 | 99,814,557 | 1401B01 | D14S0521i |
| 99,774,855 | 99,814,557 | 1401B12 | Z67128 |
| 22,619,887 | 23,215,702 | 1503D06 | D15S0006i |
| 22,619,887 | 23,215,702 | 1505B09 | D15S0228i |
| 22,619,887 | 23,215,702 | 1505C09 | D15S0230i |
| 22,619,887 | 23,215,702 | 1506E01 | D15S122 |


| 22,619,887 | 23,215,702 | 1502G05 | D15S128 |
| :---: | :---: | :---: | :---: |
| 31,945,720 | 32,118,595 | 1505D04 | D15S0304i |
| 31,945,720 | 32,118,595 | 1504E03 | D15S0477i |
| 31,945,720 | 32,118,595 | 1503 A02 | D15S1040 |
| 32,869,723 | 32,875,181 | 1504G07 | D15S0246i |
| 32,869,723 | 32,875,181 | 1504D04 | D15S0470i |
| 36,567,590 | 36,644,224 | 1501C10 | D15S0487i |
| 37,660,572 | 37,676,960 | 1504F10 | D15S0084i |
| 37,660,572 | 37,676,960 | 1501A07 | D15S0247i |
| 38,366,448 | 38,387,330 | 1501G07 | D15S994 |
| 38774661 | 38811646 | RAD51 | new design |
| 39,008,839 | 39,018,529 | 1503F08 | D15S0093i |
| 39,008,839 | 39,018,529 | 1505G10 | D15S0265i |
| 39,412,361 | 39,460,538 | 1505G11 | D15S0289i |
| 39,739,902 | 39,849,433 | 1505D09 | D15S0231i |
| 39,739,902 | 39,849,433 | 1506D12 | D15S0236i |
| 41,612,949 | 41,769,525 | 1504D10 | D15S0232i |
| 41,612,949 | 41,769,525 | 1505A07 | D15S0354i |
| 41825882 | 41852096 | PDIA3 | new design |
| 42,790,977 | 42,797,649 | 1504A03 | D15S0312i |
| 42,790,977 | 42,797,649 | 1505E01 | D15S0315i |
| 47,502,751 | 47,566,815 | 1504C10 | D15S0253i |
| 47,502,751 | 47,566,815 | 1506D07 | D15S0259i |
| 48,321,436 | 48,345,218 | 1505A02 | D15S0115i |
| 49,288,961 | 49,418,086 | 1506H07 | D15S0139i |
| 49,288,961 | 49,418,086 | 1504E04 | D15S0447i |
| 50,098,739 | 50,145,754 | $1503 \mathrm{HO1}$ | D15S982 |
| 50,189,114 | 50,192,264 | 1505E12 | D15S0302i |
| 54,998,125 | 55,368,008 | 463E02 | chr15.fa.O7frz. 53612467 |
| 54,998,125 | 55,368,008 | 1503C12 | D15S0180i |
| 54,998,125 | 55,368,008 | 1506H10 | D15S0446i |
| 54,998,125 | 55,368,008 | 1503B05 | D15S648 |
| 54,998,125 | 55,368,008 | 9907G10 | Z67030 |
| 56,675,802 | 56,829,469 | 1505C10 | D15S0255i |
| 56,675,802 | 56,829,469 | 1501F02 | D15S148 |
| 57,184,612 | 57,204,536 | 1504C12 | D15S0027i |
| 57,718,358 | 57,736,991 | 1506D09 | D15S0393i |
| 58,084,427 | 58,085,434 | 1506G06 | D15S0158i |
| 58,084,427 | 58,085,434 | 1503 F 05 | HUMUT1232 |
| 58,426,642 | 58,477,477 | 1506A11 | D15S0479i |
| 58,576,755 | 59,308,794 | 1502G07 | D15S0042i |
| 58,576,755 | 59,308,794 | 1502G11 | D15S0154i |
| 58,576,755 | 59,308,794 | 1503A12 | D15S0176i |
| 58,576,755 | 59,308,794 | 1505E09 | D15S0234i |
| 58,576,755 | 59,308,794 | 1506G12 | D15S0240i |
| 58,576,755 | 59,308,794 | 1506D03 | D15S970 |
| 58,576,755 | 59,308,794 | 104G09 | Z67427 |
| 61,986,288 | 62,125,574 | 1503 A05 | D15S644 |
| 61,986,288 | 62,125,574 | 1502H06 | D15S993 |
| 61,986,288 | 62,125,574 | $738 \mathrm{A07}$ | DIJ28_10022026 |
| 62,235,067 | 62,242,407 | 1506B09 | D15S0227i |
| 62,995,046 | 63,038,086 | 1501D07 | D15S0013i |
| 62,995,046 | 63,038,086 | 1506D01 | D15S1009 |
| 63,196,770 | 63,213,227 | 1501B07 | D15S0182i |
| 64,466,674 | 64,570,936 | 1505C01 | D15S0168i |
| 64,466,674 | 64,570,936 | 1505F02 | D15S0177i |


| 64,466,674 | 64,570,936 | 1506H08 | HUMUT5980 |
| :---: | :---: | :---: | :---: |
| 65,145,249 | 65,274,587 | 1502G12 | D15S0156i |
| 65,145,249 | 65,274,587 | 1502B06 | D15S988 |
| 65,622,075 | 65,886,506 | 1506H04 | D15S0269i |
| 65,622,075 | 65,886,506 | 1503 A03 | D15S1015 |
| 65,622,075 | 65,886,506 | 1503D04 | Z66922 |
| 66,381,096 | 66,511,546 | 1504G09 | D15S0187i |
| 66,381,096 | 66,511,546 | 1504G02 | D15S0386i |
| 66,381,096 | 66,511,546 | 1502A11 | D15S0455i |
| 67,532,177 | 67,534,939 | 1502 E 12 | D15S0371i |
| 68,127,597 | 68,177,310 | 1502D03 | D15S650 |
| 68,127,597 | 68,177,310 | T003G12 | D15S977 |
| 69,889,948 | 69,897,654 | 1506B05 | D15S0033i |
| 69,889,948 | 69,897,654 | T001B03 | D15S0327i |
| 70,364,122 | 70,455,868 | 1505B08 | D15S0282i |
| 71,639,410 | 71,712,806 | 1506 A07 | D15S0423i |
| 71,763,675 | 71,793,912 | 1504B10 | HUMUT744 |
| 72,417,157 | 72,447,134 | T001C05 | D15S0332i |
| 72,417,157 | 72,447,134 | 1506A10 | Z67571 |
| 72,798,943 | 72,804,930 | 1505C11 | D15S0280i |
| 72,798,943 | 72,804,930 | 1504H04 | D15S0392i |
| 73,546,515 | 73,658,680 | 1502A01 | D15S0481i |
| 74,020,333 | 74,091,842 | 1506C10 | D15S0104i |
| 74,020,333 | 74,091,842 | 1504 A 07 | D15S0352i |
| 75,074,609 | 75,116,727 | 328H08 | AC051643.2_53971 |
| 75,074,609 | 75,116,727 | 1502H10 | D15S0350i |
| 77,001,162 | 77,024,475 | 1505D11 | D15S0281i |
| 77,001,162 | 77,024,475 | 1506F04 | D15S1023 |
| 78,040,290 | 78,050,698 | 1503B04 | D15S1005 |
| 79,262,148 | 79,392,157 | 1503G12 | D15S0200i |
| 79,262,148 | 79,392,157 | 1504C11 | D15S0207i |
| 79,262,148 | 79,392,157 | 1502E08 | D15S1041 |
| 83,578,821 | 84,093,590 | 464A06 | chr15.fa.O7frz. 83916193 |
| 83,578,821 | 84,093,590 | 1506A06 | D15S0272i |
| 83,578,821 | 84,093,590 | 045A06 | D15S972 |
| 83,578,821 | 84,093,590 | 1501A02 | D15S999 |
| 83,578,821 | 84,093,590 | 100B03 | HUMUT1211 |
| 83,578,821 | 84,093,590 | 1502D07 | Z66702 |
| 86,983,039 | 87,000,684 | 9907F09 | D15S0049i |
| 86,983,039 | 87,000,684 | 1501G08 | D15S0506i |
| 88,129,130 | 88,159,072 | 9907G11 | D15S0329i |
| 88,129,130 | 88,159,072 | 1506A04 | HUMUT644B |
| 89,061,606 | 89,159,688 | 1505F08 | D15S0202i |
| 89,061,606 | 89,159,688 | 9907G09 | D15S0273i |
| 89,061,606 | 89,159,688 | 1503 H 03 | D15S127 |
| 89,212,889 | 89,227,691 | 9907F12 | D15S0399i |
| 94,674,950 | 94,683,048 | 1501H01 | D15S0106i |
| 94,674,950 | 94,683,048 | 9907A10 | D15S0306i |
| 97,010,288 | 97,319,034 | 1504D09 | D15S0215i |
| 97,010,288 | 97,319,034 | 1501H03 | D15S0433i |
| 97,010,288 | 97,319,034 | 1502D10 | D15S120 |
| 97,010,288 | 97,319,034 | 1502C03 | Z67468 |
| 1,696,222 | 1,760,319 | 1603B11 | D16S0217i |
| 1,968,919 | 1,971,441 | 1606F08 | D16S0494i |
| 10,867,648 | 10,926,341 | 046B02 | D16S414 |
| 10,867,648 | 10,926,341 | 1606C03 | D16S497 |


| 11,255,775 | 11,257,540 | 1603H09 | D16S0155i |
| :---: | :---: | :---: | :---: |
| 11,255,775 | 11,257,540 | 1606D07 | HUMUT1334 |
| 11,549,357 | 11,588,823 | 1602G01 | D16S0316i |
| 11,549,357 | 11,588,823 | 1604C04 | D16S0512i |
| 11966465 | 11969426 | TNFRSF171 | new design |
| 11966465 | 11969426 | TNFRSF172 | new design |
| 15,704,493 | 15,858,388 | 1603 D 05 | D16S0011i |
| 15,704,493 | 15,858,388 | 9908F03 | D16S0032i |
| 15,704,493 | 15,858,388 | 1602G02 | D16S3060 |
| 15,950,935 | 16,143,774 | 1604C07 | D16S0180i |
| 162,875 | 163,708 | 1601F11 | D16S0102i |
| 2,145,800 | 2,168,131 | 1601E07 | D16S0452i |
| 21,559,426 | 21,571,473 | 1604A11 | D16S0160i |
| 21,559,426 | 21,571,473 | 1603H10 | D16S0179i |
| 23,597,692 | 23,609,189 | 1606G10 | D16S0444i |
| 23,597,692 | 23,609,189 | 1603B03 | D16S417 |
| 23,754,823 | 24,139,358 | 9908B02 | D16S0020i |
| 23,754,823 | 24,139,358 | 1603E10 | D16S0169i |
| 23,754,823 | 24,139,358 | 1604D11 | D16S0178i |
| 23,754,823 | 24,139,358 | 1601C11 | D16S0558i |
| 23,754,823 | 24,139,358 | 1605D03 | D16S0560i |
| 23,754,823 | 24,139,358 | 1602H04 | D16S420 |
| 27,232,752 | 27,283,600 | 1605C07 | D16S0091i |
| 27,232,752 | 27,283,600 | 1601H07 | D16S0519i |
| 27,379,436 | 27,468,775 | 1601 D 08 | D16S0048i |
| 277,441 | 342,465 | $682 \mathrm{C07}$ | DISO7_12395103 |
| 28,418,184 | 28,425,656 | 663D02 | chr16.fa.O7frz. 38545966 |
| 28,418,184 | 28,425,656 | 1604C11 | D16S0112i |
| 29,581,801 | 29,589,688 | 1606F12 | D16S0531i |
| 29,731,591 | 29,766,842 | 1603B06 | D16S0030i |
| 3,010,343 | 3,012,385 | 1605B07 | D16S0007i |
| 3,010,343 | 3,012,385 | 1603F01 | D16S3082 |
| 3,232,029 | 3,246,628 | 1603F06 | D16S0040i |
| 3,232,029 | 3,246,628 | 1604E08 | D16S0211i |
| 3,531,826 | 3,567,290 | 1605C12 | D16S0002i |
| 3,531,826 | 3,567,290 | 1602C08 | D16S0103i |
| 3,630,847 | 3,654,064 | 1606B08 | D16S0120i |
| 3,716,568 | 3,870,723 | 1605C11 | D16S0023i |
| 3,716,568 | 3,870,723 | 1603C08 | D16S0101i |
| 3,716,568 | 3,870,723 | 1601B06 | D16S3065 |
| 30,362,453 | 30,364,725 | 1602E07 | D16S0552i |
| 30,391,551 | 30,442,007 | 1606G11 | D16S0462i |
| 30,815,429 | 30,822,382 | 1601F05 | D16S0562i |
| 31178789 | 31251714 | ITGAM1 | new design |
| 31178789 | 31251714 | ITGAM2 | new design |
| 31,274,010 | 31,301,819 | 1604F03 | D16S0471i |
| 4,466,426 | 4,500,349 | 1604E12 | D16S0328i |
| 45,746,798 | 46,052,519 | 1604B01 | D16S0508i |
| 45,746,798 | 46,052,519 | 1602G12 | D16S0517i |
| 45,746,798 | 46,052,519 | 1602G07 | D16S3044 |
| 46,758,323 | 46,838,806 | 9908A08 | D16S0425i |
| 46,758,323 | 46,838,806 | 1601H10 | D16S0427i |
| 49,288,551 | 49,324,488 | 1604G06 | D16S0350i |
| 49,288,551 | 49,324,488 | 1601F09 | D16S3035 |
| 52082693 | 52094671 | AKTIP1 | new design |
| 52082693 | 52094671 | AKTIP2 | new design |


| 54,070,589 | 54,098,104 | 493H04 | chr16.fa.O7frz.65771871 |
| :---: | :---: | :---: | :---: |
| 54,070,589 | 54,098,104 | 9908A06 | D16S0335i |
| 54,394,264 | 54,424,576 | 1603A11 | D16S0198i |
| 55,180,768 | 55,182,501 | 9908H04 | D16S0287i |
| 55,180,768 | 55,182,501 | 1603D02 | D16S3071 |
| 55,581,018 | 55,673,941 | 1606A04 | D16S0356i |
| 55,581,018 | 55,673,941 | 1605 E 10 | D16S0377i |
| 55,581,018 | 55,673,941 | 1601A06 | D16S0380i |
| 55,950,219 | 55,957,602 | 1605F06 | D16S0364i |
| 55,996,180 | 56,007,475 | 1601B10 | D16S0158i |
| 56,616,783 | 56,638,306 | 9908E04 | D16S0281i |
| 56,616,783 | 56,638,306 | 1601H11 | HUMUT5103 |
| 64,958,064 | 64,996,190 | 1602C06 | D16S0343i |
| 64,958,064 | 64,996,190 | 1601G10 | D16S0541i |
| 65,143,967 | 65,170,463 | 1604E11 | D16S0353i |
| 65,620,551 | 65,692,462 | 1601B03 | D16S0422i |
| 65,620,551 | 65,692,462 | 1606E07 | Z66903 |
| 65,745,605 | 65,751,306 | 1606H11 | D16S0424i |
| 65,765,371 | 65,767,127 | 1606D01 | D16S421 |
| 66,525,908 | 66,528,254 | 1602F03 | D16S0298i |
| 66,525,908 | 66,528,254 | 1601D12 | D16S0386i |
| 66,676,845 | 66,818,338 | 1604A04 | D16S0313i |
| 66,902,446 | 66,948,663 | T001E06 | D16S0381i |
| 67,018 | 75,845 | 1601C05 | D16S521 |
| 67,328,696 | 67,426,945 | 1604C06 | D16S0412i |
| 68,156,498 | 68,296,054 | 1605A07 | D16S0344i |
| 68,156,498 | 68,296,054 | 1604E06 | D16S0556i |
| 68,890,573 | 68,925,232 | 1604C12 | D16S0246i |
| 70,685,116 | 70,704,312 | 1603D07 | D16S0079 |
| 70,685,116 | 70,704,312 | 047H01 | D16S3106 |
| 71,378,456 | 71,639,775 | 1603C06 | D16S0034i |
| 71,378,456 | 71,639,775 | 1603 E 06 | D16S0039i |
| 71,378,456 | 71,639,775 | 1603D08 | D16S0105i |
| 71,378,456 | 71,639,775 | 1602H03 | D16S0113i |
| 71,378,456 | 71,639,775 | 1603A10 | D16S0159i |
| 71,378,456 | 71,639,775 | 9908B05 | D16S0299i |
| 73,885,109 | 74,024,888 | 9908D06 | D16S0339 |
| 73,885,109 | 74,024,888 | 1606E10 | D16S0362i |
| 73,885,109 | 74,024,888 | 1604H01 | D16S0544i |
| 76,613,992 | 76,623,499 | 1606E06 | D16S518 |
| 78,185,732 | 78,192,112 | 9908G06 | D16S0375i |
| 78,185,732 | 78,192,112 | 1603B01 | D16S3040 |
| 84,268,781 | 84,280,089 | 1601H09 | D16S0565i |
| 84,490,275 | 84,513,713 | 1601G09 | D16S0414i |
| 84,490,275 | 84,513,713 | 1601A01 | D16S0487i |
| 85,101,659 | 85,105,548 | 1601A02 | D16S0526i |
| 85,101,659 | 85,105,548 | 1606F05 | D16S520 |
| 86,421,130 | 86,460,615 | 1602C05 | D16S0554i |
| 86,421,130 | 86,460,615 | 1602E01 | D16S413 |
| 87,232,502 | 87,234,383 | 9908F02 | D16S0251i |
| 88,512,527 | 88,529,713 | 1604B08 | D16S0252i |
| 1,483,902 | 1,495,792 | 1706D12 | D17S0145i |
| 1,483,902 | 1,495,792 | 1702B07 | D17S0440i |
| 1,909,888 | 2,220,160 | 1701G08 | D17S0205i |
| 1,909,888 | 2,220,160 | 1704E11 | D17S0208i |
| 1,909,888 | 2,220,160 | 1703D10 | D17S0211i |


| 1,909,888 | 2,220,160 | 1702A12 | D17S0496i |
| :---: | :---: | :---: | :---: |
| 10,365,192 | 10,393,704 | 1705D12 | D17S0038i |
| 10,365,192 | 10,393,704 | 1706D04 | D17S1852 |
| 11,864,860 | 11,987,865 | 1702E05 | D17S0111i |
| 15,616,046 | 15,629,130 | 1702E04 | D17S0372i |
| 15,616,046 | 15,629,130 | 1704F01 | D17S0374i |
| 15,875,983 | 16,059,570 | 1701G03 | D17S0001i |
| 15,875,983 | 16,059,570 | 1701F01 | D17S1843 |
| 16,259,613 | 16,281,042 | 1701B01 | D17S1857 |
| 16,783,123 | 16,816,127 | 1704E03 | D17S0425i |
| 17,655,794 | 17,681,050 | 1702D07 | D17S0428i |
| 17,932,008 | 17,952,017 | 1704F03 | D17S0430i |
| 17,932,008 | 17,952,017 | 1701B12 | D17S0525i |
| 18,028,014 | 18,053,993 | 1705G04 | D17S0483i |
| 18,516,347 | 18,516,964 | 9909F02 | D17S0459i |
| 19,221,659 | 19,227,445 | 1705F09 | D17S0433i |
| 2,443,686 | 2,535,638 | 1704A09 | D17S0041i |
| 2,443,686 | 2,535,638 | 1704E12 | D17S0360i |
| 2,443,686 | 2,535,638 | 1705E04 | D17S0484i |
| 21,128,581 | 21,159,118 | 1705F03 | D17S0154i |
| 22,980,951 | 23,000,711 | 1704C03 | D17S0441i |
| 23,107,919 | 23,151,682 | 1704E09 | D17S0024i |
| 23,107,919 | 23,151,682 | 1704C05 | D17S0472i |
| 23,718,425 | 23,721,844 | 9909C02 | D17S0404i |
| 23,875,086 | 23,889,302 | 1702H03 | D17S0401i |
| 24,424,663 | 24,531,556 | 1702D06 | D17S0402i |
| 24,424,663 | 24,531,556 | 9908E09 | D17S0406i |
| 24,424,663 | 24,531,556 | 1706B05 | D17S0522i |
| 24,424,663 | 24,531,556 | 1706H05 | D17S841 |
| 25,549,032 | 25,586,831 | 1705B01 | D17S0153i |
| 25,549,032 | 25,586,831 | 1701E04 | Z67368 |
| 26,133,828 | 26,175,826 | 1705G05 | D17S0196i |
| 26,133,828 | 26,175,826 | 1702A05 | D17S0465i |
| 29,606,409 | 29,608,335 | 1706H06 | D17S1293 |
| 29,707,584 | 29,709,742 | 1706B04 | D17S0493i |
| 29,711,512 | 29,714,365 | $331 \mathrm{G12}$ | AC011193.2_31574 |
| 3,415,491 | 3,459,454 | 1701B06 | D17S0309i |
| 3,415,491 | 3,459,454 | 1701D09 | D17S829 |
| 3,486,522 | 3,511,585 | 1701G07 | D17S0090i |
| 3,564,671 | 3,660,578 | 1701B04 | D17S1298 |
| 3,746,634 | 3,766,709 | 1701E06 | D17S1828 |
| 31,116,989 | 31,146,753 | 1701E01 | D17S1833 |
| 31,222,611 | 31,231,490 | 1703 A07 | D17S0086i |
| 31,811,186 | 31,816,297 | 1702F11 | D17S0438i |
| 32,516,040 | 32,841,015 | 1705A03 | D17S0107i |
| 32,516,040 | 32,841,015 | 1706D09 | D17S0265i |
| 32,516,040 | 32,841,015 | 1704D11 | D17S0272i |
| 32,924,064 | 32,947,709 | 1701B02 | D17S0124i |
| 33,046,526 | 33,077,600 | 1703 E 07 | D17S0095i |
| 35,097,919 | 35,138,441 | 1703B07 | D17S0089 |
| 35,097,919 | 35,138,441 | 1705D02 | D17S0260i |
| 35,425,214 | 35,427,592 | 303B05 | AC007776.1_62510 |
| 35,472,589 | 35,503,646 | 303 C 05 | AC007776.1_75537 |
| 35,502,567 | 35,510,499 | 9908H11 | D17S0330i |
| 35,502,567 | 35,510,499 | 1706D11 | D17S0452i |
| 35,718,972 | 35,767,420 | 1706E09 | D17S0280i |


| 35,798,321 | 35,827,695 | 1704G10 | D17S0273i |
| :---: | :---: | :---: | :---: |
| 35,963,547 | 35,975,250 | 1706D08 | D17S0285i |
| 35,963,547 | 35,975,250 | 1703E04 | HUMUT186 |
| 36,787,447 | 36,792,181 | 1703 A08 | D17S0108i |
| 36,886,467 | 36,891,194 | 1705D08 | D17S0053i |
| 36,886,467 | 36,891,194 | 1702F02 | D17S0530i |
| 37,222,727 | 37,232,995 | 1706B11 | D17S0368i |
| 37,222,727 | 37,232,995 | 1706B01 | HUMUT8182 |
| 37,506,952 | 37,518,277 | 1703 E 12 | D17S0266i |
| 37,506,952 | 37,518,277 | 1702F01 | HUMUT8184 |
| 37,604,721 | 37,681,950 | 1706C05 | D17S0161i |
| 37,604,721 | 37,681,950 | 048G02 | D17S1801 |
| 37,604,721 | 37,681,950 | 1706G10 | D17S1802 |
| 38,084,961 | 38,087,371 | 1701C11 | D17S0521i |
| 38,238,949 | 38,249,303 | 1702C05 | G10143 |
| 38,916,860 | 38,957,206 | 1704H09 | D17S0170i |
| 38,916,860 | 38,957,206 | 1706G11 | D17S0426i |
| 38,916,860 | 38,957,206 | 092E08 | HUMUT573 |
| 39,199,015 | 39,211,872 | 1705G11 | D17S0007i |
| 39,199,015 | 39,211,872 | 1706G03 | D17S0173i |
| 39,199,015 | 39,211,872 | 1702F06 | D17S951 |
| 39,509,647 | 39,556,540 | 1703A09 | D17S0134i |
| 39,509,647 | 39,556,540 | 1706B10 | D17S0320i |
| 39,682,566 | 39,700,993 | 1702A11 | D17S1860 |
| 39,805,076 | 39,822,399 | 1703D08 | D17S0122i |
| 4,480,963 | 4,491,709 | 1705A01 | D17S0238i |
| 4,560,533 | 4,571,544 | 1701A02 | D17S0350i |
| 4,646,397 | 4,648,756 | 607C09 | chr17.fa.O7frz.5865115 |
| 4,683,351 | 4,742,135 | 331 A03 | AC015913.4_60812 |
| 4,776,372 | 4,779,067 | 571C11 | chr17.fa.O7frz. 5607732 |
| 4,789,692 | 4,793,067 | 1704H04 | D17S0432i |
| 40,392,587 | 40,401,170 | 1701G01 | D17S0506i |
| 40,392,587 | 40,401,170 | 738D08 | DIJ28_10035753 |
| 40,696,278 | 40,750,148 | 1704C09 | D17S0025i |
| 40,696,278 | 40,750,148 | 1705C11 | D17S0142i |
| 40,696,278 | 40,750,148 | 1702D05 | D17S950 |
| 41,217,449 | 41,268,973 | 9908C10 | D17S0253i |
| 41,217,449 | 41,268,973 | 1703F12 | D17S0267i |
| 41,217,449 | 41,268,973 | 1701 E 10 | D17S0480i |
| 42,196,855 | 42,251,081 | 9908C12 | D17S0339 |
| 42,196,855 | 42,251,081 | 1702C12 | D17S791 |
| 42,196,855 | 42,251,081 | 1703D02 | D17S920 |
| 42,686,207 | 42,745,076 | 332C05 | AC064817.4_80427 |
| 42,686,207 | 42,745,076 | 1706A03 | D17S0092i |
| 42,686,207 | 42,745,076 | 9909B01 | D17S0373i |
| 43,165,609 | 43,178,484 | 1703E08 | D17S0123i |
| 43,165,609 | 43,178,484 | 1702D10 | D17S0527i |
| 43,165,609 | 43,178,484 | 1706F12 | D17S806 |
| 43,480,720 | 43,493,841 | 1705D09 | D17S0281i |
| 43,565,804 | 43,862,551 | 9908F10 | D17S0286i |
| 43,565,804 | 43,862,551 | 1704G07 | D17S0453i |
| 43,565,804 | 43,862,551 | 1703B02 | D17S958 |
| 44,007,868 | 44,010,742 | 1704D01 | D17S1827 |
| 44,927,654 | 44,947,360 | 1702G03 | D17S797 |
| 45,422,368 | 45,427,587 | 1705F05 | D17S0188i |
| 45,616,456 | 45,633,992 | 1703B09 | D17S0135i |


| 45,616,456 | 45,633,992 | 1704B09 | D17S0358i |
| :---: | :---: | :---: | :---: |
| 46,585,919 | 46,604,103 | 1705B02 | D17S0004i |
| 5,276,823 | 5,283,195 | 1704H11 | D17S0113i |
| 5,284,956 | 5,312,905 | 1701B10 | D17S0224i |
| 5,343,472 | 5,428,553 | 1702E08 | D17S0517i |
| 50,697,370 | 50,755,886 | 1706C06 | D17S0021i |
| 50,697,370 | 50,755,886 | 1704C06 | D17S1799 |
| 52,320,269 | 52,346,408 | 1702E06 | D17S0105i |
| 52,320,269 | 52,346,408 | 1703A10 | D17S0168i |
| 53,625,088 | 53,636,783 | 1702F09 | D17S0193i |
| 53702201 | 53713295 | MPO1 | new design |
| 53702201 | 53713295 | MPO2 | new design |
| 54,997,668 | 55,040,484 | $9908 \mathrm{B12}$ | D17S0336i |
| 54,997,668 | 55,040,484 | 9909G01 | D17S0391i |
| 58,058,494 | 58,124,629 | 1702C03 | D17S1835 |
| 58,058,494 | 58,124,629 | 1701H10 | D17S794 |
| 58,058,494 | 58,124,629 | 1701A07 | D17S808 |
| 58,908,166 | 58,952,935 | 1705A06 | D17S0349 |
| 59,205,299 | 59,250,409 | 1704G02 | D17S0271i |
| 59,348,294 | 59,349,930 | 1705D06 | D17S0278i |
| 59,754,142 | 59,817,723 | 1704D06 | D17S0439 |
| 59,926,200 | 59,932,872 | 1706F11 | D17S0390i |
| 6,840,108 | 6,856,220 | 1705A02 | D17S0158i |
| 6,918,580 | 6,924,324 | 465 E 03 | chr17.fa.O7frz.7725213 |
| 60,437,295 | 60,483,216 | 1702B05 | D17S1792 |
| 60,437,295 | 60,483,216 | 1703A03 | D17S1825 |
| 60,955,143 | 60,988,227 | 1701D12 | D17S0486i |
| 60,955,143 | 60,988,227 | 1702G04 | D17S0519i |
| 61,729,388 | 62,237,324 | 150E09 | AC006263.1_34791 |
| 61,729,388 | 62,237,324 | 1704E08 | D17S0011i |
| 61,729,388 | 62,237,324 | 1703G06 | D17S0075i |
| 61,729,388 | 62,237,324 | 1704C07 | D17S0454i |
| 61,729,388 | 62,237,324 | 1703B04 | D17S1291 |
| 61,729,388 | 62,237,324 | 048C10 | D17S1816 |
| 61,729,388 | 62,237,324 | 1701E03 | D17S942 |
| 64,922,433 | 65,051,067 | 1705G09 | D17S0337i |
| 64,922,433 | 65,051,067 | 1701A09 | D17S1786 |
| 68,672,755 | 68,679,689 | 467B04 | chr17.fa.O7frz. 78835314 |
| 68,672,755 | 68,679,689 | 9908 E 12 | D17S0343i |
| 68,672,755 | 68,679,689 | 1706D05 | D17S0399 |
| 69,974,117 | 69,992,528 | 1704F10 | D17S0297i |
| 69,974,117 | 69,992,528 | 9908F12 | D17S0345i |
| 7,156,702 | 7,173,362 | 1701H07 | D17S0478i |
| 7,283,413 | 7,288,980 | 1704A01 | D17S0479i |
| 7,392,932 | 7,405,649 | 1701C05 | Z67321 |
| 7,393,099 | 7,405,649 | 1704C12 | D17S0018i |
| 7,883,083 | 7,893,177 | 1706F05 | D17S0446i |
| 70,048,842 | 70,053,877 | 1706E01 | HUMUT7429 |
| 70,087,099 | 70,100,017 | 1703B05 | HUMUT1523 |
| 70,202,047 | 70,220,712 | 1702D04 | D17S0394i |
| 70,780,669 | 70,797,109 | 1702F07 | D17S0356i |
| 70,780,669 | 70,797,109 | 1705H03 | D17S0383i |
| 71,229,111 | 71,265,494 | 1706F07 | D17S1839 |
| 71,644,009 | 71,648,966 | 303G08 | AC015801.21_117665 |
| 71,644,009 | 71,648,966 | 303H08 | AC015801.21_98885 |
| 71,892,297 | 71,895,536 | 049G01 | D17S1817 |


| 71,892,297 | 71,895,536 | 1703G03 | D17S785 |
| :---: | :---: | :---: | :---: |
| 72,220,514 | 72,234,158 | 9908A11 | D17S0302i |
| 73,721,872 | 73,733,311 | 1705E03 | D17S0049i |
| 73,721,872 | 73,733,311 | 1704B04 | D17S0529i |
| 73,864,454 | 73,867,753 | T002C05 | D17S0219i |
| 73,864,454 | 73,867,753 | 1703G04 | HUMUT952 |
| 74,181,727 | 74,289,971 | 1704E05 | D17S0410i |
| 74,181,727 | 74,289,971 | 1704G04 | D17S0463i |
| 74,360,654 | 74,433,067 | 1703A11 | D17S0222i |
| 75,723,612 | 75,735,533 | 1706H11 | D17S0294i |
| 75,723,612 | 75,735,533 | 1704B06 | D17S0471i |
| 76,580,274 | 76,588,528 | 1705B11 | D17S0223i |
| 76,705,160 | 76,754,467 | 1704A05 | D17S0531i |
| 77,091,594 | 77,094,422 | 1706B03 | D17S0407i |
| 77,418,886 | 77,422,527 | 9908H08 | D17S0215i |
| 77,866,035 | 77,868,769 | 1702E12 | D17S928 |
| 77,872,189 | 77,884,930 | 1701E11 | D17S0537i |
| 78,070,883 | 78,153,743 | 1704A07 | D17S0411i |
| 78,070,883 | 78,153,743 | 1701D04 | D17S0516i |
| 8,722,953 | 8,756,559 | 1704D09 | D17S0019i |
| 8,722,953 | 8,756,559 | 1704B11 | D17S0020i |
| 8,722,953 | 8,756,559 | 1701A01 | D17S786 |
| 11,679,263 | 11,871,922 | 1804E09 | D18S0049 |
| 11,679,263 | 11,871,922 | 1805C07 | D18S0278i |
| 11,679,263 | 11,871,922 | 1805C10 | D18S0344i |
| 11,679,263 | 11,871,922 | 9901C03 | D18S0507i |
| 11,679,263 | 11,871,922 | 1804D07 | D18S482 |
| 12,775,480 | 12,874,334 | 573D07 | chr18.fa.O7frz. 16363530 |
| 12,775,480 | 12,874,334 | 1803F04 | D18S0050i |
| 13,716,680 | 13,754,554 | 1806G09 | D18S0055i |
| 13,716,680 | 13,754,554 | 1803 A03 | Z67649 |
| 13,815,543 | 13,816,861 | 1801D09 | Z67345 |
| 16,787,533 | 16,944,869 | 1803D05 | D18S0073i |
| 16,787,533 | 16,944,869 | 1806D06 | D18S0554i |
| 18,003,414 | 18,036,225 | 1801B01 | Z67399 |
| 19,523,560 | 19,789,028 | 1802H07 | D18S0204i |
| 19,523,560 | 19,789,028 | 1804H04 | D18S0285i |
| 19,523,560 | 19,789,028 | 1801C03 | D18S0448i |
| 20,294,591 | 20,313,919 | 1803E01 | D18S1107 |
| 20,294,591 | 20,313,919 | 1802F09 | D18S1108 |
| 204,522 | 258,049 | 1801D01 | D18S0041i |
| 23,784,933 | 24,011,189 | 1803A06 | D18S0087i |
| 23,784,933 | 24,011,189 | 1803B06 | D18S0090i |
| 23,784,933 | 24,011,189 | 1805H02 | D18S0226i |
| 23,784,933 | 24,011,189 | 1805A04 | D18S0410i |
| 23,784,933 | 24,011,189 | 1804H01 | D18S0479i |
| 3,402,072 | 3,448,409 | 123D10 | AC006211.1_151493 |
| 3,402,072 | 3,448,409 | $113 \mathrm{B09}$ | AC006211.1_168801 |
| 3,402,072 | 3,448,409 | 1804B03 | D18S0455i |
| 309,356 | 490,685 | 1803B10 | D18S0192i |
| 309,356 | 490,685 | 1804H09 | D18S0429i |
| 37,789,197 | 37,915,446 | 1803D08 | D18S0155i |
| 37,789,197 | 37,915,446 | 1803H10 | D18S0206i |
| 37,789,197 | 37,915,446 | 1802E08 | D18S0312i |
| 41,558,155 | 41,585,297 | 1803E07 | D18S0129i |
| 41,558,155 | 41,585,297 | 1805H03 | D18S0144i |


| 41,659,543 | 41,678,045 | 1805F11 | D18S0385i |
| :---: | :---: | :---: | :---: |
| 44,700,221 | 44,731,079 | 1803D11 | D18S0223i |
| 44,700,221 | 44,731,079 | 1805F05 | D18S0297i |
| 46,340,482 | 46,512,194 | 1804G12 | D18S0119i |
| 46,340,482 | 46,512,194 | 1802D08 | D18S0490i |
| 46,340,482 | 46,512,194 | 1803D02 | D18S479 |
| 51,045,967 | 51,406,858 | 418H01 | AC018994.3_111068 |
| 51,045,967 | 51,406,858 | 1804B08 | D18S0003i |
| 51,045,967 | 51,406,858 | 1806D10 | D18S0382i |
| 51,045,967 | 51,406,858 | 1806G06 | HUMUT7024B |
| 54,489,598 | 54,568,350 | 1802D12 | D18S0565i |
| 54,489,598 | 54,568,350 | 1802C12 | D18S0566i |
| 55,085,251 | 55,091,605 | 1805D08 | D18S0317i |
| 55,085,251 | 55,091,605 | 1804E01 | D18S0518i |
| 55,148,088 | 55,177,463 | 1801A05 | D18S1155 |
| 56,189,564 | 56,190,562 | 575A04 | chr18.fa.O7frz. 64410150 |
| 56,189,564 | 56,190,562 | 1802D01 | D18S0244i |
| 58,143,500 | 58,205,872 | 1803H08 | D18S0168i |
| 58,143,500 | 58,205,872 | 1802B02 | D18S0441i |
| 58,941,559 | 59,137,593 | 1803A12 | D18S0241i |
| 58,941,559 | 59,137,593 | 1804A02 | D18S0321i |
| 58,941,559 | 59,137,593 | 1803H03 | HUMUT574 |
| 59,705,922 | 59,722,100 | 1806F01 | D18S68 |
| 59,767,574 | 59,779,093 | 9901E05 | HUMUT2039 |
| 6,931,885 | 7,107,813 | 1806E11 | D18S0258i |
| 6,931,885 | 7,107,813 | 1805D04 | D18S0350i |
| 6,931,885 | 7,107,813 | 1805B03 | D18S0352i |
| 647,619 | 663,492 | 573C03 | chr18.fa.O7frz. 941566 |
| 65,681,172 | 65,775,140 | 1805B09 | D18S0327i |
| 65,681,172 | 65,775,140 | 1803 HO 2 | Z67518 |
| 65,681,172 | 65,775,140 | 1801D07 | Z67555 |
| 66,107,243 | 66,145,329 | 1804C08 | D18S0407i |
| 66,107,243 | 66,145,329 | 1806C07 | D18S0510i |
| 7,557,817 | 8,396,854 | 1805E01 | D18S0045i |
| 7,557,817 | 8,396,854 | 1803E04 | D18S0047i |
| 7,557,817 | 8,396,854 | 1804E06 | D18S0193i |
| 7,557,817 | 8,396,854 | 1803C10 | D18S0194i |
| 7,557,817 | 8,396,854 | 1806E02 | D18S0196i |
| 7,557,817 | 8,396,854 | 1803 E 10 | D18S0199 |
| 7,557,817 | 8,396,854 | 1801A08 | D18S0202i |
| 7,557,817 | 8,396,854 | 9901B06 | D18S0478i |
| 7,557,817 | 8,396,854 | 1801E06 | D18S1163 |
| 711,592 | 802,547 | 1801H06 | D18S0446i |
| 711,592 | 802,547 | 9909H05 | D18S0452i |
| 72,819,777 | 72,973,762 | 9901H05 | D18S0498i |
| 72,819,777 | 72,973,762 | 1802H04 | D18S1097 |
| 75,256,760 | 75,390,311 | 1805G11 | D18S0394i |
| 75,256,760 | 75,390,311 | 9909E05 | D18S0439 |
| 9,465,007 | 9,528,106 | 1806C11 | D18S0075i |
| 1,018,174 | 1,037,627 | 9909H06 | D19S886 |
| 1,560,293 | 1,603,328 | 1903H09 | D19S0054i |
| 1,560,293 | 1,603,328 | 9901B09 | D19S0081i |
| 1,560,293 | 1,603,328 | 1904B07 | D19S0190i |
| 10,083,197 | 10,087,065 | 1901B03 | D19S0008i |
| 10,083,197 | 10,087,065 | 1901E07 | D19S0395i |
| 10,258,650 | 10,260,198 | 1904D05 | D19S0206i |


| 10,322,205 | 10,352,211 | 1903A01 | D19S0093i |
| :---: | :---: | :---: | :---: |
| 10,457,796 | 10,475,243 | 1903B01 | HUMUT5187 |
| 10,673,106 | 10,803,579 | T001E09 | D19S0283i |
| 10,843,253 | 10,894,448 | 1901H01 | D19S0245i |
| 10,932,606 | 11,033,953 | 9909C08 | D19S0301i |
| 11,061,132 | 11,105,490 | 1902C03 | D19S0372i |
| 11,296,093 | 11,311,321 | 305C09 | AC020561.3_124319 |
| 11,348,883 | 11,356,019 | 9901E08 | D19S0139i |
| 11,477,744 | 11,500,972 | 1902A06 | D19S0364i |
| 12,763,286 | 12,765,129 | 1903A02 | D19S914 |
| 12,910,423 | 12,916,303 | 9909E06 | HUMUT8091 |
| 12,917,654 | 12,925,455 | 1903C05 | D19S0195i |
| 13,933,352 | 13,978,097 | 1901D09 | D19S0311i |
| 13,933,352 | 13,978,097 | 1902D12 | D19S0390i |
| 14,063,500 | 14,089,559 | 1904A11 | D19S0196i |
| 14,353,213 | 14,380,535 | 1902F02 | D19S0198i |
| 14,444,278 | 14,447,174 | 1901 H 10 | D19S226 |
| 14,570,918 | 14,646,810 | 1903H02 | D19S0352i |
| 14,704,205 | 14,750,353 | 1904D10 | D19S0086i |
| 15,024,015 | 15,027,900 | 1903E12 | D19S0009i |
| 15,024,015 | 15,027,900 | 1901F09 | D19S929 |
| 15,131,444 | 15,172,792 | 1901C08 | D19S0048i |
| 15,440,463 | 15,451,312 | 1901G10 | G08034 |
| 15,587,421 | 15,601,445 | 1901E08 | D19S588 |
| 15,612,707 | 15,634,634 | 1903H05 | D19S0318i |
| 15,849,834 | 15,869,885 | 1902E08 | D19S0050i |
| 15,884,181 | 15,906,326 | 9901C11 | D19S0354i |
| 16,105,838 | 16,130,381 | 467D12 | chr19.fa.O7frz.17685171 |
| 16,105,838 | 16,130,381 | 1901D08 | D19S0248i |
| 16,105,838 | 16,130,381 | 1901D03 | D19S885 |
| 16,296,648 | 16,299,345 | 9901D09 | D19S917 |
| 16,489,705 | 16,514,248 | 1904A02 | D19S0114i |
| 17,203,694 | 17,217,151 | 1902E10 | D19S410 |
| 17,203,694 | 17,217,151 | 1901H08 | D19S593 |
| 17,374,755 | 17,377,457 | 1904F08 | D19S0151i |
| 17,374,755 | 17,377,457 | 1903A10 | D19S0162i |
| 17,766,658 | 17,785,385 | 1902C08 | D19S0109i |
| 17,766,658 | 17,785,385 | 1902D02 | D19S0159i |
| 17,788,322 | 17,819,800 | 1901F12 | D19S0022i |
| 18,031,371 | 18,058,702 | 1902H04 | D19S0051i |
| 18,125,016 | 18,142,343 | 1901C11 | D19S212 |
| 18,357,968 | 18,360,987 | 1902F12 | D19S0215i |
| 18,357,968 | 18,360,987 | 1901G06 | D19S898 |
| 18,503,568 | 18,515,383 | 1903G09 | D19S0284i |
| 18,891,494 | 18,900,436 | $1903 \mathrm{H06}$ | D19S0013i |
| 19,164,008 | 19,173,678 | 9901A08 | D19S0087i |
| 2,427,135 | 2,429,257 | 1904A07 | D19S0370i |
| 2,427,135 | 2,429,257 | 1901H07 | D19S565 |
| 3,545,504 | 3,557,658 | 9901D11 | D19S0106i |
| 3,927,054 | 3,936,461 | 1903B07 | D19S0366i |
| 3,958,748 | 3,990,383 | 1902F06 | D19S0328i |
| 34,994,741 | 35,007,059 | 1902B01 | D19S0044i |
| 34,994,741 | 35,007,059 | 1901D05 | G08036 |
| 37,763,944 | 37,770,171 | 1903B05 | D19S0035i |
| 38,482,776 | 38,485,160 | 9909H08 | D19S0332i |
| 38,482,776 | 38,485,160 | 674H06 | DISO7_10007438 |


| 38,482,776 | 38,485,160 | 093E04 | HUMUT1974 |
| :---: | :---: | :---: | :---: |
| 4,041,319 | 4,075,126 | 1902A02 | D19S0032i |
| 4,180,495 | 4,188,525 | 1903D05 | D19S0123i |
| 4,180,495 | 4,188,525 | 1901D07 | D19S0137i |
| 4,608,557 | 4,621,415 | 9901F10 | D19S0040i |
| 4,608,557 | 4,621,415 | 1901B11 | D19S0131i |
| 4,766,944 | 4,782,716 | 9901G10 | D19S0060i |
| 4,766,944 | 4,782,716 | 9909D07 | D19S0098i |
| 40,451,721 | 40,462,558 | 1902A08 | D19S0085i |
| 40,451,721 | 40,462,558 | 1904F02 | D19S0273i |
| 40,895,670 | 40,899,780 | 1902F01 | D19S0384i |
| 41,070,983 | 41,085,025 | 1903C10 | D19S0002i |
| 41,070,983 | 41,085,025 | 1904F01 | D19S0227i |
| 41,070,983 | 41,085,025 | 1901B07 | D19S876 |
| 41,191,863 | 41,196,981 | 1902B09 | D19S224 |
| 43,770,121 | 43,800,471 | T002D09 | D19S0380i |
| 43,770,121 | 43,800,471 | $1901 \mathrm{E12}$ | D19S422 |
| 43,984,155 | 43,995,422 | 1901F01 | D19S0361i |
| 44,082,455 | 44,091,374 | $9901 \mathrm{E11}$ | D19S0108i |
| 44,082,455 | 44,091,374 | 1902C04 | D19S417 |
| 44,082,455 | 44,091,374 | 1902H07 | D19S881 |
| 44,426,033 | 44,427,609 | 1901F07 | D19S0193i |
| 44,450,997 | 44,452,572 | 9909G07 | D19S0285i |
| 44,589,293 | 44,591,885 | 1902B08 | D19S0375i |
| 44,785,004 | 44,789,954 | 1902A03 | D19S0260i |
| 44,886,786 | 44,891,928 | 1902B02 | D19S0275i |
| 44,913,735 | 44,920,508 | 1901G08 | D19S0083i |
| 447,490 | 456,342 | 1902G10 | D19S0073i |
| 447,490 | 456,342 | 1903 A04 | D19S0338i |
| 45,389,491 | 45,413,314 | 1902H08 | D19S0226i |
| 45,389,491 | 45,413,314 | 1902E07 | HUMUT5576 |
| 45,389,491 | 45,413,314 | $9901 \mathrm{B10}$ | HUMUT6385 |
| 45,645,541 | 45,663,516 | 1904H05 | D19S0210i |
| 45,645,541 | 45,663,516 | 1903F05 | D19S0276i |
| 46,041,284 | 46,226,008 | 9901F08 | D19S223 |
| 46,041,286 | 46,226,008 | 1901B10 | D19S0262i |
| 46,041,286 | 46,226,008 | 1901G07 | HUMUT5036 |
| 46,390,955 | 46,405,284 | 1904C05 | D19S0214i |
| 46528254 | 46551656 | TGFB11 | new design |
| 46528254 | 46551656 | TGFB12 | new design |
| 46,904,377 | 46,925,686 | 239F08 | AC005794.1_22677 |
| 46,904,377 | 46,925,686 | 1904D09 | D19S0223i |
| 46,992,381 | 47,007,431 | 576F01 | chr19.fa.O7frz. 51175481 |
| 46,992,381 | 47,007,431 | 051H06 | D19S423 |
| 47,394,592 | 47,416,115 | 1902A10 | D19S0094i |
| 47,394,592 | 47,416,115 | 9909A08 | D19S0293i |
| 47,577,500 | 47,579,250 | 1902C10 | D19S872 |
| 47,703,298 | 47,724,479 | 1902G05 | D19S0111i |
| 47,776,235 | 47,790,890 | 1904E07 | D19S0103i |
| 47,776,235 | 47,790,890 | 1903F08 | D19S0126i |
| 48,063,198 | 48,075,711 | 239F05 | AC005260.1_494 |
| 48,063,198 | 48,075,711 | 9901B07 | D19S211 |
| 48,549,651 | 48,559,368 | 1904G12 | D19S0280i |
| 48,549,651 | 48,559,368 | 1903F02 | D19S0350i |
| 48,739,032 | 48,771,998 | 1901A08 | D19S0295i |
| 48,739,032 | 48,771,998 | 9909A07 | D19S408 |


| 48,842,088 | 48,866,539 | 9901C08 | D19S0107i |
| :---: | :---: | :---: | :---: |
| 48,912,078 | 48,916,013 | 1903F09 | D19S0029i |
| 48,912,078 | 48,916,013 | 9909B07 | D19S217 |
| 49,839,066 | 49,858,690 | 1904A03 | D19S0229i |
| 49,839,066 | 49,858,690 | 1903F04 | D19S574 |
| 49,943,820 | 49,955,140 | 9901H08 | D19S0164i |
| 49,943,820 | 49,955,140 | 1901A03 | HUMUT7544 |
| 5,157,379 | 5,237,399 | 1904E06 | D19S0265i |
| 5,157,379 | 5,237,399 | 1901F10 | D19S0362i |
| 5,793,902 | 5,802,482 | 1904F10 | D19S0130i |
| 5,793,902 | 5,802,482 | 9901B11 | D19S0325i |
| 5,944,175 | 6,061,554 | 1903D01 | D19S0303i |
| 5,944,175 | 6,061,554 | 1901B01 | D19S0304i |
| 50,196,539 | 50,233,292 | 468B12 | chr19.fa.O7frz.55431947 |
| 50,546,686 | 50,565,669 | 1903 E 07 | D19S0105i |
| 50,546,686 | 50,565,669 | 9901D12 | D19S0317i |
| 50,604,712 | 50,619,017 | 9901D08 | D19S0077i |
| 50,784,865 | 50,797,294 | 1903G08 | D19S0001i |
| 51,059,358 | 51,068,895 | 1904D11 | D19S0221i |
| 51,059,358 | 51,068,895 | 1901A11 | D19S0387i |
| 51,059,358 | 51,068,895 | 1901E02 | D19S0393i |
| 51,214,255 | 51,218,163 | 1904 E10 | D19S0110i |
| 51,869,413 | 51,911,597 | 1904B04 | D19S0253i |
| 52,356,000 | 52.367 .000 | 9901A12 | D19S0181i |
| 52,415,921 | 52,427,863 | 1902F04 | D19S0264i |
| 52,504,971 | 52,517,173 | 1904H08 | D19S0185i |
| 52,544,386 | 52,577,795 | 1901A02 | D19S606 |
| 53,310,515 | 53,365,372 | 1904C01 | D19S0160i |
| 53,403,325 | 53,450,955 | 1904D01 | D19S0287i |
| 53,520,441 | 53,525,623 | 1902H05 | D19S0171i |
| 53,814,360 | 53,825,474 | 1904D07 | D19S0271i |
| 53,943,080 | 53,950,459 | 9901A11 | D19S0166i |
| 53,950,628 | 53,953,395 | 1904G10 | D19S0173i |
| 53,950,628 | 53,953,395 | 1904H04 | D19S0180i |
| 54,149,929 | 54,156,867 | 1903F07 | D19S0179i |
| 54,211,049 | 54,212,159 | 1904A09 | D19S0186i |
| 54,217,939 | 54,244,212 | 1904G05 | D19S0289i |
| 54,280,277 | 54,303,682 | 1901C01 | D19S0297i |
| 54,530,240 | 54,535,675 | 1903C08 | D19S0072i |
| 54,530,240 | 54,535,675 | 1902D06 | D19S604 |
| 54,530,240 | 54,535,675 | 1901D02 | HUMUT2523 |
| 54708304 | 54721402 | FCGRT1 | new design |
| 54708304 | 54721402 | FCGRT2 | new design |
| 54,750,780 | 54,775,626 | 1903H03 | D19S0300i |
| 55,084,723 | 55,124,598 | 1902D07 | D19S0337i |
| 55,571,515 | 55,578,051 | 469B02 | chr19.fa.O7frz.63753376 |
| 55,571,515 | 55,578,051 | 9901H06 | D19S0165i |
| 56,319,977 | 56,325,379 | 154C08 | AF135024.2_4066 |
| 56,319,977 | 56,325,379 | $9901 \mathrm{G12}$ | D19S0353i |
| 56,605,087 | 56,612,869 | 1901A06 | D19S0254i |
| 56,714,795 | 56,726,922 | 1902A09 | D19S0324i |
| 56,806,996 | 56,831,696 | 1903F12 | D19S0068i |
| 56,940,839 | 56,946,962 | 1903 A07 | D19S0027i |
| 56,955,995 | 56,965,591 | 9901C10 | D19S0014i |
| 58,484,666 | 58,486,687 | 1903B11 | D19S0291i |
| 58,484,666 | 58,486,687 | 1902C05 | D19S921 |


| 58,988,650 | 59,019,409 | 9909G06 | D19S0058i |
| :---: | :---: | :---: | :---: |
| 58,988,650 | 59,019,409 | 1904E05 | D19S0288i |
| 59,077,279 | 59,102,713 | 1904H09 | D19S0244i |
| 59,077,279 | 59,102,713 | 670G07 | DISO7_10004464 |
| 59,289,745 | 59,297,806 | 1902G12 | D19S0170i |
| 59,289,745 | 59,297,806 | 1903D03 | D19S0355i |
| 59,412,549 | 59,418,709 | 1904G11 | D19S0230i |
| 59,491,666 | 59,496,077 | 1903H07 | D19S0038i |
| 59,557,047 | 59,568,533 | 1904G06 | D19S0235i |
| 59,618,417 | 59,639,882 | 1902F05 | D19S0239 |
| 59,738,595 | 59,748,862 | 1904F04 | D19S0169i |
| 59,911,791 | 59,916,501 | 664A03 | DISO7_10004466 |
| 6,235,811 | 6,344,184 | 1902C06 | D19S0334i |
| 6,323,444 | 6,326,040 | 1901B12 | D19S0274i |
| 6,482,037 | 6,486,933 | 1904D06 | D19S0391i |
| 6,536,850 | 6,542,163 | 608H08 | chr19.fa.O7frz. 7699148 |
| 6,628,878 | 6,671,660 | 663A12 | chr19.fa.O7frz. 7846308 |
| 6,628,878 | 6,671,660 | 9901F09 | D19S0146i |
| 6,723,722 | 6,808,371 | 9901G11 | D19S0132i |
| 6,838,577 | 6,891,464 | 1901E09 | D19S0144i |
| 6,838,577 | 6,891,464 | 9909D08 | D19S0313i |
| 60,077,361 | 60,095,055 | 1903G03 | D19S926 |
| 60,168,465 | 60,204,318 | 9909D06 | D19S0453i |
| 60,384,428 | 60,412,654 | 1903F01 | D19S605 |
| 60,911,610 | 60,941,580 | 1904G09 | D19S0241i |
| 60,911,610 | 60,941,580 | 1901G05 | Z66860 |
| 61,099,123 | 61,135,489 | 1902G03 | D19S0389 |
| 63,754,745 | 63,758,298 | 1903D11 | D19S0290i |
| 7,067,049 | 7,245,045 | 305B05 | AC010606.5_70425 |
| 7,067,049 | 7,245,045 | 1903C01 | D19S0340i |
| 7,067,049 | 7,245,045 | 1903G04 | D19S406 |
| 7,659,662 | 7,673,032 | 1904A01 | D19S0117i |
| 7,659,662 | 7,673,032 | 9901C07 | D19S905 |
| 7,734,081 | 7,740,491 | 1901C06 | D19S912 |
| 7,874,728 | 7,885,363 | 1903F03 | D19S0351i |
| 776,097 | 783,017 | 1902D05 | D19S0342i |
| 8,023,934 | 8,033,547 | 1902F07 | D19S922 |
| 8,491,689 | 8,548,330 | 1902A11 | D19S0343i |
| 810,665 | 814,624 | 1903C09 | D19S0011i |
| 9,806,999 | 9,821,358 | 1902G06 | D19S0279i |
| 9,931,237 | 9,982,147 | 1902A05 | D19S583 |
| 1,614,666 | 1,727,298 | 0201A07 | D2S0869i |
| 10,101,133 | 10,112,414 | 0205D06 | D2S0137i |
| 101,680,920 | 101,877,584 | 0207D10 | D2S2264 |
| 101,974,738 | 102,011,317 | 0220C03 | HUMUT1265 |
| 102,125,678 | 102,162,766 | 0217B09 | D2S1275i |
| 102,169,865 | 102,222,243 | $0217 \mathrm{B11}$ | D2S1036i |
| 102294394 | 102334929 | IL1RL11 | new design |
| 102,345,529 | 102,381,650 | 0211B05 | D2S0556i |
| 102,401,686 | 102,435,457 | 0208E03 | D2S373 |
| 102,602,598 | 102,694,241 | 0208E11 | D2S0123i |
| 108,360,853 | 108,370,702 | 0216C04 | D2S0211i |
| 108,360,853 | 108,370,702 | 0205B09 | D2S1889 |
| 108,702,369 | 108,767,683 | 0201A02 | D2S0205i |
| 108,877,361 | 108,972,260 | 0216C11 | D2S1281i |
| 11,239,229 | 11,402,162 | 0206B05 | D2S1586i |


| 11,239,229 | 11,402,162 | 0205G04 | D2S168 |
| :---: | :---: | :---: | :---: |
| 11,239,229 | 11,402,162 | 0202G05 | Z67467 |
| 111,597,781 | 111,641,058 | 0205G06 | D2S0198i |
| 111,597,781 | 111,641,058 | 0208A05 | D2S1892 |
| 112,372,662 | 112,503,416 | 377B11 | AC067761.2_149902 |
| 112,372,662 | 112,503,416 | 0221H11 | D2S0532i |
| 113,247,963 | 113,259,442 | $0212 \mathrm{G11}$ | D2S1051i |
| 113303808 | 113310827 | IL1B1 | new design |
| 113303808 | 113310827 | IL1B2 | new design |
| 113303808 | 113310827 | IL1B3 | new design |
| 113387017 | 113392930 | IL1F71 | new design |
| 113387017 | 113392930 | IL1F72 | new design |
| 113387017 | 113392930 | IL1F73 | new design |
| 113,452,077 | 113,459,698 | 0202E11 | D2S1276i |
| 113,591,941 | 113,608,064 | 0201D01 | D2S0193i |
| 113,591,941 | 113,608,064 | 0210G06 | D2S0321i |
| 118,288,725 | 118,306,425 | 0217C04 | D2S0886i |
| 119,416,215 | 119,468,706 | 0215E06 | D2S0233i |
| 119,416,215 | 119,468,706 | 0202B11 | D2S0627i |
| 119,630,289 | 119,632,941 | 0211A06 | D2S0582i |
| 119,630,289 | 119,632,941 | 0205C04 | D2S2254 |
| 119,905,950 | 119,911,486 | 0202A10 | D2S0218i |
| 119,905,950 | 119,911,486 | 0216B01 | D2S0592i |
| 120,233,677 | 120,451,507 | 0218G01 | D2S0897i |
| 120,233,677 | 120,451,507 | 0212H06 | D2S0911i |
| 120,233,677 | 120,451,507 | 0209A04 | D2S1533i |
| 120,819,469 | 120,825,444 | 0202F10 | D2S2329 |
| 120,819,469 | 120,825,444 | 0204A07 | D2S2341 |
| 121,266,327 | 121,466,321 | 0203G09 | D2S2212 |
| 121,266,327 | 121,466,321 | 0206B07 | D2S2258 |
| 121,266,327 | 121,466,321 | 0208B04 | D2S283 |
| 121,266,327 | 121,466,321 | 0207C09 | Z67547 |
| 121,811,825 | 122,123,522 | 278D01 | AC013399.2_117318 |
| 121,811,825 | 122,123,522 | 378G07 | AC018737.3_67168 |
| 121,811,825 | 122,123,522 | 502H02 | chr2.fa.O7frz. 125811119 |
| 121,811,825 | 122,123,522 | 0217D12 | D2S1064i |
| 121,811,825 | 122,123,522 | 0221E05 | D2S343 |
| 127,130,154 | 127,170,716 | 0211E06 | D2S0593i |
| 127,130,154 | 127,170,716 | 0211C07 | D2S0617i |
| 127,130,154 | 127,170,716 | 0219H11 | D2S1467i |
| 127,778,609 | 127,817,240 | 0209E07 | D2S1631i |
| 127,778,609 | 127,817,240 | 0206B04 | D2S2271 |
| 127,892,486 | 127,903,288 | 0218D08 | D2S0214i |
| 127,892,486 | 127,903,288 | 0216D05 | D2S0596i |
| 130,830,088 | 130,848,614 | 0209F03 | D2S1522i |
| 130,830,088 | 130,848,614 | 0209D04 | D2S1536i |
| 136,313,666 | 136,350,481 | 0201H08 | D2S0632i |
| 136,313,666 | 136,350,481 | 0201G05 | Z67485 |
| 136,705,639 | 136,709,450 | 0203E04 | D2S1714i |
| 136,705,639 | 136,709,450 | 0221C09 | D2S2196 |
| 138,438,278 | 138,490,404 | 0213D08 | D2S1302i |
| 138,438,278 | 138,490,404 | 0206F10 | D2S1810i |
| 15,648,753 | 15,688,676 | 0219E10 | D2S1383i |
| 15,648,753 | 15,688,676 | 0218C05 | D2S1389i |
| 15,998,134 | 16,004,580 | 0210A09 | D2S0381i |
| 15,998,134 | 16,004,580 | 0203C08 | D2S1828i |


| 151,835,231 | 151,854,620 | 0214E05 | D2S0657i |
| :---: | :---: | :---: | :---: |
| 151,835,231 | 151,854,620 | 0214D12 | D2S1334i |
| 153,216,334 | 153,283,014 | 0211F07 | D2S0624i |
| 153,216,334 | 153,283,014 | 0204D02 | D2S2299 |
| 156,889,194 | 156,897,474 | 0202B07 | D2S1807i |
| 157,979,377 | 158,008,850 | 0216D06 | D2S0717i |
| 157,979,377 | 158,008,850 | 0211D12 | D2S0724i |
| 160,277,256 | 160,333,330 | 0221D09 | D2S0708i |
| 160,277,256 | 160,333,330 | 0206G12 | D2S156 |
| 160,277,256 | 160,333,330 | 009G10 | D2S306 |
| 160,368,118 | 160,469,493 | 0213E02 | D2S1128i |
| 160,505,506 | 160,627,367 | 0201H05 | D2S1122i |
| 160,664,438 | 160,765,009 | 0213C11 | D2S1403i |
| 161,701,712 | 161,800,928 | 0217B03 | D2S0677i |
| 161,701,712 | 161,800,928 | 0211 E 12 | D2S0725i |
| 162,557,001 | 162,639,298 | 0218F11 | D2S0929i |
| 162,557,001 | 162,639,298 | 0214G04 | D2S0932i |
| 162,831,835 | 162,883,285 | 0215E09 | D2S0917i |
| 170,149,096 | 170,202,500 | 0216G02 | D2S0649i |
| 170,149,096 | 170,202,500 | 0213E08 | D2S1313i |
| 172,487,204 | 172,556,846 | 9902F08 | D2S0689i |
| 172,487,204 | 172,556,846 | 0214H01 | D2S1444i |
| 173,000,616 | 173,079,256 | 0210A02 | D2S0048i |
| 173,129,025 | 173,172,108 | 0211H05 | D2S0581i |
| 174,481,504 | 174,538,676 | 381G03 | AC055875.2_105198 |
| 174,481,504 | 174,538,676 | 504B05 | chr2.fa.O7frz. 181842884 |
| 174,921,124 | 174,968,689 | 0201D08 | D2S0320i |
| 174,921,124 | 174,968,689 | 0201E02 | D2S1731i |
| 175,132,548 | 175,255,873 | 0211G11 | D2S0705i |
| 175,132,548 | 175,255,873 | 0209H08 | D2S1665i |
| 176,689,738 | 176,692,916 | 0210G01 | D2S0046i |
| 176,689,738 | 176,692,916 | 0219H07 | D2S0260i |
| 177,965,731 | 178,112,411 | 0211F10 | D2S0687i |
| 177,965,731 | 178,112,411 | 0213B11 | D2S1399i |
| 177,965,731 | 178,112,411 | 0203 E 07 | D2S2173 |
| 182,029,864 | 182,110,719 | 0210G05 | D2S0253i |
| 182,029,864 | 182,110,719 | 0215D10 | D2S1147i |
| 183,406,982 | 183,439,743 | 0216A09 | D2S0735i |
| 183,651,732 | 183,673,616 | 0215D06 | D2S1380i |
| 183,651,732 | 183,673,616 | 0220E12 | D2S1462i |
| 187,163,045 | 187,253,873 | 279F08 | AC017026.5_11120 |
| 187,163,045 | 187,253,873 | 381G05 | AC017026.6_146393 |
| 187,163,045 | 187,253,873 | 381A06 | AC017101.8_55453 |
| 189,547,344 | 189,585,717 | 381G07 | AC066694.2_218537 |
| 189,547,344 | 189,585,717 | 429C05 | chr2.fa.O7frz. 197184084 |
| 189,547,344 | 189,585,717 | 0219D03 | D2S0774i |
| 190,133,561 | 190,153,858 | 0215H10 | D2S0739i |
| 190,133,561 | 190,153,858 | 0219B09 | D2S0746i |
| 191,542,121 | 191,587,181 | 0212H02 | D2S0775i |
| 191,542,121 | 191,587,181 | 0215G02 | D2S1178i |
| 191,602,551 | 191,724,539 | 0214H03 | D2S0745i |
| 197,336,917 | 197,372,670 | 0209H06 | D2S1614i |
| 197,336,917 | 197,372,670 | 0207F02 | D2S1621i |
| 198,059,553 | 198,073,243 | 0214C08 | D2S1664i |
| 20,264,039 | 20,288,675 | 0210C08 | D2S0359i |
| 201,443,924 | 201,462,244 | 0204E09 | D2S116 |


| 201,443,924 | 201,462,244 | 0207D04 | D2S1707i |
| :---: | :---: | :---: | :---: |
| 201,756,100 | 201,802,372 | 0209E04 | D2S1539i |
| 201,806,396 | 201,854,521 | 0212G01 | D2S0749i |
| 202,949,916 | 203,140,719 | 0217C11 | D2S0201i |
| 202,949,916 | 203,140,719 | 0218G11 | D2S1768i |
| 203,811,658 | 203,878,579 | 0217B01 | D2S1174i |
| 203,811,658 | 203,878,579 | 0213E04 | D2S1183i |
| 204,279,443 | 204,310,801 | 0219F06 | G09915 |
| 204,279,443 | 204,310,801 | 0208F10 | HUMUT426 |
| 204,440,754 | 204,446,928 | 169G12 | AF225900.1_4678 |
| 204,440,754 | 204,446,928 | 0219C01 | D2S1188i |
| 204,440,754 | 204,446,928 | 0221H04 | D2S307 |
| 207,653,323 | 207,738,859 | 0208A11 | D2S0057i |
| 207,653,323 | 207,738,859 | 0215A11 | D2S0116i |
| 207,653,323 | 207,738,859 | 0213H11 | D2S1424i |
| 208,102,931 | 208,171,818 | 0221A07 | D2S1701i |
| 211,050,678 | 211,252,076 | 340F03 | AC021150.7_98846 |
| 211,050,678 | 211,252,076 | 0220A12 | D2S0094i |
| 211,050,678 | 211,252,076 | 0221D05 | D2S0298i |
| 211,050,678 | 211,252,076 | 0213A12 | D2S1430i |
| 215,933,409 | 216,009,041 | 0214G12 | D2S0265i |
| 215,933,409 | 216,009,041 | 0210H06 | D2S0323i |
| 216,680,435 | 216,779,248 | 0221F07 | D2S0052i |
| 216,680,435 | 216,779,248 | 0219G07 | D2S0226i |
| 216,680,435 | 216,779,248 | 0205G10 | D2S0230i |
| 216,985,441 | 217,056,021 | 0204D05 | D2S0809i |
| 216,985,441 | 217,056,021 | 0206B06 | D2S1655i |
| 216,985,441 | 217,056,021 | 0219C03 | D2S1659i |
| 217,206,372 | 217,237,404 | 0217D03 | D2S0278i |
| 217,206,372 | 217,237,404 | 0219F12 | D2S1476i |
| 218,698,991 | 218,710,220 | 0204C02 | D2S1206i |
| 218,955,161 | 218,968,994 | 0219D05 | D2S0791i |
| 218,955,161 | 218,968,994 | 0220A10 | D2S1538i |
| 219,354,745 | 219,388,259 | 0219F02 | D2S0828i |
| 219,354,745 | 219,388,259 | 011A01 | D2S2250 |
| 219,354,745 | 219,388,259 | 0221D02 | G08149 |
| 219,628,173 | 219,633,433 | 0210A07 | D2S0336i |
| 219,628,173 | 219,633,433 | 0210B07 | D2S0340i |
| 219,628,173 | 219,633,433 | 0205G11 | D2S0790i |
| 219,822,677 | 219,826,882 | 0220B12 | D2S0090i |
| 220,145,161 | 220,148,679 | 0204H01 | D2S1338i |
| 227,578,168 | 227,737,519 | 0214E03 | D2S0276i |
| 227,578,168 | 227,737,519 | 0221G02 | D2S1349 |
| 227,737,525 | 227,887,751 | 0201C03 | D2S1802i |
| 228,045,286 | 228,130,548 | 0204E11 | D2S1673i |
| 228,386,814 | 228,390,494 | 280F06 | AC068692.3_9554 |
| 228,386,814 | 228,390,494 | 0214F12 | HUMUT8098 |
| 230,741,896 | 230,792,932 | 0220G02 | D2S172 |
| 230,741,896 | 230,792,932 | 0220F06 | D2S1735i |
| 230,741,896 | 230,792,932 | 0206B11 | D2S1813i |
| 231,681,199 | 231,698,068 | 0215H09 | D2S0292i |
| 231,681,199 | 231,698,068 | 0215G01 | D2S0794i |
| 233,633,433 | 233,781,288 | 0221A06 | D2S331 |
| 233,633,433 | 233,781,288 | 0220D03 | HUMUT8067 |
| 234,191,030 | 234,346,695 | 0220E08 | D2S0042i |
| 234,191,030 | 234,346,695 | 0218C03 | D2S0839i |


| 234,191,030 | 234,346,695 | 0216F12 | D2S1220i |
| :---: | :---: | :---: | :---: |
| 234,191,030 | 234,346,695 | 082D06 | Z67659 |
| 237,143,182 | 237,155,730 | 0203C10 | D2S0288i |
| 237,143,182 | 237,155,730 | 0214B03 | D2S1219i |
| 237,897,401 | 237,987,559 | 0206A04 | D2S1796i |
| 237,897,401 | 237,987,559 | 0207 F 06 | D2S1806i |
| 238,893,821 | 238,972,536 | 383C04 | AC013400.4_10628 |
| 238,893,821 | 238,972,536 | 506H10 | chr2.fa.O7frz. 250246308 |
| 238,893,821 | 238,972,536 | 0207H02 | D2S0335i |
| 239,000,365 | 239,025,630 | 0220D04 | D2S0841i |
| 239,000,365 | 239,025,630 | 0204C09 | D2S1221i |
| 239,635,319 | 239,987,580 | 0213C05 | D2S1202i |
| 239,635,319 | 239,987,580 | 0205H02 | D2S1205i |
| 239,635,319 | 239,987,580 | 0213F05 | D2S1214i |
| 239,635,319 | 239,987,580 | 0202H10 | D2S1704i |
| 24,126,075 | 24,140,055 | 0219E03 | D2S0149i |
| 24,126,075 | 24,140,055 | 0207A05 | D2S1779i |
| 241,148,144 | 241,152,104 | 0219G05 | D2S0338i |
| 241,148,144 | 241,152,104 | 0209D07 | D2S1629i |
| 241,903,396 | 241,942,115 | 0210C01 | D2S0033i |
| 241,903,396 | 241,942,115 | 0203H09 | D2S1565i |
| 242,146,865 | 242,162,226 | 0220G05 | D2S1540i |
| 242,440,711 | 242,449,731 | 0209G07 | D2S1637i |
| 25,237,226 | 25,245,063 | 0210D09 | D2S0397i |
| 25,237,226 | 25,245,063 | 0204C04 | D2S171 |
| 27,383,769 | 27,384,634 | 0219A11 | D2S1352i |
| 29,269,144 | 29,997,936 | 0214G01 | D2S0965i |
| 29,269,144 | 29,997,936 | 0216E02 | D2S1255i |
| 29,269,144 | 29,997,936 | 0202A04 | D2S146 |
| 29,269,144 | 29,997,936 | 0218A05 | D2S1687i |
| 29,269,144 | 29,997,936 | 0209E10 | D2S1765i |
| 29,269,144 | 29,997,936 | 0204E02 | D2S2383 |
| 31,410,691 | 31,491,117 | 0215F01 | D2S0437i |
| 31,410,691 | 31,491,117 | 0210H11 | D2S0446i |
| 31,410,691 | 31,491,117 | 0202H07 | D2S2203 |
| 31,410,691 | 31,491,117 | 0203H01 | D2S352 |
| 32,303,022 | 32,344,427 | 0218E10 | D2S0143i |
| 32,303,022 | 32,344,427 | 0217C12 | D2S1405i |
| 32,435,234 | 32,697,470 | 0220B04 | D2S0152i |
| 32,435,234 | 32,697,470 | 0215D07 | D2S0311i |
| 33,025,896 | 33,478,080 | 0210G03 | D2S0151i |
| 33,025,896 | 33,478,080 | 0210D08 | D2S0363i |
| 33,025,896 | 33,478,080 | 0210F08 | D2S0371i |
| 33,025,896 | 33,478,080 | 0210C09 | D2S0391i |
| 33,025,896 | 33,478,080 | 0205C12 | D2S0453i |
| 33,025,896 | 33,478,080 | 0212B09 | D2S0985i |
| 33,025,896 | 33,478,080 | 0212D09 | D2S0992i |
| 33,025,896 | 33,478,080 | 0206B03 | D2S2325 |
| 33,025,896 | 33,478,080 | 0205E12 | D2S2347 |
| 33,514,920 | 33,643,162 | 0215D08 | D2S0384i |
| 37,331,149 | 37,398,541 | 0215H01 | D2S0873i |
| 37,331,149 | 37,398,541 | 0214A09 | D2S0877i |
| 37,331,149 | 37,398,541 | 0209E02 | D2S1501i |
| 38,148,154 | 38,156,796 | 0215H06 | D2S0459i |
| 38,148,154 | 38,156,796 | 0201A08 | D2S0967i |
| 38,878,375 | 38,956,525 | 0210F10 | D2S0416i |


| 38,878,375 | 38,956,525 | 0214G11 | D2S0980i |
| :---: | :---: | :---: | :---: |
| 38,878,375 | 38,956,525 | 0208C01 | D2S2331 |
| 39,066,469 | 39,201,067 | 0202D03 | D2S0425i |
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| 39,329,911 | 39,517,946 | 0210D01 | D2S0036i |
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| 43,717,916 | 43,848,630 | 0202F06 | D2S119 |
| 45,732,547 | 46,268,633 | 0217F04 | D2S0434i |
| 45,732,547 | 46,268,633 | 0220B11 | D2S1482i |
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| 45,732,547 | 46,268,633 | T002B12 | D2S2291 |
| 46,661,580 | 46,697,708 | 0210A11 | D2S0423i |
| 46,661,580 | 46,697,708 | 0220D06 | D2S1719i |
| 46,779,595 | 46,843,431 | 0220A09 | D2S1717i |
| 48,395,374 | 48,459,938 | 0215E08 | D2S0988i |
| 48,395,374 | 48,459,938 | 0209B06 | D2S1595i |
| 48,767,471 | 48,836,321 | 0218B03 | D2S0420i |
| 49,043,156 | 49,235,134 | 0209D02 | D2S1500i |
| 49,043,156 | 49,235,134 | 0209A03 | D2S1509i |
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| 58,127,224 | 58,240,510 | 376B06 | AC068193.5_165968 |
| 58,127,224 | 58,240,510 | 0220C10 | D2S0443i |
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| 6,935,247 | 6,955,821 | 0215D11 | D2S0351i |
| 6,935,247 | 6,955,821 | 0212E07 | D2S0936i |
| 60,962,254 | 61,003,682 | 0211E03 | D2S0524i |
| 64,173,499 | 64,225,062 | 0212G09 | D2S1003i |
| 64,173,499 | 64,225,062 | 0212H09 | D2S1008i |
| 68,203,572 | 68,341,866 | 0214D03 | D2S0182i |
| 68,203,572 | 68,341,866 | 0217G12 | D2S0305i |
| 68,203,572 | 68,341,866 | 0202E04 | D2S0484i |
| 69,995,707 | 70,023,581 | 376H04 | AC019206.3_137449 |
| 69,995,707 | 70,023,581 | 0221C01 | D2S0183i |
| 69,995,707 | 70,023,581 | 0216B12 | D2S0313i |
| 70,527,924 | 70,634,438 | 0214D07 | D2S0039i |
| 70,527,924 | 70,634,438 | 0206F08 | D2S0076i |
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| 70,910,855 | 70,916,461 | 0202D01 | D2S1464i |
| 72,209,875 | 72,228,471 | 0208E02 | D2S2112 |
| 73,842,837 | 73,860,756 | 0201A12 | D2S1715i |
| 74,598,766 | 74,606,826 | 0205B12 | D2S0066i |
| 74,598,766 | 74,606,826 | 0209A10 | D2S1745i |
| 74,634,795 | 74,638,181 | 0220E03 | D2S0062i |
| 85,737,951 | 85,748,823 | 0214B09 | D2S0171i |
| 85,737,951 | 85,748,823 | 0219B08 | D2S0315i |
| 85,919,782 | 85,969,648 | 0216E03 | D2S0575i |
| 85,919,782 | 85,969,648 | 0208C03 | D2S388 |
| 86,521,954 | 86,573,350 | 377E03 | AC068288.3_48784 |
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| 86,865,239 | 86,871,638 | 0214B05 | D2S0560i |
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| 97,696,461 | 97,722,755 | 0221B09 | D2S2222 |
| 99,225,141 | 99,238,002 | 0215D04 | D2S1049i |
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| 1,297,622 | 1,321,806 | 2005H10 | D20S0258i |
| 1,399,386 | 1,420,233 | 2004B05 | D20S0382i |
| 1,491,568 | 1,548,655 | 2004E05 | D20S0171i |
| 1,491,568 | 1,548,655 | 2001E04 | D20S906 |
| 1,822,813 | 1,868,543 | 2001B11 | D20S0035i |
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| 10,566,334 | 10,602,636 | 2005A09 | D20S0238i |
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| 2,224,647 | 2,269,725 | 2001A09 | D20S0434i |
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| 2,769,366 | 2,967,320 | 2005D03 | D20S0271i |
| 2,769,366 | 2,967,320 | 2002G03 | D20S0337i |
| 22,509,643 | 22,514,102 | 2001B05 | D20S0179 |
| 22,974,270 | 22,978,301 | 2002D06 | D20S0180i |
| 22,974,270 | 22,978,301 | 2003A10 | D20S0197i |
| 23,007,995 | 23,014,977 | 2002F06 | D20S0077i |
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| 29,565,892 | 29,621,031 | 2004C01 | D20S0416i |
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| 29,996,419 | 30,003,556 | 2001B09 | D20S863 |
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| 3,724,386 | 3,734,762 | 2001E02 | D20S0131i |
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| 31,541,589 | 31,701,503 | 2005E06 | D20S0007i |
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| 31,862,780 | 31,905,831 | 2002B04 | D20S878 |
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| 33,212,131 | 33,228,828 | 2001A02 | D20S0376i |
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| 34,953,761 | 35,013,590 | 2004B04 | D20S0396i |
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| 37,024,409 | 37,101,778 | 2003F10 | D20S0215i |
| 39,199,291 | 39,237,775 | 2004B01 | D20S0418i |
| 4,614,996 | 4,630,236 | 2004F09 | D20S0053i |
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| 40,134,806 | 41,252,024 | 243F09 | AL117374.39_58894 |
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| 40,134,806 | 41,252,024 | 2003E09 | D20S0191i |
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| 42,681,577 | 42,713,797 | 2003F04 | D20S0030i |
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| 43,028,529 | 43,142,014 | 2005A10 | D20S0353i |
| 43,028,529 | 43,142,014 | 2004H03 | D20S119 |
| 43,028,529 | 43,142,014 | 240D10 | Z93016.2_66966 |
| 43,185,480 | 43,186,520 | 9909B10 | D20S481 |
| 43,269,052 | 43,271,827 | 2005G06 | D20S0335i |
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| 43,387,342 | 43,410,478 | 2005G04 | D20S0300i |
| 43,903,768 | 43,919,442 | 246E05 | AL050348.21_45960 |
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| 44,180,313 | 44,366,257 | 2002G09 | D20S0390i |
| 44,180,313 | 44,366,257 | 2004A01 | D20S836 |
| 47,553,818 | 47,618,114 | 2003F11 | D20S0241i |
| 47,553,818 | 47,618,114 | 2005B10 | D20S0247i |
| 47,553,818 | 47,618,114 | 2004H11 | D20S0349 |
| 47,862,657 | 47,942,179 | 2005F11 | D20S0299i |
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| 48,560,294 | 48,634,706 | 2004C06 | D20S0024i |
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| 49,441,083 | 49,592,665 | 2004D07 | D20S0056i |
| 49,441,083 | 49,592,665 | 2005C10 | D20S0363i |
| 49,441,083 | 49,592,665 | 2003C02 | D20S857 |
| 54,257,195 | 54,258,278 | 2002D04 | D20S0333i |
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| 54,637,765 | 54,647,746 | 2005C12 | D20S0301i |


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| 56,990,597 | 57,015,697 | 2005C06 | D20S0082i |
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| 57,308,877 | 57,334,442 | 2002D08 | D20S171 |
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| 6,696,311 | 6,708,927 | 2003B10 | D20S0199i |
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| 61,337,680 | 61,342,299 | 2005C05 | D20S0067i |
| 61,589,810 | 61,600,949 | 2004D04 | D20S0387i |
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| 62,181,932 | 62,215,047 | 2002H03 | D20S0432i |
| 10,042,713 | 10,120,798 | 2004D08 | D20S0005i |
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| 36,679,559 | 36,710,995 | 2102A01 | D21S0090i |
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| 37,661,729 | 37,809,347 | $2101 \mathrm{E12}$ | D21S0183i |
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| 38,675,671 | 38,955,488 | 2103F03 | D21S0025i |
| 38,675,671 | 38,955,488 | 2102E10 | D21S0200i |
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| 40,306,213 | 41,140,909 | 2101D12 | D21S0209i |
| 40,306,213 | 41,140,909 | 2103E11 | D21S0213i |
| 40,306,213 | 41,140,909 | 2101F07 | D21S1887 |
| 40,306,213 | 41,140,909 | 2101F03 | D21S1906 |
| 41,655,820 | 41,702,739 | 2102H05 | D21S0088i |
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| 41,720,024 | 41,753,008 | 2102D06 | D21S0228i |
| 42,599,751 | 42,608,775 | 2102F03 | D21S0151i |
| 43,963,406 | 44,006,608 | 2101F02 | D21S0020i |
| 43,963,406 | 44,006,608 | 2103B11 | D21S0205i |
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| 45,130,296 | 45,173,181 | 2101C05 | D21S0241i |
| 45737914 | 45786779 | SLC19A1 | new design |
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| 15,945,849 | 15,971,405 | 2202F03 | D22S0091i |
| 16,501,485 | 16,593,383 | 9910C08 | D22S0210i |
| 16,591,460 | 16,631,812 | 2202F12 | D22S0026i |
| 16,591,460 | 16,631,812 | 2201C06 | D22S0231i |
| 17,141,172 | 17,159,474 | 9910D07 | D22S0216i |
| 17,403,798 | 17,489,962 | 2201A09 | D22S0151i |
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| 18,091,066 | 18,092,297 | 9910E02 | D22S0041i |
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| 19,108,875 | 19,122,146 | 2202B03 | D22S0089 |
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| 20,326,542 | 20,328,588 | 2201G07 | D22S446 |
| 20,443,946 | 20,551,970 | 2201E11 | D22S0109i |
| 20,929,200 | 20,929,926 | 2202A11 | D22S0119i |
| 21,220,123 | 21,231,768 | 2201B10 | D22S0232i |
| 21,852,552 | 21,990,224 | 9910C03 | D22S0055i |
| 21,852,552 | 21,990,224 | 103C02 | HUMUT8144 |
| 22,245,312 | 22,252,495 | 9910F03 | D22S0068i |
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| 22,706,141 | 22,714,271 | 2202F02 | D22S0169i |
| 22,737,765 | 22,904,596 | 9910E06 | D22S0184i |
| 22,737,765 | 22,904,596 | 2202F01 | D22S1174 |
| 23,153,537 | 23,168,325 | 2201B06 | D22S0176i |
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| 26,474,265 | 26,527,486 | 2202D06 | D22S0063i |
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| 28,966,441 | 28,972,748 | 2202E07 | D22S0018i |


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| 30,125,539 | 30,160,438 | 9910H06 | D22S0193i |
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| 35,906,152 | 35,914,276 | 2202F05 | D22S0148i |
| 35,951,238 | 35,970,251 | 2202G08 | D22S0171i |
| 36,216,346 | 36,245,193 | 2201C10 | D22S0135i |
| 36,216,346 | 36,245,193 | 9910B06 | D22S0177i |
| 36,837,448 | 36,907,763 | 2202C11 | D22S0099i |
| 36,837,448 | 36,907,763 | 2201F10 | D22S0106i |
| 37,209,389 | 37,232,262 | 2202G12 | D22S0092i |
| 37,209,389 | 37,232,262 | 9910F05 | D22S0166i |
| 37,460,681 | 37,481,928 | 2101D09 | D21S1919 |
| 37,460,681 | 37,481,928 | 053B05 | D21S267 |
| 37,949,310 | 37,971,006 | 2202F09 | D22S0037i |
| 38,125,692 | 38,163,078 | 2202G10 | D22S0002i |
| 38,246,515 | 38,248,637 | 2201B01 | D22S428 |
| 38,627,032 | 38,698,204 | 2201G09 | D22S0108i |
| 38,627,032 | 38,698,204 | 2202G04 | D22S284 |
| 39,817,736 | 39,906,024 | 2202C06 | D22S0021i |
| 40,664,687 | 40,673,094 | 2202B11 | D22S0007i |
| 40,664,687 | 40,673,094 | 2202A10 | D22S0064i |
| 40,852,445 | 40,856,827 | 2201D03 | D22S0227i |
| 40,885,963 | 40,941,389 | 9910B07 | D22S0199i |
| 41,108,917 | 41,158,340 | 2202H04 | D22S0200i |
| 41,108,917 | 41,158,340 | 2202H03 | D22S0201i |
| 41,309,671 | 41,340,906 | 252A05 | Z93241.11_118962 |
| 41,418,071 | 41,446,820 | 2201D06 | HUMUT1091 |
| 41,836,701 | 41,855,662 | 2201F05 | D22S1151 |
| 41,836,701 | 41,855,662 | 054F01 | D22S1179 |
| 44,925,163 | 45,018,317 | 2202F04 | D22S1149 |
| 45,394,963 | 45,454,352 | 2201H01 | D22S0202i |
| 45,394,963 | 45,454,352 | 9910E08 | D22S0225i |
| 45,458,971 | 45,512,816 | 2202F10 | D22S0173i |
| 49,311,047 | 49,315,321 | 2202F07 | D22S0096i |
| 49,385,997 | 49,396,843 | 9910D08 | D22S0222i |
| 49,459,936 | 49,518,507 | 728A10 | - |
| 10,181,563 | 10,260,427 | 0304D05 | D3S1329i |
| 103,029,547 | 103,062,556 | 0301F10 | D3S1228i |
| 103,029,547 | 103,062,556 | 632E08 | DISO7_10001184 |
| 106,568,403 | 106,778,434 | 0313F05 | D3S1080i |
| 106,568,403 | 106,778,434 | 0310D06 | D3S1083i |
| 106,568,403 | 106,778,434 | 0315A01 | D3S1591 |
| 106,859,799 | 107,070,577 | 0310G09 | D3S0029i |


| 106,859,799 | 107,070,577 | 0309D06 | D3S0668i |
| :---: | :---: | :---: | :---: |
| 106,859,799 | 107,070,577 | 0301G07 | D3S1493i |
| 109,244,631 | 109,292,625 | 0310A05 | D3S0548i |
| 109,244,631 | 109,292,625 | 0309C01 | D3S0552i |
| 109,244,631 | 109,292,625 | 0302C06 | Z67497 |
| 11,269,400 | 11,279,415 | 0314A07 | D3S0216i |
| 11,269,400 | 11,279,415 | 0301D05 | D3S0835i |
| 110,024,321 | 110,056,542 | 0303H10 | D3S1302 |
| 112,273,555 | 112,395,063 | 0307C11 | D3S0116i |
| 112,273,555 | 112,395,063 | 0313B10 | G08281 |
| 112,743,546 | 112,853,906 | 0309F05 | D3S0644i |
| 112,743,546 | 112,853,906 | 0302E07 | D3S1443i |
| 112,743,546 | 112,853,906 | 0306H04 | D3S1572 |
| 113,522,943 | 113,564,349 | 0307C06 | D3S1359i |
| 113,522,943 | 113,564,349 | 0315D12 | Z66960 |
| 113,667,463 | 113,701,066 | 0301F03 | Z67725 |
| 114,017,246 | 114,047,487 | 0312E12 | D3S0657i |
| 114,017,246 | 114,047,487 | 0304C11 | D3S1410i |
| 114,122,746 | 114,176,650 | 0308F03 | D3S0290i |
| 12,169,568 | 12,175,851 | 393G02 | AC015546.16_112255 |
| 12,169,568 | 12,175,851 | 396A05 | AC026166.2_143575 |
| 12,169,568 | 12,175,851 | 0317H08 | D3S1127i |
| 12,304,359 | 12,450,843 | 0304H07 | D3S0036i |
| 12,304,359 | 12,450,843 | 0311H05 | D3S0208i |
| 12,304,359 | 12,450,843 | 0317C09 | D3S1133i |
| 120,102,167 | 120,347,588 | 0303G01 | D3S0934i |
| 120,102,167 | 120,347,588 | 0315H02 | D3S3515 |
| 120,102,167 | 120,347,588 | 729D03 | DIJ28_10010297 |
| 120,725,832 | 120,761,139 | 285H01 | AC069519.8_29705 |
| 120,725,832 | 120,761,139 | 013E09 | D3S3513 |
| 120,792,984 | 120,831,342 | 398B01 | AC073352.7_5796 |
| 120,982,021 | 121,020,022 | 0306B04 | D3S3620 |
| 121,028,233 | 121,295,954 | 0314D03 | D3S0118i |
| 121,937,926 | 121,984,605 | 0315B05 | D3S1059i |
| 123,256,911 | 123,322,673 | 0310G01 | D3S1225i |
| 123,256,911 | 123,322,673 | 0317A01 | D3S1779i |
| 123,256,911 | 123,322,673 | 0303B04 | D3S3720 |
| 124,813,833 | 125,085,839 | 0316C06 | D3S1322i |
| 124,813,833 | 125,085,839 | 0306E04 | D3S3552 |
| 125,964,485 | 126,088,842 | 0308E05 | D3S0359i |
| 125,964,485 | 126,088,842 | 0308G08 | D3S0446i |
| 125,964,485 | 126,088,842 | 0315D07 | D3S0698i |
| 128,190,192 | 128,238,922 | 0307B03 | D3S1258i |
| 128,190,192 | 128,238,922 | 0304D07 | D3S1459i |
| 128,799,943 | 128,823,969 | 0310H09 | D3S0356i |
| 128,799,943 | 128,823,969 | 013F11 | D3S3607 |
| 129,253,902 | 129,273,216 | 0313A04 | D3S0790i |
| 129,253,902 | 129,273,216 | 0304C01 | Z67486 |
| 129,680,960 | 129,694,718 | 0315F04 | D3S0688i |
| 130,262,300 | 130,263,941 | 0313 E 12 | D3S0360i |
| 130,756,708 | 130,808,351 | 0311 E 07 | D3S0958i |
| 130,756,708 | 130,808,351 | 0307G02 | D3S1250i |
| 131,176,253 | 131,179,470 | 0314E02 | D3S0200i |
| 131,176,253 | 131,179,470 | 0312B11 | D3S0348i |
| 131,176,253 | 131,179,470 | 0302B07 | D3S1176i |
| 131,880,468 | 131,948,340 | 0309F10 | D3S0809i |


| 131,880,468 | 131,948,340 | 0305B06 | D3S1053i |
| :---: | :---: | :---: | :---: |
| 133,798,784 | 133,804,072 | 0302G03 | D3S1306i |
| 135,996,950 | 136,461,999 | 390E07 | AC016951.9_56629 |
| 135,996,950 | 136,461,999 | 0308D08 | D3S0442i |
| 135,996,950 | 136,461,999 | 0311C06 | D3S0959i |
| 135,996,950 | 136,461,999 | 0316 E 05 | D3S1590 |
| 135,996,950 | 136,461,999 | 0315E10 | D3S3641 |
| 135,996,950 | 136,461,999 | 0316A04 | D3S3696 |
| 135,996,950 | 136,461,999 | $0303 A 05$ | Z67391 |
| 138,159,397 | 138,212,610 | 0308D05 | D3S0357i |
| 138,159,397 | 138,212,610 | 0301A12 | D3S3617 |
| 139,856,921 | 139,960,875 | 0315F01 | D3S0555i |
| 139,856,921 | 139,960,875 | 0307E07 | D3S1383i |
| 14,161,648 | 14,195,143 | 0312A05 | D3S0854i |
| 14,964,240 | 15,065,784 | 0312G02 | D3S0836i |
| 14,964,240 | 15,065,784 | 0301H08 | D3S0841i |
| 140,145,756 | 140,148,491 | 0309D02 | D3S0565i |
| 140,145,756 | 140,148,491 | 0304C04 | G09845 |
| 142,939,741 | 142,947,933 | 0308A05 | D3S0345i |
| 142,939,741 | 142,947,933 | 0301F12 | D3S1451i |
| 143,078,160 | 143,128,072 | 0305F10 | D3S1061i |
| 144,466,754 | 145,049,979 | 386E07 | AC026673.12_150939 |
| 144,466,754 | 145,049,979 | 386D07 | AC026673.12_52550 |
| 144,466,754 | 145,049,979 | 398G05 | AC073242.3_44298 |
| 144,466,754 | 145,049,979 | 515D08 | chr3.fa.O7frz. 163941355 |
| 144,466,754 | 145,049,979 | 0305H03 | D3S1171i |
| 144,466,754 | 145,049,979 | 0307B04 | D3S1276i |
| 144,466,754 | 145,049,979 | 0317A12 | D3S1281i |
| 144,466,754 | 145,049,979 | 0305D08 | D3S1536i |
| 144,466,754 | 145,049,979 | 014E03 | D3S3599 |
| 149,898,355 | 149,943,478 | 390F08 | AF245699.1_60143 |
| 149,898,355 | 149,943,478 | 0312D03 | D3S0592i |
| 149,898,355 | 149,943,478 | 0317A09 | D3S1131i |
| 150,230,604 | 150,287,007 | 0313D08 | D3S0585i |
| 150,230,604 | 150,287,007 | 0303G02 | D3S1555 |
| 150,330,061 | 150,373,995 | 0317H01 | D3S0362i |
| 152,412,595 | 152,478,847 | 387F12 | AC024886.10_174359 |
| 152,412,595 | 152,478,847 | 0310E04 | D3S1000i |
| 152,637,167 | 152,659,187 | 0301B10 | D3S1499i |
| 154,035,426 | 154,038,535 | 0302G04 | D3S0145i |
| 154,035,426 | 154,038,535 | 0301F01 | D3S1193i |
| 155,476,152 | 155,524,971 | 0301D08 | D3S3677 |
| 155,476,152 | 155,524,971 | 0305H05 | D3S3710 |
| 156,280,153 | 156,384,186 | 0312E11 | D3S0134i |
| 156,280,153 | 156,384,186 | 0312G05 | D3S1004i |
| 158,637,301 | 158,644,071 | 0307 A 05 | D3S1311i |
| 16,949,586 | 17,107,089 | 0303C06 | D3S1429i |
| 16,949,586 | 17,107,089 | 0304D08 | D3S1452i |
| 161,189,323 | 161,196,500 | 388C05 | AC026118.8_8957 |
| 161,189,323 | 161,196,500 | 0311D01 | D3S2442 |
| 161,189,323 | 161,196,500 | 0303B05 | D3S3580 |
| 161,427,938 | 161,650,320 | 0306C10 | D3S0398i |
| 161,427,938 | 161,650,320 | 0315C07 | D3S0693i |
| 161,427,938 | 161,650,320 | 0301 E 11 | D3S1482i |
| 162,284,365 | 162,305,854 | 0307B06 | D3S1354i |
| 168,884,388 | 168,935,345 | 0311G10 | D3S0989i |


| 168,884,388 | 168,935,345 | 0303D02 | D3S1494i |
| :---: | :---: | :---: | :---: |
| 170,285,244 | 170,347,054 | 0308E07 | D3S0395i |
| 170,285,244 | 170,347,054 | T003G02 | D3S1282 |
| 170,965,092 | 170,965,542 | 0309H08 | D3S0726i |
| 170,965,092 | 170,965,542 | 0302H04 | D3S3523 |
| 171,422,919 | 171,506,459 | 0305D04 | D3S1013i |
| 171,422,919 | 171,506,459 | 0315B06 | D3S1289i |
| 173,706,158 | 173,723,963 | 0306B09 | D3S0009i |
| 173,706,158 | 173,723,963 | 0315C10 | D3S0162i |
| 178,221,867 | 178,397,734 | 0312D09 | D3S1017i |
| 178,221,867 | 178,397,734 | 0305A12 | D3S1092i |
| 178,221,867 | 178,397,734 | 0311D03 | D3S1522i |
| 180,349,005 | 180,435,194 | 0317F06 | D3S1052i |
| 180,349,005 | 180,435,194 | 0315D01 | Z66727 |
| 184,322,697 | 184,363,317 | 0303C09 | D3S1295i |
| 184,322,697 | 184,363,317 | 0317 D 12 | D3S1301i |
| 185,572,467 | 185,578,626 | 389H02 | AC068634.11_165297 |
| 185,572,467 | 185,578,626 | 0303D05 | D3S3578 |
| 186,563,664 | 186,683,322 | 0317B09 | D3S1132i |
| 187,917,814 | 187,944,437 | 0311B11 | D3S1037i |
| 187,917,814 | 187,944,437 | 0306B12 | D3S1093i |
| 188,131,210 | 188,279,035 | 128G07 | AC007488.15_56318 |
| 188,131,210 | 188,279,035 | 287H05 | AC034190.2_77059 |
| 188,131,210 | 188,279,035 | 0311C01 | D3S1030i |
| 188,131,210 | 188,279,035 | 0305D10 | D3S1397i |
| 188,418,632 | 188,492,446 | 0316D10 | D3S3600 |
| 188,568,862 | 188,572,066 | 0306E10 | D3S0421i |
| 188,568,862 | 188,572,066 | 706B03 | DID22N_0041504 |
| 188,869,388 | 188,870,895 | 0311B08 | D3S1021i |
| 188,869,388 | 188,870,895 | 0315H11 | D3S1293i |
| 188,869,388 | 188,870,895 | 634E03 | DISO7_10001334 |
| 190,831,910 | 191,097,759 | 389G03 | AC063939.9_142349 |
| 190,831,910 | 191,097,759 | 0314C07 | D3S0402i |
| 190,831,910 | 191,097,759 | 0310H11 | D3S1101i |
| 191,714,585 | 191,858,537 | 0314A03 | D3S0024i |
| 191,714,585 | 191,858,537 | 0303B11 | D3S0027i |
| 191,714,585 | 191,858,537 | 0302F01 | D3S1404i |
| 193,342,413 | 193,928,066 | 389H04 | AC026671.12_22466 |
| 193,342,413 | 193,928,066 | 389F04 | AC027042.2_47409 |
| 193,342,413 | 193,928,066 | 0314H08 | D3S0146i |
| 193,342,413 | 193,928,066 | 0305C09 | D3S0409i |
| 193,342,413 | 193,928,066 | 0309D03 | D3S0601i |
| 193,342,413 | 193,928,066 | 0309F04 | D3S0626i |
| 193,342,413 | 193,928,066 | 0311F11 | D3S1036i |
| 193,342,413 | 193,928,066 | 0311F12 | D3S1038i |
| 193,342,413 | 193,928,066 | 076H08 | D3S2418 |
| 193,342,413 | 193,928,066 | 015B03 | D3S3663 |
| 193,342,413 | 193,928,066 | 0316F03 | D3S3669 |
| 195,595,348 | 195,601,523 | 0311G12 | D3S1039i |
| 197,260,553 | 197,293,343 | 0305G02 | D3S0165i |
| 197,951,312 | 198,043,756 | 0313E04 | D3S0161i |
| 197,951,312 | 198,043,756 | 0310C11 | D3S0163i |
| 198,214,553 | 198,241,043 | 0313F11 | D3S0158i |
| 198,214,553 | 198,241,043 | 0302H08 | D3S1464i |
| 213,650 | 426,098 | $0311 \mathrm{E11}$ | D3S0597i |
| 213,650 | 426,098 | 0317C07 | D3S1067i |


| 213,650 | 426,098 | 0307C01 | D3S1206i |
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| 23,908,576 | 23,933,541 | 0310F04 | D3S0837i |
| 23,961,810 | 23,996,241 | 0311F04 | D3S0472i |
| 24,134,709 | 24,511,317 | 357H09 | AC069214.5_236353 |
| 24,134,709 | 24,511,317 | 0307D10 | D3S0003i |
| 24,134,709 | 24,511,317 | 0312E08 | D3S0048i |
| 24,134,709 | 24,511,317 | 0314E04 | D3S0223i |
| 24,134,709 | 24,511,317 | 0307C08 | D3S1469i |
| 25,190,893 | 25,614,424 | 0313B08 | D3S0234i |
| 25,190,893 | 25,614,424 | 0317G01 | D3S0238i |
| 25,190,893 | 25,614,424 | 0308A10 | D3S0479i |
| 25,190,893 | 25,614,424 | 0308D10 | D3S0486i |
| 25,190,893 | 25,614,424 | 0301C01 | D3S1583 |
| 27,732,872 | 27,738,807 | 0315D06 | D3S1439i |
| 3,086,421 | 3,127,031 | 0307F12 | D3S0205i |
| 3,086,421 | 3,127,031 | 0310C06 | D3S0213i |
| 30,622,998 | 30,710,638 | 0314B04 | D3S0204i |
| 30,622,998 | 30,710,638 | 0314E07 | D3S0228i |
| 31,549,495 | 31,652,560 | 0308A12 | D3S0518i |
| 32,255,175 | 32,386,817 | $0314 \mathrm{B07}$ | D3S0752i |
| 32,255,175 | 32,386,817 | 0305F08 | D3S1150i |
| 32,255,175 | 32,386,817 | 0301B04 | D3S1531i |
| 32,408,167 | 32,471,337 | 384E06 | AC026763.10_85474 |
| 32,408,167 | 32,471,337 | 384A06 | AC046140.9_14929 |
| 32,497,808 | 32,519,869 | 393 A 07 | AC010742.3_125628 |
| 32,497,808 | 32,519,869 | 384C06 | AC046140.9_123741 |
| 32,968,070 | 32,972,840 | 0315B10 | D3S1457i |
| 33,512,741 | 33,734,852 | 0313B04 | D3S0254i |
| 33,814,561 | 33,886,198 | 282A02 | AC078780.1_24631 |
| 33,814,561 | 33,886,198 | 084G01 | Z66622 |
| 37,468,817 | 37,836,285 | 0315F10 | D3S0073i |
| 37,468,817 | 37,836,285 | 0304A01 | D3S0246i |
| 37,468,817 | 37,836,285 | T001F01 | D3S0438i |
| 37,468,817 | 37,836,285 | 0315C04 | D3S0453i |
| 37,468,817 | 37,836,285 | 0315H04 | D3S0776i |
| 37,468,817 | 37,836,285 | 0313D07 | D3S1400i |
| 37,468,817 | 37,836,285 | 0303H02 | D3S3623 |
| 37,878,129 | 38,000,964 | 0317G08 | D3S1123i |
| 37,878,129 | 38,000,964 | 9902F12 | D3S1349i |
| 38155009 | 38159517 | MyD881 | new design |
| 38155009 | 38159517 | MyD882 | new design |
| 39,279,989 | 39,298,190 | 0303H03 | D3S0070i |
| 39,279,989 | 39,298,190 | 0306F01 | D3S3527 |
| 39,279,989 | 39,298,190 | 0303F12 | D3S3593 |
| 39,346,219 | 39,351,077 | 0303H11 | D3S0063i |
| 4,510,136 | 4,864,081 | 0306E09 | D3S0180i |
| 4,510,136 | 4,864,081 | 0311G03 | D3S0214i |
| 4,510,136 | 4,864,081 | 0309D11 | D3S0846i |
| 4,510,136 | 4,864,081 | 0303A02 | D3S1122i |
| 4,510,136 | 4,864,081 | 0304F04 | D3S1470i |
| 4,510,136 | 4,864,081 | 0304H11 | Z67067 |
| 41,216,004 | 41,256,938 | 0301C04 | D3S1526i |
| 41,216,004 | 41,256,938 | 0303G05 | HUMUT2543 |
| 42,617,151 | 42,665,237 | 0302C09 | D3S3687 |
| 42,825,980 | 42,883,779 | 0312F12 | D3S1081i |
| 42,888,688 | 42,892,637 | 0314E08 | D3S0061i |


| 45,098,773 | 45,162,918 | 0307F10 | D3S0012i |
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| 45,098,773 | 45,162,918 | 0312D07 | D3S0065i |
| 45,098,773 | 45,162,918 | 9903A01 | D3S1392i |
| 45,240,962 | 45,242,758 | 0310D07 | D3S1151i |
| 45,903,023 | 45,919,671 | 0313A06 | D3S0445i |
| 46,037,295 | 46,043,983 | 0312E10 | D3S0869i |
| 46,218,204 | 46,224,836 | 0303B09 | Z67705 |
| 46,227,186 | 46,283,166 | 0313C02 | D3S0004i |
| 46,423,725 | 46,426,018 | 0307E10 | D3S0010i |
| 46,452,500 | 46,481,657 | 0309F03 | D3S0605i |
| 46,894,240 | 46,926,585 | 0313G02 | D3S0884i |
| 46,894,240 | 46,926,585 | 0306E07 | G08276 |
| 47,397,528 | 47,429,935 | 0316D11 | D3S0236i |
| 47,819,625 | 47,866,687 | 0317F02 | D3S0840i |
| 48,420,266 | 48,446,464 | 0301A03 | D3S1313i |
| 49,369,613 | 49,370,795 | 0306C07 | D3S2409 |
| 49,371,582 | 49,424,530 | 0301D04 | D3S1507i |
| 49,482,595 | 49,548,048 | 0315F03 | D3S0075i |
| 49,482,595 | 49,548,048 | 0316C09 | D3S3629 |
| 49,696,391 | 49,701,099 | 0304A11 | D3S1378i |
| 49,899,439 | 49,916,074 | 0317E01 | D3S0077i |
| 49,899,439 | 49,916,074 | 0315C03 | D3S2449 |
| 49,899,439 | 49,916,074 | 0304F07 | D3S3667 |
| 52,059,799 | 52,065,329 | 581B10 | chr3.fa.O7frz. 59298456 |
| 52,059,799 | 52,065,329 | 611B01 | chr3.fa.O7frz. 59596281 |
| 52,230,138 | 52,248,223 | 0317G04 | D3S0058i |
| 52,230,138 | 52,248,223 | 0304C07 | D3S3561 |
| 52,504,396 | 52,533,551 | 0305E10 | D3S1159i |
| 52,504,396 | 52,533,551 | 0317G12 | D3S1341i |
| 53,170,263 | 53,201,773 | 0305F02 | D3S0882i |
| 53,855,612 | 53,874,867 | 0315G12 | D3S1117i |
| 53,855,612 | 53,874,867 | 0305G12 | D3S1449i |
| 55,474,783 | 55,496,371 | 0301B09 | D3S3719 |
| 55,517,376 | 56,477,431 | 341H03 | AC021129.3_44166 |
| 55,517,376 | 56,477,431 | 0313C08 | D3S0261i |
| 55,517,376 | 56,477,431 | 0309H03 | D3S0609i |
| 55,517,376 | 56,477,431 | 0309B12 | D3S0885i |
| 55,517,376 | 56,477,431 | 0310D08 | D3S1416i |
| 55,517,376 | 56,477,431 | 0303E05 | D3S3588 |
| 55,517,376 | 56,477,431 | 0316B06 | D3S3721 |
| 55,517,376 | 56,477,431 | 0304G05 | D3S3724 |
| 55,517,376 | 56,477,431 | 076B07 | G08284 |
| 55,517,376 | 56,477,431 | 0315H10 | Z67483 |
| 57,103,316 | 57,179,374 | 0308C11 | D3S0511i |
| 57,103,316 | 57,179,374 | 0312H08 | D3S0866i |
| 57,103,316 | 57,179,374 | 0315G09 | D3S1173i |
| 61,522,285 | 62,255,613 | 0306C09 | D3S0016i |
| 61,522,285 | 62,255,613 | 0308D02 | D3S0251i |
| 61,522,285 | 62,255,613 | 0308B03 | D3S0264i |
| 61,522,285 | 62,255,613 | 0310H05 | D3S0637i |
| 61,522,285 | 62,255,613 | 0311D09 | D3S0760i |
| 61,522,285 | 62,255,613 | 0313G01 | D3S0763i |
| 61,522,285 | 62,255,613 | 0317B06 | D3S1044i |
| 61,522,285 | 62,255,613 | 0301F11 | D3S1045i |
| 69,216,780 | 69,237,929 | 0315E12 | D3S0263i |
| 69,871,323 | 70,100,177 | 611H02 | chr3.fa.O7frz.81730591 |


| 69,871,323 | 70,100,177 | 0306D04 | D3S1296 |
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| 69,871,323 | 70,100,177 | 0316G02 | D3S1366i |
| 69,871,323 | 70,100,177 | $0307 \mathrm{B07}$ | D3S1373i |
| 71,087,426 | 71,715,830 | 283C05 | AC069204.3_182407 |
| 71,087,426 | 71,715,830 | 0316G01 | D3S0080i |
| 71,087,426 | 71,715,830 | 0311H11 | D3S0095i |
| 71,087,426 | 71,715,830 | 0313A01 | D3S0098i |
| 71,087,426 | 71,715,830 | 0309G05 | D3S0645i |
| 71,087,426 | 71,715,830 | 0314G12 | D3S0651i |
| 71,087,426 | 71,715,830 | 0309D12 | D3S0887i |
| 71,087,426 | 71,715,830 | 0309G12 | D3S0892i |
| 71,087,426 | 71,715,830 | T003A02 | D3S1562 |
| 71,087,426 | 71,715,830 | 0302B09 | D3S3516 |
| 71,087,426 | 71,715,830 | 0303E09 | D3S3568 |
| 85,858,322 | 86,200,641 | 430G11 | chr3.fa.O7frz. 92701960 |
| 85,858,322 | 86,200,641 | 430H11 | chr3.fa.O7frz. 92800740 |
| 85,858,322 | 86,200,641 | 0312E03 | D3S0089i |
| 85,858,322 | 86,200,641 | 9903F01 | D3S1472i |
| 85,858,322 | 86,200,641 | 0313H07 | D3S1595 |
| 87,359,140 | 87,387,339 | 0312C11 | D3S0289i |
| 87,359,140 | 87,387,339 | 0315E08 | D3S0894i |
| 87,359,140 | 87,387,339 | 0301C12 | D3S1513i |
| 9,919,150 | 9,933,086 | 0301 E 06 | D3S0050i |
| 9,933,782 | 9,950,314 | 0311A03 | D3S0179i |
| 95,074,647 | 95,175,412 | 0305G09 | D3S1471i |
| 1,764,832 | 1,780,396 | 0415D12 | D4S1055i |
| 101,088,265 | 101,090,535 | 0411D02 | D4S1218i |
| 102,163,610 | 102,487,376 | 343G09 | AP001963.1_82726 |
| 102,163,610 | 102,487,376 | 0402G08 | D4S0206i |
| 102,163,610 | 102,487,376 | 0411B03 | D4S0343i |
| 102,163,610 | 102,487,376 | 0415F08 | D4S0796i |
| 102,163,610 | 102,487,376 | 0413D09 | D4S0802i |
| 102,163,610 | 102,487,376 | 0414H05 | D4S0966i |
| 102,930,919 | 103,214,918 | 0410C06 | D4S0099i |
| 102,930,919 | 103,214,918 | 0409E09 | D4S0514i |
| 102,930,919 | 103,214,918 | 0406D09 | D4S2961 |
| 103,641,518 | 103,757,507 | 0404D06 | D4S0013i |
| 103,641,518 | 103,757,507 | 0403C10 | D4S0107i |
| 107,456,302 | 107,489,097 | 0415A12 | D4S0535i |
| 107,456,302 | 107,489,097 | 0416C09 | D4S0886i |
| 109,072,166 | 109,094,062 | 0404F07 | D4S0088i |
| 109,072,166 | 109,094,062 | 0410A11 | D4S0516i |
| 11,009,086 | 11,040,487 | 0415G12 | D4S0064i |
| 11,009,086 | 11,040,487 | 0405G12 | D4S2949 |
| 110,881,301 | 110,942,590 | 0416E04 | D4S0738i |
| 111,053,499 | 111,152,868 | 0411B01 | D4S0023i |
| 111,053,499 | 111,152,868 | 0405E10 | D4S0035i |
| 111,053,499 | 111,152,868 | 0410F08 | D4S0091i |
| 111,053,499 | 111,152,868 | 0411F06 | D4S1217i |
| 111,616,697 | 111,702,872 | 0406F09 | D4S0877i |
| 111,616,697 | 111,702,872 | 0407B11 | D4S0961i |
| 114,190,319 | 114,524,337 | 0406H08 | D4S0048i |
| 114,190,319 | 114,524,337 | 0401E03 | D4S0096i |
| 114,190,319 | 114,524,337 | 0409A10 | D4S0524i |
| 114,190,319 | 114,524,337 | 0415H11 | D4S1250i |
| 114,593,021 | 114,902,177 | 172G04 | AC004168.2_12092 |


| 114,593,021 | 114,902,177 | 0401H01 | D4S0003i |
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| 114,593,021 | 114,902,177 | 0408F09 | D4S0066i |
| 114,593,021 | 114,902,177 | 9911E01 | D4S0112i |
| 114,593,021 | 114,902,177 | 0402B05 | D4S1611 |
| 121,200,029 | 121,207,411 | 0406A02 | D4S0548i |
| 121,200,029 | 121,207,411 | 0409B11 | D4S0559i |
| 122,808,598 | 122,837,626 | 0405C05 | D4S0358i |
| 122,957,975 | 122,964,505 | 0403B08 | D4S1246i |
| 123,592,075 | 123,597,339 | 0405G03 | D4S1051i |
| 123,753,221 | 123,761,662 | 0410A05 | D4S0753i |
| 123,967,313 | 124,038,840 | 583E05 | chr4.fa.O7frz.130586887 |
| 123,967,313 | 124,038,840 | 0406D05 | D4S0540i |
| 129,021,551 | 129,039,377 | 0416B06 | D4S0804i |
| 14,950,658 | 15,056,887 | 0408H03 | D4S1104i |
| 14,950,658 | 15,056,887 | 0402G05 | D4S2362 |
| 140,156,393 | 140,186,543 | 0415A01 | D4S0352i |
| 142,777,204 | 142,874,062 | 0411A07 | D4S0680i |
| 142,777,204 | 142,874,062 | 0414D10 | D4S0778i |
| 144,654,066 | 144,694,017 | 0416D04 | D4S0734i |
| 144,654,066 | 144,694,017 | 0404B05 | D4S1292i |
| 145,136,707 | 145,159,946 | 0401G07 | D4S1279i |
| 145,249,906 | 145,281,294 | 0404G04 | D4S1223i |
| 146,622,401 | 146,699,778 | 0403F04 | Z67626 |
| 148,646,691 | 148,647,812 | 0416A01 | - |
| 148,646,691 | 148,647,812 | 0403F02 | D4S0821i |
| 148,646,691 | 148,647,812 | 0415G09 | D4S1131i |
| 149,219,370 | 149,582,973 | 0412A09 | D4S0381i |
| 149,219,370 | 149,582,973 | 0409H03 | D4S0386i |
| 149,219,370 | 149,582,973 | 0412G03 | D4S0658i |
| 149,219,370 | 149,582,973 | 0411G08 | D4S0681i |
| 149,219,370 | 149,582,973 | 0413E04 | D4S0687i |
| 15,313,738 | 15,343,508 | 0414G06 | D4S1084i |
| 15,313,738 | 15,343,508 | 0406F04 | Z67830 |
| 15,388,999 | 15,460,167 | 0407E01 | D4S2960 |
| 15,578,955 | 15,686,664 | 129E03 | AC005598.6_64773 |
| 15,578,955 | 15,686,664 | 0402D10 | D4S0060i |
| 153,461,860 | 153,675,622 | 0408B12 | D4S0270i |
| 153,461,860 | 153,675,622 | 0401E09 | D4S0599i |
| 153,461,860 | 153,675,622 | 0401B11 | D4S0610i |
| 153,461,860 | 153,675,622 | 0406H01 | D4S1057i |
| 154,824,891 | 154,846,693 | 0408C05 | D4S1132i |
| 154,824,891 | 154,846,693 | 0402D08 | D4S1179i |
| 155,703,596 | 155,711,688 | 0416C03 | D4S0101i |
| 156,349,231 | 156,357,678 | 0404D12 | D4S0183i |
| 159,849,729 | 159,864,002 | 0415A07 | D4S2997 |
| 159,849,729 | 159,864,002 | 129F08 | D63861.1_4468 |
| 166,519,538 | 166,638,926 | 0414F01 | D4S1252i |
| 166,519,538 | 166,638,926 | 0415G01 | D4S2952 |
| 175,647,955 | 175,680,213 | 292F09 | AC009887.5_114826 |
| 175,647,955 | 175,680,213 | 292A10 | AC021528.3_90278 |
| 175,647,955 | 175,680,213 | 0411C03 | D4S0773i |
| 175,647,955 | 175,680,213 | 0414E03 | HUMUT880B |
| 177,841,685 | 177,950,889 | 0413D08 | D4S0874i |
| 185,545,909 | 185,632,697 | 0402F07 | D4S1535 |
| 185,545,909 | 185,632,697 | 0415A04 | D4S3047 |
| 185,545,909 | 185,632,697 | 695E08 | DISD22_0004409 |


| 187,227,303 | 187,243,246 | 0416E08 | D4S0875i |
| :---: | :---: | :---: | :---: |
| 187,349,668 | 187,371,606 | 0411G02 | D4S0225i |
| 187,349,668 | 187,371,606 | 0411D01 | D4S0824i |
| 2,440,605 | 2,487,382 | 0416H09 | D4S0908i |
| 2,815,382 | 2,901,587 | 0402A11 | D4S0001i |
| 24,138,185 | 24,195,282 | 0401D06 | D4S0045i |
| 24,138,185 | 24,195,282 | 0409C06 | D4S0460i |
| 24,405,153 | 24,411,562 | 0415H01 | D4S0009i |
| 24,405,153 | 24,411,562 | 0414H03 | D4S0039i |
| 24,844,751 | 24,889,811 | T003B03 | D4S2970 |
| 25,930,430 | 26,045,851 | 343F03 | AC044869.2_137286 |
| 25,930,430 | 26,045,851 | 0404G01 | D4S0451i |
| 25,930,430 | 26,045,851 | 0407F06 | Z67448 |
| 25,930,430 | 26,045,851 | 0415F02 | Z67691 |
| 38450255 | 38460984 | TLR101 | new design |
| 38450255 | 38460984 | TLR102 | new design |
| 38,504,618 | 38,507,555 | 0405F09 | D4S1050i |
| 38,965,471 | 39,044,390 | 0402H07 | D4S0279i |
| 38,965,471 | 39,044,390 | 0402H02 | D4S0805i |
| 38,965,471 | 39,044,390 | 0414B11 | D4S1181i |
| 39,874,965 | 39,922,663 | 0413G01 | D4S0288i |
| 39,874,965 | 39,922,663 | 0403H10 | D4S0292i |
| 47,762,988 | 47,831,030 | 0403B03 | D4S0441i |
| 47,762,988 | 47,831,030 | 0411H11 | D4S0449i |
| 47,762,988 | 47,831,030 | 0416B08 | D4S0864i |
| 5,067,214 | 5,072,100 | 518E04 | chr4.fa.O7frz. 5109115 |
| 5,067,214 | 5,072,100 | 0415C05 | HUMUT5936 |
| 54,790,204 | 54,859,171 | 0411 E10 | D4S0176i |
| 54,790,204 | 54,859,171 | 0402D06 | D4S1630 |
| 55,218,842 | 55,301,638 | 0413A08 | D4S0005i |
| 55,218,842 | 55,301,638 | 0405G02 | D4S428 |
| 55,218,842 | 55,301,638 | 0401C09 | G08377 |
| 55,639,401 | 55,686,519 | 0406C03 | D4S1274i |
| 68,107,041 | 68,155,206 | 0404H09 | D4S0180i |
| 68,107,041 | 68,155,206 | 0412H01 | D4S0481i |
| 68,285,688 | 68,304,399 | 0414H02 | D4S3018 |
| 68,369,189 | 68,432,311 | 0409E01 | D4S0303i |
| 68,369,189 | 68,432,311 | 0410F09 | D4S0332i |
| 69,085,497 | 69,116,840 | 0414C07 | D4S1323i |
| 69,996,782 | 70,013,293 | 0408C07 | D4S1288i |
| 70,180,783 | 70,323,496 | 0414E07 | D4S0082i |
| 70,180,783 | 70,323,496 | 0414B07 | D4S1245i |
| 70,928,761 | 70,936,836 | 0414G12 | D4S0723i |
| 70,928,761 | 70,936,836 | 694C08 | DISD22_0003744 |
| 71,740,548 | 71,751,128 | 0406F06 | D4S0330i |
| 74825139 | 74828297 | IL81 | new design |
| 74,921,277 | 74,923,341 | 0408E11 | D4S0196i |
| 75,065,660 | 75,066,541 | 0406F05 | D4S0494i |
| 75,121,170 | 75,123,354 | 9903B03 | D4S1308i |
| 75,449,724 | 75,473,341 | 0409A07 | D4S0471i |
| 75,529,717 | 75,539,590 | 0413D04 | D4S0475i |
| 75,529,717 | 75,539,590 | 0416E10 | D4S0942i |
| 75,889,001 | 75,938,853 | 0407F10 | D4S0951i |
| 75,889,001 | 75,938,853 | 0414G02 | D4S1558 |
| 77,141,523 | 77,147,665 | 0407D02 | D4S3042 |
| 77,298,918 | 77,354,059 | 0403E10 | D4S1291i |


| 77,298,918 | 77,354,059 | 0407H01 | D4S2990 |
| :---: | :---: | :---: | :---: |
| 78,651,931 | 78,752,010 | 0407H10 | D4S0955i |
| 78,651,931 | 78,752,010 | 0407A11 | D4S0959i |
| 78,651,931 | 78,752,010 | 0402A08 | D4S1165i |
| 81,406,766 | 81,431,195 | 0413C05 | D4S0175i |
| 81,406,766 | 81,431,195 | 0415C11 | D4S0937i |
| 82,171,143 | 82,193,749 | 0410A06 | D4S0187i |
| 82,171,143 | 82,193,749 | 0409E07 | D4S0484i |
| 84,432,639 | 84,475,330 | 0407H02 | D4S395 |
| 87,156,656 | 87,593,307 | 0401E04 | D4S0197i |
| 87,156,656 | 87,593,307 | 0409F09 | D4S0517i |
| 87,156,656 | 87,593,307 | 0416F12 | D4S1110i |
| 87,156,656 | 87,593,307 | 9903A03 | D4S1255i |
| 87,156,656 | 87,593,307 | 0415F10 | Z67188 |
| 87,734,909 | 87,955,326 | 0408F10 | D4S0152i |
| 87,734,909 | 87,955,326 | 0406G11 | D4S1542 |
| 87,734,909 | 87,955,326 | 0408A08 | G08379 |
| 89,115,826 | 89,123,592 | 0412B08 | D4S0314i |
| 89,115,826 | 89,123,592 | 0403E03 | D4S1158i |
| 89,230,440 | 89,299,035 | 0415F07 | D4S0898i |
| 89,518,915 | 89,583,272 | 0415A09 | D4S0900i |
| 89,518,915 | 89,583,272 | 0414D05 | D4S0901i |
| 89,518,915 | 89,583,272 | 0413 F07 | D4S1171i |
| 9,055,358 | 9,061,338 | 0411H02 | D4S0709i |
| 95,438,730 | 95,483,050 | 0407H11 | D4S0976i |
| 95,898,151 | 96,295,099 | 0407C07 | D4S0054i |
| 95,898,151 | 96,295,099 | 0413H02 | D4S0198i |
| 95,898,151 | 96,295,099 | 0409A09 | D4S0506i |
| 95,898,151 | 96,295,099 | 0405G06 | D4S0960i |
| 95,898,151 | 96,295,099 | 0406A08 | D4S0982i |
|  |  | 0514C01 | D5S1480 |
|  |  | 0504A06 | D5S647 |
| 1,306,282 | 1,348,162 | 0515A11 | D5S2005 |
| 1,306,282 | 1,348,162 | 018D04 | D5S678 |
| 10,732,343 | 10,814,344 | 0513D06 | D5S0309i |
| 10,732,343 | 10,814,344 | 0507B01 | D5S432 |
| 110,433,677 | 110,441,623 | 0503D08 | D5S0383i |
| 110,433,677 | 110,441,623 | 0509F12 | D5S0676i |
| 110,587,968 | 110,858,483 | 0506H08 | D5S0122i |
| 110,587,968 | 110,858,483 | 0512F03 | D5S0345i |
| 110,587,968 | 110,858,483 | 0509C11 | D5S0643i |
| 110,587,968 | 110,858,483 | 0511B01 | D5S0649i |
| 112,101,483 | 112,209,835 | 0502C01 | D5S0048i |
| 112,101,483 | 112,209,835 | 0509G12 | D5S0677i |
| 114,942,247 | 114,989,610 | 0514F09 | D5S0658i |
| 114,942,247 | 114,989,610 | 0510F07 | D5S0873i |
| 115,168,329 | 115,180,304 | 0508B12 | D5S0358i |
| 122,386,977 | 122,400,324 | 9903E11 | D5S1319i |
| 122,386,977 | 122,400,324 | 0502E08 | Z67521 |
| 131,424,121 | 131,426,796 | 0501A03 | D5S1174i |
| 131,658,035 | 131,707,798 | 0509C01 | D5S0386i |
| 131,658,035 | 131,707,798 | 0515E01 | D5S1984 |
| 131,658,035 | 131,707,798 | 661B02 | DISO7_10004993 |
| 131,733,343 | 131,759,205 | 0512B12 | D5S0822i |
| 131,845,200 | 131,854,389 | 0511 E 05 | D5S0707i |
| 131,920,529 | 132,007,651 | 0513B07 | D5S0868i |


| 132,235,913 | 132,238,637 | 0507F07 | D5S0004i |
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| 132,235,913 | 132,238,637 | 0507C11 | D5S1011i |
| 132,415,561 | 132,468,608 | 0501B03 | D5S1016i |
| 134,122,360 | 134,194,710 | 0506H11 | D5S0021i |
| 134,122,360 | 134,194,710 | 0501G10 | D5S458 |
| 134,934,274 | 134,942,868 | 0501H01 | D5S0829i |
| 135,255,834 | 135,259,415 | 0508F06 | D5S0085i |
| 135,255,834 | 135,259,415 | 0513B12 | D5S0780i |
| 135,255,834 | 135,259,415 | 0504A05 | D5S816 |
| 137,503,358 | 137,542,257 | 0514E07 | D5S0182i |
| 137,503,358 | 137,542,257 | 0503B06 | D5S0421i |
| 137,503,358 | 137,542,257 | 0501F02 | D5S414 |
| 137,829,080 | 137,832,903 | 0507A12 | D5S1032i |
| 137,829,080 | 137,832,903 | 0502C06 | D5S500 |
| 139,207,444 | 139,403,063 | 0513D01 | D5S0272i |
| 139,207,444 | 139,403,063 | 0509B02 | D5S0415i |
| 139,207,444 | 139,403,063 | 0511H07 | D5S0694i |
| 139,692,612 | 139,706,359 | 0501A06 | D5S0273i |
| 139,692,612 | 139,706,359 | 0503C09 | D5S1305i |
| 139,991,501 | 139,993,439 | 0504G03 | D5S0084i |
| 140,005,142 | 140,007,424 | 0511D07 | D5S0704i |
| 140,494,984 | 140,497,888 | 0506C07 | D5S658 |
| 140,541,164 | 140,545,980 | 0504F08 | D5S0116i |
| 140,980,627 | 140,996,596 | 0506D11 | D5S1267i |
| 141,951,927 | 142,046,134 | 368A06 | AC016560.6_172920 |
| 141,951,927 | 142,046,134 | 0506 E 12 | D5S0125i |
| 142,637,689 | 142,795,270 | 0514B11 | D5S0041i |
| 142,637,689 | 142,795,270 | 0510C02 | D5S0500i |
| 143,171,919 | 143,180,477 | 0509E05 | D5S0505i |
| 143,171,919 | 143,180,477 | 0511C05 | D5S0699i |
| 148,734,023 | 148,739,031 | 347 A 08 | AC012613.5_58125 |
| 148,734,023 | 148,739,031 | 0506C06 | D5S1196i |
| 148,855,038 | 148,911,200 | 0514C02 | D5S0717i |
| 149,413,051 | 149,473,128 | 0514B07 | D5S0162i |
| 149,413,051 | 149,473,128 | 0508F09 | D5S0264i |
| 149,473,595 | 149,515,615 | 0508E08 | D5S0157i |
| 149,473,595 | 149,515,615 | 0503D05 | D5S2015 |
| 149,761,393 | 149,772,685 | 0514H09 | D5S0879i |
| 150,207,879 | 150,260,488 | 0502E10 | D5S1023i |
| 150,380,112 | 150,388,747 | 0503E02 | D5S0266i |
| 156,445,421 | 156,468,716 | 0506H05 | D5S0445i |
| 156,540,432 | 156,614,687 | 369H02 | AC009185.4_71677 |
| 156,540,432 | 156,614,687 | 0514G07 | D5S0196i |
| 158,058,006 | 158,459,347 | 254D05 | AC007200.1_36545 |
| 158,058,006 | 158,459,347 | 369D03 | AC011376.2_79673 |
| 158,058,006 | 158,459,347 | 0505E11 | D5S0002i |
| 158,058,006 | 158,459,347 | 0508B07 | D5S0097i |
| 158,058,006 | 158,459,347 | 0515G06 | D5S2038 |
| 158,058,006 | 158,459,347 | 0506F06 | D5S412 |
| 158,674,369 | 158,690,059 | 295 G 10 | AC011376.2_147332 |
| 158,674,369 | 158,690,059 | 0511A09 | D5S0722i |
| 158,674,369 | 158,690,059 | 058F03 | Z67033 |
| 159,707,339 | 159,730,207 | 0508A03 | D5S1138i |
| 159,707,339 | 159,730,207 | 0501A01 | D5S403 |
| 162,797,155 | 162,804,600 | 0507G05 | D5S2093 |
| 162,820,241 | 162,851,525 | 0513A12 | D5S0091i |


| 168,996,871 | 169,442,959 | 0514A07 | D5S0160i |
| :---: | :---: | :---: | :---: |
| 168,996,871 | 169,442,959 | 0510D07 | D5S0277i |
| 168,996,871 | 169,442,959 | 0508F11 | D5S0341i |
| 168,996,871 | 169,442,959 | 0511C06 | D5S0726i |
| 168,996,871 | 169,442,959 | 0505C10 | D5S1456 |
| 168,996,871 | 169,442,959 | 0504B01 | D5S1961 |
| 168,996,871 | 169,442,959 | 0505H02 | D5S1973 |
| 168,996,871 | 169,442,959 | 0506E04 | D5S504 |
| 169,465,495 | 169,469,305 | 0506A03 | D5S0063i |
| 169,607,666 | 169,657,400 | 0513H10 | D5S0169i |
| 169,607,666 | 169,657,400 | 0506H01 | HUMUT7277 |
| 172,127,707 | 172,130,809 | 0511E06 | D5S0736i |
| 172,504,130 | 172,523,989 | 0511G07 | D5S0747i |
| 172,504,130 | 172,523,989 | 0514C05 | D5S1088i |
| 172,504,130 | 172,523,989 | 0503D09 | D5S394 |
| 175,017,637 | 175,045,847 | 0514D06 | D5S0066i |
| 175,017,637 | 175,045,847 | 0508C08 | D5S0155i |
| 176,446,493 | 176,457,733 | 0513H04 | D5S0448i |
| 176,871,184 | 176,876,573 | 0513F09 | D5S0512i |
| 176,871,184 | 176,876,573 | 0506D09 | D5S1078i |
| 179,058,536 | 179,091,248 | 021C07 | D5S2073 |
| 179,595,388 | 179,640,218 | 0504E02 | D5S0752i |
| 179,945,812 | 180,009,172 | 348F06 | AC022095.4_12069 |
| 179,945,812 | 180,009,172 | 0513A04 | D5S0755i |
| 179,945,812 | 180,009,172 | 021D07 | D5S408 |
| 180,458,383 | 180,460,484 | 9903F04 | D5S1069i |
| 180,458,383 | 180,460,484 | 0501E06 | Z66649 |
| 324,739 | 488,225 | 672C05 | chr5.fa.O7frz. 1514538 |
| 324,739 | 488,225 | 0514G10 | D5S1178i |
| 324,739 | 488,225 | 0514A01 | D5S392 |
| 34,022,040 | 34,160,396 | 0510G04 | D5S0562i |
| 34,022,040 | 34,160,396 | 0511B11 | D5S0798i |
| 34,022,040 | 34,160,396 | 0508B01 | D5S1079i |
| 35,084,621 | 35,266,334 | 0504H10 | D5S0580i |
| 35,084,621 | 35,266,334 | 0508F01 | D5S1093i |
| 35,084,621 | 35,266,334 | 0506 E 10 | D5S1213i |
| 35,084,621 | 35,266,334 | 0503C11 | D5S493 |
| 35,892,748 | 35,915,462 | 0508H03 | D5S1173i |
| 35,892,748 | 35,915,462 | 0515F05 | D5S2025 |
| 36,187,946 | 36,219,904 | 0508H10 | D5S0326i |
| 36,187,946 | 36,219,904 | 0506H06 | D5S1268i |
| 36,187,946 | 36,219,904 | 0504F02 | Z66972 |
| 38,510,822 | 38,631,253 | 0501A09 | D5S1071i |
| 38,510,822 | 38,631,253 | 0511E02 | D5S1197i |
| 38,510,822 | 38,631,253 | 0503C05 | Z67436 |
| 39,141,114 | 39,255,432 | 0507H07 | D5S0146i |
| 39,141,114 | 39,255,432 | 0508G08 | D5S0217i |
| 39,141,114 | 39,255,432 | 0505D06 | D5S0573i |
| 39,320,061 | 39,400,412 | 0510C03 | D5S1073i |
| 40,715,789 | 40,729,594 | 0505A02 | D5S0954i |
| 40,715,789 | 40,729,594 | 0505C08 | D5S1061i |
| 40,877,043 | 40,896,025 | 0513C07 | D5S0566i |
| 40,945,356 | 41,018,798 | 0507E06 | D5S1457 |
| 41,178,093 | 41,297,297 | 597A07 | chr5.fa.O7frz.46786737 |
| 41,178,093 | 41,297,297 | 0515A07 | D5S0081i |
| 42,459,783 | 42,757,736 | 0509H07 | D5S0583i |


| 42,459,783 | 42,757,736 | 0507E08 | D5S0927i |
| :---: | :---: | :---: | :---: |
| 42,459,783 | 42,757,736 | 0505H03 | D5S1264i |
| 42,459,783 | 42,757,736 | 0506A10 | D5S1297i |
| 43,229,915 | 43,448,250 | 0514G11 | D5S0180i |
| 43,229,915 | 43,448,250 | 9903A09 | D5S0986i |
| 43,229,915 | 43,448,250 | 0514F05 | D5S1251i |
| 44,340,854 | 44,424,541 | 0513G03 | D5S0483i |
| 44,340,854 | 44,424,541 | 0502D10 | D5S2063 |
| 52,119,531 | 52,285,242 | 0515H02 | D5S1202i |
| 52,119,531 | 52,285,242 | 0503F10 | D5S623 |
| 52,119,531 | 52,285,242 | 667F02 | DISO7_10995141 |
| 52,321,014 | 52,423,947 | 0501E09 | D5S1184i |
| 52,321,014 | 52,423,947 | 0505F03 | D5S1239i |
| 54,355,838 | 54,366,155 | 0501H05 | D5S1191i |
| 54,434,230 | 54,441,837 | 667B03 | DISO7_11025103 |
| 54,587,830 | 54,639,278 | 0508A01 | D5S1056i |
| 55,069,609 | 55,148,362 | 0508H08 | D5S0225i |
| 55,069,609 | 55,148,362 | 0513H06 | D5S0578i |
| 55,069,609 | 55,148,362 | 0503F03 | D5S664 |
| 55,183,091 | 55,248,922 | 0504A01 | D5S645 |
| 55,266,680 | 55,326,529 | 0502A11 | D5S1340i |
| 56,146,022 | 56,227,736 | 0509D08 | D5S0591i |
| 56,146,022 | 56,227,736 | 0505A09 | D5S1237i |
| 58,302,468 | 59,320,301 | 293D08 | AC026095.3_41930 |
| 58,302,468 | 59,320,301 | 526B08 | chr5.fa.O7frz. 64839304 |
| 58,302,468 | 59,320,301 | 0508B05 | D5S0012i |
| 58,302,468 | 59,320,301 | 0508E06 | D5S0079i |
| 58,302,468 | 59,320,301 | 0513E01 | D5S0585i |
| 58,302,468 | 59,320,301 | 0510F04 | D5S0785i |
| 58,302,468 | 59,320,301 | 0511F12 | D5S0843i |
| 58,302,468 | 59,320,301 | 019A06 | D5S2000 |
| 58,302,468 | 59,320,301 | 0502E03 | D5S2080 |
| 58,302,468 | 59,320,301 | 0515B04 | D5S2091 |
| 58,302,468 | 59,320,301 | 0501F04 | D5S431 |
| 58,302,468 | 59,320,301 | 0507B02 | D5S468 |
| 58,302,468 | 59,320,301 | 0504E06 | HUMUT2092 |
| 58,302,468 | 59,320,301 | 0504G04 | HUMUT5438 |
| 66,513,872 | 66,528,368 | 0512C10 | D5S0188i |
| 66,513,872 | 66,528,368 | 0508D11 | D5S0336i |
| 66,513,872 | 66,528,368 | 0503F08 | D5S1210i |
| 67,547,360 | 67,633,405 | 0512B10 | D5S0235i |
| 67,547,360 | 67,633,405 | 0509E08 | D5S0593i |
| 68,498,593 | 68,509,828 | 367 A02 | AC010273.3_156905 |
| 68,498,593 | 68,509,828 | 294A03 | AC022107.5_95947 |
| 68,498,593 | 68,509,828 | 0501E07 | HUMUT1151 |
| 68,682,567 | 68,701,596 | 9903G08 | D5S0934i |
| 68,682,567 | 68,701,596 | 0514E05 | D5S1177i |
| 7,922,217 | 7,954,237 | 0507B09 | D5S0943i |
| 7,922,217 | 7,954,237 | 0514B05 | D5S0949i |
| 72,777,839 | 72,780,108 | 0509H04 | D5S0493i |
| 72,777,839 | 72,780,108 | 0504F06 | D5S1140i |
| 73,958,990 | 73,973,005 | 0514D09 | D5S0610i |
| 73,958,990 | 73,973,005 | 0509A10 | D5S0618i |
| 73,958,990 | 73,973,005 | 0505G07 | D5S0623i |
| 74,668,790 | 74,693,685 | 0513C04 | D5S0144i |
| 74,668,790 | 74,693,685 | 0510E04 | D5S0148i |


| 74,702,684 | 74,843,719 | 0501G04 | D5S0840i |
| :---: | :---: | :---: | :---: |
| 75,947,063 | 75,954,996 | 0508D08 | D5S0156i |
| 75,947,063 | 75,954,996 | 0512B06 | D5S0338i |
| 76,047,542 | 76,067,054 | 0503F05 | D5S1284i |
| 76,150,610 | 76,166,896 | 0507E03 | D5S424 |
| 77,816,794 | 77,841,979 | 0508F07 | D5S0124i |
| 77,816,794 | 77,841,979 | 0509D10 | D5S0629i |
| 79,957,801 | 79,986,556 | 0510G12 | D5S1057i |
| 79,957,801 | 79,986,556 | 0502C07 | D5S1120i |
| 82,803,339 | 82,912,737 | 0513F03 | D5S0194i |
| 82,803,339 | 82,912,737 | 9903D06 | D5S0778i |
| 82,803,339 | 82,912,737 | 9903C05 | Z67581 |
| 86,599,461 | 86,723,489 | $0511 \mathrm{E11}$ | D5S0615i |
| 86,599,461 | 86,723,489 | 0513G02 | D5S1205i |
| 86,599,461 | 86,723,489 | 0515D09 | D5S1248i |
| 92,944,799 | 92,956,077 | 0508G11 | D5S0342i |
| 96,122,277 | 96,169,559 | 0514G12 | D5S0053i |
| 96,122,277 | 96,169,559 | 0511F01 | D5S0633i |
| 96,122,277 | 96,169,559 | 0509H10 | D5S0638i |
|  |  | 0606D01 | D6S426 |
|  |  | 0606F12 | D6S461 |
| 1,257,675 | 1,259,983 | 9904D01 | D6S0367i |
| 1,257,675 | 1,259,983 | 0609F06 | D6S0630i |
| 1,335,068 | 1,340,831 | 0605B06 | Z67528 |
| 1,555,206 | 1,559,131 | 0603H06 | D6S0303i |
| 1,569,040 | 2,190,845 | 0614H01 | D6S0280i |
| 1,569,040 | 2,190,845 | 0611G01 | D6S0641i |
| 1,569,040 | 2,190,845 | 0609 A07 | D6S0645i |
| 1,569,040 | 2,190,845 | 0613H03 | D6S0649i |
| 10,636,575 | 10,737,587 | 0605F02 | D6S0656i |
| 10,636,575 | 10,737,587 | 0605H07 | D6S1025i |
| 100,939,606 | 101,019,494 | 0602F10 | D6S0792i |
| 100,939,606 | 101,019,494 | 0612F12 | D6S0996i |
| 101,953,385 | 102,624,651 | 0602B03 | D6S1642 |
| 105,282,661 | 105,414,867 | 0609F11 | D6S0772i |
| 105,282,661 | 105,414,867 | 0605D02 | D6S1254i |
| 106,640,888 | 106,664,507 | 0614C09 | D6S1049i |
| 106,640,888 | 106,664,507 | 0606G07 | D6S1241i |
| 106,739,044 | 106,880,388 | 0612B08 | D6S0031i |
| 106,739,044 | 106,880,388 | 0606G05 | D6S0779i |
| 107,918,010 | 108,089,195 | 0608B04 | D6S0085i |
| 107,918,010 | 108,089,195 | 0613B04 | D6S0999i |
| 108,593,955 | 108,616,706 | 0613D02 | D6S0043i |
| 108,593,955 | 108,616,706 | 023G09 | D6S1594 |
| 108,593,955 | 108,616,706 | 0604A01 | Z67377 |
| 108,987,719 | 109,108,661 | 0607F07 | D6S0497i |
| 108,987,719 | 109,108,661 | 0605D07 | D6S1167i |
| 109,794,412 | 109,810,353 | 0607H09 | D6S1050i |
| 109,820,624 | 109,868,524 | 0604H03 | D6S0905i |
| 11,291,517 | 11,490,535 | 531B10 | chr6.fa.O7frz. 12034704 |
| 11,291,517 | 11,490,535 | 0608C04 | D6S0089i |
| 11,291,517 | 11,490,535 | 0603E10 | D6S0103i |
| 11,291,517 | 11,490,535 | 0610F08 | D6S0283i |
| 11,291,517 | 11,490,535 | 0608F12 | D6S0386i |
| 110,527,715 | 110,607,900 | 0609G05 | D6S0605i |
| 110,527,715 | 110,607,900 | 0602E08 | HUMUT893 |


| 111,409,984 | 111,453,487 | 0603 A07 | D6S1698 |
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| 112,088,228 | 112,301,348 | 0612G11 | D6S0766i |
| 112,088,228 | 112,301,348 | 0601A09 | D6S0768i |
| 112,088,228 | 112,301,348 | 023D10 | D6S302 |
| 112,088,228 | 112,301,348 | 175G07 | Z97989.1_71410 |
| 112,536,654 | 112,682,605 | 0608B06 | D6S0172i |
| 112,536,654 | 112,682,605 | 0612E09 | D6S0394i |
| 112,536,654 | 112,682,605 | 0612E02 | D6S0775i |
| 114,368,571 | 114,399,029 | 0609G11 | D6S0776i |
| 114,368,571 | 114,399,029 | 0601B03 | D6S401 |
| 116,369,386 | 116,488,614 | 0605E06 | D6S0428i |
| 116,369,386 | 116,488,614 | 0602F07 | D6S0878i |
| 117,305,068 | 117,360,008 | 0606A01 | D6S1206i |
| 118,892,932 | 119,137,924 | 0614G08 | D6S0282i |
| 118,892,932 | 119,137,924 | 0602D01 | D6S0548i |
| 118,892,932 | 119,137,924 | 0612B04 | D6S0784i |
| 118,892,932 | 119,137,924 | 0601H06 | D6S0790i |
| 119,540,965 | 119,712,625 | 0607G06 | D6S0273i |
| 119,540,965 | 119,712,625 | 0609G02 | D6S0461i |
| 12,398,582 | 12,405,413 | 0602E02 | D6S0100i |
| 12,398,582 | 12,405,413 | 0614F03 | D6S0144i |
| 12,825,819 | 13,396,624 | 0601D05 | D6S0371i |
| 124,166,768 | 125,188,502 | 0602D11 | D6S0559i |
| 126,144,000 | 126,293,950 | 0612D02 | D6S0930i |
| 127,813,023 | 127,879,540 | 0613G06 | D6S0573i |
| 128,331,625 | 128,883,453 | 0613F06 | D6S0077i |
| 128,331,625 | 128,883,453 | 0613E04 | D6S0270i |
| 128,331,625 | 128,883,453 | 0603D01 | D6S0536i |
| 128,331,625 | 128,883,453 | 0605F03 | D6S0822i |
| 128,331,625 | 128,883,453 | 0607A05 | D6S1030 |
| 128,331,625 | 128,883,453 | 0604D06 | D6S1033 |
| 129,246,035 | 129,879,407 | 0612B06 | D6S0589i |
| 129,246,035 | 129,879,407 | 0604B10 | D6S0798i |
| 129,246,035 | 129,879,407 | 0611A06 | D6S0833i |
| 129,246,035 | 129,879,407 | 0601H03 | D6S1047i |
| 129,246,035 | 129,879,407 | 0601E03 | D6S1075i |
| 129,246,035 | 129,879,407 | 0605C03 | D6S1226i |
| 129,246,035 | 129,879,407 | 0602G07 | D6S1244i |
| 131,202,180 | 131,426,017 | 0607B08 | D6S0834i |
| 131,202,180 | 131,426,017 | 0607A04 | D6S1572 |
| 131,508,154 | 131,646,366 | 0612E11 | D6S0068i |
| 131,508,154 | 131,646,366 | 0609D12 | D6S0799i |
| 131,935,977 | 131,947,165 | 9903 D 12 | D6S1622i |
| 131,935,977 | 131,947,165 | 024D02 | D6S457 |
| 131,935,977 | 131,947,165 | 0605F01 | HUMUT5174B |
| 132,000,135 | 132,110,243 | 207D10 | AC005587.1_83579 |
| 132,000,135 | 132,110,243 | 183 F 08 | AL135904.11_38895 |
| 132,310,199 | 132,314,206 | 0612F06 | D6S0028i |
| 132,310,199 | 132,314,206 | 0614D03 | D6S0041i |
| 132,658,887 | 132,764,357 | 0608A09 | D6S0271i |
| 132,658,887 | 132,764,357 | 0614H07 | D6S0998i |
| 133,044,422 | 133,076,881 | 0605C07 | D6S0407i |
| 133,044,422 | 133,076,881 | 0613C10 | D6S1171i |
| 134,532,081 | 134,680,889 | 0614C07 | D6S0807i |
| 134,532,081 | 134,680,889 | 0601G06 | D6S270 |
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| 136,705,565 | 136,913,485 | 0611B10 | D6S0835i |
| :---: | :---: | :---: | :---: |
| 136,919,878 | 137,155,349 | 0604A08 | D6S0422i |
| 136,919,878 | 137,155,349 | T001A12 | D6S0478i |
| 136,919,878 | 137,155,349 | 0602E11 | D6S0800i |
| 136,919,878 | 137,155,349 | 0613B07 | D6S1003i |
| 137,362,801 | 137,407,991 | 0613 F 11 | D6S0808i |
| 137,362,801 | 137,407,991 | 0606A06 | D6S1009 |
| 137,506,650 | 137,536,478 | 0611G10 | D6S0813i |
| 138,230,274 | 138,246,142 | 0610B03 | D6S0907i |
| 138,230,274 | 138,246,142 | 0611A05 | D6S0915i |
| 139,158,950 | 139,266,900 | 0609B05 | D6S0590i |
| 139,158,950 | 139,266,900 | 0605F05 | D6S0801i |
| 139,497,942 | 139,543,639 | 0601C06 | D6S0561i |
| 139,497,942 | 139,543,639 | 0604A12 | D6S0564i |
| 139,735,089 | 139,737,478 | 0604A05 | D6S1109i |
| 14,225,715 | 14,245,128 | 0602A09 | D6S0879i |
| 14,225,715 | 14,245,128 | 0614A02 | D6S429 |
| 143,114,297 | 143,308,031 | 0608G07 | D6S0229i |
| 143,114,297 | 143,308,031 | 0603H11 | D6S0337i |
| 143,114,297 | 143,308,031 | 094F12 | HUMUT525 |
| 143,114,297 | 143,308,031 | 087H02 | HUMUT7700 |
| 143,788,765 | 143,813,517 | 0605H11 | D6S0094i |
| 143,788,765 | 143,813,517 | 0610G11 | D6S0560i |
| 143,971,010 | 144,194,014 | 0608F07 | D6S0228i |
| 143,971,010 | 144,194,014 | 0603B06 | D6S0552i |
| 143,971,010 | 144,194,014 | 0614A01 | D6S1704 |
| 144,513,356 | 144,551,200 | 0613D08 | D6S0830i |
| 144,513,356 | 144,551,200 | 0614G07 | D6S0987i |
| 144,513,356 | 144,551,200 | 0601 E 10 | D6S1003 |
| 144,654,566 | 145,215,863 | 0603G05 | D6S0836i |
| 147,566,565 | 147,748,588 | 0609D06 | D6S0626i |
| 147,566,565 | 147,748,588 | 0609E06 | D6S0627i |
| 147,566,565 | 147,748,588 | 0607G07 | D6S0628i |
| 148,313 | 151,392 | 0603H10 | D6S1086i |
| 148,313 | 151,392 | 0610A01 | D6S1139i |
| 149,680,756 | 149,774,442 | 9903A12 | - |
| 149,680,756 | 149,774,442 | $187 \mathrm{B08}$ | AL031056.1_26387 |
| 149,680,756 | 149,774,442 | 183B02 | AL031056.1_52627 |
| 149,680,756 | 149,774,442 | 0611H09 | D6S0203i |
| 149,680,756 | 149,774,442 | 0606G03 | D6S0241i |
| 149,680,756 | 149,774,442 | 024D06 | D6S1553 |
| 149,867,324 | 149,908,864 | 9904A01 | D6S0831i |
| 150,112,273 | 150,174,249 | 0610B02 | D6S1117i |
| 150,251,294 | 150,253,863 | 0607G05 | HUMUT5779 |
| 150,304,829 | 150,312,064 | 0601A08 | D6S0841i |
| 151,603,202 | 151,719,602 | 0612E04 | D6S0933i |
| 151,603,202 | 151,719,602 | 0601H08 | D6S476 |
| 152,170,379 | 152,466,099 | 0603C04 | D6S0027i |
| 152,170,379 | 152,466,099 | 0609G01 | D6S0433i |
| 152,170,379 | 152,466,099 | 0609A03 | D6S0468i |
| 152,170,379 | 152,466,099 | 0610C09 | D6S0846i |
| 152,484,515 | 153,000,227 | 0610C11 | D6S0117i |
| 152,484,515 | 153,000,227 | 0605G05 | D6S0226i |
| 152,484,515 | 153,000,227 | 0611 E12 | D6S0853i |
| 152,484,515 | 153,000,227 | 0611A12 | D6S0935i |
| 153,113,626 | 153,122,593 | 0613B10 | D6S0593i |


| 154,402,136 | 154,609,693 | 0611H10 | D6S0346i |
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| 155,620,488 | 155,686,932 | 0601B01 | D6S0937i |
| 155,620,488 | 155,686,932 | 0604A11 | D6S1162i |
| 155,758,194 | 155,818,729 | 183G06 | AL133474.9_33793 |
| 155,758,194 | 155,818,729 | 176F12 | AL133474.9_63821 |
| 155,758,194 | 155,818,729 | 0609D02 | D6S0453i |
| 157,140,756 | 157,572,094 | 0605A09 | D6S0470i |
| 157,140,756 | 157,572,094 | 0610A03 | D6S0838i |
| 158,164,282 | 158,286,097 | 0613F01 | D6S0044i |
| 158,164,282 | 158,286,097 | 0606C02 | D6S0582i |
| 158,509,372 | 158,535,008 | 586F01 | chr6.fa.O7frz. 168445622 |
| 158,509,372 | 158,535,008 | 537F05 | chr6.fa.O7frz. 168477401 |
| 158,509,372 | 158,535,008 | 0611C07 | D6S0863i |
| 16,407,322 | 16,869,700 | 0609D04 | D6S0516i |
| 160,020,138 | 160,034,343 | 0611E08 | D6S0970i |
| 160,066,607 | 160,097,341 | 0601G02 | D6S1581 |
| 160,310,121 | 160,447,573 | 0606F04 | D6S0005i |
| 160,310,121 | 160,447,573 | 0601E08 | D6S0260i |
| 160,310,121 | 160,447,573 | 0605C10 | D6S0279i |
| 160,462,853 | 160,499,740 | 0604G11 | D6S0445i |
| 160,462,853 | 160,499,740 | 0602B01 | D6S0553i |
| 160,462,853 | 160,499,740 | 0614D07 | D6S0854i |
| 161,471,047 | 161,615,097 | 0608C10 | D6S0324i |
| 161,471,047 | 161,615,097 | 0606E02 | D6S0330i |
| 161,471,047 | 161,615,097 | 0602G08 | D6S0351i |
| 161,688,442 | 163,068,793 | T001G10 | D6S1011i |
| 163,068,154 | 163,656,514 | $0611 \mathrm{G12}$ | D6S0934i |
| 165,660,766 | 165,995,578 | 0602G06 | D6S0459i |
| 166,742,844 | 167,195,761 | 0601C05 | D6S1585 |
| 167,332,660 | 167,473,174 | 0613F09 | D6S0086i |
| 167,332,660 | 167,473,174 | 0606B11 | D6S0566i |
| 167,332,660 | 167,473,174 | 0612F07 | D6S0859i |
| 167,928,066 | 167,940,388 | 0608H03 | D6S0075i |
| 167,928,066 | 167,940,388 | 0606E10 | D6S0218i |
| 170,457,769 | 170,556,162 | 0608G04 | D6S0105i |
| 170,457,769 | 170,556,162 | 0608G10 | D6S0331i |
| 170,457,769 | 170,556,162 | $0601 \mathrm{E11}$ | D6S1590 |
| 170,686,134 | 170,704,312 | 0606D10 | D6S0204i |
| 170,686,134 | 170,704,312 | 0612B01 | D6S0249i |
| 18,236,521 | 18,263,353 | 0613H08 | D6S0037i |
| 2,710,665 | 2,731,926 | 0607F09 | D6S1039i |
| 2,832,499 | 2,848,513 | 0612B11 | D6S0116i |
| 20,642,667 | 21,340,614 | 0603B10 | D6S0153i |
| 22,395,459 | 22,405,709 | 0602C09 | D6S0373i |
| 22,395,459 | 22,405,709 | 0607C05 | Z67232 |
| 22,677,657 | 22,679,871 | 0605B12 | D6S0136i |
| 22,677,657 | 22,679,871 | 0611F05 | D6S0958i |
| 237,053 | 296,355 | 0611C05 | D6S0887i |
| 237,053 | 296,355 | 0601E05 | D6S1090i |
| 24,758,184 | 24,775,240 | 135B03 | AL031230.1_102019 |
| 24,758,184 | 24,775,240 | 0601B07 | D6S0676i |
| 25,387,285 | 25,728,737 | 0606D09 | G08569 |
| 26,195,427 | 26,205,038 | T002G03 | D6S0284i |
| 26,510,460 | 26,523,445 | 0607H08 | D6S1026i |
| 26,510,460 | 26,523,445 | 0602G12 | D6S1248i |
| 27,323,487 | 27,332,327 | 0603C01 | D6S0243i |


| 27,323,487 | 27,332,327 | 0613A05 | D6S1155i |
| :---: | :---: | :---: | :---: |
| 28,217,695 | 28,233,215 | 0611G03 | D6S0111i |
| 28,217,695 | 28,233,215 | 0610F02 | D6S0660i |
| 28,301,046 | 28,309,239 | 0606C06 | D6S0147i |
| 29,631,368 | 29,778,041 | 181A05 | AC006162.1_38368 |
| 29,631,368 | 29,778,041 | 186A07 | AL050328.24_38345 |
| 29,732,755 | 29,748,128 | 174B01 | AL050328.24_40038 |
| 29,798,531 | 29,803,052 | 180G06 | AF055066.1_308034 |
| 29799096 | 29803052 | D6S2770 | D6S2770 |
| 29799096 | 29803052 | D6S2872 | D6S2872 |
| 29799096 | 29803052 | D6S2910 | D6S2910 |
| 29799096 | 29803052 | D6S2911 | D6S2911 |
| 29,802,425 | 29,802,895 | $613 \mathrm{G10}$ | chr6.fa.O7frz. 33382117 |
| 29,902,723 | 30,021,633 | 0610D04 | D6S0668i |
| 3,009,212 | 3,060,420 | 0607A11 | D6S1088i |
| 30018310 | 30021633 | D6S2704 | D6S2704 |
| 30018310 | 30021633 | D6S2707 | D6S2707 |
| 30018310 | 30021633 | D6S2838 | D6S2838 |
| 30018310 | 30021633 | D6S2847 | D6S2847 |
| 30018310 | 30021633 | 697B02 | DISD22_0011489 |
| 30018310 | 30021633 | 022G02 | - |
| 30018310 | 30021633 | 186B05 | - |
| 30,030,962 | 30,031,390 | 0607C02 | D6S265 |
| 30,107,469 | 30,109,633 | 0603F12 | D6S0509i |
| 30,212,487 | 30,224,491 | 733H11 | DIJ28_10010194 |
| 30,227,701 | 30,236,690 | 0610D05 | D6S0014i |
| 30,420,877 | 30,422,649 | 0609C04 | D6S0505i |
| 30,561,651 | 30,562,700 | 0612D01 | D6S0493i |
| 30565250 | 30569077 | C3_2_11, D6S2840 | C3_2_11, D6S2840 |
| 30565250 | 30569077 | D6S2799 | D6S2799 |
| 30,621,633 | 30,632,987 | 0609G04 | D6S0525i |
| 30,775,563 | 30,793,645 | 0605H05 | D6S1124i |
| 30888622 | 30906415 | C4_2_12, D6S2827 | C4_2_12, D6S2827 |
| 30,983,956 | 30,989,859 | 0607D12 | D6S1128i |
| 31,059,474 | 31,065,654 | 0606D05 | D6S0076i |
| 31344505 | 31432935 | C1_2_5, D6S2811 | C1_2_5, D6S2811 |
| 31344505 | 31432935 | C1_2_A, D6S2793 | C1_2_A, D6S2793 |
| 31344505 | 31432935 | C1_4_3, D6S2930 | C1_4_3, D6S2930 |
| 31344505 | 31432935 | C1_4_4, D6S2931 | C1_4_4, D6S2931 |
| 31344505 | 31432935 | D6S2792 | D6S2792 |
| 31429628 | 31432914 | C1_3_2a, D6S2902 | C1_3_2a, D6S2902 |
| 31429628 | 31432914 | C2_4_3, D6S2938 | C2_4_3, D6S2938 |
| 31429628 | 31432914 | C4_2_7, D6S2825 | C4_2_7, D6S2825 |
| 31,475,540 | 31,491,069 | 186H03 | AB031008.1_2319 |
| 31,538,938 | 31,541,565 | 0602G01 | D6S0517i |
| 31,604,718 | 31,605,987 | 0612H04 | D6S1220i |
| 31,648,042 | 31,650,080 | TNFa | Udalova |
| 31,648,042 | 31,650,080 | TNFb | Udalova |
| 31,648,042 | 31,650,080 | TNFc | Udalova |
| 31,648,042 | 31,650,080 | TNFd | Udalova |
| 31,648,042 | 31,650,080 | TNFe | Udalova |
| 31648072 | 31650077 | S2780, D6S1615, D6S2 | 6S2780, D6S1615, D6S278 |
| 31651329 | 31654091 | D6S2924 | D6S2924 |
| 31,690,984 | 31,692,781 | 022F03 | D6S273 |
| 31696429 | 31713533 | BAT2CA, D6S2787 | BAT2CA, D6S2787 |
| 31,752,440 | 31,759,796 | 186A04 | AF134726.1_179024 |
| 31,129,963 | 31,135,632 | 134E07 | - |


| 32003473 | 32021428 | D6S2740 | D6S2740 |
| :---: | :---: | :---: | :---: |
| 32003473 | 32021428 | D6S2913 | D6S2913 |
| 32084175 | 32185131 | 3-3, D6S2920 | 3-3, D6S2920 |
| 32,204,462 | 32,206,045 | 0603B07 | D6S1255i |
| 32,266,521 | 32,299,822 | $0613 \mathrm{G11}$ | D6S0267i |
| 32,266,521 | 32,299,822 | 079C09 | D6S1014 |
| 32270598 | 32299822 | D6S2894 | new design |
| 32368453 | 32460310 | 3-7, D6S2892 | 3-7, D6S2892 |
| 32,368,464 | 32,447,662 | 0609D03 | D6S0483i |
| 32,515,597 | 32,520,943 | 0610H04 | D6S0518i |
| 32515625 | 32520801 | DRA_CA, D6S2883 | DRA_CA, D6S2883 |
| 32,654,524 | 32,665,603 | 310G03 | AC026010.3_134784 |
| 32,713,112 | 32,719,407 | 022B04 | D6S1666 |
| 32735222 | 32754296 | D6S2818 | D6S2818 |
| 32735222 | 32754296 | G511525 | new design |
| 32,735,225 | 32,742,572 | 9903 E 12 | D6S0067i |
| 32,831,445 | 32,839,446 | 697A03 | DISD22_0000187 |
| 32,888,518 | 32,892,803 | 132A02 | X87344.1_184861 |
| 32,897,588 | 32,914,525 | 0602A04 | D6S1104i |
| 33,044,415 | 33,057,075 | 0614C04 | D6S0274i |
| 33,188,206 | 33,204,868 | 0602F11 | D6S0512i |
| 33238447 | 33268223 | M2_2_9, D6S2731 | M2_2_9, D6S2731 |
| 33238447 | 33268223 | M2_4_25, D6S2822 | M2_4_25, D6S2822 |
| 33238447 | 33268223 | 186C02 | - |
| 33,276,631 | 33,280,192 | 0609A04 | D6S0498i |
| 33,326,027 | 33,347,640 | 186G05 | Z97183.1_16345 |
| 33,375,449 | 33,390,142 | 0608H06 | D6S0196i |
| 33,648,307 | 33,655,997 | 133 F 12 | Z93017.6_2832 |
| 33,696,500 | 33,772,329 | 0601H07 | D6S1165i |
| 34,541,883 | 34,610,984 | 0601A01 | HUMUT2253 |
| 34,541,883 | 34,610,984 | 0607 F 05 | HUMUT6615 |
| 35,418,313 | 35,503,933 | 0613A02 | D6S0990i |
| 35,649,345 | 35,804,338 | 0614D11 | D6S1211i |
| 35,649,345 | 35,804,338 | 0603G11 | D6S1645 |
| 35,908,789 | 35,996,942 | 0608H02 | D6S0036i |
| 35,908,789 | 35,996,942 | 0601G04 | D6S0664i |
| 36,103,551 | 36,186,513 | 0602C10 | G10173 |
| 36,129,769 | 36,215,820 | 0611B04 | D6S0035i |
| 36,129,769 | 36,215,820 | T002C03 | D6S0078i |
| 36,569,647 | 36,623,234 | 0611H01 | D6S0234i |
| 36,569,647 | 36,623,234 | 0605A03 | D6S0992i |
| 36,754,413 | 36,763,094 | 0605C05 | D6S0335i |
| 36,754,413 | 36,763,094 | 0605B02 | D6S1051 |
| 36,930,581 | 36,950,778 | 0603H03 | D6S0508i |
| 36,930,581 | 36,950,778 | 9904H01 | D6S1010i |
| 37,245,957 | 37,251,182 | 0609A08 | D6S0682i |
| 37,895,285 | 38,230,375 | 0601A12 | D6S0233i |
| 37,895,285 | 38,230,375 | 0603C11 | D6S0312i |
| 37,895,285 | 38,230,375 | 0612F05 | D6S0670i |
| 38,250,711 | 38,673,848 | 0613D06 | D6S0347i |
| 38,250,711 | 38,673,848 | 0609H07 | D6S0679i |
| 38,792,313 | 39,106,545 | 0611A09 | D6S0017i |
| 38,792,313 | 39,106,545 | 0608B12 | D6S0370i |
| 38,792,313 | 39,106,545 | 0610E02 | D6S0524i |
| 38,792,313 | 39,106,545 | 0614F07 | D6S0895i |
| 39,868,120 | 39,980,622 | 0605E03 | D6S1168i |


| 39,868,120 | 39,980,622 | 0604D12 | D6S1641 |
| :---: | :---: | :---: | :---: |
| 41,102,749 | 41,114,906 | 0614E03 | D6S0047i |
| 41,102,749 | 41,114,906 | 0603A10 | D6S0302i |
| 41,224,979 | 41,230,048 | 0607D02 | D6S1575 |
| 41,284,270 | 41,298,360 | 9904A02 | D6S1012i |
| 41,411,505 | 41,426,603 | 0608G08 | D6S0263i |
| 41,622,142 | 41,678,100 | 0602D03 | D6S0379i |
| 41,622,142 | 41,678,100 | 0609B08 | D6S0683i |
| 41,622,142 | 41,678,100 | 0608H01 | D6S1672 |
| 42,010,649 | 42,124,404 | 0608C12 | D6S0372i |
| 42,010,649 | 42,124,404 | 0604H07 | D6S1552 |
| 42,300,647 | 42,527,767 | 0608A10 | D6S0320i |
| 42,300,647 | 42,527,767 | 0612H08 | D6S0671i |
| 42,300,647 | 42,527,767 | 0603C02 | D6S1237i |
| 42,300,647 | 42,527,767 | 0602A07 | HUMUT2081 |
| 43,039,586 | 43,054,936 | 0601G12 | D6S0294i |
| 43,039,586 | 43,054,936 | 0606H05 | D6S1209i |
| 43,152,007 | 43,237,435 | 0605D03 | D6S1582 |
| 43,845,924 | 43,862,202 | 0609E07 | D6S0662i |
| 43,845,924 | 43,862,202 | 0610H01 | D6S1031i |
| 44,322,802 | 44,329,598 | 0607A09 | D6S1027i |
| 44,322,802 | 44,329,598 | 0607 A 02 | D6S1650 |
| 44,333,881 | 44,341,503 | 0607G08 | D6S1024i |
| 45,404,032 | 45,626,797 | 0611D06 | D6S0918i |
| 46,625,404 | 46,728,482 | 0608C03 | D6S0060i |
| 46,625,404 | 46,728,482 | 0603D10 | D6S1541 |
| 46,625,404 | 46,728,482 | 0610D01 | D6S1638 |
| 46,779,897 | 46,811,389 | 0604B11 | D6S0691i |
| 47,307,227 | 47,385,639 | 0605G11 | D6S0290i |
| 47,307,227 | 47,385,639 | 0604G10 | D6S0489i |
| 47,307,227 | 47,385,639 | 0604H02 | Z66926 |
| 47,553,899 | 47,702,620 | 0611G07 | D6S0874i |
| 47,553,899 | 47,702,620 | 0604E03 | D6S0978i |
| 47,953,998 | 48,144,384 | 0609C08 | D6S0686i |
| 47,953,998 | 48,144,384 | 0610E07 | D6S0715i |
| 47,953,998 | 48,144,384 | 0604B07 | HUMUT596 |
| 49,680,830 | 49,712,511 | 0613H10 | D6S0299i |
| 49,680,830 | 49,712,511 | 0613F02 | HUMUT6326 |
| 50,035,964 | 50,039,777 | 0606D11 | D6S0598i |
| 50,084,810 | 50,097,607 | 0605D01 | D6S0602i |
| 51,588,104 | 52,060,382 | 0608F10 | D6S0327i |
| 51,588,104 | 52,060,382 | 0603C07 | D6S0487i |
| 51,588,104 | 52,060,382 | 0611H05 | D6S0919i |
| 51,588,104 | 52,060,382 | 0605H09 | HUMUT7510 |
| 52,159,144 | 52,163,395 | 0612D03 | D6S0702i |
| 52,209,438 | 52,217,257 | 0605H02 | D6S0250i |
| 52,209,438 | 52,217,257 | 0604F04 | D6S1195i |
| 52,392,953 | 52,468,540 | 0609F02 | D6S0460i |
| 52,470,159 | 52,549,821 | 0610H02 | D6S0709i |
| 52,764,183 | 52,776,616 | 0606B02 | D6S0115i |
| 53,470,098 | 53,517,790 | 0610D03 | D6S0692i |
| 53,470,098 | 53,517,790 | 0606C07 | D6S1623 |
| 55,300,226 | 55,375,250 | 0606D06 | D6S1636 |
| 55,300,226 | 55,375,250 | 0607H02 | D6S294 |
| 55,726,402 | 55,848,334 | 134G03 | AL137178.7_78870 |
| 55,726,402 | 55,848,334 | 0602A06 | D6S1236i |


| 55,726,402 | 55,848,334 | T003B05 | D6S1661 |
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| 55,726,402 | 55,848,334 | 058C03 | G10114 |
| 56,029,347 | 56,366,851 | 181F03 | AL034452.8_92704 |
| 56,029,347 | 56,366,851 | 0614F04 | D6S0369i |
| 56,029,347 | 56,366,851 | 0605A10 | D6S1189i |
| 56,927,732 | 57,000,099 | 0602H10 | D6S1115i |
| 57,019,470 | 57,143,057 | 0613H01 | D6S0309i |
| 57,145,083 | 57,157,694 | 0603F10 | D6S0710i |
| 6,533,340 | 6,600,215 | 174G08 | AL031123.14_110913 |
| 6,533,340 | 6,600,215 | 0602C02 | D6S0311i |
| 6,533,340 | 6,600,215 | 0613E01 | D6S0579i |
| 62,447,824 | 63,054,091 | 0614G09 | D6S1212i |
| 63,964,538 | 63,980,909 | 0607D11 | D6S1093i |
| 63,964,538 | 63,980,909 | 0601E02 | D6S1120i |
| 64,339,879 | 64,351,448 | 0607A03 | D6S1658 |
| 66,095,895 | 66,473,839 | 0608A05 | D6S0119i |
| 69,401,980 | 70,156,124 | 0609B03 | D6S0469i |
| 7,486,869 | 7,531,945 | 133 H 08 | AL390026.1_29062 |
| 7,486,869 | 7,531,945 | 598G03 | chr6.fa.O7frz. 8505863 |
| 7,486,869 | 7,531,945 | 0605E10 | D6S0377i |
| 7,672,009 | 7,826,752 | 0603G03 | D6S0225i |
| 7,672,009 | 7,826,752 | 0609B07 | D6S0647i |
| 73,388,241 | 73,965,295 | 0611F04 | D6S0927i |
| 74,161,192 | 74,184,013 | 0609H05 | D6S0609i |
| 74,161,192 | 74,184,013 | 0610E01 | D6S1228i |
| 74,282,194 | 74,288,344 | 0612G07 | D6S0726i |
| 74,282,194 | 74,288,344 | $0613 \mathrm{B11}$ | D6S0882i |
| 74,462,548 | 74,591,509 | 0605D09 | D6S0741i |
| 74,462,548 | 74,591,509 | 0614G05 | D6S1596 |
| 74,462,548 | 74,591,509 | 0603D04 | D6S406 |
| 78,228,641 | 78,229,900 | 0601D09 | D6S1066i |
| 78,228,641 | 78,229,900 | 0602G10 | Z66785 |
| 79,633,908 | 79,665,039 | 0606C04 | D6S1192i |
| 79,633,908 | 79,665,039 | 0603F05 | D6S1243i |
| 79,967,681 | 80,001,174 | 585F07 | chr6.fa.O7frz. 86184415 |
| 79,967,681 | 80,001,174 | 0607C08 | D6S0920i |
| 80,873,083 | 81,112,706 | 0610G04 | D6S0574i |
| 80,873,083 | 81,112,706 | 0609E09 | D6S0728i |
| 80,873,083 | 81,112,706 | 0602A08 | D6S1181i |
| 86,216,528 | 86,262,215 | 0601A04 | D6S0734i |
| 86,216,528 | 86,262,215 | 0612G08 | D6S1141i |
| 87,597,028 | 87,709,921 | 0608C09 | D6S0297i |
| 87,597,028 | 87,709,921 | 0608H11 | D6S0357i |
| 87,851,935 | 87,861,569 | 0608C01 | D6S1182i |
| 9,813,644 | 10,168,908 | 0607H07 | D6S0639i |
| 90,199,616 | 90,400,123 | 0603B12 | D6S0248i |
| 90,199,616 | 90,400,123 | 0605A04 | D6S0254i |
| 90,199,616 | 90,400,123 | 0611C12 | D6S0883i |
| 90,596,349 | 90,640,876 | 0607F03 | D6S1613 |
| 91,280,013 | 91,353,628 | 0612E06 | D6S0738i |
| 91,280,013 | 91,353,628 | 0607H03 | D6S1570 |
| 94,007,860 | 94,185,993 | 0612F03 | D6S0584i |
| 94,007,860 | 94,185,993 | 0614B03 | D6S1056 |
| 97,479,217 | 97,695,351 | 0608B09 | D6S0288i |
| 97,479,217 | 97,695,351 | $0611 \mathrm{E07}$ | D6S0884i |
| 97,479,217 | 97,695,351 | 0601B05 | D6S1246i |


| 100,156,359 | 100,159,259 | 0710D03 | D7S0053i |
| :---: | :---: | :---: | :---: |
| 100325551 | 100331651 | ACHE1 | new design |
| 100325551 | 100331651 | ACHE2 | new design |
| 100325551 | 100331651 | ACHE3 | new design |
| 100,557,172 | 100,569,026 | 0714E11 | D7S1170i |
| 100,557,172 | 100,569,026 | 0704D01 | D7S1273i |
| 101,715,172 | 101,748,898 | 0710E04 | D7S0272i |
| 101,883,690 | 101,892,293 | 0712D11 | D7S0888i |
| 101,883,690 | 101,892,293 | 026D09 | D7S2448 |
| 102,899,473 | 103,417,199 | 0711C12 | D7S0034i |
| 102,899,473 | 103,417,199 | 0707B11 | D7S0064i |
| 102,899,473 | 103,417,199 | 0710H02 | D7S1125i |
| 102,899,473 | 103,417,199 | T003G05 | D7S2504 |
| 102,899,473 | 103,417,199 | 0701G09 | D7S818 |
| 102,899,473 | 103,417,199 | 0706D06 | Z66799 |
| 104,544,059 | 104,816,577 | 0711F08 | D7S0303i |
| 104,544,059 | 104,816,577 | 0708D07 | D7S0399i |
| 104,544,059 | 104,816,577 | 0702A04 | D7S0732i |
| 104,544,059 | 104,816,577 | 0705B08 | D7S1841 |
| 104,544,059 | 104,816,577 | 0706G04 | D7S2545 |
| 105,677,892 | 105,712,603 | 0702A08 | D7S0255i |
| 105,677,892 | 105,712,603 | 0711H05 | D7S0358i |
| 105,677,892 | 105,712,603 | 0702F12 | D7S1083i |
| 106,292,977 | 106,334,828 | 0712A03 | D7S0621i |
| 106,292,977 | 106,334,828 | 0705H07 | D7S0909i |
| 107,351,499 | 107,431,040 | 206C07 | AC005048.2_36985 |
| 107,351,499 | 107,431,040 | 0709H10 | D7S0882i |
| 107,451,232 | 107,558,036 | 0701B12 | D7S0018i |
| 111,850,462 | 111,903,483 | 0702H02 | D7S0267i |
| 111,850,462 | 111,903,483 | 0709B08 | D7S0811i |
| 113,842,288 | 114,117,391 | 0706F08 | D7S0095i |
| 113,842,288 | 114,117,391 | 0701F03 | D7S0693i |
| 113,842,288 | 114,117,391 | 0713F02 | D7S0983i |
| 113,842,288 | 114,117,391 | 0707C08 | D7S1244i |
| 115,952,075 | 115,988,466 | 137G11 | AJ133269.1_180046 |
| 115,952,075 | 115,988,466 | 0708C09 | D7S0455i |
| 115,952,075 | 115,988,466 | 0709D01 | D7S0554i |
| 116,099,695 | 116,225,676 | 0701D08 | D7S0551i |
| 116,099,695 | 116,225,676 | 0709C06 | D7S0741i |
| 116,099,695 | 116,225,676 | 0703E09 | D7S2460 |
| 121,300,395 | 121,489,326 | 139C09 | AC006020.2_128904 |
| 121,300,395 | 121,489,326 | 0702E11 | D7S0384i |
| 121,300,395 | 121,489,326 | 0705G12 | D7S0404i |
| 121,300,395 | 121,489,326 | 0701D09 | D7S0566i |
| 124,250,549 | 124,357,110 | 0702H07 | D7S0017i |
| 124,250,549 | 124,357,110 | 0708B05 | D7S0322i |
| 124,250,549 | 124,357,110 | 0710H04 | D7S0939i |
| 127,668,567 | 127,684,917 | 0705B05 | D7S1166i |
| 127,668,567 | 127,684,917 | 0707G06 | D7S1171i |
| 128,365,230 | 128,377,325 | 0704B10 | D7S1076i |
| 128,365,230 | 128,377,325 | 0705G02 | D7S1278i |
| 128,615,949 | 128,640,622 | 0712D01 | D7S0936i |
| 128,615,949 | 128,640,622 | 0712B09 | D7S1068i |
| 129,038,791 | 129,184,158 | 541E08 | chr7.fa.O7frz. 134016965 |
| 129,038,791 | 129,184,158 | 0712C01 | D7S0929i |
| 129,038,791 | 129,184,158 | 0706A06 | D7S530 |


| 13,897,379 | 13,995,289 | 0707D12 | D7S0119i |
| :---: | :---: | :---: | :---: |
| 13,897,379 | 13,995,289 | 0702D09 | D7S0199i |
| 13,897,379 | 13,995,289 | 0702H04 | D7S0211i |
| 13,897,379 | 13,995,289 | 0702B05 | D7S0679i |
| 130,663,175 | 130,831,931 | 0713 E 11 | D7S0800i |
| 130,663,175 | 130,831,931 | 0705E02 | D7S0816i |
| 130,663,175 | 130,831,931 | 0701B02 | D7S1235i |
| 130,663,175 | 130,831,931 | 0707F08 | D7S1255i |
| 138,786,805 | 138,818,998 | 0713H09 | D7S0374i |
| 139,124,668 | 139,366,560 | 0713F06 | D7S0049i |
| 139,124,668 | 139,366,560 | 0714G01 | D7S0143i |
| 139,124,668 | 139,366,560 | 0713D02 | D7S0323i |
| 139,124,668 | 139,366,560 | 0704A03 | D7S0327i |
| 139,124,668 | 139,366,560 | 0711B10 | D7S0995i |
| 139,124,668 | 139,366,560 | 0705F10 | D7S1003i |
| 139,753,916 | 139,772,419 | 0707C04 | D7S1107i |
| 141,273,626 | 141,293,252 | 0712C04 | D7S0948i |
| 141,273,626 | 141,293,252 | 0711B11 | D7S0950i |
| 141,645,314 | 142,221,097 | 0713E03 | D7S0604i |
| 141,645,314 | 142,221,097 | 0709G05 | D7S0727i |
| 141,645,314 | 142,221,097 | 0711E09 | D7S0972i |
| 141,645,314 | 142,221,097 | 0707F04 | D7S1113i |
| 141,645,314 | 142,221,097 | 0701C09 | D7S1193i |
| 141,645,314 | 142,221,097 | 0711C03 | D7S1199i |
| 141,645,314 | 142,221,097 | 0704G05 | D7S2473 |
| 142,348,323 | 142,369,625 | 0711C10 | D7S0957i |
| 142,348,323 | 142,369,625 | 0704F09 | D7S1104i |
| 142,695,524 | 142,714,907 | 0714 D 07 | D7S0996i |
| 142,798,327 | 142,816,107 | 0711H10 | D7S1004i |
| 148,135,408 | 148,212,347 | 0712D02 | D7S0960i |
| 148,135,408 | 148,212,347 | 9904G04 | D7S1052i |
| 148,135,408 | 148,212,347 | 0712D07 | D7S1056i |
| 149,666,351 | 149,669,696 | 9904E04 | D7S0793i |
| 149,666,351 | 149,669,696 | 0712F08 | D7S0812i |
| 150,319,080 | 150,342,609 | 542A05 | chr7.fa.O7frz. 157560056 |
| 150,319,080 | 150,342,609 | T003A06 | D7S636 |
| 150,381,832 | 150,385,929 | 0705C05 | D7S1277i |
| 155,288,319 | 155,297,728 | 0701 D 07 | D7S0446i |
| 155,288,319 | 155,297,728 | 0706C06 | D7S550 |
| 157,024,516 | 158,073,179 | 0710A07 | D7S0968i |
| 157,024,516 | 158,073,179 | 0711A05 | D7S1039i |
| 157,024,516 | 158,073,179 | 0707D06 | D7S1159i |
| 157,024,516 | 158,073,179 | 0707F06 | D7S1167i |
| 157,024,516 | 158,073,179 | 0704H03 | D7S1196i |
| 157,024,516 | 158,073,179 | 0704A05 | D7S1232i |
| 157,024,516 | 158,073,179 | 0701H08 | D7S2423 |
| 18,501,894 | 19,003,518 | 204F10 | AC002433.1_139651 |
| 18,501,894 | 19,003,518 | 0707E11 | D7S0073i |
| 18,501,894 | 19,003,518 | 0710G04 | D7S0086i |
| 18,501,894 | 19,003,518 | 0701D02 | D7S0163i |
| 18,501,894 | 19,003,518 | 0714H01 | D7S0186i |
| 18,501,894 | 19,003,518 | 0705C09 | D7S0578i |
| 18,501,894 | 19,003,518 | 0709D04 | D7S0651i |
| 18,501,894 | 19,003,518 | 0709E04 | D7S0655i |
| 18,501,894 | 19,003,518 | 0709A09 | D7S0832i |
| 18,501,894 | 19,003,518 | 0702B07 | D7S638 |


| 19,121,616 | 19,123,820 | 0708F10 | D7S0488i |
| :---: | :---: | :---: | :---: |
| 19,121,616 | 19,123,820 | 025E01 | D7S2495 |
| 2,912,308 | 3,050,025 | 0701D10 | D7S2484 |
| 2,912,308 | 3,050,025 | 0701D04 | D7S2521 |
| 20,337,250 | 20,421,907 | 0707G12 | D7S0132i |
| 20,337,250 | 20,421,907 | 0714D08 | D7S0200i |
| 20,337,250 | 20,421,907 | 0713A03 | D7S0208i |
| 22,732,028 | 22,738,141 | 0703F04 | D7S0667i |
| 22,732,028 | 22,738,141 | 0701C07 | D7S629 |
| 23,252,841 | 23,281,254 | 0712H01 | D7S0842i |
| 24,290,332 | 24,298,002 | 0706E08 | D7S0088i |
| 24,290,332 | 24,298,002 | 0714C11 | D7S0644i |
| 25,124,800 | 25,131,480 | 0703E04 | D7S1791 |
| 27,147,521 | 27,149,812 | 0708B10 | D7S0477i |
| 27,147,521 | 27,149,812 | 0708E10 | D7S0486i |
| 30,430,672 | 30,484,833 | 0708H08 | D7S0440i |
| 30,658,725 | 30,706,244 | 0713F04 | D7S0077i |
| 30,658,725 | 30,706,244 | 0714C03 | G09471 |
| 30,917,993 | 30,931,656 | 0705H03 | D7S0658i |
| 30,917,993 | 30,931,656 | 0706G01 | D7S526 |
| 32,963,577 | 33,013,067 | 0709E06 | D7S0749i |
| 32,963,577 | 33,013,067 | 0713D07 | D7S0843i |
| 38,246,150 | 38,374,181 | 0710E10 | D7S0036i |
| 38,246,150 | 38,374,181 | 0706G08 | D7S0106i |
| 38,246,150 | 38,374,181 | 0706H08 | D7S0176i |
| 38,246,150 | 38,374,181 | 0705D09 | D7S2497 |
| 4,688,456 | 4,777,600 | 0712A06 | D7S1022i |
| 4,688,456 | 4,777,600 | 0705C08 | D7S511 |
| 41,695,126 | 41,709,231 | 9904A03 | - |
| 41,695,126 | 41,709,231 | 0706H02 | D7S2548 |
| 41,970,196 | 42,241,712 | 0705A12 | D7S0643i |
| 41,970,196 | 42,241,712 | 0709C04 | D7S0647i |
| 41,970,196 | 42,241,712 | 0705G05 | D7S0797i |
| 41,970,196 | 42,241,712 | 0702E08 | D7S671 |
| 43,589,251 | 43,632,247 | 0708F05 | D7S0336i |
| 43,589,251 | 43,632,247 | 0708D10 | D7S0484i |
| 43,589,251 | 43,632,247 | 0709C10 | D7S0866i |
| 44,150,395 | 44,195,563 | 0701H07 | D7S1249i |
| 44,150,395 | 44,195,563 | 0701C05 | G09840 |
| 44,571,928 | 44,581,175 | 0708H01 | D7S0167i |
| 44,571,928 | 44,581,175 | 0704G06 | D7S2488 |
| 44,802,777 | 44,809,240 | 0704H05 | D7S0180i |
| 44,802,777 | 44,809,240 | 0706A03 | D7S478 |
| 44,968,786 | 44,985,203 | 0703D03 | D7S2427 |
| 5,533,312 | 5,536,747 | 0706 E 07 | G08627 |
| 50,314,924 | 50,438,053 | 0701A11 | D7S1189i |
| 50,625,259 | 50,828,652 | 0704F10 | D7S0624i |
| 50,625,259 | 50,828,652 | 0709F07 | D7S0785i |
| 50,625,259 | 50,828,652 | 0707F01 | D7S0786i |
| 55,054,219 | 55,242,525 | 0714A08 | D7S0988i |
| 55,716,261 | 55,748,439 | 0710H01 | D7S1153i |
| 6,380,651 | 6,410,123 | 0710A02 | D7S1024i |
| 6,380,651 | 6,410,123 | 0701C01 | D7S1186i |
| 65,063,110 | 65,084,635 | 0707B07 | D7S1181i |
| 65,063,110 | 65,084,635 | 0706B03 | D7S2549 |
| 7,643,100 | 7,724,763 | 0709H04 | D7S0680i |


| 7,643,100 | 7,724,763 | 0702F03 | G10238 |
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| 72,486,045 | 72,488,386 | 0705D10 | Z67541 |
| 72,821,263 | 72,822,536 | 0710C03 | D7S1124i |
| 73,262,023 | 73,282,100 | 0712C10 | D7S1213i |
| 73,262,023 | 73,282,100 | 0706B08 | HUMUT623 |
| 73,341,741 | 73,458,201 | 0712A12 | D7S0104i |
| 73,341,741 | 73,458,201 | 0711F03 | D7S0116i |
| 73,506,056 | 73,654,853 | 0713C03 | D7S0360i |
| 73,506,056 | 73,654,853 | 0706H04 | D7S2472 |
| 73,709,966 | 73,812,958 | 0706E03 | D7S1870 |
| 73,826,245 | 73,841,595 | 138B07 | AC004883.2_46325 |
| 73,848,420 | 73,905,777 | 9904G03 | D7S1175i |
| 74,975,005 | 74,995,389 | 0712G09 | D7S0670i |
| 74,975,005 | 74,995,389 | 0713A02 | D7S1122i |
| 75,236,778 | 75,257,150 | 0708B11 | D7S0501i |
| 75,236,778 | 75,257,150 | 0713C08 | HUMUT533B |
| 75,769,859 | 75,771,548 | 0711G01 | D7S1218i |
| 76,662,535 | 76,667,080 | 0713 E 12 | D7S0900i |
| 76,662,535 | 76,667,080 | 0701C03 | D7S1077i |
| 77,004,351 | 77,107,324 | 0706C09 | D7S0240i |
| 77,004,351 | 77,107,324 | 0707E07 | D7S1210i |
| 79,602,076 | 79,686,661 | 0708H11 | D7S0519i |
| 79,602,076 | 79,686,661 | 0702D06 | D7S2443 |
| 80,069,459 | 80,144,262 | 0711G12 | D7S0999i |
| 80,069,459 | 80,144,262 | 0711B09 | D7S1007i |
| 81,166,258 | 81,237,388 | 0705H02 | D7S0495i |
| 81,166,258 | 81,237,388 | 0709F04 | D7S0674i |
| 82,831,158 | 83,116,260 | 0714A12 | D7S0227i |
| 82,831,158 | 83,116,260 | 0714F08 | D7S0232i |
| 82,831,158 | 83,116,260 | 0704A08 | D7S0249i |
| 82,831,158 | 83,116,260 | 0704F12 | D7S0415i |
| 82,831,158 | 83,116,260 | 0714B02 | D7S0597i |
| 82,831,158 | 83,116,260 | 0714G05 | D7S0894i |
| 82,831,158 | 83,116,260 | 0705D06 | D7S1015i |
| 82,831,158 | 83,116,260 | 0705G10 | D7S2540 |
| 86,970,884 | 87,180,500 | 0702H08 | D7S0121i |
| 86,970,884 | 87,180,500 | 0708B04 | D7S0275i |
| 86,970,884 | 87,180,500 | 0713G07 | D7S1228i |
| 86,970,884 | 87,180,500 | 713C11 | DIJ28_10004693 |
| 91,579,402 | 91,601,946 | 0714C04 | D7S0248i |
| 91,579,402 | 91,601,946 | 0708F03 | D7S0250i |
| 91,579,402 | 91,601,946 | 0705B11 | D7S0677i |
| 92,072,171 | 92,301,148 | 0714D04 | D7S0251i |
| 92,072,171 | 92,301,148 | 0704B06 | D7S0278i |
| 92,072,171 | 92,301,148 | 0714H04 | D7S0522i |
| 93,861,809 | 93,898,480 | 0710G11 | D7S0123i |
| 93,861,809 | 93,898,480 | 0703H05 | D7S0302i |
| 93,861,809 | 93,898,480 | 0711E01 | D7S0640i |
| 96,156,015 | 96,177,139 | 586G07 | chr7.fa.O7frz. 98288632 |
| 96,156,015 | 96,177,139 | 0702B01 | D7S1274i |
| 97,199,311 | 97,207,720 | 0706G11 | D7S0543i |
| 97,199,311 | 97,207,720 | 0703A11 | HUMUT901 |
| 97,760,007 | 97,868,316 | 0710C04 | D7S0022i |
| 97,760,007 | 97,868,316 | 0710G07 | D7S0057i |
| 989,361 | 995,802 | 0703C09 | D7S1263i |
| 989,361 | 995,802 | 0703D12 | D7S2474 |


| 99,083,437 | 99,219,744 | 0712C12 | D7S0916i |
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| 99,083,437 | 99,219,744 | 0705C10 | D7S647 |
| 99,263,572 | 99,302,109 | 540B09 | chr7.fa.O7frz. 102029376 |
| 99,499,406 | 99,517,299 | 698C04 | DISD22_0005257 |
| 99,771,673 | 99,803,388 | 0710A06 | D7S0755i |
| 99,771,673 | 99,803,388 | 0702B11 | D7S2480 |
| 6,825,000 | 6,900,000 | 0804G07 | D8S1706 |
| 6,825,000 | 6,900,000 | 0801D07 | D8S1819 |
| 10,659,883 | 10,734,796 | 0805A11 | D8S0730i |
| 10,659,883 | 10,734,796 | 0811 A07 | D8S0894i |
| 10,659,883 | 10,734,796 | 0802G05 | D8S520 |
| 102,000,090 | 102,034,745 | 350B05 | AC027373.2_36983 |
| 102,000,090 | 102,034,745 | 0808G03 | D8S0336i |
| 103,730,188 | 103,737,128 | 0807F02 | D8S0305i |
| 103,730,188 | 103,737,128 | 0803B02 | D8S1834 |
| 104,222,097 | 104,311,709 | 0809A10 | D8S0324i |
| 104,222,097 | 104,311,709 | 0811D06 | D8S0643i |
| 105,421,228 | 105,438,092 | 0802B08 | D8S1738 |
| 105,421,228 | 105,438,092 | 0808D03 | HUMUT5342 |
| 108,330,886 | 108,579,459 | 0804E04 | D8S0323i |
| 108,330,886 | 108,579,459 | 0807E12 | D8S0566i |
| 108,330,886 | 108,579,459 | 0801H01 | D8S0781i |
| 108,330,886 | 108,579,459 | 9904E09 | D8S0787i |
| 108,330,886 | 108,579,459 | 0804A11 | D8S0797i |
| 11,388,919 | 11,459,522 | 0804A05 | D8S1695 |
| 11,388,919 | 11,459,522 | 0805C01 | D8S1759 |
| 11,388,919 | 11,459,522 | 0801D03 | D8S265 |
| 11,599,122 | 11,654,920 | 0806E02 | D8S0813i |
| 11,868,871 | 11,869,517 | 0811G09 | D8S1033i |
| 110,168,900 | 110,200,989 | 0801F10 | D8S0907i |
| 110,168,900 | 110,200,989 | 0808C03 | D8S0923i |
| 12,212,843 | 12,220,196 | 0807C07 | D8S0411i |
| 120,004,977 | 120,033,492 | 0801C12 | D8S0421i |
| 120,004,977 | 120,033,492 | 9904F09 | D8S0793i |
| 120,497,882 | 120,505,776 | 0802B04 | D8S0048i |
| 120,497,882 | 120,505,776 | 0805F04 | D8S1823 |
| 121,206,533 | 121,453,454 | 9904G09 | D8S0809i |
| 121,206,533 | 121,453,454 | 0808B03 | D8S0937i |
| 121,206,533 | 121,453,454 | 0806G06 | D8S1000i |
| 128,817,498 | 128,822,856 | 0810G01 | D8S0020i |
| 128,817,498 | 128,822,856 | 0811F10 | D8S0988i |
| 141,590,586 | 141,596,434 | 0808G02 | D8S0596i |
| 141,590,586 | 141,596,434 | 0808D05 | D8S0705i |
| 141,590,586 | 141,596,434 | 030H03 | D8S1717 |
| 141,737,683 | 142,080,514 | T002F06 | D8S0710i |
| 141,737,683 | 142,080,514 | 0803F07 | D8S1035i |
| 141,737,683 | 142,080,514 | 0803F06 | D8S1704 |
| 142,501,189 | 142,510,802 | 0803F08 | D8S0822i |
| 143,950,775 | 143,958,238 | 0811D08 | D8S0811i |
| 144,171,274 | 144,175,199 | 0808C05 | D8S1011i |
| 144,870,498 | 144,876,619 | 0811D03 | D8S0722i |
| 16,009,761 | 16,094,595 | 0806C12 | D8S0211i |
| 16,009,761 | 16,094,595 | 0801C07 | D8S0449i |
| 16,009,761 | 16,094,595 | 733H01 | DIJ28_10036995 |
| 16,894,049 | 16,904,061 | 256D03 | AB020858.1_88789 |
| 16,894,049 | 16,904,061 | 405D08 | AC072058.1_86060 |


| 16,894,049 | 16,904,061 | 645F10 | DISO7_10002436 |
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| 18,293,035 | 18,303,003 | 0810E12 | D8S0455i |
| 18,293,035 | 18,303,003 | T002A03 | D8S1145 |
| 21,822,330 | 21,827,151 | 0805H08 | D8S0073i |
| 21,822,330 | 21,827,151 | 0802D07 | D8S1025i |
| 21,955,883 | 21,962,266 | 0811F05 | D8S0467i |
| 21,955,883 | 21,962,266 | 0803E08 | HUMUT5312 |
| 22,354,541 | 22,454,583 | 0806D11 | D8S0186i |
| 22,354,541 | 22,454,583 | 0806D03 | D8S0839i |
| 22,601,117 | 22,606,760 | 0805A09 | D8S0190i |
| 22,601,117 | 22,606,760 | 0803B12 | D8S0654i |
| 22,601,117 | 22,606,760 | 0803F05 | D8S1733 |
| 22,933,591 | 22,982,637 | 0801A07 | D8S0005i |
| 22,933,591 | 22,982,637 | 0802C01 | D8S1008i |
| 23016377 | 23030895 | TNFRSF10C | new design |
| 23049046 | 23077488 | TNFRSF10D1 | new design |
| 23049046 | 23077488 | TNFRSF10D2 | new design |
| 23104009 | 23138584 | TNFRSF10A1 | new design |
| 23104009 | 23138584 | TNFRSF10A2 | new design |
| 23,157,114 | 23,175,452 | 0808F06 | D8S0452i |
| 25,332,693 | 25,338,087 | 0806D12 | D8S0213i |
| 25,757,490 | 25,958,292 | 0806 E 11 | D8S0187i |
| 25,757,490 | 25,958,292 | 0802F05 | D8S0929i |
| 25,757,490 | 25,958,292 | 0804B09 | D8S1031i |
| 25,757,490 | 25,958,292 | 713C09 | DIJ28_10000970 |
| 26,296,331 | 26,326,562 | 0806A04 | D8S0850i |
| 27,224,916 | 27,372,824 | 0806B11 | D8S0182i |
| 27,224,916 | 27,372,824 | 0803H08 | D8S0438i |
| 27,224,916 | 27,372,824 | 0809H04 | D8S0457i |
| 27,510,351 | 27,528,288 | 0807G08 | D8S0461i |
| 27,510,351 | 27,528,288 | 0802H02 | D8S1839 |
| 27,547,304 | 27,590,211 | 9904A09 | D8S0690i |
| 27,783,655 | 27,906,117 | 0806E03 | D8S0841i |
| 29,249,530 | 29,264,104 | 0811C03 | D8S0718i |
| 29,249,530 | 29,264,104 | 0809A02 | D8S0891i |
| 30,555,422 | 30,635,274 | 9904A06 | - |
| 30,555,422 | 30,635,274 | 543G11 | chr8.fa.O7frz. 33849047 |
| 30,555,422 | 30,635,274 | 0810C11 | D8S0636i |
| 33,568,393 | 33,577,043 | 544G01 | chr8.fa.O7frz. 37049102 |
| 33,568,393 | 33,577,043 | 028F04 | D8S1845 |
| 38,153,263 | 38,189,966 | 0811H03 | D8S0008i |
| 38,153,263 | 38,189,966 | 0811A02 | D8S1791 |
| 38,389,406 | 38,445,296 | 406A04 | AC011237.4_133993 |
| 38,389,406 | 38,445,296 | 0804H08 | D8S1038i |
| 38,389,406 | 38,445,296 | 0811B08 | D8S1821 |
| 38,734,008 | 38,829,703 | 349E03 | AC016813.3_111110 |
| 38,734,008 | 38,829,703 | 0811A11 | G10158 |
| 39,890,485 | 39,905,120 | 0810F05 | D8S0218i |
| 39,890,485 | 39,905,120 | 0809B01 | D8S0924i |
| 41,629,901 | 41,873,437 | 614G08 | chr8.fa.O7frz. 46429060 |
| 41,629,901 | 41,873,437 | 0808G04 | D8S0083i |
| 41,629,901 | 41,873,437 | 0809H02 | D8S0240i |
| 41,629,901 | 41,873,437 | 0808C02 | D8S0687i |
| 42,247,986 | 42,309,130 | 0808H08 | D8S0250i |
| 42,247,986 | 42,309,130 | 0802A05 | D8S0742i |
| 48,848,222 | 49,035,296 | 0804A12 | D8S0897i |


| 54,300,829 | 54,326,747 | 0811B03 | D8S0712i |
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| 54,300,829 | 54,326,747 | 0811F11 | D8S0716i |
| 56,954,926 | 57,086,493 | 587G05 | chr8.fa.O7frz.62681686 |
| 56,954,926 | 57,086,493 | 0802D09 | D8S0902i |
| 56,954,926 | 57,086,493 | 028C09 | D8S1828 |
| 59,565,292 | 59,575,275 | 9904F08 | D8S0253i |
| 59,565,292 | 59,575,275 | 0805F02 | D8S1723 |
| 59,658,617 | 59,734,940 | 0808C08 | D8S0948i |
| 6,344,580 | 6,408,338 | 543B01 | chr8.fa.O7frz. 7433403 |
| 6,344,580 | 6,408,338 | 0809G08 | D8S0184i |
| 65,671,246 | 65,873,902 | 0810F07 | D8S0493i |
| 65,671,246 | 65,873,902 | 0801F03 | D8S544 |
| 65,671,246 | 65,873,902 | 064D05 | G08709 |
| 67,251,166 | 67,253,380 | 0806F10 | D8S0174i |
| 71,648,227 | 71,683,158 | 0811E09 | D8S0714i |
| 75,066,141 | 75,103,859 | 0802G10 | D8S0070i |
| 75,066,141 | 75,103,859 | 0803C06 | D8S526 |
| 76,482,826 | 76,641,623 | 0811A05 | D8S0294i |
| 76,482,826 | 76,641,623 | 0801C09 | D8S0746i |
| 76,482,826 | 76,641,623 | 0805E12 | D8S0765i |
| 78,057,713 | 78,074,994 | 0806E05 | D8S0898i |
| 78,057,713 | 78,074,994 | 0802G06 | D8S0899i |
| 79,807,560 | 79,880,313 | 407D09 | AC048363.3_2317 |
| 79,807,560 | 79,880,313 | T001A11 | D8S0679i |
| 79,807,560 | 79,880,313 | 056 A 04 | Z67101 |
| 8,597,076 | 8,599,026 | 0802G07 | D8S0827i |
| 8,597,076 | 8,599,026 | 0808C07 | D8S0830i |
| 82,042,600 | 82,186,858 | 0806H08 | D8S0101i |
| 82,042,600 | 82,186,858 | 0808C11 | D8S0296i |
| 82,355,326 | 82,359,563 | 0806C08 | D8S0062i |
| 82,355,326 | 82,359,563 | 0809E03 | D8S0085i |
| 82,553,481 | 82,558,023 | 0808E09 | D8S0081i |
| 82,807,243 | 82,834,305 | 0807H10 | D8S0496i |
| 82,807,243 | 82,834,305 | 029C03 | D8S525 |
| 89,118,576 | 89,408,892 | 0806A09 | D8S0107i |
| 89,118,576 | 89,408,892 | 0810D07 | D8S0298i |
| 89,118,576 | 89,408,892 | 0807E02 | D8S0304i |
| 89,118,576 | 89,408,892 | 0809G02 | D8S0413i |
| 89,118,576 | 89,408,892 | 0810E02 | D8S0913i |
| 90,839,110 | 90,872,433 | 0806A01 | D8S0779i |
| 90,839,110 | 90,872,433 | 0806F01 | D8S0791i |
| 91,014,740 | 91,066,075 | 0811C11 | D8S0997i |
| 93,040,328 | 93,176,619 | 0801D02 | D8S0780i |
| 93,040,328 | 93,176,619 | 0806D01 | D8S0785i |
| 93,040,328 | 93,176,619 | 0806H01 | D8S0795i |
| 95,330,657 | 95,343,733 | 0803B05 | D8S1083 |
| 95,330,657 | 95,343,733 | 0805D07 | Z66605 |
| 95,961,628 | 95,976,660 | 0807G11 | D8S0545i |
| 95,961,628 | 95,976,660 | 0802H07 | D8S1042i |
| 97,343,340 | 97,415,950 | 314B04 | AC068091.3_100690 |
| 97,343,340 | 97,415,950 | 0804A07 | D8S0102i |
| 97,343,340 | 97,415,950 | 0804E08 | D8S0944i |
| 97,343,340 | 97,415,950 | 029 A 05 | D8S1772 |
| 97,575,058 | 97,693,213 | 0810E10 | D8S0108i |
| 97,575,058 | 97,693,213 | 0803A10 | D8S0796i |
| 97,575,058 | 97,693,213 | 698H11 | DISD22_0005839 |


| 99,199,244 | 99,239,816 | 0810H02 | D8S0560i |
| :---: | :---: | :---: | :---: |
| 99,199,244 | 99,239,816 | 0804B04 | D8S1778 |
| 99,199,244 | 99,239,816 | 0805B04 | D8S506 |
| 99,536,041 | 99,907,085 | 0810A08 | D8S0103i |
| 99,536,041 | 99,907,085 | 9904D10 | D8S0852i |
| 99,536,041 | 99,907,085 | 0806B04 | D8S0855i |
| 99,536,041 | 99,907,085 | 0811B07 | D8S0926i |
| 99,536,041 | 99,907,085 | 0803G04 | D8S1749 |
| 91,883,383 | 91,884,738 | 0901B04 | D9S1797 |
| 91,883,383 | 91,884,738 | 0902F09 | D9S1820 |
| 100,907,233 | 100,956,406 | $0904 \mathrm{E12}$ | D9S0215i |
| 100,907,233 | 100,956,406 | 0905H11 | D9S0532i |
| 101,024,380 | 101,032,722 | 0909D02 | D9S0086i |
| 101,623,958 | 101,668,994 | 0905H03 | D9S0313i |
| 101,623,958 | 101,668,994 | 0906C02 | D9S0726i |
| 103,393,718 | 103,397,104 | 0905A11 | D9S0506i |
| 103,393,718 | 103,397,104 | 9905A09 | D9S0724i |
| 107,046,724 | 107,241,273 | 9905H09 | D9S0793i |
| 107,046,724 | 107,241,273 | 0903C07 | D9S1784 |
| 107,464,599 | 107,465,214 | 0908H02 | D9S0219i |
| 110,656,692 | 110,659,068 | 0909G09 | D9S0390i |
| 110,656,692 | 110,659,068 | 0901A05 | D9S0830i |
| 111,177,800 | 111,300,407 | 0906 A07 | D9S0748i |
| 111,177,800 | 111,300,407 | 0904C04 | D9S1835 |
| 112,045,912 | 112,058,741 | 0907H11 | D9S0629i |
| 112,045,912 | 112,058,741 | 031D09 | D9S1675 |
| 112,431,057 | 112,431,557 | 0902C11 | D9S1828 |
| 113,364,678 | 113,401,917 | 0908E06 | D9S0213i |
| 113,364,678 | 113,401,917 | 9905D06 | D9S0608i |
| 113,698,867 | 113,737,470 | 0909D11 | D9S0513i |
| 113,698,867 | 113,737,470 | 9905F06 | D9S0616i |
| 113,698,867 | 113,737,470 | 0907C03 | D9S0802i |
| 114,967,620 | 115,024,010 | 9905B09 | D9S0754i |
| 114,967,620 | 115,024,010 | 0903G02 | D9S262 |
| 115,209,336 | 115,212,773 | 0904G11 | D9S0165i |
| 115,209,336 | 115,212,773 | 0907 E 10 | D9S0510i |
| 116,125,157 | 116,128,578 | 081G08 | D9S302 |
| 116,131,890 | 116,135,357 | 0903 A 01 | D9S0172i |
| 116,591,421 | 116,608,229 | 0901C10 | D9S0116i |
| 116,591,421 | 116,608,229 | 0908H11 | D9S0840i |
| 116,704,945 | 116,732,591 | 0904G07 | D9S0015i |
| 116,822,634 | 116,920,260 | 0904F02 | D9S155 |
| 116,822,634 | 116,920,260 | 0904H04 | D9S1776 |
| 119,506,405 | 119,519,589 | 9905G04 | D9S0554i |
| 119,506,405 | 119,519,589 | 0909G10 | D9S1864 |
| 122,704,492 | 122,730,868 | 0906A10 | D9S0050i |
| 122,704,492 | 122,730,868 | 0904E04 | D9S1823 |
| 122,754,434 | 122,852,375 | 0904C08 | D9S0039i |
| 124,173,050 | 124,197,802 | 0907D12 | D9S0318i |
| 124,173,050 | 124,197,802 | 9905C07 | D9S0649i |
| 126,155,565 | 126,217,542 | 0905D12 | D9S0545i |
| 126,155,565 | 126,217,542 | 0904H06 | HUMUT7968 |
| 126,283,336 | 126,309,530 | 0904B05 | D9S1840 |
| 126,319,380 | 126,573,410 | 0908E08 | D9S0176i |
| 126,319,380 | 126,573,410 | 0902G04 | Z67401 |
| 127,036,953 | 127,043,430 | 0904E05 | D9S1825 |


| 129,587,898 | 129,592,887 | 414B02 | AL162586.7_116830 |
| :---: | :---: | :---: | :---: |
| 129,587,898 | 129,592,887 | 0901F06 | D9S0133i |
| 129,617,112 | 129,656,856 | 317C02 | AL157935.6_74493 |
| 129,951,171 | 129,956,333 | 0908A09 | D9S0735i |
| 130,485,844 | 130,498,488 | 0902H06 | D9S0546i |
| 130,913,050 | 130,951,046 | 0907C01 | D9S0200i |
| 131,540,433 | 131,555,165 | 352H04 | AC007936.2_31251 |
| 131,540,433 | 131,555,165 | 0909H05 | D9S0091i |
| 131,855,526 | 131,942,264 | 0909E12 | D9S0020i |
| 131,855,526 | 131,942,264 | 0906F04 | D9S0720i |
| 131,855,526 | 131,942,264 | 9905H08 | D9S0723i |
| 132,874,325 | 132,958,267 | 0906C05 | D9S0761i |
| 132,874,325 | 132,958,267 | 0906G05 | D9S0767i |
| 133,441,978 | 133,605,282 | 0901H04 | D9S0547i |
| 133,441,978 | 133,605,282 | 0907G03 | D9S0550i |
| 134,458,205 | 134,535,609 | 0906G08 | D9S0198i |
| 134,458,205 | 134,535,609 | 9905H03 | HUMUT6781 |
| 134,895,897 | 134,923,709 | 0903H01 | D9S0032i |
| 134,895,897 | 134,923,709 | 0906C09 | D9S0045i |
| 135,120,384 | 135,140,451 | 0909E11 | D9S0620i |
| 135,276,941 | 135,314,329 | 0906E04 | D9S0785i |
| 135,276,941 | 135,314,329 | T003D07 | D9S164 |
| 135,616,837 | 135,847,547 | 0906G09 | D9S0049i |
| 135,616,837 | 135,847,547 | 9904F12 | D9S0052i |
| 135,616,837 | 135,847,547 | 9904G12 | D9S0057i |
| 135,616,837 | 135,847,547 | 9905A01 | D9S0062i |
| 135,616,837 | 135,847,547 | 0902H08 | D9S0858i |
| 136,673,473 | 136,876,510 | 0901D02 | D9S0634i |
| 136,940,837 | 136,949,630 | 9905C03 | D9S0548i |
| 138,508,717 | 138,560,135 | 0909A08 | D9S0347i |
| 138,900,786 | 138,940,888 | 0901D04 | D9S0734i |
| 139437668 | 139448679 | NOXA11 | new design |
| 139437668 | 139448679 | NOXA12 | new design |
| 15,454,064 | 15,501,017 | 0909D05 | D9S0187i |
| 15,454,064 | 15,501,017 | 0909C04 | D9S0668i |
| 2,005,342 | 2,183,624 | 9905G02 | D9S0348i |
| 2,005,342 | 2,183,624 | 0905A06 | D9S0367i |
| 2,005,342 | 2,183,624 | 0903F01 | D9S0371i |
| 2,794,152 | 2,834,095 | 0908E02 | D9S0561i |
| 2,794,152 | 2,834,095 | 0908B06 | D9S0566i |
| 21,067,104 | 21,067,962 | 0907F05 | D9S0224i |
| 21,067,104 | 21,067,962 | 0901B03 | D9S0783i |
| 21,130,213 | 21,132,144 | 0905H05 | D9S0366i |
| 21,191,234 | 21,229,990 | 9905G09 | D9S0786i |
| 21,357,423 | 21,358,961 | 0909B09 | D9S0369i |
| 21,470,838 | 21,472,312 | 0906G10 | D9S0122i |
| 21,470,838 | 21,472,312 | 0901H02 | D9S0373i |
| 21,957,751 | 21,984,490 | 0909G05 | D9S0018i |
| 21,957,751 | 21,984,490 | 0906A11 | D9S0115i |
| 21,992,902 | 21,999,312 | 0904D01 | D9S1870 |
| 26,894,081 | 26,937,461 | 0907G06 | D9S0788i |
| 27,099,236 | 27,220,173 | 9905F02 | D9S0285i |
| 27,099,236 | 27,220,173 | 0909B05 | D9S0749i |
| 27,099,236 | 27,220,173 | 0909E10 | D9S169 |
| 27,514,302 | 27,516,496 | 0906F06 | D9S0708i |
| 3,208,297 | 3,515,983 | 0906G11 | D9S0191i |


| 3,208,297 | 3,515,983 | 0908D07 | D9S0406i |
| :---: | :---: | :---: | :---: |
| 3,208,297 | 3,515,983 | 0903D08 | D9S0424i |
| 3,208,297 | 3,515,983 | T001F11 | D9S0571i |
| 3,208,297 | 3,515,983 | 0904B07 | HUMUT537 |
| 32,445,300 | 32,516,322 | 0907G02 | D9S0582i |
| 32,445,300 | 32,516,322 | 0903D09 | D9S0842i |
| 33,242,469 | 33,254,744 | 0908H09 | D9S0673i |
| 33,254,167 | 33,271,525 | 0904D12 | D9S0188i |
| 34,636,635 | 34,651,884 | 0906H03 | D9S0812i |
| 34,636,635 | 34,651,884 | 063 E 02 | Z67043 |
| 35,599,976 | 35,608,753 | 0906H07 | D9S0693i |
| 35,687,334 | 35,722,369 | 411A07 | AL133410.26_90374 |
| 35,687,334 | 35,722,369 | 0909E02 | D9S0363i |
| 36,562,873 | 36,667,679 | 470C11 | chr9.fa.O7frz. 39457705 |
| 36,562,873 | 36,667,679 | 0905C08 | D9S0435i |
| 37,909,131 | 38,059,249 | 0908D03 | D9S0445i |
| 37,909,131 | 38,059,249 | 0906H01 | D9S0683i |
| 4,975,245 | 5,118,183 | 0905H06 | D9S0413i |
| 4,975,245 | 5,118,183 | 0902B02 | D9S0838i |
| 5,440,525 | 5,460,547 | 0902A05 | D9S0755i |
| 5,440,525 | 5,460,547 | 0901G11 | D9S0762i |
| 5,500,570 | 5,561,252 | 0906H04 | D9S0768i |
| 6,205,809 | 6,247,983 | 0909B08 | D9S0235i |
| 6,205,809 | 6,247,983 | 0904A01 | D9S1852 |
| 70,817,241 | 70,818,849 | 0902C05 | D9S0147i |
| 74,956,493 | 74,975,129 | 0905H01 | D9S0260i |
| 74,956,493 | 74,975,129 | 0907C07 | D9S0584i |
| 76,302,072 | 76,491,937 | 0905E09 | D9S0465i |
| 76,302,072 | 76,491,937 | 0905F09 | D9S0469i |
| 76,302,072 | 76,491,937 | 9905F07 | D9S0656i |
| 78,824,391 | 78,825,689 | 0901H06 | D9S0489i |
| 8,304,246 | 9,008,735 | 0907D07 | D9S0242i |
| 8,304,246 | 9,008,735 | 0908H01 | D9S0247i |
| 8,304,246 | 9,008,735 | 0909F08 | D9S0408i |
| 8,304,246 | 9,008,735 | 0903B11 | D9S0417i |
| 8,304,246 | 9,008,735 | 0905E07 | D9S0420i |
| 8,304,246 | 9,008,735 | T003G06 | D9S1676 |
| 88,749,098 | 88,751,924 | 0902A09 | D9S1680 |
| 89,301,963 | 89,513,369 | 0908F04 | D9S0487i |
| 89,301,963 | 89,513,369 | 0907B08 | D9S0493i |
| 89,301,963 | 89,513,369 | 0907F12 | D9S0501i |
| 89,301,963 | 89,513,369 | 0906G01 | D9S257 |
| 91,115,925 | 91,121,438 | 412F07 | AL160054.5_63624 |
| 91,181,972 | 91,302,708 | 0901C01 | D9S906 |
| 92,603,890 | 92,700,652 | 0905F03 | D9S0311i |
| 92,603,890 | 92,700,652 | 0902C09 | D9S1836 |
| 93,211,148 | 93,225,965 | 352B02 | AL353645.3_94104 |
| 93,211,148 | 93,225,965 | 0908F11 | D9S0194i |
| 97,245,083 | 97,318,923 | 472F02 | chr9.fa.O7frz.96923615 |
| 97,245,083 | 97,318,923 | 9905G01 | D9S0585i |
| 97,245,083 | 97,318,923 | 0904D03 | D9S1816 |
| 99,477,012 | 99,499,460 | 0907G08 | D9S0800i |
| 99,655,357 | 99,658,818 | 472C04 | chr9.fa.O7frz. 100149745 |
| 99,785,462 | 99,818,046 | 0906E03 | D9S0832i |
| 100,491,091 | 100,527,839 | 2304C07 | DXS0684i |
| 100,491,091 | 100,527,839 | 2308F02 | DXS0923i |


| 103,697,652 | 104,898,478 | 2306D03 | DXS0001i |
| :---: | :---: | :---: | :---: |
| 103,697,652 | 104,898,478 | 2310B01 | DXS0151i |
| 103,697,652 | 104,898,478 | 2306H08 | DXS0208i |
| 103,697,652 | 104,898,478 | 2304G05 | DXS0433i |
| 103,697,652 | 104,898,478 | 2307D04 | DXS0438i |
| 103,697,652 | 104,898,478 | 2307E04 | DXS0442i |
| 103,697,652 | 104,898,478 | 2307H04 | DXS0450i |
| 103,697,652 | 104,898,478 | 2309D05 | DXS0629i |
| 103,697,652 | 104,898,478 | 2311B07 | DXS0741i |
| 103,697,652 | 104,898,478 | 2303A01 | DXS0964i |
| 103,697,652 | 104,898,478 | 2304D04 | DXS0979i |
| 103,697,652 | 104,898,478 | 2308C05 | DXS8112 |
| 103,697,652 | 104,898,478 | 2304E10 | Z67212 |
| 105,823,724 | 105,926,902 | 2301B07 | DXS8048 |
| 105,823,724 | 105,926,902 | 055B11 | DXS8097 |
| 105,823,724 | 105,926,902 | 2301A10 | HUMUT1690 |
| 106,843,107 | 106,905,858 | 2306G07 | DXS0175i |
| 106,843,107 | 106,905,858 | 2307F04 | DXS0443i |
| 107,103,687 | 107,105,431 | 2307G04 | DXS0447i |
| 107,285,493 | 107,569,383 | 2301H07 | DXS0140i |
| 107,285,493 | 107,569,383 | 2301D09 | DXS0141i |
| 107,285,493 | 107,569,383 | 2311D11 | DXS0640i |
| 107,285,493 | 107,569,383 | 2304B01 | DXS6797 |
| 107,569,810 | 107,827,431 | 2311H11 | DXS0641i |
| 107,569,810 | 107,827,431 | 2303D11 | DXS0716i |
| 110,226,244 | 110,350,816 | 2301G11 | DXS0968i |
| 114,144,794 | 114,159,792 | 2308A06 | DXS0727i |
| 115,216,003 | 115,220,253 | 552C03 | chrX.fa.O7frz. 114508264 |
| 117,745,563 | 117,812,530 | 2311B02 | DXS0287i |
| 117,745,563 | 117,812,530 | 2306B11 | DXS0308i |
| 119,446,367 | 119,487,189 | 2304F06 | DXS0156i |
| 119,446,367 | 119,487,189 | 2307H10 | DXS0622i |
| 12,795,123 | 12,818,420 | 2307D09 | DXS0567i |
| 12,834,679 | 12,851,209 | 2309E08 | DXS0246i |
| 12,903,148 | 12,905,267 | 2306A10 | DXS0244i |
| 122,821,558 | 122,875,510 | 2311B11 | DXS0634i |
| 122,821,558 | 122,875,510 | 2305H07 | DXS8098 |
| 123,307,875 | 123,334,686 | 2305F09 | DXS0171i |
| 123,307,875 | 123,334,686 | 2310D06 | DXS0613i |
| 128,408,159 | 128,485,158 | 2303D04 | DXS0835i |
| 128,408,159 | 128,485,158 | 2304C10 | DXS0839i |
| 128,408,159 | 128,485,158 | 2301B08 | DXS0843i |
| 128,607,006 | 128,616,595 | 2305C02 | DXS0166i |
| 128,607,006 | 128,616,595 | 2307D11 | DXS0635i |
| 129,091,018 | 129,127,489 | 2306G08 | DXS0205i |
| 129,091,018 | 129,127,489 | 2310F07 | DXS0483i |
| 130,235,161 | 130,361,358 | 2306A09 | DXS0212i |
| 130,235,161 | 130,361,358 | 2307A11 | DXS0624i |
| 134,482,215 | 134,544,100 | 2305F03 | DXS0701i |
| 134,895,264 | 134,957,089 | 9910E12 | DXS0735i |
| 134,895,264 | 134,957,089 | 2301E05 | DXS0826i |
| 135,558,002 | 135,570,215 | 2307H06 | DXS0495i |
| 135,558,002 | 135,570,215 | 2301G06 | DXS0829i |
| 137,541,401 | 137,894,912 | 2308H11 | DXS0042i |
| 137,541,401 | 137,894,912 | 2302H08 | DXS0173i |
| 137,541,401 | 137,894,912 | 2308 H 10 | DXS0184i |


| 137,541,401 | 137,894,912 | 2308G10 | DXS0199i |
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| 137,541,401 | 137,894,912 | 2309C05 | DXS0869i |
| 137,541,401 | 137,894,912 | 2306G02 | HUMUT1537 |
| 149,685,467 | 149,817,837 | 260E03 | AF002223.1_128496 |
| 149,685,467 | 149,817,837 | 2306C08 | DXS0189i |
| 15,273,639 | 15,312,498 | 721B09 | DIJ28_10038784 |
| 15,273,639 | 15,312,498 | 2307G09 | DXS0575i |
| 15,273,639 | 15,312,498 | 258F05 | U75931.1_14164 |
| 15,392,290 | 15,484,573 | 2303G09 | DXS1053 |
| 15,489,077 | 15,530,199 | 2310A08 | DXS0008i |
| 152,413,591 | 152,428,206 | 2307F07 | DXS0516i |
| 152,413,591 | 152,428,206 | 2307A08 | DXS0524i |
| 152,561,182 | 152,569,975 | 2303G03 | DXS8087 |
| 152,780,163 | 152,804,802 | 2310E04 | DXS0638i |
| 152,929,145 | 152,938,625 | 2311H05 | HUMUT2234 |
| 153,412,800 | 153,428,981 | 2302B07 | DXS1073 |
| 153,644,229 | 153,659,158 | 2309H10 | DXS0494i |
| 18,167,355 | 18,282,768 | 2305G08 | DXS0133i |
| 18,167,355 | 18,282,768 | 2304H09 | DXS0667i |
| 19,288,095 | 19,443,363 | 2310G03 | DXS0240i |
| 19,288,095 | 19,443,363 | 9911B02 | DXS0782i |
| 19,288,095 | 19,443,363 | 2305B08 | DXS7592 |
| 19,462,014 | 19,815,640 | 2305D08 | DXS0248i |
| 19,462,014 | 19,815,640 | 2308B03 | DXS0789i |
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| 2,680,115 | 2,743,968 | 9911C02 | DXS0787i |
| 22,927,999 | 22,931,627 | 2308E02 | DXS0064i |
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| 22,927,999 | 22,931,627 | 2301F09 | DXS7110 |
| 23,592,300 | 23,614,437 | 2311F06 | DXS0350i |
| 28,515,437 | 29,884,761 | 2306E04 | DXS0044i |
| 28,515,437 | 29,884,761 | 2306F05 | DXS0094i |
| 28,515,437 | 29,884,761 | 2309B08 | DXS0339i |
| 28,515,437 | 29,884,761 | 2304B12 | DXS0343i |
| 28,515,437 | 29,884,761 | 2306G12 | DXS0348i |
| 28,515,437 | 29,884,761 | 2307A01 | DXS0351i |
| 28,515,437 | 29,884,761 | 2310H06 | DXS0361i |
| 28,515,437 | 29,884,761 | 2307H09 | DXS0576i |
| 28,515,437 | 29,884,761 | 2308G07 | DXS0783i |
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| 28,515,437 | 29,884,761 | 2306H01 | Z67237 |
| 30,232,244 | 30,237,636 | 125G12 | AC005185.1_116697 |
| 30,232,244 | 30,237,636 | 2310E11 | DXS0063i |
| 30,232,244 | 30,237,636 | 257A12 | U31929.1_4333 |
| 37,429,931 | 37,476,322 | 2305G07 | DXS0078i |
| 37,524,208 | 37,557,658 | 2302D10 | DXS0873i |
| 38,305,553 | 38,433,118 | 2307H01 | DXS0369i |
| 38,305,553 | 38,433,118 | 2301G10 | DXS0539i |
| 41,077,595 | 41,108,669 | 2311B05 | DXS0705i |
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| 44,588,193 | 44,589,078 | 2309D03 | DXS0585i |
| 44,588,193 | 44,589,078 | 2302H02 | DXS0943i |
| 46,349,697 | 46,503,434 | 2308C01 | DXS0019i |
| 46,349,697 | 46,503,434 | 2301A05 | DXS1003 |
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| 47,368,557 | 47,374,648 | 2310D01 | DXS0396i |
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| 48,427,112 | 48,434,762 | 2306H03 | DXS0016i |
| 48,439,930 | 48,452,347 | 2306F04 | DXS0059i |
| 48,529,906 | 48,537,662 | 2303D07 | DXS0753i |
| 48,545,170 | 48,568,336 | 2310B07 | DXS0020i |
| 48,993,841 | 49,008,232 | 2306E01 | DXS1208 |
| 49,856,156 | 50,111,653 | 2308F07 | DXS0765i |
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| 53,128,274 | 53,134,447 | 2301A07 | DXS8017 |
| 53,128,274 | 53,134,447 | 055E03 | DXS8062 |
| 53,238,059 | 53,271,329 | 2308B07 | DXS0780i |
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| 55,052,213 | 55,074,136 | 665B12 | DISO7_10004639 |
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| 57,719,936 | 57,723,438 | 2301D01 | DXS0074i |
| 66,680,599 | 66,867,186 | 2311A06 | DXS0033i |
| 66,680,599 | 66,867,186 | 2303G11 | DXS0108i |
| 68,752,636 | 69,176,047 | 2308C10 | DXS0109i |
| 68,752,636 | 69,176,047 | 2307B04 | DXS0425i |
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| 69,394,741 | 69,396,379 | 2308A02 | DXS0724i |
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| 70,232,772 | 70,240,110 | 2302B01 | DXS0830i |
| 70,243,979 | 70,248,188 | 2310D04 | DXS0553i |
| 70,752,491 | 70,755,092 | 2310B10 | DXS0421i |
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| 71,341,232 | 71,375,602 | 2310A06 | DXS0409i |
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| 71,466,091 | 71,709,623 | 2302H01 | DXS0114i |
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| 71,466,091 | 71,709,623 | 2307F09 | DXS0574i |
| 76,596,303 | 76,598,669 | 2309F09 | DXS0281i |
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| 77,413,617 | 77,469,743 | 2309G08 | DXS0252i |
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| 9,391,369 | 9,647,778 | 2309A09 | DXS0017i |
| 9,391,369 | 9,647,778 | 2306A06 | DXS0105i |
| 9,391,369 | 9,647,778 | 2309A04 | DXS0324i |
| 9,391,369 | 9,647,778 | 2305G03 | DXS8051 |
| 95,826,365 | 96,746,652 | 2311A01 | DXS0012i |
| 95,826,365 | 96,746,652 | 2308C12 | DXS0146i |
| 95,826,365 | 96,746,652 | 2307A05 | DXS0454i |
| 95,826,365 | 96,746,652 | 2301B12 | DXS0458i |
| 95,826,365 | 96,746,652 | T001C06 | DXS0588i |
| 95,826,365 | 96,746,652 | 2304G12 | DXS0649i |
| 95,826,365 | 96,746,652 | 2305D03 | DXS0651i |
| 95,826,365 | 96,746,652 | 2303B12 | DXS0653i |
| 95,826,365 | 96,746,652 | 2307A12 | DXS0655i |
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| 99,984,969 | 100,015,990 | 2304C06 | DXS0191i |
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| 13,322,554 | 13,482,162 | 9911D06 | DYS0039i |
| 13525413 | 13541784 | DDX3Y1 | new design |
| 13525413 | 13541784 | DDX3Y2 | new design |


| $13,869,653$ | $14,101,947$ | 9911 A06 | DYS0035i |
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| $13,869,653$ | $14,101,947$ | 9911 F 11 | DYS0083i |
| $14,324,841$ | $14,327,298$ | 9911 C 12 | DYS0088i |
| $19,611,898$ | $19,614,093$ | 9911 D12 | DYS0089i |
| 2769527 | 2794997 | RPS4Y11 | new design |
| 2769527 | 2794997 | RPS4Y12 | new design |
| $20,326,689$ | $20,366,212$ | 9911 H 09 | DYS0069i |
| $6,838,727$ | $7,019,724$ | 9911 B 10 | DYS0071i |
| $6,838,727$ | $7,019,724$ | 9911 F 10 | DYS0075i |


| Marker Start Position | Marker End Position | Marker start - gene start | Marker end - gene end |
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| 1377882 | 1378152 | 117,361 | 103,529 |
| 1674131 | 1674253 | 116,794 | 51,144 |
| 1965222 | 1965610 | -6,547 | -141,084 |
| 10480824 | 10481227 | 41,658 | 26,027 |
| 100984474 | 100984814 | 26,589 | 7,625 |
| 101457925 | 101458128 | -17,107 | -21,534 |
| 107413661 | 107414087 | 12,837 | 10,648 |
| 107942872 | 107943023 | 27,567 | -366,085 |
| 108359820 | 108360046 | 444,515 | 50,938 |
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| 108101217 | 108101367 | 185,912 | -207,741 |
| 109482692 | 109482919 | 76,048 | 62,772 |
| 109925706 | 109926123 | 32,882 | -12,375 |
| 109854474 | 109854584 | -38,350 | -83,914 |
| 11024850 | 11025217 | 15,683 | -4,660 |
| 10916159 | 10916533 | -93,008 | -113,344 |
| 11206544 | 11206940 | 117,365 | -38,236 |
| 11275250 | 11275466 | 186,071 | 30,290 |
| 11136881 | 11137209 | 47,702 | -107,967 |
| 11813253 | 11813518 | 44,886 | 24,816 |
| 11843587 | 11843676 | 15,234 | 12,687 |
| 110244279 | 110244430 | -10,499 | -30,714 |
| 111259436 | 111259863 | 44,092 | 15,782 |
| 111215883 | 111216019 | 539 | -28,062 |
| 111576226 | 111576568 | 4,422 | -11,017 |
| 111951031 | 111951504 | 123,538 | 43,397 |
| 111801431 | 111801622 | -26,062 | -106,485 |
| 112076662 | 112076907 | 190,299 | 16,071 |
| 113057162 | 113057354 | 11,911 | 5,775 |
| 114042643 | 114042849 | -50,338 | -173,055 |
| 114203304 | 114203593 | 110,323 | -12,311 |
| 114891669 | 114891958 | -20,032 | -33,830 |
| 115021860 | 115022030 | -26,753 | -80,117 |
| 115380627 | 115380776 | 6,689 | 2,312 |
| 115626587 | 115626756 | -3,473 | -55,624 |
| 116813768 | 116814047 | -44,912 | -101,137 |
| 117015268 | 117015647 | 96,714 | 3,749 |
| 117126586 | 117126967 | 28,056 | 13,593 |
| 117222284 | 117222492 | -31,918 | -112,011 |
| 117358295 | 117358401 | 104,093 | 23,898 |
| 117554280 | 117554477 | 66,548 | -602 |
| 117477252 | 117477671 | -10,480 | -77,408 |
| 119888983 | 119889236 | 37,627 | 30,036 |
| 12107161 | 12107554 | 61,140 | -19,297 |
| 12156704 | 12156927 | 7,057 | -34,945 |
| 120395204 | 120395554 | 139,505 | -18,245 |
| 120698748 | 120698903 | 29754 | 38557 |
| 120724753 | 120724985 | -3749 | -12475 |
| 144243331 | 144243536 | 7,083 | -11,689 |
| 144341864 | 144342095 | -65,291 | -84,876 |
| 146398562 | 146398665 | -19,973 | -23,709 |
| 146482484 | 146482839 | -84,877 | -94,308 |
| 148224704 | 148224871 | 57,536 | 49,475 |
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| 149005432 | 149005676 | 36,257 | 619 |
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| 149074817 | 149074986 | 39,506 | 27,550 |
| 149126523 | 149126710 | 91,212 | 79,274 |
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| 149568202 | 149568358 | 37,165 | 1,543 |
| 15592470 | 15592659 | -63,341 | -97,823 |
| 15743102 | 15743307 | 87,291 | 52,825 |
| 150124005 | 150124257 | 84,641 | 53,285 |
| 150321738 | 150321938 | 50,132 | 45,803 |
| 151509676 | 151510156 | -27,286 | -39,662 |
| 151679764 | 151680119 | 66,956 | 65,370 |
| 151741386 | 151741558 | -32,313 | -33,786 |
| 151815693 | 151816092 | 32,980 | 26,856 |
| 151903033 | 151903133 | 2,128 | -7,015 |
| 151981475 | 151981706 | 80,570 | 71,558 |
| 152254939 | 152255252 | 25,086 | 24,002 |
| 152311055 | 152311181 | 81,202 | 79,931 |
| 152640133 | 152640283 | 4160 | 68267 |
| 152689113 | 152689266 | -44820 | 19284 |
| 153167546 | 153167951 | -46,207 | -50,397 |
| 153368540 | 153368689 | 1,980 | -5,321 |
| 153695008 | 153695274 | 270,084 | 265,944 |
| 154674287 | 154674388 | -25,856 | -62,856 |
| 155099951 | 155100256 | 57,292 | 46,986 |
| 155091484 | 155091672 | 48,825 | 38,402 |
| 155821590 | 155821916 | 71,799 | 32,982 |
| 155907593 | 155907785 | 97,430 | 73,291 |
| 155968421 | 155968850 | -13,724 | -44,696 |
| 156053140 | 156053528 | 70,995 | 39,982 |
| 156491097 | 156491389 | 74,736 | 70,079 |
| 156581636 | 156581790 | 55,436 | 51,746 |
| 157182878 | 157183052 | -53,504 | -108,517 |
| 157343650 | 157343800 | 107,268 | 52,231 |
| 157436866 | 157437022 | 28,843 | -5,892 |
| 157529540 | 157529738 | 3,412 | -14,900 |
| 157905533 | 157905683 | -43,170 | -45,320 |
| 158026403 | 158026552 | 9,057 | 7,595 |
| 158110214 | 158110363 | -53,239 | -71,647 |
| 158339075 | 158339223 | 11,321 | 4,120 |
| 158611823 | 158611977 | 8,342 | 2,715 |
| 158710267 | 158710590 | -11,177 | -49,086 |
| 158994467 | 158994763 | 79,307 | 46,498 |
| 159042070 | 159042242 | 9,518 | -22,427 |
| 159241305 | 159241641 | 9,680 | -33,763 |
| 159289844 | 159289978 | 58,219 | 14,574 |
| 159476005 | 159476157 | 24,312 | 19,044 |
| 159547782 | 159547902 | 81,703 | 73,312 |
| 159718090 | 159718302 | 23754 | 37682 |
| 159925743 | 159926078 | 66,133 | 58,458 |
| 16068553 | 16068848 | 21,607 | -70,694 |
| 160680728 | 160680994 | 49,048 | 32,442 |
| 163670101 | 163670279 | 33,323 | -10,778 |
| 165776202 | 165776309 | 109,701 | 21,838 |
| 165742032 | 165742474 | 75,531 | -11,997 |
| 165996572 | 165996681 | 38,740 | -30,003 |
| 166760192 | 166760479 | -16,434 | -19,380 |


| 166847127 | 166847541 | 34,792 | 29,602 |
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| 167782019 | 167782189 | 31,991 | -40,261 |
| 167839015 | 167839240 | 14,354 | -26,791 |
| 167888112 | 167888292 | 63,451 | 22,261 |
| 167934170 | 167934404 | 7,738 | -13,059 |
| 17538987 | 17539227 | 31,710 | -23,859 |
| 170879507 | 170879676 | -15,270 | -22,961 |
| 170974676 | 170975097 | 79,899 | 72,460 |
| 171297482 | 171297762 | 21,759 | 11,083 |
| 171264445 | 171264712 | -11,278 | -21,967 |
| 171384999 | 171385262 | -34,494 | -57,832 |
| 171509854 | 171509987 | 90,361 | 66,893 |
| 171743126 | 171743355 | 30,098 | 18,786 |
| 172228252 | 172228588 | 88,690 | 75,449 |
| 173372906 | 173373307 | 69,289 | -10,518 |
| 173789361 | 173789708 | 230,803 | -189,821 |
| 173663383 | 173663770 | 104,825 | -315,759 |
| 173551440 | 173551669 | -7,118 | -427,860 |
| 173979342 | 173979564 | 420,784 | 35 |
| 173943745 | 173943862 | 385,187 | -35,667 |
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| 177098271 | 177098441 | 12,978 | -8,397 |
| 177222734 | 177223120 | -95,001 | -110,533 |
| 177468156 | 177468492 | 150,421 | 134,839 |
| 178439502 | 178439869 | 48,911 | 81 |
| 178371799 | 178371991 | -18,792 | -67,797 |
| 179291903 | 179292263 | 22,141 | -49 |
| 18558257 | 18558663 | 251,430 | -18,900 |
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| 18306011 | 18306145 | -816 | -271,418 |
| 18640976 | 18641135 | 334,149 | 63,572 |
| 181023651 | 181023751 | -51,476 | -99,759 |
| 181140182 | 181140466 | 65,055 | 16,956 |
| 181453066 | 181453438 | 31,044 | -27,224 |
| 181485371 | 181485534 | 63,349 | 4,872 |
| 181778627 | 181778785 | -12,693 | -47,849 |
| 184498645 | 184499049 | -33,389 | -51,268 |
| 184973466 | 184973772 | 65,920 | 57,593 |
| 190964044 | 190964300 | 92,139 | 68,241 |
| 190826568 | 190826757 | -45,337 | -69,302 |
| 194983594 | 194984042 | 95,963 | 785 |
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| 195153322 | 195153523 | 142,751 | -863 |
| 195259589 | 195259868 | 80,069 | 64,889 |
| 196859008 | 196859237 | -15,416 | -133,798 |
| 198427146 | 198427367 | 163,793 | 14,192 |
| 198979477 | 198979894 | 119,830 | 74,145 |
| 198766750 | 198766850 | -92,897 | -138,899 |
| 2521865 | 2522093 |  |  |
| 2530030 | 2530401 | -50880 | -43788 |
| 20310948 | 20311409 | -71 | -7,228 |
| 20246759 | 20246900 | -64,260 | -71,737 |
| 20383149 | 20383327 | 72,130 | 64,690 |
| 20807260 | 20807633 | 19,232 | -10,355 |
| 20755451 | 20755747 | -32,577 | -62,241 |
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| 200450701 | 200450807 | 67,937 | 53,475 |
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| 201387330 | 201387616 | 60,925 | $-15,540$ |
| 201338325 | 201338509 | 11,920 | $-64,647$ |
| 201635202 | 201635518 | 58,827 | 48,278 |
| 201989619 | 201989975 | $-11,338$ | $-22,148$ |
| 202348103 | 202348336 | 39,237 | $-15,158$ |
| 202327292 | 202327557 | 18,426 | $-35,937$ |
| 202692591 | 202692790 | 34,212 | $-33,385$ |
| 202816750 | 202817052 | 64,616 | 23,181 |
| 203304684 | 203304960 | 25,721 | $-8,801$ |
| 203220013 | 203220180 | $-58,950$ | $-93,581$ |
| 203891616 | 203892025 | 58,286 | 23,402 |
| 204741554 | 204741703 | 31,140 | 4,857 |
| 204833379 | 204833538 | $-42,125$ | $-90,843$ |
| 205013527 | 205013705 | 88,615 | 39,454 |
|  |  | $\mathrm{~N} / \mathrm{A}$ | N |


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| 22822726 | 22822865 | -12,979 | -15,897 |
| 22897463 | 22897645 | 61,758 | 58,883 |
| 23141418 | 23141558 | 231,373 | 27,153 |
| 221394688 | 221394784 | 44,418 | 11,537 |
| 224053319 | 224053540 | -11,140 | -46,344 |
| 224133098 | 224133270 | 68,639 | 33,386 |
| 224579194 | 224579455 | -35,821 | -82,959 |
| 224715979 | 224716073 | 100,964 | 53,659 |
| 225001629 | 225002044 | 115,615 | 8,397 |
| 224922112 | 224922501 | 36,098 | -71,146 |
| 226712934 | 226713078 | 1,631 | 881 |
| 227588826 | 227589226 | -44,789 | -47,242 |
| 227637808 | 227638013 | 4,193 | 1,545 |
| 233985087 | 233985489 | 94,123 | -128,074 |
| 234092495 | 234092797 | 201,531 | -20,766 |
| 233918710 | 233919094 | 27,746 | -194,469 |
| 233960402 | 233960594 | 69,438 | -152,969 |
| 234369444 | 234369756 | 163,691 | 66,050 |
| 234215458 | 234215755 | 9,705 | -87,951 |
| 234756938 | 234757107 | 132,635 | 42,458 |
| 234635612 | 234635898 | 11,309 | -78,751 |
| 234697293 | 234697439 | 72,990 | -17,210 |
| 234988436 | 234988869 | 72,014 | -5,685 |
| 24365214 | 24365363 | 46,366 | 23,165 |
| 24284111 | 24284522 | -34,737 | -57,676 |
| 24327532 | 24327681 | 8,684 | -14,517 |
| 240059211 | 240059445 | -18,894 | -60,419 |
| 241879146 | 241879476 | 160,988 | -200,577 |
| 241929347 | 241929516 | 211,189 | -150,537 |
| 241980312 | 241980412 | 262,154 | -99,641 |
| 242132500 | 242132680 | 414,342 | 52,627 |
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| 245668817 | 245669049 | 20,843 | -9,984 |
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| 26523553 | 26523675 | 6,555 | 4,074 |
| 27063369 | 27063469 | -26,198 | -36,080 |
| 27161953 | 27162115 | 51,387 | 49,068 |
| 27284905 | 27285150 | -12,988 | -80,909 |
| 27406448 | 27406579 | 108,555 | 40,520 |
| 27826445 | 27826654 | 15,283 | -7,721 |
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| 32217659 | 32218079 | 73,050 | 41,501 |
| 32319158 | 32319636 | 67,141 | 20,599 |
| 32431626 | 32431870 | -57,854 | -92,483 |
| 32598999 | 32599175 | 109,519 | 74,822 |
| 32582337 | 32582751 | 52,063 | 10,928 |
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| 37880392 | 37880591 | 75,388 | 46,482 |
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| 51227256 | 51227512 | 549,518 | 28,988 |
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| 11230768 | 11230868 | 143,478 | -187,812 |
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| 112213182 | 112213522 | -34,404 | -47,770 |
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| 112381801 | 112381926 | 64,362 | 27,542 |
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| 12315190 | 12315371 | 103,548 | 63,405 |
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| 124863927 | 124864289 | -39,856 | -50,587 |
| 125000685 | 125000816 | 96,902 | 85,940 |
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| 127590942 | 127591112 | 76,046 | 16,095 |
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| 129430152 | 129430372 | 4,648 | 932 |
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| 131176113 | 131176333 | 20,657 | -279,025 |
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| 134980088 | 134980242 | 54,190 | 39,880 |
| 135205989 | 135206368 | 15,132 | -18,346 |
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| 15100482 | 15100739 | 121,118 | 64,302 |
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| 17365771 | 17366154 | 54,488 | 46,556 |
| 17935391 | 17935717 | 44,023 | -57,467 |
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| 26801475 | 26801569 | 34,337 | -95,169 |
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| 30856805 | 30856995 | 93,933 | 66,227 |
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| 31856227 | 31856469 | 208,797 | -2,279 |
| 31859304 | 31859717 | 211,874 | 969 |
| 31602801 | 31603020 | -44,629 | -255,728 |
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| 31787228 | 31787345 | 139,798 | -71,403 |
| 33306963 | 33307354 | 77,637 | 20,150 |
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| 59827471 | 59827581 | 62,726 | 27,066 |
| 6102858 | 6103019 | 68,518 | 42,863 |
| 6162899 | 6163377 | 70,241 | 19,083 |
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| 61519163 | 61519332 | 60,998 | -300,162 |
| 61820415 | 61820639 | 362,250 | 1,145 |
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| 72065347 | 72065494 | 38,237 | 32,973 |
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| 80698998 | 80699282 | -78,228 | -85,814 |
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| 133323661 | 133323811 | 33,266 | -3,510 |
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| 405192 | 405367 | -9476 | 2030 |
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| 4975673 | 4976118 | 230,597 | 5,883 |
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| 43927725 | 43928112 | 68,754 | 29,720 |
| 44591879 | 44592346 | 48,162 | -5,569 |
| 44536919 | 44537277 | -6,798 | -60,638 |
| 45860353 | 45860453 | -3,425 | -24,139 |
| 45933382 | 45933561 | 69,604 | 48,969 |
| 46807925 | 46808037 | 110,594 | 90,406 |
| 46634982 | 46635131 | -62,349 | -82,500 |
| 47332639 | 47332808 | 105,556 | 85,836 |
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| 47566731 | 47567132 | 23,267 | 4,442 |
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| 47658692 | 47659143 | 90,900 | 78,627 |
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| 47979419 | 47979563 | 20,730 | -166,683 |
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| 5258197 | 5258397 | 54,927 | 45,943 |
| 5640602 | 5641025 | -26,893 | -47,644 |
| 5685349 | 5685820 | 17,854 | -2,849 |
| 5776756 | 5777027 | 109,261 | 88,358 |
| 56766568 | 56766738 | 8,938 | 5,249 |
| 56813022 | 56813215 | -49,503 | -80,910 |
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| 6644378 | 6644852 | 62,838 | 56,175 |
| 60310130 | 60310329 | -64,853 | -69,691 |
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| 65854507 | 65854868 | 15,973 | 13,777 |
| 65975563 | 65975718 | -15,411 | -25,664 |
| 66132595 | 66133021 | 61,628 | 45,648 |
| 66187699 | 66187856 | 100,188 | 95,233 |
| 66600844 | 66601220 | 19,947 | 5,160 |
| 66526118 | 66526306 | -54,779 | -69,754 |
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| 66948890 | 66949039 | 60,675 | 51,257 |
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| 69324892 | 69325079 | 27,914 | 25,727 |
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| 105703670 | 105703837 | 202,507 | 23,126 |
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| 108248621 | 108248864 | 48,454 | 17,456 |
| 108195877 | 108196139 | -4,290 | -35,269 |
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| 112103089 | 112103288 | 123,044 | 83,072 |
| 112051097 | 112051240 | 71,052 | 31,024 |
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| 12146717 | 12147053 | 31,572 | -108,161 |
| 12293759 | 12293984 | 178,614 | 38,770 |
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| 124113254 | 124113485 | 115,929 | 73,865 |
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| 129286415 | 129286564 | 73,458 | 70,326 |
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| 4605075 | 4605250 | 35,570 | 11,948 |
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| 47544321 | 47544518 | 34,515 | 12,294 |
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| 55498736 | 55498853 | 106,252 | 91,605 |
| 55791940 | 55792334 | 22,783 | 16,808 |
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| 6587061 | 6587187 | 73,189 | 69,390 |
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| 8252748 | 8252849 | 85,255 | 70,379 |
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| 8933305 | 8933450 | 66,821 | 12,804 |
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| 91915912 | 91916032 | 222,655 | 68,894 |
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| 94989229 | 94989364 | 70,487 | 35,868 |
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| 20257883 | 20258180 | 82,404 | 62,943 |
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| 20643000 | 20643381 | 30,350 | 22,160 |
| 20557425 | 20557629 | -55,225 | -63,592 |
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| 21215397 | 21215842 | 72,227 | 39,205 |
| 23139163 | 23139311 | 96,440 | -8,921 |
| 23243644 | 23243795 | 200,921 | 95,563 |
| 23083142 | 23083346 | 40,419 | -64,886 |
| 25963459 | 25963901 | 237,183 | 86,526 |
| 25834146 | 25834547 | 107,870 | -42,828 |
| 25729969 | 25730139 | 3,693 | -147,236 |
| 26078404 | 26078846 | 48,564 | -82,239 |
| 26805545 | 26805847 | -91,136 | -101,976 |
| 26957206 | 26957639 | 60,525 | 49,816 |
| 27358433 | 27358626 | -75,840 | -82,691 |
| 27478401 | 27478570 | 2,990 | -94,159 |
| 27618297 | 27618554 | 142,886 | 45,825 |
| 27953530 | 27953699 | 179,740 | -13,533 |
| 27869868 | 27870023 | 96,078 | -97,209 |
| 28043073 | 28043316 | 269,283 | 76,084 |
| 29970494 | 29970820 | 39,610 | -118,909 |
| 29906396 | 29906794 | -24,488 | -182,935 |
| 30003534 | 30003738 | 72,650 | -85,991 |
| 30163702 | 30163949 | 232,818 | 74,220 |
| 30244427 | 30244579 | 36,782 | 8,023 |
| 31922785 | 31923197 | 135,168 | 51,388 |
| 35835464 | 35835578 | -69,031 | -79,430 |
| 36317134 | 36317436 | 25,795 | 15,696 |
| 40132897 | 40133162 | 105,096 | -5,572 |
| 39975108 | 39975292 | -52,693 | -163,442 |
| 41970551 | 41970720 | -64,321 | -109,428 |
| 42092311 | 42092485 | 57,439 | 12,337 |
| 44821373 | 44821567 | 228,723 | 65,330 |
| 44496186 | 44496554 | -96,464 | -259,683 |
| 44713243 | 44713628 | 120,593 | -42,609 |
| 44830757 | 44830862 | 21,749 | 17,357 |
| 46302100 | 46302455 | -3,414 | -65,724 |
| 46345241 | 46345710 | 39,727 | -22,469 |
| 47788809 | 47789010 | -94,361 | -98,937 |
| 47997390 | 47997628 | 114,220 | 109,681 |
| 48196230 | 48196355 | 17,538 | 14,856 |
| 52255232 | 52255509 | 79,832 | 43,561 |
| 94559555 | 94559705 | 89,471 | -191,983 |
| 94698049 | 94698332 | 227,965 | -53,356 |
| 94480217 | 94480614 | 10,133 | -271,074 |
| 94740327 | 94740523 | 270,243 | -11,165 |
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| 98666284 | 98666427 | -78,506 | -91,281 |
| 101709325 | 101709631 | 92,186 | 33,855 |
| 101822811 | 101823084 | 60,436 | -18,200 |
| 102307652 | 102308130 | -5,917 | -134,251 |
| 102347697 | 102347928 | 34,128 | -94,453 |
| 103022865 | 103023071 | -69,777 | -75,836 |
| 103312211 | 103312378 | 78,504 | 60,829 |
| 104341507 | 104341623 | 50,978 | 44,587 |
| 104670447 | 104670755 | 83,665 | 67,956 |


| 105529292 | 105529704 | 101,198 | 101,594 |
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| 105335154 | 105335468 | -92,940 | -92,642 |
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| 19906719 | 19906868 | 57,352 | 35,571 |
| 20276160 | 20276537 | -42,890 | -43,927 |
| 20618135 | 20618406 | 37,884 | 36,180 |
| 22182690 | 22182915 | 1,022,793 | 92,000 |
| 21479279 | 21479478 | 319,382 | -611,437 |
| 22028150 | 22028453 | 868,253 | -62,462 |
| 21328896 | 21329186 | 168,999 | -761,729 |
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| 21577795 | 21577963 | 417,898 | -512,952 |
| 21186185 | 21186349 | 26,288 | -904,566 |
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| 22280814 | 22281202 | -94,819 | -105,441 |
| 22535821 | 22536096 | 76,248 | 67,595 |
| 22656227 | 22656373 | -128 | -2,292 |
| 22814584 | 22814823 | -31,282 | -35,975 |
| 22962373 | 22962572 | 50,515 | 47,120 |
| 23630109 | 23630242 | -31,098 | -47,774 |
| 23839637 | 23839920 | 91,010 | 84,900 |
| 23943961 | 23944294 | 36,867 | 25,644 |
| 24044148 | 24044344 | -68,416 | -70,962 |
| 24121670 | 24121860 | 48330 | 51453 |
| 24192338 | 24192541 | -22338 | -19228 |
| 28330240 | 28330505 | 25,439 | 21,884 |
| 28257603 | 28257879 | -47,198 | -50,742 |
| 29435297 | 29435499 | 319,861 | -31,152 |
| 29262603 | 29262789 | 147,167 | -203,862 |
| 29526109 | 29526368 | 410,673 | 59,717 |
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| 34335036 | 34335455 | 85,638 | 81,806 |
| 34231541 | 34231758 | -17,857 | -21,891 |
| 34918147 | 34918589 | -22,321 | -25,114 |
| 35033722 | 35034192 | 93,254 | 90,489 |
| 37116005 | 37116456 | -12,935 | -17,784 |
| 37232711 | 37233184 | 103,771 | 98,944 |
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| 37824490 | 37824956 | 77,535 | 72,937 |
| 44710536 | 44710892 | 55,677 | 36,620 |
| 49191881 | 49192154 | 56,716 | 41,014 |
| 49928543 | 49928979 | -26,450 | -140,147 |
| 50152652 | 50153047 | 197,659 | 83,921 |
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| 51759902 | 51760366 | -44,279 | -52,826 |
| 51890305 | 51890758 | 39,442 | 25,684 |
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| 53436482 | 53436805 | -49,725 | -56,557 |
| 53593304 | 53593509 | 107,097 | 100,147 |
| 53938580 | 53938926 | 5,157 | -17,756 |
| 54629121 | 54629387 | 65,527 | 43,427 |
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| 55718215 | 55718408 | 63,369 | -119,376 |
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| 61143560 | 61143747 | 285,374 | 56,296 |
| 60800266 | 60800425 | -57,920 | -287,026 |
| 61065410 | 61065576 | 207,224 | -21,875 |
| 62818684 | 62818896 | -30,711 | -35,420 |
| 62892243 | 62892460 | 42,848 | 38,144 |
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| 63876107 | 63876504 | 254,719 | 1,434 |
| 63720780 | 63720959 | 99,392 | -154,111 |
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| 68543815 | 68544025 | 133,022 | 28,278 |
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| 70370762 | 70370929 | 106,157 | 25,288 |
| 74852031 | 74852209 | 36,747 | 33,524 |
| 75038385 | 75038685 | -20,152 | -44,401 |
| 75091785 | 75092149 | 33,248 | 9,063 |
| 75464406 | 75464680 | -29,789 | -52,562 |
| 75527214 | 75527586 | 33,019 | 10,344 |
| 75942526 | 75942991 | 35,047 | -93,970 |
| 76025446 | 76025619 | 117,967 | -11,342 |
| 75863464 | 75863650 | -44,015 | -173,311 |
| 77174362 | 77174469 | -34,140 | -69,640 |
| 77293976 | 77294208 | 85,474 | 50,099 |
| 80747781 | 80748039 | 256,253 | 65,640 |
| 80471349 | 80471808 | -20,179 | -210,591 |
| 80827913 | 80828075 | 111,766 | 70,747 |
| 87457690 | 87457886 | -11,421 | -71,774 |
| 87574460 | 87574634 | 105,349 | 44,974 |
| 87965948 | 87966195 | -37,919 | -124,681 |
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| 88951893 | 88952084 | 259,619 | -1,043 |
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| 88740800 | 88740999 | 48,526 | -212,128 |
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| 90693068 | 90693495 | -75,561 | -96,482 |
| 92208560 | 92209028 | -31,347 | -75,737 |
| 92301822 | 92301976 | 61,915 | 17,211 |
| 92665583 | 92665684 | -52,711 | -55,318 |
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| 93521287 | 93521445 | -65,732 | -95,866 |
| 94013907 | 94014188 | 113,503 | 100,010 |
| 93907101 | 93907324 | 6,697 | -6,854 |
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| 99275129 | 99275290 | 54,722 | 11,899 |
| 99909870 | 99910019 | 135,015 | 95,462 |
| 99765742 | 99765988 | -9,113 | -48,569 |
| 23078354 | 23078568 | 458,467 | -137,134 |
| 23155553 | 23155754 | 535,666 | -59,948 |
| 22855116 | 22855330 | 235,229 | -360,372 |
| 23231137 | 23231291 | 611,250 | 15,589 |


| 22681893 | 22682091 | 62,006 | -533,611 |
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| 32012936 | 32013103 | 67,216 | -105,492 |
| 31913669 | 31913869 | -32,051 | -204,726 |
| 32820177 | 32820560 | -49,546 | -54,621 |
| 32973028 | 32973177 | 103,305 | 97,996 |
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| 37776157 | 37776513 | 115,585 | 99,553 |
| 37660967 | 37661197 | 395 | -15,763 |
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| 38739331 | 38739549 | 35330 | 72097 |
| 39052668 | 39052839 | 43,829 | 34,310 |
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| 39544355 | 39544546 | 131,994 | 84,008 |
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| 41622051 | 41622403 | 9,102 | -147,122 |
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| 42800986 | 42801377 | 10,009 | 3,728 |
| 47550841 | 47551198 | 48,090 | -15,617 |
| 47657612 | 47658032 | 154,861 | 91,217 |
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| 49407764 | 49408106 | 118,803 | -9,980 |
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| 55368974 | 55369156 | 370,849 | 1,148 |
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| 56666269 | 56666486 | -9,533 | -162,983 |
| 56830252 | 56830385 | 154,450 | 916 |
| 57249332 | 57249720 | 64,720 | 45,184 |
| 57689592 | 57689840 | -28,766 | -47,151 |
| 58001826 | 58002255 | -82,601 | -83,179 |
| 58139383 | 58139534 | 54,956 | 54,100 |
| 58478125 | 58478377 | 51,483 | 900 |
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| 59297551 | 59297816 | 720,796 | -10,978 |
| 58782555 | 58782724 | 205,800 | -526,070 |
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| 61905971 | 61906120 | -80,317 | -219,454 |
| 62317691 | 62317867 | 82,624 | 75,460 |
| 63127006 | 63127286 | 131,960 | 89,200 |
| 62942725 | 62942862 | -52,321 | -95,224 |
| 63239929 | 63240121 | 43,159 | 26,894 |
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| 64540767 | 64540965 | 74,093 | -29,971 |
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| 65320336 | 65320572 | 175,087 | 45,985 |
| 65113637 | 65113910 | -31,612 | -160,677 |
| 65835777 | 65836140 | 213,702 | -50,366 |
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| 65669136 | 65669238 | 47,061 | -217,268 |
| 66422171 | 66422538 | 41,075 | -89,008 |
| 66518358 | 66518512 | 137,262 | 6,966 |
| 66297190 | 66297339 | -83,906 | -214,207 |
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| 68241434 | 68241634 | 113,837 | 64,324 |
| 69940758 | 69941117 | 50,810 | 43,463 |
| 69841475 | 69841628 | -48,473 | -56,026 |
| 70313328 | 70313733 | -50,794 | -142,135 |
| 71725492 | 71725884 | 86,082 | 13,078 |
| 71857694 | 71858093 | 94,019 | 64,181 |
| 72387628 | 72387823 | -29,529 | -59,311 |
| 72477905 | 72478090 | 60,748 | 30,956 |
| 72794409 | 72794608 | -4,534 | -10,322 |
| 72841471 | 72841726 | 42,528 | 36,796 |
| 73573462 | 73573611 | 26,947 | -85,069 |
| 73935516 | 73935815 | -84,817 | -156,027 |
| 74102486 | 74102604 | 82,153 | 10,762 |
| 75018571 | 75018934 | -56,038 | -97,793 |
| 75079805 | 75080007 | 5,196 | -36,720 |
| 77055934 | 77056165 | 54,772 | 31,690 |
| 76928739 | 76929010 | -72,423 | -95,465 |
| 78111951 | 78112070 | 71,661 | 61,372 |
| 79182433 | 79182589 | -79,715 | -209,568 |
| 79231537 | 79231915 | -30,611 | -160,242 |
| 79452215 | 79452352 | 190,067 | 60,195 |
| 83706160 | 83706564 | 127,339 | -387,026 |
| 84166314 | 84166691 | 587,493 | 73,101 |
| 83712649 | 83712876 | 133,828 | -380,714 |
| 84046409 | 84046579 | 467,588 | -47,011 |
| 83678213 | 83678322 | 99,392 | -415,268 |
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| 86956758 | 86957203 | -26,281 | -43,481 |
| 87096274 | 87096437 | 113,235 | 95,753 |
| 88099167 | 88099366 | -29,963 | -59,706 |
| 88188346 | 88188631 | 59,216 | 29,559 |
| 89007946 | 89008196 | -53,660 | -151,492 |
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| 89198746 | 89198873 | 137,140 | 39,185 |
| 89313939 | 89314196 | 101,050 | 86,505 |
| 94659259 | 94659444 | -15,691 | -23,604 |
| 94715440 | 94715645 | 40,490 | 32,597 |
| 96961037 | 96961440 | -49,251 | -357,594 |
| 97077023 | 97077172 | 66,735 | -241,862 |
| 97409689 | 97409850 | 399,401 | 90,816 |
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| 1715132 | 1715300 | 18,910 | -45,019 |
| 1941711 | 1941861 | -27,208 | -29,580 |
| 10960331 | 10960485 | 92,683 | 34,144 |
| 10988018 | 10988179 | 120,370 | 61,838 |


| 11253513 | 11253680 | -2,262 | -3,860 |
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| 11325274 | 11325545 | 69,499 | 68,005 |
| 11545405 | 11545633 | -3,952 | -43,190 |
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| 11952949 | 11953116 | 13516 | 16310 |
| 11968282 | 11968434 | -1817 | 992 |
| 15946157 | 15946367 | 241,664 | 87,979 |
| 15832715 | 15832958 | 128,222 | -25,430 |
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| 16039379 | 16039743 | 88,444 | -104,031 |
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| 2153935 | 2154221 | 8,135 | -13,910 |
| 21639956 | 21640295 | 80,530 | 68,822 |
| 21491146 | 21491448 | -68,280 | -80,025 |
| 23634635 | 23635014 | 36,943 | 25,825 |
| 23684673 | 23684800 | 86,981 | 75,611 |
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| 24101554 | 24101749 | 346,731 | -37,609 |
| 23742470 | 23742824 | -12,353 | -396,534 |
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| 24143725 | 24143913 | 388,902 | 4,555 |
| 27332776 | 27333207 | 100,024 | 49,607 |
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| 28446611 | 28447003 | 28,427 | 21,347 |
| 29585304 | 29585508 | 3,503 | -4,180 |
| 29717086 | 29717245 | -14,505 | -49,597 |
| 2973215 | 2973608 | -37,128 | -38,777 |
| 3057182 | 3057379 | 46,839 | 44,994 |
| 3270186 | 3270361 | 38,157 | 23,733 |
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| 3917284 | 3917686 | 200,716 | 46,963 |
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| 45855925 | 45856074 | 109,127 | -196,445 |
| 45755306 | 45755468 | 8,508 | -297,051 |
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| 46689476 | 46689676 | -68,847 | -149,130 |
| 46924887 | 46925148 | 166,564 | 86,342 |
| 49384477 | 49384823 | 95,926 | 60,335 |
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| 54106207 | 54106515 | 35,618 | 8,411 |
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| 55222408 | 55222493 | 41,640 | 39,992 |
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| 56024812 | 56025048 | 28,632 | 17,573 |
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| 56552978 | 56553146 | -63,805 | -85,160 |
| 64882417 | 64882654 | -75,647 | -113,536 |
| 65094021 | 65094170 | 135,957 | 97,980 |
| 65191351 | 65191682 | 47,384 | 21,219 |
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| 65589309 | 65589450 | -31,242 | -103,012 |
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| 67515184 | 67515304 | 186,488 | 88,359 |
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| 68218683 | 68218832 | 62,185 | -77,222 |
| 68942004 | 68942399 | 51,431 | 17,167 |
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| 70745261 | 70745460 | 60,145 | 41,148 |
| 71474329 | 71474502 | 95,873 | -165,273 |
| 71600267 | 71600426 | 221,811 | -39,349 |
| 71522743 | 71522918 | 144,287 | -116,857 |
| 71647224 | 71647436 | 268,768 | 7,661 |
| 71565950 | 71566111 | 187,494 | -73,664 |
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| 74082377 | 74082765 | 197,268 | 57,877 |
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| 78139168 | 78139367 | -46,564 | -52,745 |
| 78209120 | 78209232 | 23,388 | 17,120 |
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| 15690829 | 15691057 | 74,783 | 61,927 |
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| 16005240 | 16005438 | 129,257 | -54,132 |
| 16355951 | 16356133 | 96,338 | 75,091 |
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| 17664978 | 17665246 | 9,184 | -15,804 |
| 17974790 | 17974994 | 42,782 | 22,977 |
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| 23154137 | 23154468 | 46,218 | 2,786 |
| 23225251 | 23225444 | 117,332 | 73,762 |
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| 24434601 | 24434751 | 9,938 | -96,805 |
| 24566594 | 24566862 | 141,931 | 35,306 |
| 25564795 | 25565234 | 15,763 | -21,597 |
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| 29584279 | 29584562 | -22,130 | -23,773 |
| 29764758 | 29764899 | 57,174 | 55,157 |
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| 32534609 | 32535044 | 18,569 | -305,971 |
| 32834762 | 32835086 | 318,722 | -5,929 |
| 32711674 | 32712041 | 195,634 | -128,974 |
| 32956258 | 32956455 | 32,194 | 8,746 |
| 33158937 | 33159143 | 112,411 | 81,543 |
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| 35228291 | 35228766 | 130,372 | 90,325 |
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| 35603899 | 35604099 | 101,332 | 93,600 |
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| 36795531 | 36795960 | -90,936 | -95,234 |
| 36963799 | 36964015 | 77,332 | 72,821 |
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| 39300407 | 39300768 | 101,392 | 88,896 |
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| 39175619 | 39175792 | -23,396 | -36,080 |
| 39566265 | 39566482 | 56,618 | 9,942 |
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| 4689816 | 4690139 | 43,419 | 41,383 |
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| 4836563 | 4836811 | 60,191 | 57,744 |
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| 40429552 | 40429701 | 36,965 | 28,531 |
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| 43217422 | 43217588 | 51,813 | 39,104 |
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| 43166868 | 43167022 | 1,259 | -11,462 |
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| 45465137 | 45465591 | 42,769 | 38,004 |
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| 45676226 | 45676554 | 59,770 | 42,562 |
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| 5499787 | 5499937 | 156,315 | 71,384 |
| 50660637 | 50660937 | -36,733 | -94,949 |
| 50795628 | 50795719 | 98,258 | 39,833 |
| 52385101 | 52385333 | 64,832 | 38,925 |
| 52310353 | 52310633 | -9,916 | -35,775 |
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| 53676373 | 53676538 | 25828 | 36757 |
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| 59430647 | 59431054 | 82,353 | 81,124 |
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| 59967269 | 59967418 | 41,069 | 34,546 |
| 6878488 | 6878922 | 38,380 | 22,702 |
| 6936589 | 6936982 | 18,009 | 12,658 |
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| 62261110 | 62261431 | 531,722 | 24,107 |
| 62309476 | 62309723 | 580,088 | 72,399 |
| 61711640 | 61711743 | -17,748 | -525,581 |
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| 68732511 | 68732808 | 59,756 | 53,119 |
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| 69882746 | 69883135 | -91,371 | -109,393 |
| 70021387 | 70021603 | 47,270 | 29,075 |
| 7198529 | 7198678 | 41,827 | 25,316 |
| 7284908 | 7285058 | 1,495 | -3,922 |
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| 7489622 | 7489957 | 96,523 | 84,308 |
| 7931173 | 7931386 | 48,090 | 38,209 |
| 70073050 | 70073218 | 24,208 | 19,341 |
| 70105487 | 70105785 | 18,388 | 5,768 |
| 70222075 | 70222326 | 20,028 | 1,614 |
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| 71313458 | 71313703 | 84,347 | 48,209 |
| 71694392 | 71694802 | 50,383 | 45,836 |
| 71713182 | 71713612 | 69,173 | 64,646 |
| 71993688 | 71993888 | 101,391 | 98,352 |


| 71942968 | 71943164 | 50,671 | 47,628 |
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| 75761616 | 75761765 | 38,004 | 26,232 |
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| 77846169 | 77846317 | -19,866 | -22,452 |
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| 78124664 | 78124790 | 53,781 | -28,953 |
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| 8828321 | 8828649 | 105,368 | 72,090 |
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| 8752505 | 8752653 | 29,552 | -3,906 |
| 11711693 | 11712033 | 32,430 | -159,889 |
| 11658519 | 11658718 | -20,744 | -213,204 |
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| 11958362 | 11958526 | 279,099 | 86,604 |
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| 13777416 | 13777790 | 60,736 | 23,236 |
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| 37770662 | 37770882 | -18,535 | -144,564 |
| 37988134 | 37988309 | 198,937 | 72,863 |
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| 46428295 | 46428518 | 87,813 | -83,676 |
| 46315429 | 46315724 | -25,053 | -196,470 |
| 51332534 | 51332775 | 286,567 | -74,083 |
| 51404172 | 51404547 | 358,205 | -2,311 |
| 51024679 | 51025142 | -21,288 | -381,716 |
| 51180270 | 51180704 | 134,303 | -226,154 |
| 54396910 | 54397059 | -92,688 | -171,291 |
| 54506404 | 54506563 | 16,806 | -61,787 |
| 55129037 | 55129236 | 43,786 | 37,631 |
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| 55214674 | 55214878 | 66,586 | 37,415 |
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| 58946489 | 58946749 | 4,930 | -190,844 |
| 59205218 | 59205463 | 263,659 | 67,870 |
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| 59837986 | 59838224 | 70,412 | 59,131 |
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| 835049 | 835319 | 123,457 | 32,772 |
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| 75349212 | 75349491 | 92,452 | -40,820 |
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| 1644992 | 1645289 | 84,699 | 41,961 |
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| 10152253 | 10152466 | 69,056 | 65,401 |
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| 10423635 | 10423825 | 101,430 | 71,614 |
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| 11356276 | 11356548 | 60,183 | 45,227 |
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| 12802814 | 12802908 | 39,528 | 37,779 |
| 12933953 | 12934229 | 23,530 | 17,926 |
| 12971199 | 12971526 | 53,545 | 46,071 |
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| 14748665 | 14748915 | 44,460 | -1,438 |
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| 15021462 | 15021719 | -2,553 | -6,181 |
| 15194816 | 15195102 | 63,372 | 22,310 |
| 15543228 | 15543409 | 102,765 | 92,097 |
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| 15714939 | 15715309 | 102,232 | 80,675 |
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| 15953695 | 15953983 | 69,514 | 47,657 |
| 16193930 | 16194348 | 88,092 | 63,967 |
| 16037651 | 16037932 | -68,187 | -92,449 |
| 16070758 | 16070934 | -35,080 | -59,447 |
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| 17258423 | 17258592 | 54,729 | 41,441 |
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| 17326770 | 17327165 | -47,985 | -50,292 |
| 17390091 | 17390415 | 15,336 | 12,958 |
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| 17899279 | 17899516 | 110,957 | 79,716 |
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| 18204062 | 18204270 | 79,046 | 61,927 |
| 18434072 | 18434264 | 76,104 | 73,277 |
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| 18907905 | 18908245 | 16,411 | 7,809 |
| 19244455 | 19244606 | 80,447 | 70,928 |
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| 35108867 | 35109072 | 114,126 | 102,013 |
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| 38490143 | 38490435 | 7,367 | 5,275 |
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| 44164440 | 44164669 | 81,985 | 73,295 |
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| 44640675 | 44640827 | 51,382 | 48,942 |
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| 44908211 | 44908484 | 21,425 | 16,556 |
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| 45696050 | 45696398 | 50,509 | 32,882 |
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| 46432760 | 46433198 | 41,805 | 27,914 |
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| 6963679 | 6963891 | 28,432 | 8,070 |
| 6894594 | 6894784 | -40,653 | -61,037 |
| 61024210 | 61024456 | 61,956 | 20,774 |
| 64100707 | 64100861 | -72,792 | -124,201 |
| 64179780 | 64180013 | 6,281 | -45,049 |
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| 68188178 | 68188626 | -15,394 | -153,240 |
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| 70649299 | 70649563 | 121,375 | 15,125 |
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| 70946526 | 70946787 | 35,671 | 30,326 |
| 72233266 | 72233397 | 23,391 | 4,926 |
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| 74650343 | 74650585 | 51,577 | 43,759 |
| 74556566 | 74556774 | -42,200 | -50,052 |
| 74734382 | 74734584 | 99,587 | 96,403 |
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| 85766676 | 85767120 | 28,725 | 18,297 |
| 86014326 | 86014777 | 94,544 | 45,129 |
| 85930443 | 85930702 | 10,661 | -38,946 |
| 86481806 | 86482255 | -40,148 | -91,095 |
| 86607160 | 86607620 | 85,206 | 34,270 |
| 86662286 | 86662672 | 140,332 | 89,322 |
| 86841440 | 86841786 | -23,799 | -29,852 |
| 86908869 | 86909126 | 12,898 | -33,423 |
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| 89092632 | 89092736 | -28,204 | -28,574 |
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| 9545244 | 9545477 | 81,980 | 64,350 |
| 9770787 | 9771039 | 129,235 | 82,410 |
| 96131524 | 96131764 | -41,114 | -43,142 |
| 96391129 | 96391330 | 25,918 | -13,671 |
| 97648370 | 97648747 | -48,091 | -74,008 |
| 97665848 | 97666066 | -30,613 | -56,689 |
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| 1042062 | 1042169 | 123 | -54,853 |
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| 1414174 | 1414553 | 14,788 | -5,680 |
| 1627603 | 1628002 | 136,035 | 79,347 |
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| 1915834 | 1915981 | 93,021 | 47,438 |
| 10468619 | 10469073 | -97,715 | -133,563 |
| 10647962 | 10648232 | 81,628 | 45,596 |
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| 23011100 | 23011259 | 36,830 | 32,958 |
| 23072431 | 23072641 | 64,436 | 57,664 |
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| 23710443 | 23710602 | 34,253 | 31,028 |
| 29276254 | 29276633 | -32,874 | -34,463 |
| 29531589 | 29531787 | 64,877 | 51,143 |
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| 29817890 | 29818039 | 101,974 | 43,673 |
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| 30041477 | 30041705 | 45,058 | 38,149 |
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| 32523119 | 32523324 | 108,417 | -39,535 |
| 32619207 | 32619328 | 204,505 | 56,469 |
| 33236471 | 33236781 | 24,340 | 7,953 |
| 33292497 | 33292779 | 80,366 | 63,951 |
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| 33372401 | 33372550 | 94,306 | 44,332 |
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| 35427085 | 35427236 | 20,583 | -40,003 |
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| 37158641 | 37158802 | 134,232 | 57,024 |
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| 4706071 | 4706483 | 91,075 | 76,247 |
| 41150140 | 41150578 | 1,015,334 | -101,446 |
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| 42610838 | 42610995 | -70,739 | -102,802 |
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| 43501370 | 43501723 | 114,028 | 91,245 |
| 43413351 | 43413737 | 26,009 | 3,259 |
| 43838223 | 43838449 | -65,545 | -80,993 |
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| 44260258 | 44260495 | 79,945 | -105,762 |
| 44373780 | 44373923 | 193,467 | 7,666 |
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| 47912603 | 47913082 | 49,946 | -29,097 |
| 47824925 | 47825188 | -37,732 | -116,991 |
| 48163425 | 48163888 | 32,357 | -39,790 |
| 48255969 | 48256203 | 124,901 | 52,525 |
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| 48680433 | 48680690 | 120,139 | 45,984 |
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| 54317486 | 54317726 | 60,291 | 59,448 |
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| 55241252 | 55241401 | 64,041 | -33,690 |
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| 57058409 | 57058747 | 67,812 | 43,050 |
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| 60337796 | 60337928 | 20,286 | -37,835 |
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| 17849118 | 17849521 | -38,724 | -57,615 |
| 17983081 | 17983527 | 95,239 | 76,391 |
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| 35189018 | 35189185 | 107,050 | -154,326 |
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| 36748728 | 36748907 | 69,169 | 37,912 |
| 36580779 | 36580938 | -98,780 | -130,057 |
| 37789223 | 37789394 | 127,494 | -19,953 |
| 37615962 | 37616430 | -45,767 | -192,917 |
| 37752385 | 37752576 | 90,656 | -56,771 |
| 39018313 | 39018754 | 342,642 | 63,266 |
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| 40027996 | 40028375 | -11,208 | -67,518 |
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| 40555208 | 40555687 | 248,995 | -585,222 |
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| 41050684 | 41050855 | 744,471 | -90,054 |
| 41649699 | 41650054 | -6,121 | -52,685 |
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| 44543483 | 44543743 | 13,292 | 1,213 |
| 45169636 | 45169867 | 39,340 | -3,314 |
| 45691473 | 45691724 | 46441 | 95055 |
| 46224207 | 46224587 | -1,884 | -24,804 |
| 46145961 | 46146257 | -80,130 | -103,134 |
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| 16492436 | 16492647 | -9,049 | -100,736 |
| 16729284 | 16729487 | 137,824 | 97,675 |
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| 19572504 | 19572679 | -29,210 | -65,355 |
| 19740853 | 19741038 | 41,404 | 27,919 |
| 20349158 | 20349359 | 22,616 | 20,771 |
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| 21878997 | 21879366 | 26,445 | -110,858 |
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| 22264873 | 22265321 | 19,561 | 12,826 |
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| 29029236 | 29029413 | 62,795 | 56,665 |
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| 30176682 | 30177150 | 51,143 | 16,712 |
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| 36007306 | 36007695 | 56,068 | 37,444 |
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| 36871605 | 36871832 | 34,157 | -35,931 |
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| 38284898 | 38285044 | 38,383 | 36,407 |
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| 107048902 | 107049121 | 189,103 | -21,456 |
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| 114005884 | 114006128 | -11,362 | -41,359 |
| 114223163 | 114223318 | 100,417 | 46,668 |
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| 126085702 | 126086143 | 121,217 | -2,699 |
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| 128303650 | 128303799 | 113,458 | 64,877 |
| 128834841 | 128835209 | 34,898 | 11,240 |
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| 131959370 | 131959656 | 78,902 | 11,316 |
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| 138243465 | 138243644 | 84,068 | 31,034 |
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| 77309459 | 77309619 | 10,541 | -44,440 |
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| 81336012 | 81336161 | -70,754 | -95,034 |
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| 84465266 | 84465396 | 32,627 | -9,934 |
| 87122526 | 87122759 | -34,130 | -470,548 |
| 87451125 | 87451294 | 294,469 | -142,013 |
| 87163936 | 87164088 | 7,280 | -429,219 |
| 87243136 | 87243335 | 86,480 | -349,972 |
| 87496152 | 87496301 | 339,496 | -97,006 |
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| 87820478 | 87820651 | 85,569 | -134,675 |
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| 144123911 | 144124141 | N/A | N/A |
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| 110380391 | 110380561 | -53,286 | -61,062 |
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| 132039199 | 132039604 | 118,670 | 31,953 |


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| 176850177 | 176850409 | -21,007 | -26,164 |
| 179112656 | 179112905 | 54,120 | 21,657 |
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| 34200645 | 34200758 | 178,605 | 40,362 |
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| 38729405 | 38729644 | 218,583 | 98,391 |
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| 54521287 | 54521362 | 87,057 | 79,525 |
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| 55081967 | 55082127 | 12,358 | -66,235 |
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| 73993056 | 73993295 | 34,066 | 20,290 |
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| 74859004 | 74859218 | 156,320 | 15,499 |
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| 75979075 | 75979548 | 32,012 | 24,552 |
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| 82796665 | 82796820 | -6,674 | -115,917 |
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| 92967674 | 92967861 | 22,875 | 11,784 |
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| 96027674 | 96027845 | -94,603 | -141,714 |
| 40750185 | 40750393 | N/A | N/A |
| 23689621 | 23689880 | N/A | N/A |
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| 1959022 | 1959399 | 389,982 | -231,446 |
| 2107768 | 2107968 | 538,728 | -82,877 |
| 2213166 | 2213628 | 644,126 | 22,783 |
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| 10693184 | 10693317 | 56,609 | -44,270 |
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| 12430249 | 12430479 | 31,667 | 25,066 |
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| 127955309 | 127955758 | 142,286 | 76,218 |
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| 128589350 | 128589677 | 257,725 | -293,776 |
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| 132082141 | 132082454 | 82,006 | -27,789 |
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| 137122101 | 137122569 | 202,223 | -32,780 |
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| 143119565 | 143119738 | 5,268 | -188,293 |
| 143875476 | 143875698 | 86,711 | 62,181 |
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| 149696454 | 149696628 | 15,698 | -77,814 |
| 149647323 | 149647646 | -33,433 | -126,796 |
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| 149731357 | 149731567 | 50,601 | -42,875 |
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| 149932908 | 149933057 | 65,584 | 24,193 |
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| 150264691 | 150264844 | 13,397 | 10,981 |
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| 151757233 | 151757652 | 154,031 | 38,050 |
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| 152244388 | 152244779 | 74,009 | -221,320 |
| 152780109 | 152780421 | 295,594 | -219,806 |
| 152517174 | 152517336 | 32,659 | -482,891 |
| 152604307 | 152604506 | 119,792 | -395,721 |
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| 155893582 | 155893858 | 135,388 | 75,129 |
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| 155789784 | 155789943 | 31,590 | -28,786 |
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| 157267984 | 157268385 | 127,228 | -303,709 |
| 158254185 | 158254646 | 89,903 | -31,451 |
| 158121699 | 158121941 | -42,583 | -164,156 |
| 158584828 | 158585237 | 75,456 | 50,229 |
| 158555320 | 158555779 | 45,948 | 20,771 |
| 158484231 | 158484466 | -25,141 | -50,542 |
| 16593127 | 16593309 | 185,805 | -276,391 |
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| 160362967 | 160363122 | 52,846 | -84,451 |
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| 160409020 | 160409197 | -53,833 | -90,543 |
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| 160519187 | 160519548 | 56,334 | 19,808 |
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| 165782129 | 165782332 | 121,363 | -213,246 |
| 167030450 | 167030633 | 287,606 | -165,128 |
| 167409422 | 167409870 | 76,762 | -63,304 |
| 167355649 | 167355816 | 22,989 | -117,358 |
| 167506629 | 167507105 | 173,969 | 33,931 |
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| 167965406 | 167965699 | 37,340 | 25,311 |
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| 170485524 | 170485694 | 27,755 | -70,468 |
| 170601790 | 170602006 | 144,021 | 45,844 |
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| 170761646 | 170762073 | 75,512 | 57,761 |
| 18259657 | 18260013 | 23,136 | -3,340 |
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| 20862105 | 20862340 | 219,438 | -478,274 |
| 22419518 | 22419782 | 24,059 | 14,073 |
| 22472950 | 22473076 | 77,491 | 67,367 |
| 22713240 | 22713472 | 35,583 | 33,601 |
| 22637790 | 22637989 | -39,867 | -41,882 |
| 254547 | 254760 | 17,494 | -41,595 |
| 345685 | 345874 | 108,632 | 49,519 |
| 24666439 | 24666704 | -91,745 | -108,536 |
| 24854231 | 24854406 | 96,047 | 79,166 |
| 25405006 | 25405194 | 17,721 | -323,543 |
| 26245803 | 26246097 | 50,376 | 41,059 |
| 26411190 | 26411300 | -99,270 | -112,145 |
| 26518884 | 26519041 | 8,424 | -4,404 |
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| 27342916 | 27343329 | 19,429 | 11,002 |
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| 28200881 | 28201258 | -16,814 | -31,957 |
| 28388134 | 28388412 | 87,088 | 79,173 |
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| 29751702 | 29751902 | 120,334 | -26,139 |
| 29753266 | 29753582 | 20,511 | 5,454 |
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| 29550333 | 29550545 | -248,763 | -252,507 |
| 29476083 | 29476238 | -323,013 | -326,814 |
| 29670244 | 29670434 | -128,852 | -132,618 |
| 29644100 | 29644264 | -154,996 | -158,788 |
| 29831719 | 29832103 | 29,294 | 29,208 |
| 30027185 | 30027486 | 124,462 | 5,853 |
| 3103508 | 3103611 | 94,296 | 43,191 |
| 30284629 | 30284792 | 266,319 | 263,159 |
| 29822664 | 29822810 | -195,646 | -198,823 |
| 29930636 | 29930880 | -87,674 | -90,753 |
| 30396299 | 30396607 | 377,989 | 374,974 |
| 30238657 | 30238758 | 220,347 | 217,125 |
| 30127492 | 30127613 | 109,182 | 105,980 |
| 30170485 | 30170689 | 152,175 | 149,056 |
| 30127492 | 30127613 | 96,530 | 96,223 |
| 30170485 | 30170689 | 63,016 | 61,056 |
| 30213749 | 30213943 | 1,262 | -10,548 |
| 30321166 | 30321536 | 93,465 | 84,846 |
| 30456329 | 30456500 | 35,452 | 33,851 |
| 30519687 | 30520097 | -41,964 | -42,603 |
| 30515634 | 30515830 | -49,616 | -53,247 |
| 30583169 | 30583395 | 17,919 | 14,318 |
| 30716816 | 30716971 | 95,183 | 83,984 |
| 30889107 | 30889213 | 113,544 | 95,568 |
| 30889030 | 30889254 | 408 | -17,161 |
| 30967250 | 30967351 | -16,706 | -22,508 |
| 31145725 | 31146018 | 86,251 | 80,364 |
| 31367081 | 31367280 | 22,576 | -65,655 |
| 31579685 | 31579894 | 235,180 | 146,959 |
| 31317048 | 31317463 | -27,457 | -115,472 |
| 31312116 | 31312543 | -32,389 | -120,392 |
| 31643332 | 31643497 | 298,827 | 210,562 |
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| 31648097 | 31648514 | 43,379 | 42,527 |
| N/A | N/A | N/A | N/A |
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| 31814672 | 31814942 | 62,232 | 55,146 |
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| 32327802 | 32328124 | 324,329 | 306,696 |
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| 32153714 | 32153926 | 69,539 | -31,205 |
| 32256603 | 32256752 | 52,141 | 50,707 |
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| 32377666 | 32377882 | 9,213 | -82,428 |
| 32442933 | 32443084 | 74,469 | -4,578 |
| 32505292 | 32505607 | -10,305 | -15,336 |
| 32511329 | 32511466 | -4,296 | -9,335 |
| 32707394 | 32707854 | 52,870 | 42,251 |
| 32721587 | 32721719 | 8,475 | 2,312 |
| 32845242 | 32845375 | 110,020 | 91,079 |
| 32778014 | 32778233 | 42,792 | 23,937 |
| 32785483 | 32785908 | 50,258 | 43,336 |
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| 32961904 | 32962066 | 64,316 | 47,541 |
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| 33232052 | 33232277 | 43,846 | 27,409 |
| 33281628 | 33281840 | 43,181 | 13,617 |
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| 33281662 | 33281841 | 5,031 | 1,649 |
| 33410105 | 33410512 | 84,078 | 62,872 |
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| 33571139 | 33571410 | -77,168 | -84,587 |
| 33790555 | 33790704 | 94,055 | 18,375 |
| 34598233 | 34598378 | 56,350 | -12,606 |
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| 35347923 | 35348320 | -70,390 | -155,613 |
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| 35690784 | 35691025 | 41,439 | -113,313 |
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| 36249771 | 36250032 | 120,002 | 34,212 |
| 36631575 | 36631969 | 61,928 | 8,735 |
| 36474795 | 36474972 | -94,852 | -148,262 |
| 36777970 | 36778156 | 23,557 | 15,062 |
| 36740976 | 36741204 | -13,437 | -21,890 |
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| 37035946 | 37036145 | 105,365 | 85,367 |
| 37239036 | 37239280 | -6,921 | -11,902 |
| 37940375 | 37940615 | 45,090 | -289,760 |
| 38247153 | 38247408 | 351,868 | 17,033 |
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| 38661618 | 38661787 | 410,907 | -12,061 |
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| 39139577 | 39139747 | 347,264 | 33,202 |
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| 39725403 | 39725562 | -142,717 | -255,060 |


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| 41318854 | 41319059 | 34,584 | 20,699 |
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| 41731825 | 41732017 | 109,683 | 53,917 |
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| 43874446 | 43874606 | 28,522 | 12,404 |
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| 44355441 | 44355541 | 32,639 | 25,943 |
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| 47413291 | 47413588 | 106,064 | 27,949 |
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| 51889875 | 51890101 | 301,771 | -170,281 |
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| 52103872 | 52104329 | -55,272 | -59,066 |
| 52312123 | 52312346 | 102,685 | 95,089 |
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| 53396387 | 53396728 | -73,711 | -121,062 |
| 53513334 | 53513603 | 43,236 | -4,187 |
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| 56363233 | 56363421 | 333,886 | -3,430 |
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| 56907259 | 56907424 | -20,473 | -92,675 |
| 57111380 | 57111746 | 91,910 | -31,311 |
| 57229312 | 57229578 | 84,229 | 71,884 |
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| 6695688 | 6695945 | 162,348 | 95,730 |
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| 62715703 | 62715852 | 267,879 | -338,239 |
| 64025660 | 64025763 | 61,122 | 44,854 |
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| 64409345 | 64409563 | 69,466 | 58,115 |
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| 7870600 | 7870798 | 198,591 | 44,046 |
| 7768079 | 7768240 | 96,070 | -58,512 |
| 73622850 | 73623097 | 234,609 | -342,198 |
| 74179707 | 74179889 | 18,515 | -4,124 |
| 74098291 | 74098441 | -62,901 | -85,572 |
| 74215696 | 74216093 | -66,498 | -72,251 |
| 74307562 | 74308009 | 25,368 | 19,665 |
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| 74527430 | 74527633 | 64,882 | -63,876 |
| 78324298 | 78324448 | 95,657 | 94,548 |
| 78161782 | 78162024 | -66,859 | -67,876 |
| 79706873 | 79707064 | 72,965 | 42,025 |
| 79596279 | 79596441 | -37,629 | -68,598 |
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| 81015266 | 81015415 | 142,183 | -97,291 |
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| 86306353 | 86306776 | 89,825 | 44,561 |
| 87541626 | 87541795 | -55,402 | -168,126 |
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| 94154229 | 94154474 | 146,369 | -31,519 |
| 97699568 | 97699739 | 220,351 | 4,388 |
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| 100065689 | 100066059 | -90,670 | -93,200 |
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| 100371833 | 100372040 | -46282 | -40389 |
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| 100599084 | 100599367 | 41,912 | 30,341 |
| 100520648 | 100520797 | -36,524 | -48,229 |
| 101727115 | 101727453 | 11,943 | -21,445 |
| 101847243 | 101847689 | -36,447 | -44,604 |
| 101894447 | 101894688 | 10,757 | 2,395 |
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| 104751564 | 104751762 | 207,505 | -64,815 |
| 104851797 | 104851997 | 307,738 | 35,420 |
| 104504496 | 104504626 | -39,563 | -311,951 |
| 105597520 | 105597727 | -80,372 | -114,876 |
| 105726552 | 105726909 | 48,660 | 14,306 |
| 105778845 | 105779011 | 100,953 | 66,408 |
| 106243511 | 106243926 | -49,466 | -90,902 |
| 106396984 | 106397237 | 104,007 | 62,409 |
| 107470518 | 107470694 | 119,019 | 39,654 |
| 107378539 | 107378707 | 27,040 | -52,333 |
| 107592145 | 107592427 | 140,913 | 34,391 |
| 111824685 | 111824903 | -25,777 | -78,580 |
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| 113776818 | 113777080 | -65,470 | -340,311 |
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| 115893593 | 115893788 | -58,482 | -94,678 |
| 116023121 | 116023293 | 71,046 | 34,827 |
| 116059115 | 116059287 | -40,580 | -166,389 |
| 116097894 | 116098063 | -1,801 | -127,613 |
| 116195224 | 116195416 | 95,529 | -30,260 |
| 121567185 | 121567481 | 266,790 | 78,155 |
| 121319883 | 121320093 | 19,488 | -169,233 |
| 121255206 | 121255512 | -45,189 | -233,814 |
| 121485106 | 121485313 | 184,711 | -4,013 |
| 124262464 | 124262696 | 11,915 | -94,414 |
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| 124399535 | 124399874 | 148,986 | 42,764 |
| 127688775 | 127688924 | 20,208 | 4,007 |
| 127775947 | 127776057 | 107,380 | 91,140 |
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| 13987603 | 13987762 | 90,224 | -7,527 |
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| 13926485 | 13926682 | 29,106 | -68,607 |
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| 130901037 | 130901516 | 237,862 | 69,585 |
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| 139405578 | 139405792 | 280,910 | 39,232 |
| 139789996 | 139790107 | 36,080 | 17,688 |
| 141221537 | 141221992 | -52,089 | -71,260 |
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| 44241919 | 44242154 | 91,524 | 46,591 |
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| 44516225 | 44516452 | -55,703 | -64,723 |
| 44852100 | 44852281 | 49,323 | 43,041 |
| 44898728 | 44898851 | 95,951 | 89,611 |
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| 5597181 | 5597288 | 63,869 | 60,541 |
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| 65153022 | 65153136 | 89,912 | 68,501 |
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| 92374977 | 92375168 | 302,806 | 74,020 |
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| 120571321 | 120571545 | 73,439 | 65,769 |
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| 141631901 | 141632083 | 41,315 | 35,649 |
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| 18396794 | 18397013 | 103,759 | 94,010 |
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| 22024661 | 22024894 | 68,778 | 62,628 |
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| 65892255 | 65892470 | 221,009 | 18,568 |
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| 97358290 | 97358436 | 14,950 | -57,514 |
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| 99173118 | 99173499 | -26,126 | -66,317 |
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| 116227615 | 116227889 | 95,725 | 92,532 |
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| 129681037 | 129681437 | 93,139 | 88,550 |
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| 131034543 | 131034982 | 121,493 | 83,936 |
| 131568037 | 131568281 | 27,604 | 13,116 |
| 131540451 | 131540610 | 18 | -14,555 |
| 131981878 | 131982283 | 126,352 | 40,019 |
| 131868460 | 131868560 | 12,934 | -73,704 |
| 131813132 | 131813407 | -42,394 | -128,857 |
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| 35637621 | 35637723 | 37,645 | 28,970 |
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| 29757165 | 29757273 | 1,241,728 | -127,488 |
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| 44668655 | 44668973 | 80,462 | 79,895 |
| 44590422 | 44590571 | 2,229 | 1,493 |
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| 13814930 | 13815340 | $-54,723$ | $-286,607$ |
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| 6921204 | 6921332 | 82,477 | $-98,392$ |
| 7001434 | 7001541 | 162,707 | $-18,183$ |


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|  | all_152_indall_152_10 | all_152_20 | all_152_40 | all_156_ind | all_156_10 | all_156_20 | all156_40 |
| :--- | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| D01 | 1 | 0 | 0 | 0 | 1 | 0 | 0 |
| D24 | 1 | 0 | 0 | 0 | 0 | 0 | 0 |
| P01 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| P24 | 0 | 0 | 0 | 0 | 1 | 0 | 0 |

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|  | all_152_ind | all_152_10 | all_152_20 | all_152_40 | all_156_ind | all_156_10 | all_156_20 |
| :--- | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| all156_40 |  |  |  |  |  |  |  |
| D01 | 1 | 0 | 0 | 0 | 1 | 0 | 0 |
| D24 | 1 | 0 | 0 | 0 | 0 | 0 | 0 |
| P01 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| P24 | 0 | 0 | 0 | 0 | 1 | 0 | 0 |

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|  | all_152_indall_152_10 | all_152_20 | all_152_40 | all_156_ind | all_156_10 | all_156_20 | all156_40 |  |
| :--- | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| D01 | 1 | 0 | 0 | 0 | 1 | 0 | 0 | 0 |
| D24 | 1 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| P01 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| P24 | 0 | 0 | 0 | 0 | 1 | 0 | 0 | 0 |




all_169_indall_169_10 all_169_20 all_169_40 all_173_indall_173_10 all_173_20 all_173_40 all_177_ing

| 7 | 16.84735 | 13.25337 | 15.84178 | 27 | 37.90654 | 38.65567 |
| ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 10 | 40.50675 | 31.4152 | 30.2175 | 13 | 25.72534 | 24.55138 |
| 13 | 37.54609 | 8.151417 | 16.54119 | 24 | 31.58542 | 35.42731 |
| 5 | 19.73258 | 17.15352 | 24.05867 | 33 | 40.55213 | 42.07973 |
|  | 39.17811 | 269 | 261 |  |  |  |

all_169_indall_169_10 all_169_20 all_169_40 all_173_indall_173_10 all_173_20 all_173_40 all_177_ing
$\left.\begin{array}{|r|r|r|r|r|r|r|r|}\hline 7 & 20.81844 & 43.84921 & 22.22143 & 27 & 39.45429 & 37.7834 & 39.34286 \\ \hline 10 & 27.31143 & 19.92455 & 25.64863 & 13 & 22.42095 & 23.70511 & 21.98454 \\ \hline 13 & 37.21922 & 33.43653 & 26.8886 & 24 & 23.89866 & 31.46285 & 32.26633\end{array}\right] 269$.
all_169_indall_169_10 all_169_20 all_169_40 all_173_indall_173_10 all_173_20 all_173_40 all_177_ind

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| 7 | 30.13636 | 24.48507 | 20.83099 | 27 | 31.00988 | 31.20896 | 33.76056 | 241 |


| 10 | 27.72346 | 16.51871 | 31.69862 | 13 | 25.73305 | 24.51724 | 27.89478 |
| ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 13 | 22.4442 | 14.33121 | 16.37622 | 24 | 31.49628 | 37.83439 | 31.13788 |
| 5 | 16.28648 | 25.77068 | 18.2372 | 33 | 49.28246 | 40.81193 | 44.67279 |

066B03_H


73_20 all_177_indi all_177_20 all_181_indi all_181_20 all_185_indi all_185_20 all_189_indi all.

066B03_M


066B03_L


| all_177_10 | all_177_20 | all_177_40 | all_181_ind | all_181_10 | all_181_20 | all_181_40 | all_185_indall_185_10 |  |
| ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 230.8411 | 231.934 | 229.9199 | 150 | 143.2025 | 145.1244 | 145.8586 | 13 | 13.20249 |
| 248.0861 | 253.8296 | 256.671 | 152 | 134.3829 | 139.9165 | 137.1045 | 29 | 25.29895 |
| 245.1249 | 263.9805 | 261.4703 | 130 | 119.7665 | 125.5109 | 122.5642 | 19 | 15.97706 |
| 254.0988 | 255.8287 | 246.7669 | 151 | 138.2953 | 138.4343 | 141.1516 | 23 | 21.32122 |


| all_177_10 | all_177_20 | all_177_40 | all_181_ind | all_181_10 | all_181_20 | all_181_40 | all_185_indall_185_10 |  |
| ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 232.4166 | 223.704 | 229.3786 | 150 | 136.551 | 123.4355 | 138.9143 | 13 | 12.75969 |
| 262.2048 | 266.7847 | 264.8138 | 152 | 137.6857 | 138.961 | 137.903 | 29 | 24.37714 |
| 244.7066 | 245.6656 | 247.3752 | 130 | 125.37 | 121.5557 | 126.9526 | 19 | 18.8055 |
| 250.235 | 250.4206 | 241.9914 | 151 | 126.2491 | 132.65 | 133.2894 | 23 | 19.08989 |


| all_177_10 | all_177_20 | all_177_40 | all_181_ind | all_181_10 | all_181_20 | all_181_40 | all_185_indall_185_10 |  |
| ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: |
| 230.8271 | 237.4925 | 237.5211 | 150 | 136.0504 | 137.2687 | 137.1972 | 13 | 13.97628 |
| 256.7618 | 265.3426 | 252.7436 | 152 | 137.9064 | 142.4087 | 137.9947 | 29 | 25.87522 |
| 250.9782 | 266.2739 | 255.5613 | 130 | 130.4492 | 114.0764 | 132.163 | 19 | 14.63213 |
| 242.1821 | 247.4787 | 241.6011 | 151 | 142.9826 | 138.2792 | 145.2284 | 23 | 23.2664 |


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| all_185_20 | all_185_40 | all_189_ind all_189_ | all_189__ | all_189_ |  |
| ---: | ---: | ---: | ---: | ---: | ---: |
| 13.03248 | 12.98741 | 2 | 0 | 0 | 0 |
| 24.28739 | 25.4775 | 0 | 0 | 0 | 0 |
| 16.92987 | 16.54119 | 3 | 0 | 0 | 0 |
| 20.50382 | 22.8447 | 0 | 0 | 0 | 0 |


| all_185_20 | all_185_40 | all_189_indall_189_ | lll_189_ | all_189__ |  |
| ---: | ---: | ---: | ---: | ---: | ---: |
| 13.22784 | 12.14286 | 2 | 0 | 0 | 0 |
| 24.6247 | 23.65004 | 0 | 0 | 0 | 0 |
| 17.87926 | 16.51729 | 3 | 0 | 0 | 0 |
| 19.54739 | 19.22621 | 0 | 0 | 0 | 0 |


| all_185_20 | all_185_40 | all_189_ind | all_189_ | all_189_ | all_189__ |
| ---: | ---: | ---: | ---: | ---: | ---: |
| 11.54478 | 12.69014 | 2 | 0 | 0 | 0 |
| 25.21277 | 23.6683 | 0 | 0 | 0 | 0 |
| 17.48408 | 14.76166 | 3 | 0 | 0 | 0 |
| 21.6594 | 24.2605 | 0 | 0 | 0 | 0 |

