Advancing Animal Models of Human Type 1 Diabetes by Engraftment of Functional Human Tissues in Immunodeficient Mice

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Despite decades of studying rodent models of type 1 diabetes (T1D), no therapy capable of preventing or curing T1D has successfully been translated from rodents to humans. This inability to translate otherwise promising therapies to clinical settings likely resides, to a major degree, from significant species-specific differences between rodent and human immune systems as well as species-related variances in islets in terms of their cellular composition, function, and gene expression. Indeed, taken collectively, these differences underscore the need to define interactions between the human immune system with human β cells. Immunodeficient mice engrafted with human immune systems and human β cells represent an interesting and promising opportunity to study these components in vivo. To meet this need, years of effort have been extended to develop mice depleted of undesirable components while at the same time, allowing the introduction of constituents necessary to recapitulate physiological settings as near as possible to human T1D. With this, these so-called “humanized mice” are currently being used as a preclinical bridge to facilitate identification and translation of novel discoveries to clinical settings.

Our understanding of T1D has been influenced greatly by studies performed using rodent models. The two rodent models studied most extensively are the nonobese diabetic (NOD) mouse and the biobreeding (BB) rat (Greiner et al. 2001). These two rodent models have helped define the autoimmune response

that leads to the destruction of β cells and to provide clues into the pathogenesis of T1D. These models have noted that T1D is characterized by a T-cell-mediated immune response against islet autoantigens, that it can be transferred with autoreactive lymphocytes (i.e., T cells), and that the autoimmunity persists long after the loss of

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M.A. Brehm et al.

β cells, displaying recurrent autoimmunity when transplanted with syngeneic islets (Von and Nepom 2009). Similar patterns of pathogenesis have been observed in humans, particularly with respect to recurrent autoimmunity. A key observation made by Sutherland and colleagues (1984) showed that T1D individuals transplanted with kidneys and pancreas from identical twins retained the kidney graft, but rejected the islets in the pancreatic graft. Recurrent autoimmunity has also been observed following transplantation of allogeneic islets (Vendrame et al. 2010).

In addition to studies of T1D pathogenesis, rodent models have been used to investigate potential therapeutics for the treatment and cure of this disease (Staeve-Vieira et al. 2007). In the NOD mouse, 200 therapies have been shown to prevent diabetes (Atkinson and Leiter 1999). However, it must be noted that NOD mice are resistant to tolerance induction even to nonislet tissues and grafts (Pearson et al. 2003) and thus, their immune systems appear to differ in many respects from that of nonautoimmune mice. In the BB rat, far fewer therapies have been shown to prevent diabetes (Greiner et al. 2001). However, despite decades of studies with rat as well as mouse models of T1D, we have yet to successfully translate therapies that prevent, delay, or cure T1D in humans (Roep 2007; Staeve-Vieira et al. 2007; Couzin-Frankel 2011; Greenbaum and Atkinson 2011).

Underlying this failure is the increasing awareness that mouse and human immune systems, as well as islets, differ significantly in terms of their cell composition, function, and gene expression. These distinctive features of human immune systems and islets, combined with the need to translate emerging findings from rodent biology to human therapeutic efficacy, have formed roadblocks for translating discoveries in rodents to new approaches to prevent or delay T1D in humans.

HUMAN ISLETS DIFFER SUBSTANTIALLY FROM RODENT ISLETS

Mouse and human pancreatic islets as the target of autoimmune attack differ in many ways, including cellular architecture and composition,

proliferative capacity, susceptibility to injury, ability to form islet amyloid, and expression of heat-shock proteins, islet-enriched transcription factors, antioxidant enzymes, and the principal glucose transporter (i.e., GLUT1 vs. GLUT2) (Eizirik et al. 1994; Welsh et al. 1995; Brissova et al. 2005; Butler et al. 2007). In contrast to the more familiar rodent islet cellular architecture (characterized by non- β endocrine cells surrounding the inner β -cell mass), the endocrine cells in human islets are more intermingled (Brissova et al. 2005; Cabrera et al. 2006; Bosco et al. 2010). Furthermore, in contrast to rodent β cells that replicate or regenerate in response to a number of stimuli such as insulin resistance, β -cell ablation, and partial pancreatectomy, the human β -cell proliferative capacity appears to be very modest and often nonexistent (Butler et al. 2007).

HUMAN IMMUNE SYSTEMS DIFFER SIGNIFICANTLY FROM RODENT IMMUNE SYSTEMS

Of particular interest in the study of autoimmunity are the differences between human and mouse immune systems (Mestas and Hughes 2001). The first and foremost difference involves the major histocompatibility complex (MHC), a primary genetic factor for determining diabetes susceptibility. In the NOD mouse, the MHC class I is K^dD^b, with a unique I-A^{g7} class II molecule. The initial CD4 T-cell autoimmune response in NOD mice appears to be directed against insulin (Atkinson et al. 1990; Zhang et al. 1991) and specifically, against an epitope of insulin (B9-23) presented by the I-A^{g7} class II molecule (Abiru et al. 2001). On the basis of these observations, the NIH Diabetes Prevention Trial-1 was initiated, but this study suggested a beneficial effect (i.e., delay in developing disease) in only a small subgroup of individuals at high risk for T1D and treated with oral insulin (whole molecule); no beneficial influence was seen with prophylactic daily subcutaneous administration of this antigen (Skylar et al. 2005). In humans, the B9-23 epitope does not appear to be the dominant insulin epitope

(Skowera et al. 2008; Dromey et al. 2011). Moreover, using NOD mice deficient in mouse MHC class I and expressing human HLA-A2, it has been observed that a primary initiating autoantigenic epitope appears to be IGRP_{265–273} (Niens et al. 2011). Another key difference between the study of T1D pathogenesis in rodents and humans is their environment. Mice and rats in most animal facilities are housed under specific pathogen-free (SPF) conditions in which they are not exposed to pathogenic infectious agents. In contrast, humans are not only exposed to infectious agents, they are also immunized to these agents, leading to the generation of effector/memory immune responses that generate immune systems that may cross-react with islet autoantigens (Saruger et al. 2001; Tong et al. 2002; Chou et al. 2004).

Mice and humans also differ in their balance of leukocyte subsets, defensins, inducible nitric oxide synthase (NOS), γ/δ T cells, and natural killer (NK) inhibitory receptor families, cells, and molecules that are important players in immune and autoimmune responses (Mestas and Hughes 2001). The T-cell signaling pathways and costimulatory molecule expression of humans and rodents also shows many differences, as does their cytokine and chemokine receptor expression. Of particular interest are differences in the innate immune responses that are known to play a critical role in the generation of an adaptive immune response. For example, following IV injection of lipopolysaccharide, humans are as much as 100,000-fold more susceptible to endotoxin shock than mice (Munford 2010). In addition, the Toll-like receptor (TLR) repertoires that control innate immune responses also differ substantially. TLR11 is expressed in mice but not in humans; indeed, this gene has stop codons that prevent its expression. Conversely, TLR10 is expressed in humans, but in mice this TLR is a pseudogene. With this setting, the community of translational investigators is in great need of improved small animal models that would more effectively identify human-specific agents that will prove successful in human T1D (Greenbaum and Atkinson 2011).

SCID Mice—A Paradigm Shift in T1D Research

As described above, progress in understanding the pathogenesis, as well as analyses of the efficacy of therapeutics for T1D, has been impeded by considerable differences of rodent models and human disease. This is compounded by our inability to analyze in vivo the interaction of diabetogenic human immune cells with human islets without placing individuals at risk (Atkinson et al. 2000). One potential approach to overcome these limitations is to study this interaction in vivo by engraftment of human immune cells and islets into immunodeficient mice.

Engraftment of Human Immune Systems in Genetically Modified SCID Mice

Progress in development of effective small animal models to facilitate in vivo studies of human adaptive immune responses followed the discovery of the severe combined immunodeficiency (*Prkdc^{scid}*, abbreviated as *scid*) mutation in a colony of CB17 mice (Bosma et al. 1983). However, the utility of the CB17-*scid* model, particularly for studies of human immunity and autoimmune disease such as T1D, is limited by low levels of human hematolymphoid cell engraftment owing to host adaptive and innate immune factors, including elevated NK cell activity (Bosma et al. 1988; Shultz et al. 1995; Peled et al. 1999). Improved engraftment has been achieved in NOD-*scid* mice (Greiner et al. 1998), but these mice still express detectable NK activity and innate immunity that impedes human hematolymphoid cell engraftment. To improve human hematolymphoid cell engraftment, differentiation, and immune function in immunodeficient mice, two general approaches have been pursued: (1) additional genetic modification of the host and (2) development of new protocols for engraftment.

GENETIC MODIFICATION: INTERLEUKIN-2 RECEPTOR COMMON γ CHAIN NULL IMMUNODEFICIENT MURINE HOSTS

An alternative method for decreasing innate immunity in *scid*, *Rag1^{null}*, or *Rag2^{null}* immunodeficient mice involves genetic intercrossing

M.A. Brehm et al.

with mice carrying a targeted mutation at the IL-2 γ common chain (Shultz et al. 2007). Deficiency of IL-2 receptor γ chain causes X-linked SCID in humans (Uribe and Weinberg 1998). This molecule is indispensable for IL2, IL4, IL7, IL9, IL15, and IL21 high-affinity binding and signaling (Sugamura et al. 1996).

IL-2 γ common chain deficiency completely blocks NK cell development and causes additional defects in innate immunity (Shultz et al. 2007). Research groups have independently produced targeted IL-2 receptor γ chain mutations (Cao et al. 1995; DiSanto et al. 1995; Ohbo et al. 1996). Ito et al. (2002) backcrossed a truncated *IL2r γ* mutation onto the NOD/Shi-*scid* strain. Adult NOD/Shi-*scid* *IL2r γ ^{null}* mice injected IV with human CD34⁺ umbilical cord blood (UCB) cells results in the generation of a functional human immune system (Ito et al. 2002; Yahata et al. 2002). Traggiai et al. (2004) engrafted BALB/c-*Rag2^{null}* *IL2r γ ^{null}* mice generated using an *IL2r γ ^{null}* truncated knockout originally made by Ito et al. (2002), and injection of newborn mice with human hematopoietic stem cells (HSC) generated a functional adaptive immune system (Traggiai et al. 2004). Our laboratories have generated NOD/Lt-*scid* *IL2r γ ^{null}* (NSG) mice using a complete null mutation of the *IL2r γ* gene and have documented that both adult and newborn mice engrafted with human HSC generate functional human immune systems (Ishikawa et al. 2005; Shultz et al. 2005, 2007; Brehm et al. 2010b).

The NSG Mouse

Focusing our efforts on NSG mice (Shultz et al. 2007), we have shown that these mice are superior to all other stocks of *IL2r γ ^{null}* mice in their ability to support engraftment with a functional human immune system (Shultz et al. 2007; Pearson et al. 2008a; Brehm et al. 2010b;). Because of the ability of NSG mice to engraft at high levels with mature human lymphocytes and HSC, NSG mice are ideal hosts for investigating the in vivo function of human immune and autoimmune systems. These preclinical models can also be used to identify mechanisms by which therapeutic interventions for T1D mediate their effects.

A number of human immune engraftment models have been developed that can be used to study human immune cell function, including the Hu-PBL-SCID, Hu-SRC-SCID, and the BLT (i.e., fetal hematopoietic stem cell, liver, thymus) models (Table 1). The Hu-PBL-SCID model is based on injection of human peripheral blood leukocytes (PBL) or splenocytes into NSG mice and is used for the analyses of mature immune and autoreactive T cells and the effector arm of the immune system. This model has been used to examine both alloimmunity and autoimmunity as well as viral immunity (Shultz et al. 2007). The Hu-SRC-SCID model is based on the engraftment of HSC. This model develops functional innate and adaptive human immune systems, including all of the cells of

Table 1. Models of human immune systems in immunodeficient mice

Model	Characteristics	Uses
Hu-PBL-SCID	Immunodeficient mice engrafted with human PBMCs	Study of human-specific infectious agents, alloimmunity and autoimmunity in vivo
Hu-SRC-SCID	Immunodeficient mice engrafted with human HSCs	Study of human HSC and development of a functional human immune system
SCID-hu (BLT)	Immunodeficient mice engrafted under the renal capsule with human fetal liver/thymus and injected IV with fetal liver HSC	Study of human HSC, T-cell development and intrathymic selection and development of a functional human immune system

Abbreviations: PBL, peripheral blood leukocytes; PBMCs, peripheral blood mononuclear cell cultures; SRC, scid-repopulating cell; BLT, bone marrow/liver/thymus; HSCs, hematopoietic stem cells.

hematopoietic lineage, and has been used to study many aspects of human immunobiology (Shultz et al. 2007; Manz and Di Santo 2009). In the BLT model, mice are engrafted under the renal capsule with fragments of human fetal liver and thymus, and injected intravenously with human HSC derived from the liver of the same donor tissue (McCune et al. 1988; Nami-kawa et al. 1988; Melkus et al. 2006; Sun et al. 2007; Denton et al. 2008; Wege et al. 2008; Denton and Garcia 2009). These mice also develop functional innate and adaptive human immune systems, but in this model human T cells are educated on autologous human thymic epithelium, leading to robust HLA-restricted human immunity.

Models to Study Human Islet Function and Allo- and Autoimmunity

NSG Normoglycemic Mice

One simple model to study the function of human islets in the absence of potential effects of glucose toxicity encountered following transplantation of human islets into hyperglycemic mice involves the transplantation of human islets into normoglycemic NSG mice. We have observed that we can evaluate human islet function by their ability to secrete human insulin and C-peptide following glucose administration (unpublished observations).

The NSG model system will allow the analyses of development and function of reprogrammed, ES (embryonic stem) and induced pluripotent stem (iPS) -derived β cells in a normoglycemic environment. It should also be noted that euglycemia in a mouse is ~ 120 – 160 mg/dL, whereas the “set point” in humans is ~ 80 – 100 mg/dL, essentially exposing the human β cells to chronic low hyperglycemic stimulation even in a “normoglycemic” mouse host. Despite this, there are numerous advantages for the use of this model system. These include (1) normoglycemic NSG mice are readily available in essentially unlimited numbers, (2) transplanted cells are not exposed to high levels of glucose, avoiding the potential confounding effects of glucose toxicity, (3) fewer

cells are required for functional analyses than are required for regulation of hyperglycemia in diabetic recipients, and (4) this model has been shown to successfully support the engraftment of mouse and human islets. Finally, (5) we have published a “gold standard” protocol for how to establish human immune systems in these immunodeficient mice (Pearson et al. 2008a), permitting investigation of the function of human islets or β cells in the presence of alloreactive and, in the future, autoreactive immune systems. The disadvantages of this model include the inability to determine if the engrafted islets or β cells can function properly with respect to their ability to regulate hyperglycemia in vivo.

Chemically Induced Hyperglycemic NSG Mice

NSG mice have been induced to become hyperglycemic by the injection of streptozotocin (STZ) (King et al. 2008; Pearson et al. 2008b). Transplantation of human or mouse islets, or dissociated mouse islet insulin-positive cells can restore normoglycemia. These mice can also be engrafted with functional human immune systems following injection of human peripheral blood mononuclear cells (PBMC, the Hu-PBL-SCID model) (King et al. 2009; Pino et al. 2009) or human HSC (the Hu-SRC-SCID model) (Shultz et al. 2005; Giassi et al. 2008; Jaiswal et al. 2009; Brehm et al. 2010a). Data validating the ability of the NSG mouse model to serve as recipients of human islets and human immune systems has been published (Shultz et al. 2007). Advantages for the use of this model system include (1) readily available NSG mice in essentially unlimited numbers, (2) the ability to induce hyperglycemia at will, (3) human and mouse islets and dissociated mouse islet cells (single-cell suspensions of insulin-positive cells) can restore normoglycemia, and (4) the NSG mice can be engrafted with a functional human immune system. The disadvantages of this model are the inconsistent induction of hyperglycemia in STZ-treated mice, the potential for “reversion” of the hyperglycemia by recovery of the endogenous mouse islets, and toxicity of STZ.

M.A. Brehm et al.

Monogenic Models of Hyperglycemia

The use of chemical toxins for the induction of hyperglycemia has a number of drawbacks as described above. To address this, a number of monogenic mouse models of hyperglycemia have been described. These include, among others, the *Ins2^{Akita}* (Akita), the *Eif2ak3* (PERK) knockout, the *Fxn* (frataxin) and *Tfam* conditional knockouts, and the *Ncb5or* (NCB5OR knockout) strains (Table 2). Our laboratory has focused on the use of the *Ins2^{Akita}* model of spontaneous diabetes as a foundation strain

for the generation of a monogenic model of genetically induced spontaneous hyperglycemia.

NOD-Rag1^{null} Prf1^{null} Ins2^{Akita} Mouse Model

The dominant mutation in the murine insulin 2 gene, termed *Ins2^{Akita}*, results in spontaneous nonimmune mediated hyperglycemia (Yoshioka et al. 1997; Mathews et al. 2002). This spontaneously arising mutation replaces a cysteine at position 96 with tyrosine, and disrupts a disulfide linkage required for proper folding of

Table 2. Monogenic mouse models of insulin-dependent diabetes

Mouse model	Gene symbol	Phenotype	Human disease	References
AKITA Spontaneous mutation in insulin-2 gene	<i>Ins2^{Akita}</i>	Spontaneous insulin-dependent diabetes	Permanent neonatal/infancy-onset diabetes mellitus (PNDM)	Yoshioka et al. 1997; Wang et al. 1999; Colombo et al. 2008
PERK Phosphorylation of eukaryotic translation initiation factor 2 α kinase 3 knockout	<i>Eif2ak3</i>	PERK knockout mice show progressive loss of $\alpha\beta$ cells as well as loss in pancreatic acinar cell viability	Walcott-Rallison syndrome	Shi et al. 1998; Harding et al. 1999; Zhang et al. 2002; Cavener et al. 2010; Julier and Nicolino 2010
FRATAXIN Nuclear-encoded protein localized at the mitochondrial matrix conditional knockout	<i>Fxn</i>	Conditional disruption in β cells results in diabetes owing to loss of β -cell mass	Freidrich's ataxia	Cossee et al. 2000; Ristow et al. 2000, 2003; Pandolfo 2003; Gucev et al. 2009
TFAM Mitochondrial transcription factor A Nuclear-encoded mitochondrial protein Conditional knockout NCB50r	<i>Tfam</i>	Conditional disruption in β cells results in impaired insulin secretion and hyperglycemia by 5 wk of age	Mitochondrial diabetes	Larsson et al. 1998; Silva et al. 2000; Maassen et al. 2002; Falkenberg et al. 2007
NCB5OR Cytochrome B5 reductase 4 Soluble flavoheme NAD(P)H reductase localized in the ER protects β cells from oxidative stress	<i>Cyb5r4</i> Old symbol (<i>Ncb5or</i>)	<i>Cyb5r4</i> knockout mice develop severe hyperglycemia by 7 wk of age accompanied by hepatic lipid metabolic abnormalities	No reported association with diabetes in humans	Andersen et al. 2004; Xie et al. 2004; Xu et al. 2011

Abbreviation: ER, endoplasmic reticulum.

insulin. Improper folding of the nascent insulin 2 molecule, induction of the unfolded protein response, and finally apoptosis of β cells leads to diabetes and permanent hyperglycemia (Ron 2002; Izumi et al. 2003). We generated a strain of immunodeficient spontaneously hyperglycemic mice, the NOD-*Rag1*^{null} *Prf1*^{null} *Ins2*^{Akita} strain. The NOD-*Rag1*^{null} *Prf1*^{null} *Ins2*^{Akita} strain was based on the previously described NOD-*Rag1*^{null} *Prf1*^{null} strain that was shown to accept human islet transplants and allowed for allograft rejection following engraftment with human PBMC (Banuelos et al. 2004). NOD-*Rag1*^{null} *Prf1*^{null} *Ins2*^{Akita} mice develop spontaneous hyperglycemia, similar to *Ins2*^{Akita}-harboring strains of immunocompetent mice (Pearson et al. 2008b). There was no mononuclear cell infiltration into the islets that had spontaneously lost β cells, and human islets transplanted into diabetic NOD-*Rag1*^{null} *Prf1*^{null} *Ins2*^{Akita} mice resulted in a return to euglycemia.

Immunodeficient NOD mice harboring the *Ins2*^{Akita} mutation are an appealing host for human islet transplants and for human β stem and progenitor cells when concerns about drug-induced hyperglycemia are encountered. Advantages of this model are (1) spontaneous development of hyperglycemia in the absence of administration of toxic drugs, (2) consistent and severe hyperglycemia, (3) no “reversion” of the hyperglycemia by recovery of the endogenous mouse islets, and (4) no need for administration of exogenous insulin to prevent the development of metabolic decompensation and death. Disadvantages of this model include the inability of NOD-*Rag1*^{null} *Prf1*^{null} *Ins2*^{+/Akita} mice to support consistent levels of human PBMC or HSC engraftment and the inability of HSC engraftment to generate a fully functional human immune system, including T cells (Shultz et al. 2003; Banuelos et al. 2004; Minamiguchi et al. 2005).

NOD-*Rag1*^{null} *IL-2r γ* ^{null} *Ins2*^{Akita} Mouse Model

The NOD-*Rag1*^{null} *IL-2r γ* ^{null} *Ins2*^{Akita} (NRG-Akita) mouse model of spontaneous hyperglycemia has recently been described (Brehm et al.

2010a), and these mice become hyperglycemic with the same kinetics as do NOD-*Rag1*^{null} *Prf1*^{null} *Ins2*^{Akita} mice (Pearson et al. 2008b). Furthermore, human islets can also restore normoglycemia in diabetic NRG-Akita mice. A major advantage of these mice is that they can be engrafted with a fully functional human immune system (Brehm et al. 2010b) that can reject transplanted allogeneic human islets (Brehm et al. 2010a). This model has all the advantages of the NOD-*Rag1*^{null} *Prf1*^{null} *Ins2*^{Akita} model plus the ability to support engraftment with functional human immune systems so that alloimmunity and autoimmunity can be studied in these mice.

New Models of Hyperglycemia under Development

***NOD-Scid IL-2r γ* ^{null} *Tg(Ins-rtTA)* *Tg(TET-DTA)* Strain of Mice**

This mouse model developed by Drs. Nir, Dor, and Melton expresses diphtheria toxin A (DTA) in β cells following addition of doxycycline to the drinking water (Nir et al. 2007). Expression of a single DTA molecule is toxic to mouse β cells, and the mice have been shown to remain hyperglycemic throughout the period of doxycycline administration. Following removal of doxycycline, the β cells proliferate and restore normoglycemia (Nir et al. 2007). The generation of these mice by speed congenic backcrossing of the *Ins-rtTA* and *TET-DTA* transgenes to the NSG strain is currently under way.

***NOD-Scid IL-2r γ* ^{null} *Tg(RIP-HuDTR)* Strain of Mice**

The *Tg(RIP-HuDTR)* model system developed by Dr. Herrera (Thorel et al. 2010) is currently being backcrossed onto the NSG strain. Both X-linked and autosomal versions of the model are under development. In the X-linked model system, administration of diphtheria toxin to hemizygous female mice will lead to a loss of 50% of β cells, whereas administration of diphtheria toxin to the autosomal model will result in up to 99% elimination of the β -cell mass

M.A. Brehm et al.

(Thorel et al. 2010). These models will permit timing of the induction of hyperglycemia to be precisely controlled.

Use of Immunodeficient Mice for the In Vivo Study of Human β -Cell Proliferation

It is well known that mouse β cells can be induced to proliferation in the presence of hyperglycemia (Schuit and Drucker 2008), but it is unknown whether human β cells will also experience proliferation under similar stimulation. Based on the ability of immunodeficient mice to accept human islet grafts, two groups have recently used these model systems to study the ability of hyperglycemia to induce human β -cell proliferation in vivo. In one study, human islets were transplanted into STZ-diabetic NOD-*scid* mice, and the mice were injected with BrdU (Levitt et al. 2011). In the second study, human islets were transplanted into diabetic NRG-Akita mice (DiIorio et al. 2011). In both studies, human β cells were observed to increase their proliferation rates in a hyperglycemic environment, but the rate of proliferation was much lower than that observed in mouse islets subjected to similar hyperglycemic conditions (Schuit and Drucker 2008). These model systems can now be used for the study of human β -cell responses to experimental drugs and to human immune and autoimmune systems.

Models for the Study of Human T1D

Hu-PBL-SCID Model

One approach for the study of human T1D in immunodeficient mice is the use of the Hu-PBL-SCID model. This model has historically been used to study multiple types of autoimmunity (Tighe et al. 1990; Martin et al. 1992; Davis et al. 2002), including T1D (Petersen et al. 1993). Although the autoantibodies to islet components were detected following PBL transfer from individuals with T1D, no infiltration or β -cell destruction was observed (Petersen et al. 1993). More recently, the development of human T-cell clones with specificities for islet autoantigens has permitted the study of adop-

tive transfer of diabetes into NOD-*scid* mice. In these studies, infiltration, but not islet cell destruction was detected (van Halteren et al. 2005). The availability of newer strains of immunodeficient mice based on the *IL2r γ ^{null}* mutation and transgenic expression of human HLA molecules has provided models that now appear to permit the direct study of human autoreactive T cells in vivo. A recent manuscript has used NSG-HLA-A2 mice as recipients of HLA-A2 PBMC from T1D and non-T1D donors, and reported that in the Hu-PBL-SCID model system, peripheral blood lymphocytes from T1D patients preferentially infiltrated the islets of NSG-HLA-A2 recipients (Whitfield-Larry et al. 2011). Using splenocytes from T1D donors obtained through the nPOD program supported by JDRF, we have obtained similar results in that splenocytes from HLA-A2 T1D donors preferentially induce insulinitis in NSG-HLA-A2 recipients (unpublished observations).

Hu-SRC-SCID Model

Although the Hu-PBL-SCID model enables in vivo analyses of functional mature human T cells, other cell lineages fail to engraft efficiently following injection of PBMC (Shultz et al. 2007). In rodent models of T1D, other immune cell lineages, including macrophages, NK cells, NKT cells, dendritic cells, and B cells have been implicated in the pathogenesis of T1D (Von and Nepom 2009). These additional human cell lineages can be generated in immunodeficient mice by engraftment of HSC. However, to engraft HSC from a donor with a genetic susceptibility for T1D requires the recovery of HSC from T1D donors via bone marrow biopsies or mobilization of HSC into the peripheral blood following G-CSF treatment. Alternatively, in the virus research community, TCRs from virus-specific human T-cell clones have been generated and lentivirus constructs containing these TCRs have been used to transduce human HSC that are subsequently transplanted into immunodeficient mice. For example, HIV-specific TCR constructs were used to transduce HSC ("retrogenic" approach), and these engrafted HSCs generated mature HLA-restricted

cytotoxic CD8 T cells expressing this “transgenic” TCR (Kitchen et al. 2009). A large number of human islet autoantigenic epitopes have been identified (Di Lorenzo et al. 2007). In addition, autoreactive T-cell clones are now becoming available. These tools now permit the TCR “retrogenic” approach used successfully for the study of HIV to be applied to the study of T1D.

iPS Cells for the Study of Human T1D

Recent breakthroughs in the ability to “reprogram” cells into pluripotent stem cells that can be used via directed differentiation to generate all cell lineages have been reported (Cohen and Melton 2011). Various groups are now generating iPS cells derived from T1D donors, and are pursuing the directed differentiation of these cells into β cells, HSC, and thymic epithelium. If successful, this approach would permit T1D genetically susceptible HSC to be educated on autologous thymic epithelium with autologous β cells as targets of the autoimmune response. In rodent models, it is well known that HSC can transfer diabetes to adoptive recipients (Greiner et al. 2001; Von and Nepom 2009), and isolated case reports have suggested this is also true for humans (Lampeter et al. 1998, 1993; Mellouli et al. 2009). This system would allow the analysis of the relative importance of different diabetic genotypes, risk loci, and environmental stimuli on the etiology of T1D, and may thus identify novel points for therapeutic intervention.

CONCLUDING REMARKS

Despite many decades of studying rodent models of T1D, we have yet to translate therapies that prevent or cure T1D in rodents to the successful prevention or cure of T1D in humans. This is in part owing to the fact that rodent and human immune systems, as well as their islets, differ significantly in terms of cell composition, function, and gene expression. These differences form the basis for our need to understand interactions of human autoimmune systems with human β cells to allow successful translation of emerging findings from rodent biology to humans. The use of novel immunodeficient

Animal Models of Human Type 1 Diabetes

mouse models engrafted with functional human immune systems and islets is now providing the tools to investigate specific mechanisms by which alloimmune and diabetogenic human immune cells attack human β cells and how β cells respond to immune attack. Findings from this approach can guide the development of novel strategies to prevent and cure T1D. This information, because it is based on human cells, tissues, and immune systems, has the potential to translate directly into information that will identify targets for therapeutic intervention, guide clinical trials, and ultimately transform our understanding of human T1D.

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M.A. Brehm et al.

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M.A. Brehm et al.

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