Type 1 diabetes is characterized by destruction of insulin-producing β cells in the pancreatic islets by effector T cells. Tregs, defined by the markers CD4 and FoxP3, regulate immune responses by suppressing effector T cells and are recruited to sites of action by the chemokine CCL22. Here, we demonstrate that production of CCL22 in islets after intrapancreatic duct injection of double-stranded adeno-associated virus encoding CCL22 recruits endogenous Tregs to the islets and confers long-term protection from autoimmune diabetes in NOD mice. In addition, adenoviral expression of CCL22 in syngeneic islet transplants in diabetic NOD recipients prevented β cell destruction by autoreactive T cells and thereby delayed recurrence of diabetes. CCL22 expression increased the frequency of Tregs, produced higher levels of TGF-β in the CD4+ T cell population near islets, and decreased the frequency of circulating autoreactive CD8+ T cells and CD8+ IFN-γ-producing T cells. The protective effect of CCL22 was abrogated by depletion of Tregs with a CD25-specific antibody. Our results indicate that expression of CCL22 recruits Tregs and attenuates autoimmune destruction of β cells. CCL22-mediated recruitment of Tregs to islets may be a novel therapeutic strategy for type 1 diabetes.

Introduction

The complex pathogenesis of autoimmune diabetes in NOD mice and humans involves multiple immune components. The autoimmune response recurs when β cells are transplanted into type 1 diabetic patients or NOD mice (1, 2). Tregs are primary mediators of peripheral self-tolerance and are able to suppress the proliferative and cytokine responses of other immune cells (3). Defects in Treg number and/or function may underlie spontaneous development of autoimmune diseases (4). The chemokine receptor CCR4 is highly expressed on Tregs and mediates recruitment of Tregs in vivo by its ligand, CCL22 (5). Recruitment of Tregs to human cancers producing CCL22 suppresses tumor-specific T cell immunity and contributes to tumor growth (6, 7). To determine whether CCL22-mediated recruitment of Tregs could counter the autoimmune response to β cells in diabetes, we overexpressed CCL22 in β cells in vivo in NOD mice and in syngeneic islet transplants.

Results and Discussion

CCL22 protects pancreatic β cells from autoimmune attack in type 1 diabetes. To drive islet-specific expression of CCL22, we created double-stranded adeno-associated virus serotype 8 (dsAAV8) vectors containing mouse CCL22 or GFP cDNA, driven by the rat insulin promoter (dsAAV8-RIP/CCL22 and dsAAV8-RIP/GFP, respectively). One month after injection of 1.5 × 1011 viral genomes (vg) via the pancreatic duct, CCL22 secretion was detected in islets isolated from dsAAV8-RIP/CCL22–infected mice but not controls. Infection did not impact β cell function, since immune-deficient (NOD/SCID) mice showed normal glucose tolerance at 2 and 4 months after injection with dsAAV8-RIP/GFP (data not shown).

To determine whether β cell–specific expression of CCL22 can delay or prevent diabetes, 8-week-old NOD mice were injected intraductally with dsAAV8-RIP/CCL22, dsAAV8-RIP/GFP, or PBS. NOD mice transduced with dsAAV8-RIP/CCL22 (CCL22-NOD mice) were protected from diabetes, with 9 out of 11 mice remaining normoglycemic for up to 160 days. All NOD mice transduced with dsAAV8-RIP/GFP (6 out of 6 mice) or injected with PBS (PBS-NOD mice) (7 out of 7 mice) developed diabetes within the first 100 days (Figure 1A). CCL22-NOD mice maintained normal glucose tolerance (Figure 1B) and circulating insulin levels (data not shown). Despite extensive insulitis in both CCL22-NOD and PBS-NOD mice (Figure 1C), CCL22 expression was associated with preservation of insulin-positive cells at 15 days, 1 month, and 3 months after intraductal injection.

CCL22 enhances influx of Tregs to islets and limits islet autoimmunity. By immunostaining, we observed an apparent increase in the number of FoxP3+ cells in islets of CCL22-NOD mice at 15 days, 1 month, and 3 months after injection (Supplemental Figure 1; supplemental material available online with this article; doi:10.1172/JCI43048DS1). Flow cytometric analysis showed that the frequency of FoxP3+ cells in pancreata of CCL22-NOD mice was significantly elevated by 1 month after injection and more markedly elevated at 3 months (Figure 2A). At 1 month after injection, the absolute number of CD4+FoxP3+ cells was increased in the pancreas (Figure 2B) but not in the pancreatic lymph node (PLN) (data not shown). No significant differences were observed in the absolute numbers of CD4+ or CD8+ effector cells in either the pancreas or the PLN (data not shown). The Treg/CD4 effector T cell ratio was significantly elevated in PLNs of CCL22-NOD mice and showed a nonsignificant increase in pancreas at 1 month after
injection (Supplemental Figure 2). Administration of Treg-depleting (anti-CD25) antibody to CCL22-NOD mice reversed protection from diabetes (Figure 2C and Supplemental Figure 3, A and B). CCR4 expression on Tregs in the PLNs of CCL22-NOD mice appeared higher than that in PBS control mice by 1 month after injection (Supplemental Figure 4A), although this difference was not statistically significant. These data suggest that interaction of CCL22 with CCR4 contributes to the trafficking of Tregs to both CCL22-expressing islets and the PLNs.

CD4\(^+\) cells in the PLN and pancreas of CCL22-NOD mice produced more TGF-\(\beta\) (Figure 2D), suggesting that local CCL22 expression may suppress autoimmunity in part by increasing TGF-\(\beta\) production. CCL22-NOD mice did not differ significantly from PBS-NOD controls in the proportion of CD8\(^+\) cells in pancreatic infiltrates at 1 month after injection (Supplemental Figure 4A), although this difference was not statistically significant. These data suggest that interaction of CCL22 with CCR4 contributes to the trafficking of Tregs to both CCL22-expressing islets and the PLNs.

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CCL22 protects from recurrent autoimmunity in NOD mice. To determine whether islet CCL22 expression prevents \(\beta\) cell loss in syngeneic islet transplants in NOD mice, we created an adenovirus (Ad-CCL22) in which expression of a mouse CCL22 cDNA is driven by the CMV promoter. Transduction with Ad-CCL22 (MOI of 10) resulted in islet expression of CCL22 without affecting viability or function in vitro (data not shown). Diabetic NOD female mice were transplanted with 500 syngeneic (NOD/SCID) islets transduced ex vivo with either Ad-CCL22 or Ad-LacZ or left untreated. Survival of syngeneic islet grafts in diabetic NOD recipients was markedly improved in mice transplanted with CCL22-expressing islets (Figure 3A). Protection from recurrent autoimmunity was temporary, as hyperglycemia returned in all recipients of CCL22-expressing grafts by 60 days after transplantation, likely due to gradual loss of adenoviral CCL22 expression (Supplemental Figure 5). Immunostaining revealed many insulin-producing \(\beta\) cells in CCL22-expressing grafts but virtually none in controls (Supplemental Figure 6A). Islet CCL22 expression appeared to similarly protect \(\alpha\) cells (Supplemental Figure 6A). Ad-CCL22–transduced grafts had a greater number of FoxP3\(^+\) cells than controls, as demonstrated by both flow cytometry (Figure 3B) and immunostaining (Supplemental Figure 6, B and C). Moreover, the ratio of Tregs to CD4\(^+\) effector cells was significantly greater in CCL22-expressing grafts (Figure 3C).
More Tregs than effector CD4+ cells were CCR4-positive in both CCL22- and LacZ-expressing grafts, and Tregs present in CCL22-expressing grafts expressed higher levels of CCR4 than Tregs in controls (Supplemental Figure 4B). IGRP-reactive T cells were found in similar frequencies in both groups (Figure 3D), indicating that CCL22 expression does not prevent migration of autoreactive CD8+ cells to the graft. Administration of anti-CD25 antibody reversed protection from recurrent autoimmunity in recipients of CCL22-expressing islet transplants (Figure 3E). These results suggest that CCL22-mediated recruitment of Tregs decreases graft failure and recurrence of autoimmune diabetes.

In summary, recruitment of Tregs to islets by CCL22 expression delays both spontaneous and recurrent autoimmune diabetes in NOD mice. Although insulitis clearly persists in the face of CCL22-mediated protection from diabetes, we speculate that CCL22 may not only enhance recruitment but also functionally activate Tregs that are optimally positioned for suppression of relevant targets (as suggested by increased CTLA-4 expression after CCL22 treatment; Supplemental Figure 7). CCL22 likely recruits CCR4-expressing effector cells, perhaps increasing the opportunity for Treg/effector cell interaction. Although islet CCL22 expression did not reverse disease in diabetic mice (data not shown), it will be of interest to assess the ability of other chemokines, such as CCL17, which also interacts with CCR4 (5), to prevent or cure diabetes. Recruitment of endogenous Tregs to the site of inflammation may obviate the need for expansion and infusion of cells for Treg-based diabetes therapies (11–13). Manipulating the migration of endogenous Tregs by gene- or peptide-delivery approaches may be a novel therapeutic strategy not only for diabetes but for other autoimmune diseases and organ transplantation.
Viral vector production and amplification. Recombinant dsAAV8 vectors were generated at the Children’s Hospital of Philadelphia by triple transfection protocol as previously described (14). The dsAAV8-RIP/GFP plasmid was provided by Paul Robbins (University of Pittsburgh, Pittsburgh, Pennsylvania, USA). CCL22 cDNA was from Invitrogen. To generate the adenoviral vector Ad-CCL22, pCMV/CCL22 cDNA was cloned in pShuttle plasmid. Adenovirus was produced by cotransfection of HEK293 cells with the CCL22 vector and the Adenovirus helper plasmid (ABM Good). Adenovirus was purified with the Vivapure Adenopack 100 Kit (VivaScience) and titrated by the end-point dilution method of tissue culture infective dose. Ad-LacZ was provided by R.G. Korneluk (Children’s Hospital of Eastern Ontario Research Institute, Ottawa, Ontario, Canada). The endotoxin content of the adenoviral vectors was less than 1 endotoxin units/ml (EU/ml).

Viral vector administration and islet transplantation. NOD and NOD/SCID mice were purchased from The Jackson Laboratory and housed in the Child and Family Research Institute animal facility. Animal studies were reviewed and approved by the University of British Columbia Committee on Animal Care. Injection into the pancreatic duct was performed as described previously (15). For islet transplantation, pancreatic islets were isolated from male NOD/SCID mice by ductal collagenase (Sigma-Aldrich) injection, digestion, and purification with the filtration method as described previously (16). Islets were hand picked and incubated in complete RPMI 1640 (Gibco) with Ad-CCL22 or Ad-LacZ (MOI of 10) overnight at 37°C. Islets were transplanted into the renal subcapsular space of recipient mice under isoflurane anaesthesia. Graft failure was determined when blood glucose was greater than 15 mM on 2 consecutive days. In some experiments, CD25 antibody (PC61; AdLab, University of British Columbia; 100 μg i.p.) or rat isotype control IgG (AbLab) was injected weekly after AAV8 administration or islet transplantation.

Islet function. In vivo islet function was assessed by i.p. glucose tolerance test. Mice were fasted for 5 hours and injected with D50 glucose (2 g/kg body weight; Baxter Healthcare), and tail blood glucose was measured at 0, 15, 30, 60, and 120 minutes.

CCL22 protein determination. CCL22 expression in islets was determined by Western blot. Ten micrograms of protein from islet lysates were resolved on a 20% SDS-polyacrylamide gel at 100 V. After transfer to Immobilon-P membranes (Millipore) at 100 V for 1 hour using the Trans-Blot System (Bio-Rad), immunoblot analysis was performed using polyclonal anti-CCL22 antibody (R&D Systems) and detected by chemiluminescence (Amersham). CCL22 protein levels were measured by ELISA (R&D Systems) following the manufacturer’s instructions.

Histological analysis of the pancreas. Pancreata were fixed in 4% paraformaldehyde overnight. Paraffin sections (5-μm thick) were immunostained using guinea pig anti-insulin and rabbit anti-glucagon antibodies (Dako) as well as goat anti–guinea pig Alexa Fluor 594 and goat anti-rabbit Alexa Fluor 594 (Molecular Probes) as secondary antibodies. Leukocyte immunostaining was performed after antigen retrieval by steaming for 30 minutes in Dako Target Retrieval Solution, using rat anti-
mouse CD45 antibody (BD Pharmingen) or anti-mouse FoxP3 antibody (eBioscience) and anti-rat Alexa Fluor 488 secondary antibody (AdLab, University of British Columbia). Images were captured using an Olympus BX61 microscope and In Vivo or DP software.

Flow cytometry. Pancreata were excised and digested with 1,000 U/ml of collagenase (Sigma-Aldrich) in Hank’s buffer solution (Life Technologies) for 20 minutes at 37°C. The resulting cell homogenate was filtered through a 40-μm filter basket and washed in PBS. Harvested grafts and lymph nodes were strained through a 40-μm filter and washed before incubation with antibodies. Monoclonal antibodies to CD8 (53-6.7) and CD4 (RM4-5) were purchased from BD Biosciences. Antibodies to FoxP3 (clones MF-14 and 150D) and CCR4 (clone 2G12) were obtained from BioLegend. Antibodies to IFN-γ (XMG1.2) and CTLA4 were from eBioscience, and antibody to anti-TGF-β (1D11) was from R&D Systems. Tetramers with IGRP286-294 or Insulin B31-39 epitopes were generated and provided by the Cellular Immunology Core facility at the Child and Family Research Institute. For intracellular staining, cells were fixed and permeabilized using the Cytofix/Cytoperm Kit (BD Biosciences). For intracellular cytokine staining, cells were stimulated in culture medium containing 100 ng/ml PMA, 1 μg/ml ionomycin, and 10 μg/ml brefeldin A (Sigma-Aldrich) at 37°C for 4 hours. Absolute cell number was determined by adding 50,000 counting beads (CountBright, Invitrogen) per sample before flow cytometry. A FACSCalibur cytometer (BD Biosciences) and CellQuest software (BD Biosciences) were used for flow cytometry, and data were analyzed using FlowJo 7.2.5 software.

Statistics. Graft survival curves were generated using Kaplan-Meier life-table analysis and were compared using the log-rank test. Other statistical analyses were performed using Student’s 2-tailed t test. P < 0.05 was considered statistically significant. Data in bar graphs are represented as mean ± SEM.

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