Resident macrophages of pancreatic islets have a seminal role in the initiation of autoimmune diabetes of NOD mice

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Treatment of C57BL/6 or NOD mice with a monoclonal antibody to the CSF-1 receptor resulted in depletion of the resident macrophages of pancreatic islets of Langerhans that lasted for several weeks. Depletion of macrophages in C57BL/6 mice did not affect multiple parameters of islet function, including glucose response, insulin content, and transcriptional profile. In NOD mice depleted of islet-resident macrophages starting at 3 wk of age, several changes occurred: (i) the early entrance of CD4 T cells and dendritic cells into pancreatic islets was reduced, (ii) presentation of insulin epitopes by dispersed islet cells to T cells was impaired, and (iii) development of autoimmune diabetes was significantly reduced. Treatment of NOD mice starting at 10 wk of age, when the autoimmune process has progressed, also significantly reduced the incidence of diabetes. Despite the absence of diabetes, NOD mice treated with anti-CSF-1 receptor starting at 3 or 10 wk of age still contained variably elevated leukocytic infiltrates in their islets when examined at 20–40 wk of age. Diabetes occurred in the anti–CSF-1 receptor protected mice after treatment with a blocking antibody directed against PD-1. We conclude that treatment of NOD mice with an antibody against CSF-1 receptor reduced diabetes incidence and led to the development of a regulatory pathway that controlled autoimmune progression.

Significance

Our studies indicate that the resident macrophages of the pancreatic islets of Langerhans have a seminal role in the initiation and progression of autoimmune diabetes in NOD mice. In this study, islet macrophages were depleted by administration of a monoclonal antibody to the CSF-1 receptor. Macrophage depletion, either at the start of the autoimmune process or when diabetogenesis is active, leads to a significant reduction in diabetes incidence. Depletion of the islet macrophages reduces the entrance of T cells into islets and results in the absence of antigen presentation. Concordantly, a regulatory pathway develops that controls diabetes progression. We conclude that treatments that target the islet macrophages may have important clinical relevance for the control of autoimmune type 1 diabetes.

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chemokine receptors (6). They are the sole hematopoietic cell within the islets before 3 wk of age and are able to present unconventional insulin epitopes to autoreactive T cells (6). Altogether, this posits the intraislet macrophage as a key regulator of lymphocyte entry into the islet. We have eliminated them using a monoclonal antibody to the CSF-1 receptor (CSF-1R), AF98 (11). This monoclonal antibody, and the M279 antibody against CSF-1R, when administered in vivo resulted in the loss of macrophages from many tissues (12–14). The extent of depletion depended on the amount and frequency of administration (12). Although treatment with AF98 resulted in the loss of resident macrophages, blood monocyte counts and inflammatory responses were not affected (13). We present evidence that the anti–CSF-1R antibody AF98 depleted the islet resident macrophages in two strains of mice: the normal C57BL/6 (B6) and the diabetogenic NOD. The depletion of macrophages did not affect the metabolism or islet transcriptome of the mouse. However, diabetes progression was severely blunted.

Results

Evaluation of C57BL/6 Mice. First, we evaluated the non-diabetic B6 strain, following treatment with the monoclonal antibody against CSF-1R. Injecting B6 mice with 0.25, 0.50, and 2.0 mg of AF98 led to a dose-dependent elimination of the intraislet macrophages (Fig. 1). This reduction was observed as early as 7 d after treatment and was equally evident at day 14. The depletion was long lived. A 2.0-mg dose of AF98 resulted in elimination of the islet macrophage lasting for at least 6 wk (Fig.

Fig. 1. Treatment of C57BL6 mice with AF98 antibody depleted their macrophages. (A) Female B6 mice aged 6–8 wk were administered 0.25, 0.50, or 2.0 mg of AF98 antibody i.p. Islets were examined 7 and 14 d after injection for the presence of macrophages. Box indicates CD45+CD11c+MHCII+F4/80+CD11b+ cells as a percent of total islet cellularity. (B) Female B6 mice aged 6–8 wk were treated with 2.0 mg of AF98, and their islets were examined at the time points indicated for the presence of macrophages. Top show the CD45+ cells, and Bottom show the CD45+CD11c+MHCII+F4/80+CD11b+ cells as a percent of total islet cellularity. (C) Graph of the CD45+ cells found in mice 1–2 wk after treatment with 2.0 mg of AF98. (D) Graph of CD45+F4/80+ CD64+ islet stromal macrophage populations as a percent of CD45+ cells in control and AF98-treated mice 2 wk after treatment. (E) Mice were treated for 1–2 wk with AF98 antibody, and the percent of total macrophages in lung, liver, spleen, and pancreatic lymph nodes were determined. (F) Macrophages were examined as in E except plots show the percent of the indicated subpopulation of macrophages. For all graphs, controls were untreated age-matched mice. Flow cytometry plots in A and B are representative of individual islet, lung, liver, spleen, and lymph nodes, from two to three experiments with two to four mice per treatment group. Scatter plots in E and F were calculated from three independent experiments with two to three mice per group. P values were calculated using Mann–Whitney U test with the following style: not significant (n.s., >0.1234), *P = 0.0332, **P = 0.0021, ***P = 0.0002, ****P < 0.0001.

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Macrophages started to return to the islets by 6 wk and were at normal levels by 7 wk after antibody treatment. The depletion of macrophages in the islets was consistent over multiple experiments (Fig. 1C). The macrophages of the stroma were also affected by the AFS98 treatment. The macrophage reduction was almost complete resulting in an ~85% reduction compared with control-treated mice (Fig. 1D and Fig. S1).

In addition to islets, several other tissues were examined. In most, a set of the resident macrophages was also affected 7–14 d after antibody administration. We analyzed total macrophages by expression of F4/80 and CD64 and calculated their reduction as shown in Fig. 1E. However, it was evident that the sensitivity to AFS98 treatment was variable (Fig. 1F). In the lung, alveolar macrophages were not affected, yet the CD11b+ interstitial macrophages were depleted (15) (Fig. 1E and F). In the liver, there are two macrophage populations defined by expression of F4/80. The F4/80hi Kupffer cells were reduced by about 50% (Fig. 1E and F). The F4/80int monocyte-derived macrophages were almost completely depleted by AFS98 treatment. This last finding suggest that these macrophages most likely represent the recently identified liver capsular macrophages (16). The spleen macrophages were not affected (Fig. 1E and F). However, in the pancreatic lymph node, a set of macrophages characterized by CD11b (similar in surface marker expression to the lung interstitial macrophages) were >80% reduced. Thus, not all tissue macrophages are as sensitive to anti-CSF-1R as the ones in the islets.

In summary, treatment with AFS98 led to variable effects on the macrophages residing in various tissues, while in islets, the effects were pronounced and prolonged. We also evaluated whether AFS98 treatment could affect the number of T and B cells and their differentiation in thymus and bone marrow. We found no impairment in their numbers and differentiation patterns in thymus or bone marrow (Fig. S2).

As seen in the op/op mouse, the absence of macrophages from birth can lead to systemic disruption of mouse homeostasis. Therefore, we evaluated two basic parameters of islet function after macrophage depletion: glucose tolerance and beta-cell insulin content. The glucose tolerance test measures the ability of the beta cell to sensor and respond to glucose, release insulin, and return the mouse to euglycemia. Following treatment of the B6 mice with the AFS98 antibody, neither glucose tolerance nor insulin content was affected (Fig. 2A and B). Glucose tolerance was unimpaired even after 6 wk of macrophage depletion in the islets (Fig. 2A). Measured pancreatic insulin content was stable at 3 wk after depletion (Fig. 2B).

A more global and unbiased measure of islet health is to evaluate the whole transcriptome. B6 mice were treated with 2.0 mg of AFS98 or control IgG2a at 3 wk of age, and whole islets were isolated at 6 wk of age. Comparison of AFS98 versus IgG2a-treated mice revealed 16 differentially expressed transcripts at a twofold change and 99% confidence interval in whole islets (Fig. 2C). These 16 changes were in transcripts known to be encoded strictly by the macrophage. These included transcripts previously reported to be up-regulated during diabetes progression (9).

Therefore, the majority of the islet transcriptome was not significantly affected by macrophage depletion, and the only change was the loss of the islet macrophage. In conclusion, homeostasis of islet function was not affected in a detectable manner when macrophages were depleted several weeks after birth.

**Fig. 2.** Islet function after depletion of macrophages by AFS98. (A) B6 mice were given 2.0 mg of AFS98 i.p. at 6–8 wk of age. Glucose tolerance assays were then performed on AFS98-treated and untreated mice. After the indicated number of days, the mice were fasted for 12 h and then injected with 2.0 g/kg glucose i.p. Blood glucose (mg/dL) was measured at the indicated time points. Results are pooled from two independent experiments (n = 2–3 mice per group). (B) Nine-week-old B6 females were left untreated or administered 2.0 mg of AFS98 i.p. Four days later, the mice were placed on a 20% sucrose diet for an additional 7 d, then returned to a normal diet for 2 d. The mice were then killed, and the insulin content of their islets was measured (n = 5 mice per group). (C) Three-week-old C57BL/6 mice were left untreated or administered 2.0 mg of AFS98 i.p. At 6 wk of age, their islets were isolated, total RNA was extracted, and transcripts were analyzed by microarray. Scatter plot shows the log2 mean expression values for four control and experimental mice. The dots highlighted in blue represent genes differentially expressed between treated and control mice at 99% confidence using moderated t test with Benjamini–Hochberg false discovery rate analysis. The selected genes are plotted in the heat map using Euclidean distance and normalized global expression as indicated.
Evaluation of NOD Mice.

Islets. Similar to the results observed in B6 mice, Fig. 3A shows that treatment with 0.5 or 2.0 mg of AFS98 depleted the islet macrophages in 4- to 5-wk-old NOD mice. Comparable results were obtained following treatment of NOD.Rag1−/− mice with the AFS98 antibody (Fig. S3). An analysis of NOD mice at 3–4 wk of age showed that the initial islet infiltrating T cells were all CD4+ and mostly in contact with the intraislet macrophage (9). The islet macrophages express CD11b, CD11c, and MHC-II highly on their surface. At this 3–4 wk of age period, there are very few XCR1+ DCs in islets (10).

We confirmed the colocalization of CD4 T cells with intraislet macrophages by examining islets using two-photon microscopy. NOD mouse islets were examined at 3–4 wk of age, the earliest age where one can identify the initial infiltrating T cells. Indeed, 27% of NOD islets had CD4+ T cells, 70% of which were in contact with the F4/80+ macrophages, confirming our initial studies. In the mice that were injected with 0.5 mg of anti-CSF-1R at 2 wk of age, the islets did not harbor any myeloid cells at 4 wk of age. In these mice, only 3% of islets had a detectable CD4+ T cell.

Next, we examined insulin peptide presentation by isolating the islets, dispersing the cells, and culturing them with insulin-reactive CD4 T cell hybridomas. Depletion of islet macrophages resulted in marked reduction of presentation to two different CD4 T cell hybridomas recognizing different MHC-II epitopes of insulin (17). The addition of the cognate peptides reflects the availability of MHC-II+ presenting cells. As noted in Fig. 3A, this addition did not lead to presentation in the islets of the treated mice, reflecting the paucity of presenting cells and the inability of any other islet cell to express MHC-II or present peptide. The same findings were reproduced in B6.g7 mice (Fig. S4).

Lymph node responses. While the islets from the macrophage-depleted mice were impaired in antigen presentation, this was not the case in the peripheral lymph nodes. This was determined using two approaches: carboxyfluorescein succinimidyl ester (CFSE) dilution of transferred T cells and recall response by ELISPOT. For CFSE dilution experiments, we transferred CD4+ (BDC2.5) or CD8+ (NY8.3) T cell clones into control or AFS98-treated NOD. BDC2.5 responds to a chromogranin peptide in the context of I-A^b (18), while the NY8.3 T cell divides in response to a peptide derived from IGRP, the islet-specific glucose-6-phosphatase–dependent catalytic subunit-related protein, in the context of H-2Kd (19, 20). Fig. 4A and B shows that both T cell clones proliferated in the draining pancreatic lymph node, but not in the distant inguinal lymph node. Next, to determine if depletion of the islet macrophage affected the trafficking of diabetogenic cells to the islets, we examine the entrance of TCR transgenic T cells into them. Both BDC2.5 and NY8.3 T cells entered islets of control mice but neither entered the islets from AFS98-treated mice (Fig. 4C). While the T cells entered and reacted to antigen presented in the pancreatic lymph node, this was not the case in the islets.

In the second approach, AFS-treated mice were immunized with various autoreactive peptides in the footpads and the T cell response was tested a week later; there was no impairment of the response (Fig. 5). Immunization with the insulin B9-23 peptide elicited an IL-2 and IFN-γ response to the peptide but not to the insulin protein, as reported before (17). Immunization with peptides from IGRP known to elicit CD4+ or CD8+ T cell responses was also unaffected by AFS98 treatment. The CD4+ or CD8+ T cell responses to the foreign protein hen egg lysozyme (HEL) were also unaffected.

In brief, examination by flow cytometry, direct microscopy of islets, antigen presentation assays, and T cell migration assays shows that the lack of macrophages translates into an absence of early CD4 T cell infiltration and antigen presentation capability of the islets. In contrast, the lymph nodes in the AFS-treated mice were active in antigen presentation.

Diabetes. Female NOD mice were followed for diabetes after treatment with 0.5 or 2.0 mg of AFS98 or control rat IgG2a antibody. In two experiments, mice were treated at 2–3 wk of age, a time when the diabetogenic process is starting in limited islets. The treatment was continued for several weeks using two different concentrations of antibody. In a third experiment, NOD mice were treated starting at 10 wk of age, a time when the diabetogenic process is active and most islets are already infiltrated by both CD4 and CD8 T cells. Both treatments led to a marked reduction in diabetes incidence when the mice were followed for 40 wk. The pooled results are shown in Fig. 6A. Fig. 6C shows the results of the individual experiments. Combining the three experiments, 4 of 40 AFS98-treated mice and 24 of 39 of the control mice became diabetic

At the end of the 40–42 wk of observation, two manipulations were performed on the AFS98-treated mice. First, the mice were administered an anti–PD-1 monoclonal antibody and this led to...
Fig. 4. AFS98 treatment does not affect T cell division in lymph nodes but prevents T cell entry into islets of Langerhans. NOD mice were injected with AFS98 antibody at a dose of 0.5 mg at 2 wk of age and 2.0 mg at 4 wk of age. Two TCR transgenic T cells, the CD4+ BDC2.5 and the CD8+ NY8.3, were isolated from lymph nodes and spleens of their respective mice. T cells were then labeled with CFSE and transferred into 6-wk-old NOD mice that had either been left untreated or treated with AFS98. (A and B) Seven days after T cell transfer, the pancreatic and inguinal lymph nodes were isolated and analyzed by flow cytometry. (A) Dilution of CFSE in either inguinal (Upper) or pancreatic (Lower) lymph nodes for an individual mouse per treatment is shown. Cells were gated on forward and side-scatter, CD45, CD3, and either CD4 (BDC2.5) or CD8 (NY8.3). (B) Summary of division index and proliferation index for individual mice examined as in A. Results show three or four individual mice per group. (C) Ten days after TCR transgenic T cell transfer, islets of Langerhans were isolated and examined for entry of either the BDC2.5 or NY8.3 T cells by flow cytometry. The Left two images are gated on forward and side-scatter, CD45, and CD3. The Right images are gated on forward and side-scatter. BDC T cells were identified using a clonotypic antibody to its T cell receptor. NY8.3 T cells were identified by a CD45.2 congenic label. Numbers indicate the percent of cells in each selection as a function of CD45+ cells. T cell islet entry results are representative of two to three independent experiments with two to four individual mice per group.
observed two phenotypes: 4 of 8 mice contained ~20–25% CD45+ cells in the islets, and 4 of 8 mice contained 2–5% CD45+ cells in the islets (Fig. 7C and D). (The IgG-treated control NOD mice contained ~60–70% CD45+ cells in islets.) Corroborating the flow cytometry ~50% of the pancreata examined had a degree of peri-insulitis (Fig. 7E). However, the other 50% of pancreata examined showed little to no insulitis (Fig. 7F). Therefore, macrophage depletion is protective in NOD mice long term, despite the infiltration or expansion of leukocytes in ~50% of the mice.

In mice treated with AFS98 starting at 10 wk of age, there also was a reduction in the number of leukocytes in islets at 22 wk of age. However, by 40 wk of age, all of the mice had comparable infiltrates to a 22-wk-old NOD mouse (Fig. 8). Thus, late macrophage depletion does not stop the infiltration of islets, despite preventing the development of hyperglycemia.

These results suggest that macrophage depletion can act before and after the diabeticogenic T cell pool in the NOD has fully developed. Early AFS98 treatment blocks diabetes by eliminating a key antigen-presenting cell in islets, preventing early entry of activated T cells, and limiting the expansion of the diabeticogenic T cells. In contrast, in late-treated NOD mice, there are active diabeticogenic T cells but disease is controlled by the development of a regulatory condition through at least one mechanism, PD-1/PD-L1 interaction. In both situations, we find islet infiltration with preservation of islet function.

**Discussion**

Injection of NOD mice with antibodies directed against CSF-1R depleted the islet resident macrophages while having a variable effect on macrophages of secondary lymphoid tissues and various other organs. The treatment had a profound effect in the development of diabetes. The manner in which the islet macrophage was eliminated by the antibody treatment is not clear, although it does not involve an acute inflammatory reaction (14).

The islet transcriptome analysis displayed no up-regulation of inflammatory transcripts following macrophage depletion (Fig. 2C). CSF-1R signaling is crucial for sustaining macrophage viability, which is most likely the reason for its loss following the antibody treatment (11).

There are two distinct scenarios that likely contributed to the attenuation of the active autoimmune process: (i) the impairment of (ii) the development function and (iii) the development of a regulatory process. Concerning the islets, these studies confirm the centrality of the resident macrophage in the pathophysiology of pancreatic islets. The islet macrophage arises during embryonal development. It is required for the early development of islets and during postnatal growth as is evident in the op/op mouse that lack CSF-1 (4, 5). However, in adult mice, islet function is not impaired when the macrophage is absent (our studies in B6 mice). We have not probed other possible roles for the islet macrophage such as maintaining vessel permeability, protecting against infectious agents, or modulating the physiology of beta cells. The islet macrophage is highly activated, as evidenced by their expression of TNF and IL-1 transcript and protein, and high expression levels of MHC-II (5, 10). In the context of autoimmunity, such as in NOD mice, autoreactive T cells escape thymic control and enter islets whereupon they establish long-lived contact with the macrophage (26).

The macrophage-depleted islet per se was impaired in its capacity to receive diabeticogenic T cells. In their absence, lymphocyte entry into islets was impaired. The immunofluorescence and two-photon imaging of isolated islets showed the majority of CD4 T cells in islets in contact with the macrophage. Indeed, previous analyses showed macrophage filipodia extending into the blood vessel lumens (8). I.v. injecting 0.5-μm latex beads coated with antibodies to I-Aαβ localized the beads to the macrophage/blood vessel interface. These findings point to the macrophage filopodia as the anatomical element that captures
The diabetogenic CD4 T cells, while the high-expression MHC-II may well be the key molecular substrate (8). In sum, the present and past studies point to the macrophages as the beacon, the true gaitekeeper for entrance of lymphocytes into the islet.

Three pointed findings followed the initial depletion of macrophages. First, the effect of AFS98 treatment applied to mice already undergoing an active diabetogenic process. At 10 wk of age, the autoimmune process is progressing; our own studies showed a striking inflammatory signal at this time. However, AFS98 treatment at this time was effective. This indicates that AFS98 inhibits when there are active diabetogenic T cells. Thus, the inhibitory effect of macrophage depletion is not necessarily accounted for by a lack of T cell priming. Second is that after the period of macrophage depletion, the islet becomes progressively receptive to the entrance of T cells and new macrophages. This was evident when examining the islets by flow cytometry. Despite this renewed accumulation of leukocytes into the islets, the beta cells were preserved. Histological examination at the 42-wk period showed the preservation of beta cells, albeit many islets showed the peri-insulitic lesion next to otherwise normal beta cells. This finding suggests that the newly arriving monocyte-derived macrophages or other cells in islets or in lymph nodes are not driving forward the process and may participate by controlling it. For example, the incoming macrophages may be inhibitory in a way akin of what happens in the tumor environment. Finally, a regulatory process involving the PD-1 and PD-L1 interacting molecules was evident. The mice protected by AFS98 until over 40 wk of age developed diabetes if given anti–PD-1. At face value, this result tells us that the active process had been under checkpoint control following the initial macrophage deletion.

Future studies should examine the site of the control pathway, whether it is the lymph nodes or the islets, many of which had the peri-insulitic lesion. In NOD diabetogenicity, peripheral lymph nodes play a key role in the early program of the autoimmunity (27–29). We did show here that the lymph nodes after AFS treatment could be an effective site of presentation of either diabetogenic or foreign antigens. It is not because of the lack of presenting function in lymph nodes that diabetes is curtailed. However, the nodes could be the anatomical substrate of the regulatory process involving the PD-1/PD-L1 pathway. The lymph nodes are depleted of their resident macrophages, albeit not to the extent as islets. Thus, the presentation that results in the activation of the PD-1/PD-L1 activation could be caused by DC or B cells. Lymph node macrophages at face value would foster positive interactions in a normal situation but, in their absence, would drive the process into a regulatory pathway. In our early studies, we did find that the F4/80 positive macrophages were capable of presenting minor epitopes of insulin.

Various inflammatory processes have been studied with variable results using antibodies to transiently remove tissue macrophages (30–35). Other treatment modalities have targeted blood monocytes. Clodronate-liposome treatment depletes circulating monocytes and affected a number of inflammatory reactions (36). These treatments over repeated periods of time resulted in the reduction in diabetes in NOD mice (36–40). In all of these instances, the reduction resulted in the loss of inflammatory macrophages, i.e., macrophages derived from monocytes that participated as effectors in the diabetogenic process. A study by Calderon et al. (41) showed that resident macrophages were not affected by injections of clodronate liposomes, although diabetes induced by the transfer of the activated BDC2.5 CD4+ T cells was affected. Diabetes induced by such transfer required the presence of new macrophages as effector cells modulated by IFN-γ production from activated T cells (42).

Integrating our findings with these, it is evident that macrophages have two distinct functions in autoimmune diabetes: First, as evident here, as a cell controlling the development and progression of the diabetogenic process, and second, as an effector cell once islet inflammation is induced.
Fig. 7. Early treatment with AFS98 reduces infiltrating leukocytes in NOD mice up to 12 wk of age. (A and B) Female NOD mice were left untreated or injected with 0.5 mg of AFS98 i.p. at 2 wk of age and 2.0 mg of AFS98 i.p. at 4, 7, and 10 wk of age. The islets of mice at 6 (B), 8 (A and B), and 12 (A) wk of age were isolated and analyzed by flow cytometry. (A) Shows the flow cytometry plots of myeloid and T cell compartments. Gating is indicated over the plots. Flow cytometry plots are representative of individual islet preparation from three control or four AFS98-treated mice. (B) Summarizes the flow cytometry data for all time course experiments (three to four mice per group). Bars represent the mean ± SD for two independent experiments with two to three replicates per group. (C) Flow cytometry plots of immune cell populations in islets isolated from nondiabetic control or AFS98-treated mice examined at 40–44 wk of age. Plots were generated from individual mice. Gates are indicated on the top of each column of plots. Flow cytometry plots are representative of individual islet preparations isolated from two control and eight AFS98-treated mice. (D) Summary of the data shown in C. (E and F) Hematoxylin/eosin staining of pancreatic sections isolated from nondiabetic AFS98-treated mice taken from Fig. 6A and C. The mice were treated early, and their islets were examined at 40–44 wk of age. (Scale bars, 400 μm.)
Leukocyte infiltrates are reduced following macrophage depletion at 10 wk of age. NOD mice were either treated with AFS98 or control Rat IgG2a. Seven days later, the mice were immunized with 200 μg of antibody three times over 6 d (days 0, 3, and 6; Anti–PD-1 clone RMP1-4; Leinco Technologies). Mice were then followed for diabetes incidence. For experiments examining the lymph node response by ELSIPOT, B6.g7 or NOD mice (7–10 wk old) were treated with a single 2.0-mg dose of AFS98 or control rat IgG2a. Seven days later, the mice were immunized in the footpad with 10 nmol INS B:9–23 peptide, HEL protein (Sigma-Aldrich), or IGRP:206–214 peptide. After 7 d, the animals were killed and the popliteal lymph nodes were harvested, dispersed into single-cell populations, and stained with fluorescent antibodies for analyses by flow cytometry. For experiments dealing with anti–PD-1, AFS98-treated or control NOD mice were injected with 200 μg of antibody three times over 6 d (days 0, 3, and 6; Anti–PD-1 clone RMP1-4; Leinco Technologies). Mice were then followed for diabetes incidence. For experiments examining the lymph node response by ELSIPOT, B6.g7 or NOD mice (7–10 wk old) were treated with a single 2.0-mg dose of AFS98 or control rat IgG2a. Seven days later, the mice were immunized in the footpad with 10 nmol INS B:9–23 peptide, HEL protein (Sigma-Aldrich), or IGRP:206–214 peptide. After 7 d, the animals were killed and the popliteal lymph nodes were harvested, dispersed into single-cell suspensions, and stained with fluorescent antibodies for analyses by flow cytometry.
Histology and Imaging. Microscopy imaging was performed using anoxic E800 microscope (Nikon) equipped with CFI Plan Apo Lambda DM 20x air objective, X-Cite 120PC light source (Excellitas Technologies), Eki blue fluorescence microscopy camera, and QCapture 64-bit v2.9.13 acquisition software (Qmaging).

Islets, Pancreatic Stroma, and Lymph Node Isolation. Pancreata were perfused through the common bile duct with 5.0 mL of calcium-free HBSS supplemented with 400.0 μg/mL of collagenase. Pancreata were then removed, and digested in a 37 °C water bath followed by vigorous shaking for 90 s, washed three times in HBSS, and then passed through a 70-μm strainer to retain the islets. Cells that passed the stainer represented the pancreatic stroma and were filtered a second time through a 40.0-μm filter to prepare a single-cell suspension. Islets retained on the 70-μm filter were then flushed into a Petri dish for hand-picking using a zinc-chelating dye, Dithizone (200 μg/mL 10% DMSO PBS; Sigma), to identify the islets. Hand-picked islets were then dispersed using Cell Dissociation Solution Non-Enzymatic (Sigma) for 10 min at 37 °C. T lymph nodes and spleens were digested by incubation at 37 °C in DMEM supplemented with 10% fetal calf serum, 125 μg/mL Liberase TL (Roche Life Science), and 50 μg/mL DNase I (Roche Life Science). All single-cell suspensions were then incubated with 2.4G2 conditioned media (PBS, 1% BSA, and 50% 2.4G2 in DMEM) at 4 °C for 15 min to block FC receptors. Samples were then stained with fluorescent antibodies for flow cytometry or sorting.

Antibodies for Flow Cytometry and Sorting. Flow cytometry data were acquired on a FACsCanto II (BD Biosciences) and analyzed on FlowJo v10.2 software (FlowJo, LLC). Cell sorting was performed using a FACSAria II (BD Biosciences). The following antibodies were purchased from BioLegend: BV510 anti-CD45 (30-F11), Pacific Blue (PB) anti-I-A/I-E (pan MHC-II), FITC anti-CD3 (2C11), FITC and APC anti-F4/80 (B220), PE-Cy7 anti-CD11b (M17/10), PE-Cy7 anti-CD14 (M6/29), eFluor450 anti-CD4 (RPA-T4), PE-Cy5.5 anti-CD20 (RA3-6B2), and APC anti-CD64 (555-571). APC-eFluor780 anti-CD11c (N418), PerCP-eFluor710 anti-CD80 (56-507), and APC-FoxP3 (FJK-16) were purchased from eBioscience. For intracellular staining, the FoxP3 Fix/Perm kit (eBioscience) was used according to the manufacturer’s instructions.

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