An OGT-STAT5 Axis in Regulatory T Cells Controls Energy and Iron Metabolism

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The immunosuppressive regulatory T (Treg) cells exert emerging effects on adipose tissue homeostasis and systemic metabolism. However, the metabolic regulation and effector mechanisms of Treg cells in coping with obesogenic insults are not fully understood. We have previously established an indispensably role of the O-linked N-Acetylglucosamine (O-GlcNAc) signaling in maintaining Treg cell identity and promoting Treg suppressor function, via STAT5 O-GlcNAcylation and activation. Here, we investigate the O-GlcNAc transferase (OGT)-STAT5 axis in driving the immunomodulatory function of Treg cells for metabolic homeostasis. Treg cell-specific OGT deficiency renders mice more vulnerable to high-fat diet (HFD)-induced adiposity and insulin resistance. Conversely, constitutive STAT5 activation in Treg cells confers protection against adipose tissue expansion and impaired glucose and insulin metabolism upon HFD feeding, in part by suppressing adipose lipid uptake and redistributing systemic iron storage. Treg cell function can be augmented by targeting the OGT-STAT5 axis to combat obesity and related metabolic disorders.

Keywords: O-GlcNAc, Treg - regulatory T cell, iron, fatty acid uptake, adipose tissue

INTRODUCTION

Foxp3+ regulatory T (Treg) cells are specialized immunosuppressive lymphocytes that control self-tolerance, inflammatory responses, and tissue homeostasis (1, 2). Treg cells develop in the thymus through two distinct programs involving CD25 Foxp3+ and CD25 Foxp3+ progenitors (3). The upregulation of CD25 during Treg cell development is driven by strong T cell receptor (TCR) signals (4). The upregulation of Foxp3 is TCR-independent, but instead involves IL-2 signaling through its receptor CD25 and the downstream activation of STAT5 (5–7). Besides of their indispensability during development, both TCR and IL-2/STAT5 signaling pathways are required for the suppressive function of effector Treg cells (8–10). Treg cells are essential in enforcing peripheral tolerance and loss of them leads to a fatal scurfy phenotype that is characterized by severe inflammation in endocrine and barrier organs (11–13).

In addition to preserving tissue integrity, Treg cells have been increasingly shown to support adipose tissue function and systemic metabolism. Treg cells, along with other anti-inflammatory immune cells, are abundant in adipose tissues of lean animals (14, 15). The accumulation and...
function of adipose Treg cells depend on antigen presentation by parenchymal MHC class-II molecules and local cytokines such as IL-33 to drive a PPARγ-dependent transcriptional program (16–18). Obesity profoundly remodels the immune cell compartment of fat tissue, leading to Treg cell reduction and insulin resistance. Adaptive transfer or genetic induction of Treg cells have been shown to suppress adipose inflammation and improve insulin sensitivity in obese animals (19–21). On the other hand, loss of Treg cell function by cell depletion, PPARγ knockout, or Vα5/Vβ8.2 TCR transgene leads to no or marginal changes in adipose insulin resistance (14, 18, 22). It has been proposed that Treg cells in obesity and aging may adopt pro-inflammatory properties (15, 23); therefore, restoring Treg cell function, rather than increasing Treg cell number, may offer therapeutic opportunities for metabolic diseases. Nonetheless, mechanisms mediating the insulin-sensitizing effect of adipose Treg cells are not fully understood. Also, it has not been established if Treg cells can directly modulated lipid metabolism and energy balance.

O-linked N-Acetylgalactosamine (O-GlcNAc) modification at serine or threonine residues, termed O-GlcNAcylation (24), plays a central role in signaling pathways relevant to chronic human diseases such as diabetes, cancer, and immune disorders (25–27). O-GlcNAc transferase (OGT), using UDP-GlcNAc derived from the hexosamine biosynthetic pathway as the substrate, controls diverse biological processes such as gene transcription, protein stability, and cell signaling (28–31). O-GlcNAcase (OGA) mediates the removal of O-GlcNAcylation from proteins. Protein O-GlcNAcylation senses glucose availability (30, 32), hormonal cues (29, 33, 34), cellular stress (33, 35), and immune and bacterial signals (27, 36, 37). Recently, we found in Treg cells that TCR signaling promotes OGT-mediated O-GlcNAcylation of Foxp3 and STAT5, two transcriptional factors essential for Treg cell development and function (38). Overexpression of a constitutive active form of STAT5 rescues lineage instability and suppressor dysfunction found in O-GlcNAc deficient Treg cells, thus ameliorating systemic autoimmunity in OGT-deficient mice (38). In this study, we seek to interrogate roles of the OGT-STAT5 axis in instructing Treg cells to cope with obesity-associated remodeling of the adipose tissue and systemic metabolism.

**MATERIALS AND METHODS**

**Mice**

Foxp3^{GFP-Cre} mice (stock number 016959) and Foxp3^{GFP-Cre-EERT2} mice (stock number 016961) were purchased from the Jackson Laboratory. Rosa26\_Sta5b-CA mice were provided by Dr. Alexander Rudensky at Memorial Sloan Kettering Cancer Center (10). Ogt^{fl/fl} Mice (Jackson Laboratory stock number 004860) were kindly provided by Dr. Xiaooyong Yang at Yale University. All mice were on C57BL/6 background. Mice were free to access water and fed on a regular chow, tamoxifen food (Teklad, TD.130860), or 60% high fat diet (ResearchDiets, D12492) as indicated. All procedures were approved by the Institutional Animal Care and Use Committee at the University of Minnesota. All relevant ethical regulations for animal testing and research were complied with.

**Metabolic Assays**

Body weights were monitored every week. Body composition was assessed using an EchoMRI system (39). For food intake measurement, mice were individually housed and food consumption was weighed every morning for 7 consecutive days. For the metabolic cage study, mice were first acclimated in metabolic chambers (Columbus Instruments), and then physical activity and energy expenditure were measured continuously for at least 2 days (40). For oral glucose-, and insulin- tolerance tests, 16 h fasted mice were given glucose (1.5 g/kg body weight) by gavage; 6 h fasted mice were injected with insulin (1 U/kg body weight) intraperitoneally. Tail-vein blood collected at the designated times was used to measure blood glucose level using a Contour Glucometer (Bayer). For lipid tolerance test, mice were fasted for 4 h, gavaged with 200 ul of olive oil, and tail-vein blood was collected every 1.5 h for quantification of triglyceride using a colorimetric assay kit (Cayman #10010303). Fecal energy was determined with a bomb calorimeter at Dr. Pedro Urrio’s lab. Fecal lipid was extracted with chloroform: methanol mix (41), weighed and calculated as percentage to fecal weight.

**Flow Cytometry**

For surface markers, cells were stained in PBS containing 0.5% BSA with relevant antibodies at 4°C for 30 min. For analysis of intracellular markers, cells were first fixed with Fixation/Permeabilization buffer (ThermoFisher, catalog no. 00-5123) at 4°C for 30 min and then stained in Permeabilization Buffer (ThermoFisher, catalog no. 00-8333) with relevant antibodies at 4°C for 30 min. Flow cytometry data were acquired on BD Fortessa H0081 or X-20 and analyzed with Flowjo.

**Histology**

Mouse tissues were dissected and fixed in 10% buffered formalin. Sectioning and haematoxylin & eosin (H&E) staining were performed by the Comparative Pathology Shared Resource of UMN.

**RNA and Real-time PCR**

RNA was isolated with Trizol and reverse transcribed into cDNA with the iScript™ cDNA Synthesis Kit. Real-time RT-PCR was performed using iTaq™ Universal SYBR® Green Supermix and gene-specific primers on a Bio-Rad C1000 Thermal Cycler. All data were normalized to the expression of 18s or 36b4.

**Western Blotting**

Tissues were lysed in buffer containing 1% Nonidet P-40, 50 mM Tris 3 HCl, 0.1 mM EDTA, 150 mM NaCl, protease inhibitors and an OGT inhibitor, TMG. Equal amounts of protein lysates were electrophoresed on 4-20% SDS-PAGE gels and transferred to Nitrocellulose membranes. Primary antibodies were incubated at 4°C overnight. Western blotting was visualized by peroxidase conjugated secondary antibodies and ECL chemiluminescent.
substrate with a Bio-Rad ChemiDoc Imaging System or imaged by fluorescent IgG secondary antibodies with a LI-COR Odyssey 9120 imager.

**Statistical Analyses**
Results are shown as mean ± SEM. The comparisons were carried out using two-tailed Student’s t test, one-way ANOVA followed by multiple comparisons with the Tukey adjustment, two-way ANOVA using GraphPad Prism 7.

**RESULTS**

**OGT-insufficiency in Treg Cells Leads to Insulin Resistance**
Both the Foxp3 and Ogt genes are located on the mouse X Chromosome; about 40 centimorgans apart (Figure 1A). When we generated Treg cell-specific OGT knockout (KO) using the Foxp3<sup>YFP-Cre</sup> mice with YFP-Cre knocked into the endogenous Foxp3 locus (Figure S1A), hemizygous male KO mice developed scurfy phenotypes and died of autoimmune disease by the age of 4 weeks (38), thus preventing us to study their whole-body metabolism. Due to random inactivation of the X chromosome in females, Foxp3<sup>YFP-Cre/wtOgt<sup>F/F</sup></sup> female mice possess both OGT-sufficient (WT) and -deficient (KO) Treg cells (Figure 1B). Despite lineage instability and suppressor dysfunction of KO Treg cells, the remaining WT cells were sufficient to avert autoimmunity in mice (38). Therefore, Foxp3<sup>YFP-Cre/wtOgt<sup>F/F</sup></sup> females and their littermate controls were used to investigate metabolic outcomes. When fed with normal chow (NC), no difference in body weight was observed in 12-week-old mice (Figure 1C). Compared to controls, Foxp3<sup>YFP-Cre/wtOgt<sup>F/F</sup></sup> females had higher blood glucose levels during insulin tolerance test, indicative of impaired insulin sensitivity (Figure 1D). Glucose tolerance test revealed similar blood glucose levels between the two genotypes (Figure 1B), suggesting that pancreatic insulin secretion might be increased to compensate insulin resistance in KO mice. We then treated mice with insulin to assess its downstream signaling, and found similar AKT phosphorylation in gonadal white adipose tissue (gWAT) and liver (Figure S1C). However, insulin-stimulated Ser473 phosphorylation of AKT in brown adipose tissue (BAT) was diminished in Foxp3<sup>YFP-Cre/wtOgt<sup>F/F</sup></sup> females (Figure 1E). These results demonstrate that OGT in Treg cells controls insulin pathway activation in BAT and whole-body insulin sensitivity.

**OGT-insufficiency in Treg Cells Promotes Diet-induced Obesity**
We then fed animals with 60% high fat diet (HFD) to induce obesity. Female Foxp3<sup>YFP-Cre/wtOgt<sup>F/F</sup></sup> mice gained more weight than controls (Figure 2A). Body composition analysis showed that weight gain was due to increased fat mass (Figures 2B, C). Similar to animals fed with chow, HFD Foxp3<sup>YFP-Cre/wtOgt<sup>F/F</sup></sup> females displayed impaired insulin tolerance (Figure 2D), but normal glucose tolerance (Figure 2E).

We went on to determine what caused diet-induced adiposity in Foxp3<sup>YFP-Cre/wtOgt<sup>F/F</sup></sup> mice. Their daily intake of HFD was similar to control mice, when adjusted to body weight using ANCOVA analysis (Figure 2F). Metabolic cage study found no difference in locomotor activity (Figure 2G) or respiratory exchange ratio (Figure 2H). Rather, heat production, when...
adjusted to body weight (42), was significantly reduced in Foxp3^{YFP-Cre/wtOgtF/F} mice during both light and dark periods (Figures 2I, J). Moreover, when challenged with cold, Foxp3^{YFP-Cre/wtOgtF/F} animals showed reduced heat production than the controls (Figure 2K), indicative thermogenic defects. Together, these data demonstrate that OGT-controlled Treg cell function enables metabolic adaptation to positive energy balance.

STAT5 Activation in Treg Cells Confers Protection Against Diet-induced Obesity

The regulation of Treg cell suppressive program by OGT is dependent on STAT5 O-GlcNAcylation and activation (38). We then sought to determine whether STAT5 overactivation, to the opposite of OGT knockout, would drive Treg cell action in systemic metabolism. Foxp3^{YFP-Cre/YRosa26Stat5b-CA/wt} male mice were generated to specifically overexpress a constitutively active (CA-) form of STAT5B in Treg cells (10). Flow cytometric analyses confirmed that STAT5B-CA not only increased the number of FOXP3\(^+\) Treg cells in the lymph nodes and the activity of the Foxp3 gene promoter (Figure 3A), but also drastically expanded the population of effector Treg cells that were CD44\(^+\)KLRG1\(^+\) or CD44\(^+\)CD103\(^+\) (Figure 3B, C). When fed with a normal chow, even though with comparable body weight (Figure 3D), Foxp3^{YFP-Cre/YRosa26Stat5b-CA/wt} mice showed a slight, but statistically significant reduction in fat mass (Figure 3E). There was a declining trend of blood glucose levels during the insulin tolerance test (Figure 3F). Similar locomotor activity (Figure 3G), energy expenditure (Figure 3H), and energy or lipid excretion into the feces (Figures 3I, J) were observed between control and Foxp3^{YFP-Cre/YRosa26Stat5b-CA/wt} mice. We went on to challenge the mice with HFD. The resistant to HFD-induced weight gain could be readily seen in Foxp3^{YFP-Cre/YRosa26Stat5b-CA/wt} mice as early as 3 weeks after diet-switching (Figure 4A). The reduction in weight gain was solely because of reduced fat mass (Figure 4B). Both BAT and gWAT depots were significantly smaller in Foxp3^{YFP-Cre/YRosa26Stat5b-CA/wt} mice, while liver weight remained the same (Figure 4C). We then asked what led to the lean phenotype after STAT5B-CA expression in Treg cells.
cells. Continuous metabolic monitoring revealed a substantial reduction of food intake in Foxp3YFP-Cre/YRosa26Stat5b-CA/wt mice (Figure 4D), while energy expenditure (Figure 4E) and locomotor activity (Figure 4F) were both comparable to wildtype controls. Moreover, there was a tendency that fecal energy excretion was increased after Treg cell hyperactivation (Figure 4G). We then measured lipid content in the feces and found more lipids were excreted in Foxp3YFP-Cre/YRosa26Stat5b-CA/wt mice (Figure 4H). As expected, these HFD-fed but relatively lean Foxp3YFP-Cre/YRosa26Stat5b-CA/wt mice show much better glucose handling during the glucose tolerance test (Figure 4I). These data demonstrate that Treg cell expansion and activation stimulated by tonic STAT5 signaling restrain dietary fat intake/absorption and render mice resistant to diet-induced metabolic disturbances.

Inducible STAT5 Activation in Treg Cells Improves Systemic Glucose Metabolism

Due to stochastic activity of the Foxp3 promoter (43), constitutive overexpression of STAT5B-CA led to progressive splenomegaly and hematopoietic neoplasia in Foxp3YFP-Cre/YRosa26Stat5b-CA/wt mice (data not shown), similar to that after STAT5 activation in other hematopoietic compartments (44–46). We therefore used the inducible Foxp3eGFP-Cre-ERT2 line to generate Foxp3eGFP-Cre-ERT2/YRosa26Stat5b-CA/wt male mice, fed them with tamoxifen-containing food to induce STAT5-CA expression in Treg cells, and challenged them with HFD feeding (Figure 5A). A trending reduction in weight gain could be observed in Foxp3eGFP-Cre-ERT2/YRosa26Stat5b-CA/wt mice (Figure 5B). When compared to their control counterparts, Foxp3eGFP-Cre-ERT2/YRosa26Stat5b-CA/wt mice possessed significantly less BAT (Figure 5C) and slightly smaller WAT (Figure 5D, E). No change in muscle mass was noticed (Figure 5F). Moreover, Foxp3eGFP-Cre-ERT2/YRosa26Stat5b-CA/wt animals also displayed much improved insulin tolerance (Figure 5G). Concomitantly, insulin signaling activity, reflected by AKT Ser473 phosphorylation, was augmented in BAT of Foxp3eGFP-Cre-ERT2/YRosa26Stat5b-CA/wt mice (Figure 5H). The more profound protection observed in the former constitutive model might stem from stronger and broader expression of STAT5B-CA. Nonetheless,

FIGURE 3 | Characterization of Foxp3YFP-Cre/YRosa26Stat5b-CA/wt mice. (A–C) Flow cytometric quantification of FOXP3-YFP+CD4+ Treg cells (A), CD44+KLRG1+ (B) and CD44+CD103+ (C) effector Treg cells in lymph nodes from Foxp3YFP-Cre/YRosa26Stat5b-CA/wt mice. (D) Growth curve of Foxp3YFP-Cre/YRosa26Stat5b-CA/wt (n = 10) and Foxp3YFP-Cre/YRosa26Stat5b-CA/wt (n = 9) mice fed with NC. (E) Lean and fat mass of NC-fed Foxp3YFP-Cre/YRosa26Stat5b-CA/wt (n = 7) and Foxp3YFP-Cre/YRosa26Stat5b-CA/wt (n = 6) mice. (F) Insulin tolerance test of NC-fed Foxp3YFP-Cre/YRosa26Stat5b-CA/wt (n = 7) and Foxp3YFP-Cre/YRosa26Stat5b-CA/wt (n = 6) mice. Area under curve (AUC) is shown to the right. (G) Locomotor activity and (H) body weight-adjusted heat production of NC-fed Foxp3YFP-Cre/YRosa26Stat5b-CA/wt (n = 7) and Foxp3YFP-Cre/YRosa26Stat5b-CA/wt (n = 6) mice, determined by metabolic cages. (I) Fecal energy and (J) fecal lipid percentage of NC-fed Foxp3YFP-Cre/YRosa26Stat5b-CA/wt (n = 7) and Foxp3YFP-Cre/YRosa26Stat5b-CA/wt (n = 6) mice. Data are presented as mean ± SEM. n.s., not significant; *, p < 0.05; ***, p < 0.001 by unpaired student’s t-test or two-way ANOVA (D, F).
using two complementary models, we demonstrate that STAT5 activation in Treg cells is sufficient to protect mice from diet-induced adiposity and insulin resistance.

**Treg Cell Activation Suppresses Lipid Uptake by Adipose Tissue**

We then inquired into the causes of metabolic protection after STAT5-dependent Treg cell overactivation. We first interrogated the involvement of BAT thermogenesis. BAT histology clearly revealed smaller adipocytes and less lipid deposition in Foxp3\textsuperscript{YFP-Cre/Y Rosa26\textsuperscript{Stat5b-CA/wt}} mice (Figure 6A). However, the expression of thermogenic genes remained unchanged in BAT (Figure 6B), or even reduced in inguinal WAT (iWAT) (Figure 6C). This suggests that reduced weight gain by Treg cell activation was not a result of increased thermogenesis.

Not only was gWAT mass reduced in Foxp3\textsuperscript{YFP-Cre/Y Rosa26\textsuperscript{Stat5b-CA/wt}} mice (Figures 4B, C), but adipocyte size was also much smaller (Figure 6D). Crown-like structures were more evident in Foxp3\textsuperscript{YFP-Cre/Y Rosa26\textsuperscript{Stat5b-CA/wt}} gWAT (Figure 6D), suggesting active clearing of dying adipocytes by macrophages (47). Interestingly, the expression of both C\textsuperscript{d36} (Figure 6E) and L\textsuperscript{lpl} (Figure 6F), two important genes required for cellular uptake of free fatty acid (48), were substantially ablated in gWAT and iWAT of Foxp3\textsuperscript{YFP-Cre/Y Rosa26\textsuperscript{Stat5b-CA/wt}} mice. There seemed to be no defect in dietary intake of lipid in the intestine, as intestinal C\textsuperscript{d36} expression was intact (Figure 6G) and blood triglyceride levels after an oral gavage of olive oil were even higher in Foxp3\textsuperscript{YFP-Cre/Y Rosa26\textsuperscript{Stat5b-CA/wt}} mice (Figure 6H). Taking in consideration of the elevated fecal lipid content (Figure 4H), we postulate that Treg cell activation suppresses lipid uptake into adipose tissues, leading to more energy excretion and less weight gain during HFD feeding.

**Treg Cell Activation Remodels Systemic Iron Metabolism**

Iron is critically important for adipose tissue function and systemic metabolism. Dietary iron supplementation augments thermogenesis, reduces fat mass, and protects mice from diet-induced adiposity and insulin resistance.
induced weight gain (49, 50). On the other hand, nonanemic iron deficiency impairs thermogenesis and promotes visceral obesity in mice (51). Lowering adipocyte iron uptake via genetic deletion of transferrin receptor 1 (TFR1) leads to impaired thermogenesis, increased insulin resistance, and inflammation (52). We then inquired if the metabolic protection after STAT5-dependent Treg cell activation involves iron. Ferritin is a protein complex that stores cellular iron and its protein levels were elevated in BAT, gWAT and iWAT of HFD-fed Foxp3YFP-Cre/YRosa26Stat5b-CA/wt mice, compared to Cre-only controls (Figures 7A–C), indicative of excess iron storage in fat tissues. In Foxp3YFP-Cre/YRosa26Stat5b-CA/wt adipose tissues, there was a trending increase in Tfr1 gene expression (Figure 7D) and TFR1 protein levels (Figures 7A–C). Conversely, expression of the Fpn1 gene (encoding iron export protein Ferroportin) was significantly downregulated (Figure 7E). In contrast to adipose tissue, both liver and spleen showed substantial reduction in Ferritin protein levels (Figures 7F, G), accompanied with increased TFR1 expression. The presumptive impairment in hepatic iron storage was correlated with reduced expression of iron-responsive Hamp and Hfe genes (Figure 7H), which encode Hepcidin and Homeostatic iron regulator, respectively. As a result of reduced Heparin, intestinal genes involved in iron uptake including Dmt1, Cyp3a11, Fn1 displayed varying degrees of upregulation (Figure 7I). Collectively, STAT5B-CA-mediated Treg cell overactivation redistributes the iron pool from liver and spleen to adipose tissues, which may contribute to the alleviation of metabolic stress associated with diet-induced obesity.

**DISCUSSION**

In effector Treg cells, protein O-GlcNAcylation induced by the TCR signaling, modifies and activates STAT5 for Treg cell homeostasis and function (38). In the study, we investigate whether and how the OGT-STAT5 regulatory axis dictates Treg cell function during metabolic adaptation to diet-induced obesity. Taking advantage of random X chromosome inactivation and mosaic OGT-defi ciency in Foxp3 YFP-Cre/wtOgtF/F female mice, we establish that functional deficiency in about half the Treg cells renders animals more prone to diet-induced adiposity and insulin resistance. Our model is more advantageous than other Treg cell depletion models, such as the Foxp3-DTR mice (14), since the absence of systemic inflammation in Foxp3YFP-Cre/wtOgtF/F females allowed us to perform long-term studies and avoided side effects and caveats of tissue inflammation (38). This is possibly also the reason why we observed obesity and thermogenic defects in Foxp3YFP-Cre/wtOgtF/F mice, which have not been reported in other loss-of-function animals (14, 18, 22). We speculate that other heterozygous loss of Treg function models would show similar metabolic phenotypes as the Foxp3YFP-Cre/wtOgtF/F mice, provided that systemic inflammation is avoided. In addition,
we do not imply that Treg function in coping with obesity is specifically controlled by OGT, but rather OGT is essential for most if not all Treg functions, including their regulation of insulin sensitivity and systemic metabolism. For instance, we previously demonstrated that Blimp1+ effector Treg cells are ablated in OGT knockout mice (38). Similar to OGT knockout, Blimp1-deficient mice do not have effector Tregs (53). Recently, it was shown that Blimp1+ Tregs are also involved in the homeostasis of adipose tissue (54).

It is important to note that BAT harbors Treg cell populations that have shared and distinct gene signatures compared to WAT-resident Treg cells (55). Cold exposure or β3-adrenergic stimuli enhance Treg induction that is linked by a T cell-specific STAT6-Pten axis (56). When OGT is deficient in Treg cells, BAT insulin resistance and impaired thermogenesis could be observed, contributing to diet-induced obesity. Nonetheless, it is unclear if cold or β3-adrenergic activation could directly modulate the hexosamine biosynthetic pathway, OGT activity, and protein O-GlcNAcylation in Treg cells. Other work in our laboratory has revealed a pivotal role of O-GlcNAcylation on STAT6 signaling and type 2 immunity (57). It is warranted in the future to determine if OGT also regulates the STAT6-Pten axis in tissue-specific Treg cells and adipose tissue function.

IL-2 and IL-2R-dependent STAT5 activation are required for Treg cell development (5, 58). Foxp3<sup>YFP-Cre</sup>-mediated STAT5a/b knockout in male mice causes scurfy phenotypes and death by the age of 2 months (data not shown). We also generated mosaic Foxp3<sup>YFP-Cre/YRosa26Stat5b-CA/wt</sup> female mice avoiding fatal inflammation. However, we did not observe evident metabolic perturbations in these animals (data not shown), differently from Foxp3<sup>YFP-Cre/YRosa26OgtF/F</sup> females. It indicates that the metabolic effects of Treg cell O-GlcNAcylation might not be mediated just by STAT5, and other mechanisms such as STAT6 could compensate for the loss of STAT5. Nevertheless, STAT5B-CA overexpression is sufficient to booster Treg cell number and immunosuppression (10). Using these Foxp3<sup>YFP-Cre/Rosa26Stat5b-CA/wt</sup> mice, we demonstrate that Treg cell activation renders mice resistant to diet-induced weight gain and insulin resistance, in part by suppressing fatty acid uptake in adipose tissue and remodeling systemic iron metabolism.
that STAT5-independent, but maybe STAT6-dependent mechanisms in Treg cells control adipose tissue lipolysis and thermogenesis (56).

While it has been widely accepted that Treg cells are important contributors to metabolic homeostasis, the regulatory mechanisms are not fully understood. Previous studies have revealed inflammation, lipolysis, and thermogenesis as tissue-specific targets of Treg cells. Our current study suggests two additional metabolic actions of Treg cells – fatty acid uptake and iron distribution. While intestinal lipid absorption is not perturbed, STAT5B-CA-driven Treg cell activation suppresses the expression of key genes involved in adipose fatty acid uptake – Lpl and Cd36. It would be important in the future to determine if the obesity protection in STAT5B-CA overexpression mice is dependent on LPL/CD36 downregulation. Meanwhile, the adipose iron pool seems to be expanded in STAT5B-CA overexpression, which we believe contributes to thermogenic activation and fat mass reduction. However, we are unclear whether parenchymal adipocytes or resident immune cells such as macrophages play a predominant role in the change in iron content. We also lack a mechanistic understanding of how Treg cells differentially regulate iron metabolism across different tissues (adipose vs. liver and kidney). Finally, it would be interesting to test if the suppression of lipid uptake is linked to iron redistribution in adipose tissues. In summary, our study has revealed new biology of Treg cells in metabolic regulation and the OGT-STAT5 axis as a target to harness Treg cell function for obesity and diabetes therapeutics.

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/Supplementary Material. Further inquiries can be directed to the corresponding author.

ETHICS STATEMENT

The animal study was reviewed and approved by University of Minnesota.

AUTHOR CONTRIBUTIONS

ZZ and MF generated and analyzed STAT5B-CA mice. OS and KH did flow cytometry. BL analyzed OGT-KO mice. ZM helped with Western blot. H-BR conceived the project, analyzed data and wrote...
the manuscript. All authors contributed to the article and approved the submitted version.

**FUNDING**

This work was supported by NIH grant R56 AI162791 to H-BR and R01 AI139420 to H-BR and MF.

**ACKNOWLEDGMENTS**

We thank Dr. Alexander Rudensky for providing the Rosa26\textsuperscript{Stat5b-CA} mice and Dr. Xiaoyong Yang for providing the Ogt\textsuperscript{fl/fl} mice. We thank Drs. Alessandro Bartolomucci, Maria Razzoli, and Pilar Ariza Guzman at the IBP Phenotyping Core of University of Minnesota for helping with the metabolic cages and EchoMRI analyses. We thank Dr. Pedro Urriol for providing us access to their bomb caloriometer.

**SUPPLEMENTARY MATERIAL**

The Supplementary material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fimmu.2022.874863/full#supplementary-material
32. Hardiville S, Hart GW. Nutrient Regulation of Signaling, Transcription, and Cell Physiology by O-GlcNAcylation. Cell Metab (2014) 20:208–13. doi: 10.1016/j.cmet.2014.07.014
33. Ruan HB, Ma Y, Torres S, Zhang B, Periodo C, Heck RM, et al. Calcium-Dependent O-GlcNAc Signaling Drives Liver Autophagy in Adaptation to Starvation. Genes Dev (2017) 31:1655–65. doi: 10.1101/gad.305441.117
34. Whelan SA, Lane MD, Hart GW. Regulation of the O-Linked Beta-N-Acetylgalcosamine Transferease by Insulin Signaling. J Biol Chem (2008) 283:21411–7. doi: 10.1074/jbc.M800677200
35. Martinez MR, Dias TB, Natov PS, Zachara NE. Stress-Induced O-GlcNAcylation: An Adaptive Process of Injured Cells. Biochem Soc Trans (2017) 45:237–49. doi: 10.1042/BST20160153
36. Zhao M, Ren K, Xiong X, Cheng M, Zhang Z, Huang Z, et al. Protein O-GlcNAc Modification Links Dietary and Gut Microbial Cues to the Differentiation of Enterodendocrine I. Cells. Cell Rep (2020) 32:108013. doi: 10.1016/j.celrep.2020.108013
37. Zhao M, Xiong X, Ren K, Xu B, Cheng M, Sahu C, et al. Deficiency in Intestinal Epithelial O-GlcNAcylation Predisposes to Gut Inflammation. EMBO Mol Med (2018) 10(8): e8736. doi: 10.15252/emmm.201708736
38. Liu B, Salgado OC, Singh S, Hippen KL, Maynard JC, Burlingame AL, et al. The Lineage Stability and Suppressive Program of Regulatory T Cells Require Protein O-GlcNAcylation. Nat Commun (2019) 10:354. doi: 10.1038/s41467-019-08300-3
39. Wang Q, Tang J, Jiang S, Huang Z, Song A, Hou S, et al. Inhibition of PPARgamma, Adipogenesis and Insulin Sensitivity by MAGED1. J Endocrinol (2018) 239:167–80. doi: 10.1530/JOE-18-0349
40. Huang Z, Ruan HR, Xian L, Chen W, Jiang S, Song A, et al. The Stem Cell Factor/Kit Signalling Pathway Regulates Mitochondrial Function and Energy Expenditure. Nat Commun (2014) 5:4282. doi: 10.1038/ncomms5282
41. Kraus D, Yang Q, Kahn BB. Lipid Extraction From Mouse Feces. Bio Protoc (2015) 5(1):e1375. doi: 10.1529/bioprotoc.1375
42. Tschop MH, Speckman JR, Arch JR, Auwert J, Bruning JC, Chan L, et al. A Guide to Analysis of Mouse Energy Metabolism. Nat Methods (2012) 9:57–63. doi: 10.1038/nmeth.1806
43. Bittner-Eddy PD, Fischer LA, Costalongo M. Cre-loxP Reporter Mouse. Front Immunol (2019) 10:2228. doi: 10.3389/fimmu.2019.02228
44. Burchill MA, Goetz CA, Prlic M, O’Neill JJ, Harmon IR, Bensinger SJ, et al. Distinct Effects of STAT5 Activation on CD4+ and CD8+ T Cell Homeostasis: Development of CD4+CD25+ Regulatory T Cells Versus CD8+ Memory T Cells. J Immunol (2003) 171:5853–64. doi: 10.4049/jimmunol.171.11.5853
45. Pham HTT, Maurer B, Prchal-Murphy M, Grausenburger R, Grundschobor E, Javaheri T, et al. STAT5BN642H is a Driver Mutation for T Cell Neoplasia. J Clin Invest (2018) 128:387–401. doi: 10.1172/JCI94509
46. Maurer B, Novartha H, Wingelhofer B, Pham HTT, Schlederer M, Supe T, et al. High Activation of STAT5A Drives Peripheral T-Cell Lymphoma and Leukemia. Hauematologica (2020) 105:435–47. doi: 10.3324/haematol.2019.216986
47. Murano I, Barbutelli G, Parisani V, Latini C, Muzzonigro G, Castellucci M, et al. Dead Adipocytes, Detected as Crown-Like Structures, are Prevalent in Visceral Fat Depots of Genetically Obese Mice. J Lipid Res (2008) 49:1562–8. doi: 10.1194/jlr.M800019-JLR200
48. Goldberg IE, Eckel RH, Abumrad NA. Regulation of Fatty Acid Uptake Into Tissues: Lipoprotein Lipase- and CD36-Mediated Pathways. J Lipid Res (2009) 50 Suppl:S86–90. doi: 10.1194/jlr.R800085-JLR200
49. Romero AR, Mu A, Ayres JS. Fat Specific Adipose Triglyceride Lipase is Necessary for Iron-Mediated Lipolysis and Lipid Mobilization in Response to Negative Energy Balance. JCI Insight (2021) 2021:08. doi: 10.1111/jcic.2021.08.05.455308
50. Kitamura N, Yokoyama Y, Taoka H, Nagano U, Hosoda S, Tawornwat T, et al. Iron Supplementation Regulates the Progression of High Fat Diet Induced Obesity and Hepatic Steatosis via Mitochondrial Signaling Pathways. Sci Rep (2021) 11:10753. doi: 10.1038/s41598-021-89673-8
51. Yook JS, Thomas SS, Toney AM, You M, Kim YC, Liu Z, et al. Dietary Iron Deficiency Modulates Adipocyte Iron Homeostasis, Adaptive Thermogenesis, and Obesity in C57BL/6 Mice. J Nutr (2021) 151:2967–75. doi: 10.1093/jn/ nxab22
52. Li J, Pan X, Pan G, Song Z, He Y, Zhang S, et al. Transferrin Receptor 1 Regulates Thermogenic Capacity and Cell Fate in Brown/Beige Adipocytes. Adv Sci (Weinh) (2020) 7:1903366. doi: 10.1002/advs.201903366
53. Cretney E, Xin A, Shi W, Minnich M, Masson F, Musari M, et al. The Transcription Factors Blimp-1 and IRF4 Jointly Control the Differentiation and Function of Effector Regulatory T Cells. Nat Immunol (2011) 12:304–11. doi: 10.1038/ni.2006
54. Beppu LY, Mooli RGR, Xu Q, Marrero GJ, Finley CA, Fooks AN, et al. Tregs Facilitate Obesity and Insulin Resistance via a Blimp-1/IL-10 Axis. JCI Insight (2021) 6(3):e140644. doi: 10.1172/jci.insight.140644
55. Medrikova D, Sijmonsma TP, Sovodnik K, Richards DM, Delacher M, Sticht C, et al. Brown Adipose Tissue Harbors a Distinct Sub-Population of Regulatory T Cells. PLoS One (2015) 10:e0118534. doi: 10.1371/journal.pone.0118534
56. Kalin S, Becker M, Ott VB, Serr I, Hosp F, Mollah MMH, et al. A Stat6/Pten Axis Links Regulatory T Cells with Adipose Tissue Function. Cell Metab (2017) 26:475–492 e7. doi: 10.1016/j.cmet.2017.08.008
57. Zhao M, Ren K, Xiong X, Xin Y, Zou Y, Maynard JC, et al. Epithelial STAT6 O-GlcNAcylates Drives a Concerted Anti-Helminth Alarmin Response Dependent on Tuft Cell Hyperplasia and Gasdermin C. Immunity (2022) 55:623–638 e5. doi: 10.1016/j.immuni.2022.03.009
58. Yao Z, Kanno Y, Kerenyi M, Stephens G, Durant L, Watford WT, et al. Nonredundant Roles for Stat5a/b in Directly Regulating Fopx3. Blood (2007) 109:A368–75. doi: 10.1182/blood-2006-11-055756

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