**Regulation of ILC2/eosinophil by sympathetic tone**

**Research**

**IL-33-driven ILC2/eosinophil axis in fat is induced by sympathetic tone and suppressed by obesity**

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**Abstract**

Group 2 innate lymphoid cells (ILC2s) in white adipose tissue (WAT) promote WAT browning and assist in preventing the development of obesity. However, how ILC2 in adipose tissue is regulated remains largely unknown. Here, our study shows that ILC2s are present in brown adipose tissue (BAT) as well as subcutaneous and epididymal WAT (sWAT and eWAT). The fractions of ILC2s, natural killer T (NKT) cells and eosinophils in sWAT, eWAT and BAT are significantly decreased by high-fat-diet (HFD) feeding and leptin deficiency-induced obesity. Consistent with this, the adipose expression and circulating levels of IL-33, a key inducing cytokine of ILC2, are significantly downregulated by obesity. Furthermore, administration of IL-33 markedly increases the fraction of ILC2 and eosinophil as well as the expression of UCP1 and tyrosine hydroxylase (TH), a rate-limiting enzyme in catecholamine biosynthesis, in adipose tissue of HFD-fed mice. On the other hand, cold exposure induces the expression levels of IL-33 and UCP1 and the population of ILC2 and eosinophil in sWAT, and these promoting effects of cold stress are reversed by neutralization of IL-33 signaling in vivo. Moreover, the basal and cold-induced IL-33 and ILC2/eosinophil pathways are significantly suppressed by sympathetic denervation via local injection of 6-hydroxydopamine (6-OHDA) in sWAT. Taken together, our data suggest that the ILC2/eosinophil axis in adipose tissue is regulated by sympathetic nervous system and obesity in IL-33-dependent manner, and IL-33-driven ILC2/eosinophil axis is implicated in the development of obesity.

**Key Words**

- ILC2
- eosinophil
- sympathetic tone
- obesity
- IL-33

**Introduction**

Beige or brite adipocytes, which are enriched in subcutaneous fat, have high thermogenic capacity and have become a new therapeutic target for the treatment of obesity and its related diseases (Akbari et al. 2003, Petrovic et al. 2010, Bostrom et al. 2012, Fisher et al. 2012). Type 2 immune responses are defined by the recruitment of
CD4+ T helper type 2 (Th2) cells and the production of type 2 cytokines such as interleukin (IL)-4, IL-5, IL-9 and IL-13. An adipose tissue-specific type 2 immune response is characterized by infiltration of eosinophils, group 2 innate lymphoid cells (ILC2s), IL-4- and/or IL-13-induced M2 macrophages; production of type 2 cytokines; and plays a critical role in regulating browning of white adipose tissue (WAT), energy expenditure and glucose homeostasis (Molofsky et al. 2013, Brestoff et al. 2014, Khalifeh-Soltani et al. 2014, Rao et al. 2014). Type 2 cytokines including IL4, IL5 and IL13 are the key players in type 2 immunity and mediate its specific role in WAT browning. Deficiency of IL-4 and IL-13 impairs cold-induced WAT browning and exacerbates diet-induced obesity and insulin resistance (Rao et al. 2014). Macrophages are alternatively activated by IL-4 and IL-13 (IL-4/13) and mediate IL-4/13-induced browning by producing norepinephrine in adipose tissue (Wu et al. 2011, Van Dyken & Locksley 2013). However, the production and regulation of type 2 cytokines in adipose tissue remains largely unknown.

Innate lymphoid cells are present in adipose tissue and promote type 2 inflammation and browning of fat (Brestoff et al. 2014, Lee et al. 2015). ILC2s, which were first discovered in the lung (Price et al. 2010), are activated by epithelial cell-derived cytokines IL-33 and IL-25 as well as thymic stromal lymphopoietin in response to allergens. ILC2s orchestrate type 2 innate and adaptive immune responses (Koyasu & Moro 2013, Licona-Limon et al. 2013, Cayrol & Girard 2014). Very recently, ILC2s were identified in murine and human adipose tissue as a conserved characteristic of obesity, and have been shown to drive WAT browning and prevent the development of obesity (Brestoff et al. 2014, Lee et al. 2015). On one hand, ILC2s produce type 2 cytokines IL-5 and IL-13, where IL-5 stimulates the maturation and infiltration of eosinophils and IL-13 activates M2 macrophages: eosinophils and M2 macrophages subsequently promote the browning effect in adipose tissue (Molofsky et al. 2013, Nussbaum et al. 2013, Khalifeh-Soltani et al. 2014, Lee et al. 2015). On the other hand, IL-33-elicited ILC2s themselves produce methionine-enkephalin peptides, which directly target adipocytes and promote browning and thermogenesis (Brestoff et al. 2014). In addition, natural killer T (NKT) cells found in human adipose tissue have been shown to produce and secrete IL-4, IL-5 and IL-13 (Akbari et al. 2003, Lynch et al. 2009, 2012). Invariant NKT cells are decreased in human obesity, and confer protection against the development of the metabolic syndrome and inflammation induced by high-fat-diet (HFD) feeding through anti-inflammatory cytokines such as IL-4 and IL-10 (Lynch et al. 2009, 2012). Adipose-resident eosinophils also produce type 2 cytokines, including IL-4, IL-5 and IL-13, and sustain adipose alternatively activated macrophages, thereby enhancing glucose tolerance (Wu et al. 2011). In contrast, γδ T cells, another type of lymphocytic cells present in adipose tissue, are associated with diet-induced obesity, and have been shown to promote diet-induced inflammation and insulin resistance (Mehta et al. 2015). Although some studies suggest that lymphocytes in adipose tissue are involved in the development of obesity-induced inflammation and insulin resistance (Chatzigeorgiou et al. 2012, Mehta et al. 2015), whether various lymphocyte or lymphocyte-like subsets in brown and beige adipose tissue play different roles in thermogenesis and obesity remains to be established. In addition, the β3-adrenoceptor and its downstream protein kinase A (PKA) signaling pathway have been shown to drive the ‘browning’ effect in WAT in rodents (Cao et al. 2004). However, whether the β3-adrenoceptor and its downstream signaling pathway play a role in regulating the recruitment and activation of ILC2s, NKT and γδ T cells remains unclear.

Here, we show that the frequency of ILC2s, eosinophils and NKT cells in the stromal vascular fraction (SVF) is significantly lower in brown adipose tissue (BAT) and subcutaneous WAT (sWAT) compared with epididymal WAT (eWAT). Although the cell number of total lymphocytes and macrophages was increased by HFD feeding and leptin deficiency, the fractions of ILC2s, NKT cells and eosinophils in adipose tissue were significantly decreased by 12 weeks of HFD feeding and leptin deficiency. In support of this, the circulating and adipose expression levels of ILC2-eliciting factor IL-33 were significantly downregulated by HFD feeding and leptin deficiency-induced obesity. Moreover, IL-33 administration dramatically upregulated the frequency of ILC2s and eosinophils and expression levels of UCP1 and tyrosine hydroxylase (TH) in sWAT of HFD-fed mice. Consistently, blocking IL-33 signaling with ST2 (IL-33 receptor) antibody suppressed the promoting effects of cold on ILC2/eosinophil axis and expression of UCP1 and TH in vivo. In addition, our data also suggest that sympathetic tone is required for cold-induced IL-33 production and recruitment of ILC2s and eosinophils in sWAT. Our data strongly suggest that the IL-33-driven ILC2/eosinophil axis in adipose tissue is a key pathway in the onset of obesity and may play an important role in regulating energy homeostasis.
Methods

Mice Lepr<sup>ob</sup> (commonly referred to as ob/ob) heterozygous mice were purchased from Jackson Laboratory and intercrossed to generate ob/ob homozygous mice and wild-type littermates. The lean wild-type mice were used as the control for ob/ob homozygous mice. The ob/ob heterozygous mice were only used for the generation of experimental mice but not for the following experiments. Six-week-old male C57BL/6 mice were fed either a normal chow diet (NCD) or HFD (45% kcal from fat, D12451; Research Diets Inc, New Brunswick, NJ, USA) for 1, 12, 20 or 32 weeks. All the animals were housed in the specific pathogen-free barrier facility with a 12h light:12h darkness cycle with free access to food and water. All animal experimental protocols were reviewed and approved by the Animal Care Committee of the University of New Mexico Health Sciences Center.

Denervation of inguinal fat pads

Seven-week-old male C57BL/6 mice were anesthetized using constant flow isoflurane inhalation and a small incision was made in the abdominal skin. One of the two inguinal fat (sWAT) pads was denervated and the other pad, used as the control, was injected with the vehicle. 6-hydroxydopamine (6-OHDA, Sigma) was dissolved in PBS containing 1% ascorbic acid (9 mg/mL), and injected into the denervated pad using a 25 μL Hamilton syringe (2 μL for each injection, total 24 μL/pad) within 10 min. The same volume of vehicle, PBS containing 1% ascorbic acid, was injected into the control fat pad. The needle was kept in place for 10 s after each injection to minimize reflux from the site of injection. The skin incision was closed with stainless steel wound clips and covered with nfz Puffer to protect the wound from infection after injection. The mice were then administered with 1 mg/kg of buprenorphine by intraperitoneal (i.p.) injection. Two weeks post denervation, half of operated mice were individually housed in cages with free access to food and water continuously and exposed to cold (6°C) for 2 days and all were then killed. Inguinal fat was isolated for flow cytometry analysis.

Isolation of SVF

The inguinal, epididymal and interscapular brown fat pads were collected to exclude lymph nodes, and 0.4 g inguinal fat, 0.5 g epididymal fat and 0.2 g brown fat were weighed and used for the isolation of SVF. The adipose tissues from different mice were individually digested with collagenase A (Roche), and the samples were filtered through 100 μM cell strainers (Falcon) and washed with RPMI-1640 medium. The cell pellet was collected as SVF by centrifugation and resuspended for FACS analysis.

Flow cytometry

The suspended SVF from adipose depots was fixed, blocked and stained with conjugated antibodies including anti-CD45, anti-Siglec-5, anti-CD11b and anti-CD206 (eBioscience and BioLegend) to identify macrophage subsets. To detect ILC2, the SVF cells were stimulated with a PMA/ionomycin/menonsin mixture and then incubated with antibodies to PE-conjugated anti-CD45, PerCP-conjugated anti-LIN (CD3e, CD11b, B220, CD11c, Gr-1, IgE) and Alexa 488-conjugated anti-IL-13 (eBioscience and BioLegend) after fixation and permeabilization. The T cell subpopulations were blocked and labeled with antibodies to PerCP-conjugated anti-CD3, APC-conjugated anti-γδTCR (eBioscience and BioLegend) and PE-conjugated anti-mCD1d/PBS-57 tetramer (NIH tetramer core facility) followed by fixation. FACS analysis was performed on a FACS Calibur (BD Pharmingen, San Diego, CA, USA), and the data were analyzed with FlowJo software as described previously (Yang et al. 2013).

ELISA analysis of norepinephrine and IL-33

Enzyme-linked immunosorbent assay (ELISA) of tissue samples for norepinephrine was performed following the manufacturer’s protocol using a noradrenaline research ELISA kit (LDN Labor Diagnostika Nord GmbH & Co. KG, Nordhorn, Germany). In brief, fat tissue was weighed and homogenized with 0.01 M HCl in the presence of 1 mM EDTA and 4 mM sodium metabisulphite. The norepinephrine was extracted and acylated from the tissue homogenate followed by enzyme conversion, and then measured by ELISA. And the ELISA analysis of IL-33 in serum and inguinal fat was performed following the manufacturer’s protocol using a mouse IL-33 DuoSet ELISA kit (R&D Systems).

Real-time PCR analysis

Mouse inguinal fat samples were homogenized and the total RNA was isolated with the RNeasy Lipid Tissue Mini Kit (Qiagen). RNA (1 μg) was reverse-transcribed using cDNA kit (Qiagen). PCR amplification was detected using SYBR Green PCR master mixture on Roche 480 Real-time PCR system (Roche).
Figure 1

Differential abundance of innate/innate-like lymphocytes, eosinophils and monocytes in three types of fat. The SVF from 10-week-old male C57Bl/6 mice (n = 16) was isolated, suspended, fixed, blocked, surface-stained, permeabilized and labeled by incubation with conjugate antibodies. The labeled cells were selected by size and granularity, and then gated by cell-surface markers using flow cytometry. (A) Definition of gate R1 (SSClowFScow) cells, representative FACS plots and percentage of FScowCD45+ and SSClowCD3+ cells in gate R1 in brown, epididymal and inguinal fat pads. The dot plots depict SSC and FSC, and the sequential gating strategy for analysis of CD45+ and CD3+ cells are presented. (B) Representative FACS plots of Lin-IL-13+ ILC2s in gate R1 and percentage of ILC2s in CD45+ cells from gate R1 in three fat pads. (C) Representative FACS plots of CD1d-CD3+ NKT cells in gate R1 and percentage of NKT cells in CD45+ cells from gate R1 in three fat pads. (D) Representative FACS plots of γδ T cells in gate R1 and percentage of γδ T cells in CD45+ cells from gate R1 in three fat pads. (E) Definition of gate R2 (SSChighFScow) cells, representative FACS plots and percentage of CD45+ cells in gate R2 in three fat pads. (F) Representative dot plots and percentage of Siglec-5-C11b+ macrophages from CD45+ cells from gate R2. (G) Representative dot plots and percentage of Siglec-5-C11b+CD206+ to Siglec-5-C11b+ macrophages from gate R2 in three fat pads. The data were presented as the mean ± s.e.m. *P<0.05, **P<0.01. A full colour version of this figure is available at http://dx.doi.org/10.1530/JOE-16-0229.
The primer sequences were as follows: Arginase 1 forward primer 5′-CAGCTACCTGCTGGGAAGGAAGA-3′ and reverse primer 5′-CCAAGAGTTGGTTACCTCCA-3′; CD206 forward primer 5′-TGTTGGTGAAGCTGAAAGGTA-3′ and reverse primer 5′-CAGGTGTGGGCGGACGTAG-3′; F4/80 forward primer 5′-CTTTGGCTATGGGCGG-3′ and reverse primer 5′-GAACCTCCGCATTAATGG-3′; Nr3c2 forward primer 5′-GAAGAGTGGGAGTTGCTGTTG-3′ and reverse primer 5′-CGTGCCGCCTGGAGAACC-3′ and reverse primer 5′-CTTTGGCTATGGGCGG-3′ and reverse primer 5′-GAACCTCCGCATTAATGG-3′. Each primer pair amplifies products spanning several exons, distinguishing spliced mRNA from genomic DNA contamination. The relative expression of target genes was normalized to GAPDH.

Administration of IL-33

Six-week-old male C57BL/6 mice were fed with HFD described above for 11 weeks and then administered with recombinant murine IL-33 (carrier-free, BioLegend) at 100 µg/mice/day by i.p. injection under cold exposure (6°C) for 2 days. The mice were killed, and adipose tissue was collected for flow cytometry and Western blot analyses.

Administration of IL-33 receptor antibody

Seven-month-old male C57BL/6 mice were administered with or without mouse ST2/IL-1 R4 antibody (R&D Systems) at 100 µg/mice/day by i.p. injection under cold exposure (6°C) for 2 days. The mice were killed, and adipose tissues were collected for flow cytometry analysis and Western blotting.

Western blot analysis

Fat tissue was homogenized, and the protein was normalized and separated with SDS-PAGE gel followed by immunoblots as described previously (Liu et al. 2012, 2014). The rabbit polyclonal anti-UCP1 and mouse monoclonal anti-tubulin antibodies were from Abcam. The phospho-PKA substrate and TH antibodies were purchased from cell signaling. The secondary antibodies including anti-rabbit IgG (H+L) HRP conjugate and anti-mouse IgG (H+L) AP conjugate were from Promega.

Data analysis

Statistical analysis was performed using a two-tailed Student’s t-test between two groups or one-way ANOVA among three different groups. All the results were presented as the mean ± S.E.M., and P value of <0.05 was considered to be statistically significant.

Results

The abundance of ILC2s, NKT cells and γδ T cells is lower in BAT and sWAT compared with eWAT

To investigate if lymphocyte population in adipose tissue is correlated with thermogenesis, we performed flow cytometry analysis for SVF isolated from interscapular brown, epididymal and inguinal fat pads. The viable cells in SVF were selected based on size, and gate R1 (SSClow FSC low) was defined for analysis of lymphocytes (Fig. 1A). The fraction of CD45+ cells (nucleated cells of hematopoietic origin) accounted for 48, 78 and 58% of cells in gate R1 for brown, epididymal and inguinal fat, respectively (Fig. 1A). Consistently, the fraction of CD3+ cells (T lymphocytes) within gate R1 was also significantly lower in brown (25%) and inguinal (28%) fat compared with epididymal fat (37%) (Fig. 1A), suggesting that lymphocytes and T lymphocytes are enriched in eWAT. Since innate or innate-like lymphocytes including ILC2s, NKT and γδ T cells reside in adipose tissue and have been shown to regulate energy or glucose homeostasis in WAT (Lynch et al. 2009, 2012, Brestoff et al. 2014, Lee et al. 2015, Mehta et al. 2015), we further examined the abundance of these lymphocytes in brown fat. LinIL-13+(ILC2s), CD1d tetramer (CD1d tet)+ CD3+ (NKT) and γδTCR+CD3+ (γδ T) cells accounted for 3, 5 and 8% of total CD45+ cells within gate R1 in epididymal fat (Fig. 1B, C and D), suggesting the presence of an innate immune system within adipose tissue. In addition, the fractions of ILC2s, NKT and γδ T cells were significantly lower in brown and inguinal fat compared with epididymal fat (Fig. 1B, C and D). There was no significant difference in ILC2s, NKT and γδ T cell fractions between brown and inguinal fat (Fig. 1B, C and D).

Given that ILC2s and NKT cells as well as Th2 cells produce type 2 cytokines such as IL-4 and IL-13, well-known factors of eosinophil and M2 macrophage activation and promoters of WAT browning (Lynch et al. 2009, 2012, Brestoff et al. 2014, Han et al. 2014, Lee et al. 2015), we further investigated whether the macrophage...
subset differs between brown, epididymal and inguinal fat. The cells in gate R2 (granulocytes) were defined and used for flow cytometry analysis of macrophage and eosinophil subsets (Fig. 1E). CD45+ cells accounted for 16, 34 and 24% of cells in gate R2 for brown, epididymal and inguinal fat, respectively (Fig. 1E). The Siglec-5+CD11b+ (eosinophils, 37%) and Siglec-5-CD11b+ (macrophages, 39%) cells consisted mostly of CD45+ cells within gate R2 of epididymal fat (Fig. 1F). The percentage of eosinophil, the downstream effectors of ILC2 and NKT, was significantly greater in inguinal fat compared with brown and epididymal fat. Although the fraction of Siglec-5-CD11b+ macrophages within gate R2 was decreased in inguinal fat compared with epididymal fat, the percentage of Siglec-5-CD11b+CD206+ (M2) to Siglec-5-CD11b+ macrophages was greater in inguinal compared with brown fat and epididymal fat (Fig. 1F and G). Taken together, our results suggest that ILC2s, NKT cells and γδ T cells are present in BAT and sWAT as well as eWAT, and may play an important role in regulating the biology and function of adipose tissue.

The frequency of ILC2s and eosinophils is driven by sympathetic tone in beige fat

Activation of ILC2/eosinophil/macrophage axis is induced by cold exposure and promotes browning effect in WAT (Nguyen et al. 2010, Lee et al. 2015). Given that sympathetic tone mediates the cold-induced browning effect through norepinephrine in WAT (Farmer 2008, Barbatelli et al. 2010), we then examined whether the presence of these types of immune cells is driven by sympathetic innervation. We performed the denervation of inguinal fat using a local injection of 6-OHDA in 7-week-old male C57BL/6 mice followed by 2-week recovery and subsequent cold exposure. Pretreatment with 6-OHDA led to a marked reduction in basal and cold-induced norepinephrine production (Fig. 2A) as well as a robust downregulation of UCP1 expression in inguinal fat in vivo (Fig. 2B). Moreover, sympathetic denervation significantly suppressed basal and cold-induced fractions of ILC2s and eosinophils, while denervation had no significant effect on NKT and γδ T cell fractions (Fig. 2C). Consistent with this, inhibiting sympathetic tone significantly decreased basal and cold-induced IL-33 levels in inguinal fat (Fig. 2D). These data suggest that sympathetic tone plays a critical role in regulating IL-33 production and the ILC2/eosinophil axis in beige fat. Notably, sympathetic denervation increased the basal and cold-induced CD11b+ macrophages in beige fat despite a little effect on the fraction of CD206+ M2 macrophages under cold stress condition (Fig. 2E), suggesting that CD206+ M2 macrophages are not sensitive to sympathetic tone, and other M2 macrophage markers in addition to CD206 should be considered in the future study. These results indicate that IL-33 production and ILC2/eosinophil axis activation in beige fat are driven by norepinephrine, whereas M2 macrophages activation is not significantly altered by denervation.

The abundance of ILC2s, eosinophils and NKT cells in SVF of adipose tissue is decreased by HFD feeding in vivo

Previous studies show that the expression levels of UCP1 were significantly downregulated in sWAT by chronic HFD feeding (Fromme & Klingenspor 2011, Shen et al. 2014). In contrast to sWAT, BAT UCP1 had been shown to be induced by HFD feeding (Fromme & Klingenspor 2011, Shen et al. 2014). To further investigate the effect of HFD on UCP1 expression in BAT and sWAT, we performed time course study with HFD feeding for 1, 12, 20 and 32 weeks. We found that 1–12 weeks of HFD feeding upregulated the expression of UCP1 in BAT (Fig. 3B). Although UCP1 expression was slightly downregulated by 20-weeks HFD feeding, this downregulation was not significant until 32-weeks HFD feeding (Fig. 3B). Different from BAT, UCP1 expression in sWAT was not significantly affected in 1 week but significantly suppressed from 12 to 32 weeks, suggesting that UCP1 expression in adipose tissue is regulated by overnutrition in duration-dependent manner, and the response in WAT is distinct from BAT. (Fig. 3A and B). To investigate the importance of innate/innate-like lymphocytes in adipose tissue under HFD feeding conditions, we analyzed brown, epididymal and inguinal fat from mice fed with NCD or HFD for 12 weeks for the presence of ILC2s, NKT and γδ T cells using flow cytometry. The body mass was increased by 28% under HFD feeding (NCD mice average body weight 28.1 and 35.9 g for the mice fed with HFD (data not shown)). The fraction of CD45+ cells in gate R1 was also significantly increased by HFD feeding in all three types of fat (Fig. 3C), suggesting that lymphocyte abundance is positively correlated with the development of obesity. Consistent with this, the fraction of CD3+ cells in gate R1 was also slightly increased by HFD feeding in all three types of fat, although these differences did not reach statistical significance (Fig. 3D). Importantly, the fractions of ILC2s and NKT cells, but not γδ T cells, in CD45+ cells of gate R1 were markedly decreased in all
three types of fat upon HFD feeding (Fig. 3E, F and G), suggesting that HFD feeding may selectively suppress the recruitment of ILC2s and NKT cells rather than γδ T cells. In support of this, the percentage of eosinophils in CD45+ cells within gate R2 was also significantly decreased by HFD feeding despite an increase in total CD45+ cells in gate R2 in both brown and epididymal fat (Fig. 3H and I). These data point to a negative correlation between obesity and eosinophil abundance in adipose tissue. Notably, the infiltration of total macrophages into all three fat pads was markedly increased by HFD feeding (Fig. 3J), while the percentage of Siglec-5−CD11b+CD206+ macrophage was not significantly affected in any of three fat pads upon HFD feeding (Fig. 3K). Consistently, RT-PCR results show that the mRNA levels of macrophage marker F4/80, M1 marker Nr3c2 and M2 markers Arginase and CD206 are also induced by 12 weeks of HFD feeding (Fig. 3L), suggesting that macrophage differentiation and polarization may be associated with the stage of obesity development.

ILC2 and NKT cell fractions in SVFs are decreased in adipose tissue of ob/ob mice

To further evaluate the effect of obesity on the adipose innate immune system, we isolated adipose SVFs from 9-week-old male lean and ob/ob mice and performed flow cytometry analysis. The body mass was increased by 38% in ob/ob mice (31.4 g) compared with lean mice (22.7 g) (data not shown). Intriguingly, expression levels of UCP1 were downregulated in inguinal fat and in BAT of ob/ob mice compared with lean mice (Fig. 4A, B and C).

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in the development of obesity. In addition, the fraction of ILC2s and NKT cells rather than γδ T cells in gate R1 was significantly decreased in epididymal and inguinal fat pads of ob/ob mice compared with lean mice (Fig. 4F, G and H). ILC2 and NKT cell populations were slightly decreased in brown fat in ob/ob mice, but the difference did not reach statistical significance (Fig. 4F, G and H). Similar to HFD feeding conditions, where the CD45+ cell fraction in gate R2 was increased, leptin deficiency resulted in a significant decrease in the eosinophil fraction in all three fat pads (Fig. 4I and J). The infiltration of Siglec-5−CD11b+ macrophages was increased (Fig. 4K), while the ratio of Siglec-5−CD11b+CD206+ to Siglec-5−CD11b+ macrophages was not significantly affected in any of the fat pads in ob/ob mice compared with lean mice (Fig. 4L). Taken together, these data suggest a negative correlation between activation of ILC2/eosinophil axis, NKT cell infiltration and obesity.

IL-33 administration improves ILC2/eosinophil pathway and the expression of UCP1 and TH in HFD-fed mice

Next, we dissect the role of IL-33 in the suppression of ILC2/eosinophil axis and UCP1 by obesity given that IL-33 is a key inducting factor of ILC2s and WAT browning (Brestoff et al. 2014). We found that similar to UCP1 and ILC2/eosinophil pathway, mRNA and protein levels as well as circulating levels of IL-33 in inguinal WAT was induced by cold exposure and suppressed by HFD feeding and leptin deficiency-induced...
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Figure 4

The fractions of NKT and ILC2s in SVF were decreased in brown, epididymal and inguinal fat in ob/ob mice. Brown, inguinal and epididymal fat pads were collected from 9-week-old lean (n = 12) and ob/ob (n = 14) mice, and SVF from fat was isolated and used for flow cytometry analysis. The expression levels of UCP1 in inguinal (A) and brown (B) fat were downregulated in ob/ob mice compared with lean mice. Tubulin was used as the loading control. (C) The data in Fig. 4A and B were quantified and analyzed. The percentage of FSC<sup>−</sup>CD45<sup>+</sup> cells (D) but not SSC<sup>−</sup>CD3<sup>+</sup> cells (E) in gate R1 in three fat pads was significantly increased in ob/ob mice compared with lean mice. The percentage of ILC2s (F) and NKT cells (G) but not γδT (H) in CD45<sup>+</sup> cells from gate R1 in three fat pads was significantly decreased by leptin deficiency. (I) The percentage of SSC<sup>−</sup>FSC<sup>−</sup>CD45<sup>+</sup> cells in gate R2 in three fat pads was significantly increased in ob/ob mice compared with lean mice. (J) The fraction of Siglec-5<sup>−</sup>CD11b<sup>+</sup> eosinophils was decreased by leptin deficiency. The population of Siglec-5<sup>−</sup>CD11b<sup>+</sup> macrophage (K) but not Siglec-5<sup>−</sup>CD11b<sup>−</sup>CD206<sup>+</sup> (M2) to Siglec-5<sup>−</sup>CD11b<sup>+</sup> macrophage (L) was increased by leptin deficiency. The data were presented as the mean ± s.e.m. *P < 0.05, **P < 0.01.

Figure 5

Obesity (Fig. 5A, B and C), indicating that IL-33 downregulation may mediate the suppressing effect of obesity on ILC2/eosinophil axis and UCP1 in WAT. To address this question, 6-week-old male mice were fed with HFD for 11 weeks and then injected with vehicle or IL-33 for 7 days. The flow cytometry analysis showed that IL-33 administration notably increased the fraction of ILC2 in sWAT and eWAT about 13- and 9-folds, respectively, in HFD-fed mice (Fig. 5D and E). Consistent with this, IL-33 treatment significantly upregulated the expression levels of UCP1 in sWAT and eWAT (Fig. 5F). In agreement with this, IL-33 induced the expression of TH, a key enzyme of catecholamine biosynthesis and downstream of ILC2/eosinophil pathway in sWAT and eWAT of HFD-fed mice (Fig. 5G and H) (Tsao et al. 1998, Brestoff et al. 2014, Rao et al. 2014). Moreover, the phosphorylation of PKA substrates was enhanced by IL-33 administration in eWAT and sWAT (Fig. 5G and H).

These results suggest that IL-33 treatment greatly induces the WAT browning. However, IL-33 treatment had no significant effect on PKA substrate phosphorylation and UCP1 expression in BAT despite the increased ILC2 frequency (data not shown). Taken together, these results suggest that IL-33 plays a mediatory role in suppression of ILC2s/eosinophils and WAT browning by obesity, and downregulation of IL-33 in WAT is a potential marker for obesity.

Cold-induced ILC2/eosinophil axis and expression of UCP1 and TH were suppressed by neutralization of IL-33 signaling

To further determine the role of IL-33 in regulation of catecholamine production and thermogenesis, 7-month-old male mice were injected with vehicle or ST2 antibody under cold exposure condition for 2 days.
Consistent with a previous finding of significant decrease in CD90^+ CD25^+ ILCs in both frequency and total cell number in the lungs by ST2 antibody administration (Monticelli et al. 2011), blocking IL-33 signaling decreased the fraction of ILC2 and eosinophil in sWAT and eWAT under cold stress condition (Fig. 6A and B). Moreover, neutralization of IL-33 signaling significantly downregulated the expression levels of UCP1 (Fig. 6C). In line with this, the expression levels of TH as well as PKA substrate phosphorylation were suppressed by the administration of ST2 antibody in sWAT and eWAT under cold stress condition (Fig. 6D and E). However, ST2 antibody administration had no inhibitory effect on PKA substrate phosphorylation and UCP1 expression despite a decrease in ILC2 fraction in BAT (data not shown). These results suggest that the induction of IL-33 is required for catecholamine production and WAT browning. However, the functional role of IL-33-driven ILC2 pathway in BAT is distinct from that in WAT.

**Discussion**

Type 2 inflammatory responses have been shown to play an important role in regulating browning of WAT through M2 macrophage activation and type 2 cytokines release in eWAT and sWAT (Molofsky et al. 2013, Nussbaum et al. 2013, Brestoff et al. 2014, Khalifeh-Soltani et al. 2014, Rao et al. 2014). ILC2s reside in adipose tissue and drive the browning of WAT by producing type 2 cytokines including IL-5 and IL-13 in WAT (Lee et al. 2015). However, how the recruitment and activation of adipose-resident ILC2s are regulated remains unclear. In this study, we show that ILC2s are present in BAT and sWAT, and enriched in eWAT (Fig. 1). Moreover, the fraction of ILC2s is significantly decreased in BAT, sWAT and eWAT by HFD feeding and leptin deficiency (Figs 3 and 4). In addition, the abundance of ILC2/eosinophil axis as well as the levels of IL-33 and UCP1 are induced by cold exposure and suppressed by obesity in sWAT (Figs 2, 3 and 4). Given that ILC2s and eosinophils have been shown to promote the browning of WAT under cold stress (Brestoff et al. 2014, Khalifeh-Soltani et al. 2014, Rao et al. 2014, Lee et al. 2015), our data suggest that an ILC2/eosinophil axis exists in BAT as well as WAT and is regulated by both sympathetic tone and obesity.

IL-33 is a critical factor for the maintenance and stimulation of ILC2s in adipose tissue (Brestoff et al. 2014), and the circulating level of IL-33 has been shown to be downregulated by obesity, and IL-33 administration...
Improves obesity and obesity-induced insulin resistance in vivo (Miller et al. 2010, Hasan et al. 2014, Han et al. 2015), implying that obesity may decrease ILC2 frequency in adipose tissue through downregulation of IL-33 production. Our study shows that IL-33 administration improves the percentage of ILC2s and eosinophils, catecholamine production and thermogenic gene expression in sWAT and eWAT of HFD-fed mice (Fig. 5). Our study suggests that ILC2-inducing cytokine IL-33 mediates the suppressing effect of obesity on ILC2/eosinophil axis and thermogenic gene expression in WAT, and is a potential marker of obesity. Consistent with this, neutralization of IL-33 signaling suppressed ILC2/eosinophil axis as well as catecholamine production and UCP1 expression in WAT under cold stress condition (Fig. 6). However, some beneficial effects of IL-33 are limited to WAT given that the promoting effects of IL-33 on catecholamine production and thermogenic gene expression are not observed in BAT despite the increased fraction of ILC2s in HFD-fed mice (Fig. 5 and data not shown). Our data are consistent with Brestoff’s report that IL-33 treatment has no promoting effect on UCP1 expression, while increases ILC2 fraction in BAT (Brestoff et al. 2014). Although ILC2/eosinophil axis is suppressed by obesity in all three types of fat, IL-33/ILC2 pathway may have distinct effects in BAT compared with those in WAT. Our study also suggests that in addition to ILC2/eosinophil axis, many other factors may be involved in the regulation of BAT activation. Along with this, adipose UCP1 expression is affected by HFD feeding in duration and tissue-dependent manners, and this response in WAT is distinguished from BAT (Fig. 3A and B).

Although IL-33 has been defined as an epithelial cell-derived cytokine, recent studies also report that IL-33 as well as its receptor ST2, a protein that is encoded by the IL1RL1 gene in human adipose tissue, is predominantly detectable in endothelial cells (Price et al. 2010, Zeyda et al. 2013). It remains to be clarified if obesity suppresses IL-33 production through targeting of epithelial cells or endothelial cells. On the other hand, alternative macrophage activation is a downstream event of ILC2/eosinophil pathway. However, the ratio of Siglec-5-CD11b-CD206- M2 macrophages to Siglec-5-CD11b+ macrophages is not significantly affected by HFD feeding.

Figure 6
Administration of ST2 antibody downregulated ILC2/eosinophil pathway and expression levels of TH and UCP1 in WAT. The percentages of ILC2s (A) and eosinophils (B) in CD45+ cells from gate R1 and gate R2, respectively, in sWAT and eWAT were significantly decreased by ST2 antibody injection. (C) The protein levels of UCP1 in sWAT and eWAT were downregulated by ST2 antibody administration. The expression levels of TH and phosphorylation of PKA substrates in sWAT (D) and eWAT (E) were downregulated by ST2 antibody administration. Tubulin was used as the loading control. The data in Fig. 6A and B were presented as the mean ± s.e.m. *P<0.05, **P<0.01.
and leptin deficiency (Figs 3 and 4). Moreover, the mRNA levels of both M1 and M2 markers are induced by HFD (Fig. 3L). Given that obesity leads to a shift in the activation state of adipose tissue macrophages from an M2-polarized state to an M1 proinflammatory state that contributes to insulin resistance (Lumeng et al. 2007), our data suggest that the effect of HFD feeding on differentiation and polarization of macrophage may be dynamic and needs to be further clarified. In addition, other markers of M2 macrophage such as galactose (Mgl-1) receptors, resistin-like molecule (RELM)-α and chitinases Ym1 and Ym2 should be included for the determination of M2 macrophage activation in future studies (Claria et al. 2011). Similar to HFD-fed mice, leptin deficiency leads to a decrease in ILC2/eosinophil activity axis and a downregulation of UCP1 expression in adipose tissue. Since leptin itself has been shown to positively regulate sympathetic tone and energy expenditure (Myers et al. 2009), the decreased browning and ILC2/eosinophil axis in fat depots of ob/ob mice may be resulted from leptin deficiency itself as well as leptin deficiency-induced obesity as the secondary effect. In agreement with this, the ILC2 population is decreased in epididymal fat of obese rodents and human subjects (Brestoff et al. 2014). The browning of WAT is controlled by sympathetic innervation in a β3-adrenoceptor signaling pathway-dependent manner (Farmer 2008, Barbatelli et al. 2010). However, whether the abundance of ILC2s is regulated by norepinephrine remains unknown. Our study shows that inhibiting sympathetic tone by denervation impairs the abundance of ILC2s and expression of UCP1 in sWAT (Fig. 2), suggesting that β3-adrenoceptor signaling pathway plays an important role in regulating ILC2s in adipose tissue. Consistently, it has been shown that sympathetic tone significantly suppresses the level of IL-33 in sWAT (Fig. 2D). It has been shown that cytokines IL-33 and IL-25 as well as cold stress promotes the recruitment of ILC2s and browning effect in WAT (Hams et al. 2013, Brestoff et al. 2014, Lee et al. 2015). In addition, Lee and coworkers reported that environmental cold induces the recruitment of ILC2s via IL-33 (Lee et al. 2015). Our data suggest that activated sympathetic tone may promote ILC2 recruitment by upregulating the production of IL-33 in epithelial cells or other types of cells in response to cold or other environmental stimuli. In agreement with this, loss of IL-33 signaling downregulates ILC2/eosinophil pathway, norepinephrine biosynthesis and thermogenic gene expression in WAT under cold stress condition (Fig. 6). However, whether IL-33 in response to cold or overnutrition is locally produced in fat or enters through the circulation needs to be established in the future. While activation of the ILC2–eosinophil axis has been shown to drive M2 macrophage polarization and browning (Lee et al. 2015), our data show that sympathetic tone plays a critical role in cold-induced browning and ILC2–eosinophil axis activation, but not M2 macrophage polarization, suggesting that ILC2/eosinophil activation and M2 macrophage activation may be disassociated and mediated through two independent mechanisms under some particular conditions. Consistent with this, Brestoff and coworkers reported that ILC2s themselves promote the browning of fat through self-derived factor enkephalin, but do not confer browning through macrophage-dependent mechanisms (Brestoff et al. 2014).

In summary, our study identifies sympathetic tone as a key regulator of ILC2/eosinophil axis in beige fat. In addition, inactivation of IL-33/ILC2/eosinophil pathway is a novel hallmark of obesity, and this pathway may play an important role in regulating thermogenesis and obesity development.

Declaration of interest
The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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