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Spermidine/spermine N1-acetyltransferase-mediated polyamine catabolism regulates beige adipocyte biogenesis

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Abstract

Objective: Cold and β 3-adrenergic receptor (AR) agonists activate beige adipocyte biogenesis in white adipose tissue (WAT). The two stimuli also induce expression of inflammatory cytokines in WAT. The low-grade inflammation may further promote WAT browning. However, the mechanisms to reconcile these two biological processes remain to be elucidated. In this study, we aim to investigate the roles of the rate-limiting polyamine catabolic enzyme spermidine/spermine N1-acetyltransferase (SAT1) in regulating beige adipocyte biogenesis and inflammation.

Methods: Adipose-specific SAT1 knockout mice (SAT1-aKO) were generated by crossing adiponectin-cre to SAT1-lox/lox mice. Metabolic phenotype was investigated. Primary preadipocytes were isolated from inguinal WAT (iWAT) and differentiated to adipocytes for studying beige adipocyte biogenesis.

Result: The expression and enzymatic activity of SAT1 were up-regulated in iWAT upon cold and β 3-AR stimulation. SAT1-aKO mice developed late-onset obesity on a high-fat diet with impaired cold-induced beige adipocyte biogenesis and energy expenditure. RNA-seq analysis of iWAT from cold-challenged SAT1-aKO mice revealed that, in addition to beige adipocyte

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Author contributions

FY, LZ, YC, WG, CZ and YF conducted experiments and analyzed results; KZ and MS generated SAT1 floxed mice. FY and QY wrote the manuscript; XL, ZF and QY conceived the experimental design and supervised the studies.

†Fang Yuan and Lin Zhang contributed equally to this work.

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Conflict of Interest

The authors have no conflict of interest to declare.

biogenesis signatures, the immune response markers were highly enriched among reduced genes. In cultured adipocytes, SAT1 overexpression or pharmacological activation with N1, N11-diethylnorspermine (DENSp_m) elevated oxygen consumption and increased the expression of beige adipocyte marker UCP1 and PGC-1 α . DENSp_m treatment of adipocytes also increased the expression of inflammatory genes. SAT1 activation enhanced hydrogen peroxide production in adipocytes. Antioxidant N-acetylcysteine abrogated the elevated UCP1 expression and reversed some inflammatory genes induced by SAT1 activation.

Conclusions: SAT1 activation plays a key role in cold and β 3-AR agonist-induced beige adipocyte biogenesis and low-grade inflammation.

Keywords

Polyamine metabolism; beige adipocytes; inflammation; obesity; energy expenditure

1. Introduction

Polyamines including putrescine, spermidine and spermine are small polycations that are essential for multiple cellular functions affecting cells growth, cancer and ageing [1–3]. Polyamine metabolism is tightly regulated by several rate-limiting enzymes for synthesis and catabolism. Specifically, the polyamine synthesis is modulated by ornithine decarboxylase (ODC) and adenosylmethionine decarboxylase (AMD1). The catabolism is controlled by spermidine—spermine N1-acetyltransferase (SAT1), which acetylates spermidine and spermine to generate N1-acetylspermine, N1,12-diacetylspermine, and N1-acetylspermidine. These acetylated polyamines are either excreted intact in urine or oxidized by acetyl-polyamine oxidase (APAOX) [2, 4]. The oxidation of acetylpolyamines generates putrescine and spermidine as well as hydrogen peroxide (H₂O₂) [2, 4]. The putrescine and spermidine are then returned to the polyamine metabolic flux.

Initial polyamine research was mostly focused on cancer biology since ODC and SAT1 were found to be overexpressed in multiple cancers [2, 4–6]. Subsequent studies on genetic mouse models, especially SAT1 transgenic and knockout mice, revealed that polyamine metabolism plays a major role in regulating energy expenditure and adiposity [7–9]. Whole body SAT1 knockout mice develop late-onset obesity while systemic and adipose-specific SAT1 overexpression prevent diet-induced obesity. PGC-1 α expression and AMPK activity are altered in white adipose tissue (WAT) with SAT1 overexpression or knockdown [7–9], but it is unclear to what extent they contribute to the SAT1-regulated energy metabolism and adiposity.

Brown adipose tissue (BAT) expresses high levels of uncoupling protein 1 (UCP1) in order to dissipate chemical energy to heat, therefore, plays a major role in thermogenesis. WAT is traditionally considered to be an energy storage organ. However, it is becoming clear that clusters of adipocytes in WAT undergo browning or beiging with induced expression of UCP1 in response to various environmental, hormonal or pharmacological stimuli such as cold and β 3-adrenergic receptor (AR) activation [10–13]. Interestingly, cold and β 3-AR also induce expression of inflammatory cytokines in adipocytes and recruit immune cells to WAT [14–16]. Although adipose inflammation has been linked to insulin resistance [17, 18], low

grade inflammation appears to be required for adipose tissue remodeling and beige adipocyte biogenesis [19–21]. Therefore, cold and β 3-AR agonist exert direct and indirect effects on activating WAT browning. However, the downstream signals for the dual effects remain to be fully elucidated. In the current paper, we show that adipose SAT1 is induced by cold and β 3-AR agonist stimulation. Importantly, SAT1 appears to be a key mechanistic node for both beige adipocyte biogenesis and low-grade inflammation induced by cold and β 3-AR stimulation.

2. Materials and Methods

2.1. Animal Studies

To generate adipose-specific SAT1 knockout mice, SAT1-flox/flox mice [22] were crossed to adiponectin-Cre mice (Jackson Lab). For diet-induced obesity, mice were fed a high fat diet containing 54.8% fat calories, 24.0% carbohydrate calories, and 21.2% protein calories (4.8 kcal/gram) (stock number TD.93075; Envigo Inc.) [23]. Body composition was analyzed using an EchoMRI 3-in-1 instrument (Echo Medical Systems, Houston, TX). Metabolic cage studies were performed as described previously [23].

For the cold-induced thermogenesis studies, mice were housed in a cold chamber (4°C) for 4, 12 or 72 hours. For the β 3-adrenergic agonist treatment, mice received intraperitoneal injections of CL316, 432 (C5976; Sigma) at 1mg/kg per day for 4 days. Control mice received same volumes of phosphate-buffered saline.

Glucose tolerance test was performed by injecting glucose (1g/kg) intraperitoneally to mice with food removal for 5 hours. Blood glucose levels were measured at 0, 15, 30, 60, and 120 min. Mice were maintained under a 12hr light/12hr dark cycle at constant temperature (23°C) with free access to food and water, except for the cold exposure experiments. All mouse studies were conducted in accordance with federal guidelines and were approved by the Institutional Animal Care and Use Committee of University of California, Irvine.

2.2. Isolation and Culture of Primary Adipocytes

Isolation and culture of primary adipocytes were described previously [24]. Inguinal white adipose tissue excised from 4 to 5-week-old male C57BL/6 mice was minced and digested in a PBS digestion buffer containing 2.5 U/mL collagenase D (11088882001; Sigma), 2.4 U/mL Dispase II (4942078001; Sigma), and 10 mM CaCl₂ at 37°C in a shaking water bath for 18 minutes. The digested tissue was filtered through 100 μ m nylon cell strainers (BD Biosciences). The stromal vascular fraction cells containing preadipocytes were separated from floating primary adipocytes by centrifugation at 600g for 5 minutes. The resulting pellets were resuspended and further filtered using 40 μ m nylon cell strainers (BD Biosciences). Stromal vascular fraction cells were maintained in DMEM/F12 GlutaMAX (Invitrogen) containing 15% FBS, 100U/mL penicillin, and 100 μ g/mL streptomycin. For differentiation, confluent preadipocytes were treated with medium containing 15% FBS, 0.5 mM isobutylmethylxanthine (I7018; Sigma), 1 μ M dexamethasone (D4902; Sigma), 2 μ g/mL insulin (I0546; Sigma), and 1 μ M rosiglitazone (R2408; Sigma) for 48 hours. Adipocytes were then maintained in medium containing 15% FBS and 2 μ g/mL insulin. At the 7 to 8-

day of differentiation, cells were incubated with CL 316,243 (C5976; Sigma) at 10 μ M for up to 8hrs or treated with N1, N11-Diethylnorspermine tetrahydrochloride (4A/194732; Tocris Bioscience) for 24 hours. For experiments of the PKA/CREB inhibitor, the cells were treated with 50 μ M H-89 (S1582; Selleckchem) for 1 hour prior to the stimulation of the CL316, 432. In the antioxidant experiment, adipocytes were pre-treated with 100 μ M N-acetylcysteine (NAC) (Sigma) for 30 mins before DENSp treatment. Lentiviral overexpression of SAT1 was performed as previously described [25]. Oxygen consumption rate (OCR) of adipocytes was determined using an XF24 Extracellular Flux Analyzer (Seahorse Bioscience).

2.3. SAT1 Activity Assay

Snap-frozen tissues (50–100mg) were homogenized in Tris/EDTA (pH 7.0) buffer. The soluble supernatants after centrifugation were used in activity assay. SAT1 activity was assayed radiochemically as described [5, 23] and expressed as pmol of N1-[14C]-acetylspermidine (AcSpd) generated/10 min/mg of tissue.

2.4. RNA extraction and quantitative PCR

Total RNA was extracted using the RNeasy kit (QIAGEN). cDNA was synthesized using the SuperScript III first-strand synthesis supermix for quantitative RT-PCR (qRT-PCR; Invitrogen) and used in real-time PCRs with Power SYBR Green PCR master mix (Applied Biosystems) on a 7900HT real-time PCR system (Applied Biosystems). The relative gene expression levels were calculated by the 2Ct method using Tata-binding protein (Tbp) as an internal control. Primer sequences are shown in Supplemental Table 1.

2.5. Total reactive oxygen species (ROS) and hydrogen peroxide measurements

Intracellular ROS production was measured based on the oxidation of 2',7'-dichlorofluoresce diacetate (Invitrogen) as previously described [26]. Adipocytes were seeded into a black 96-well plate at a density of 1 \times 10⁵/well. After treatment, the cells were incubated with 2',7'-dichlorofluoresce diacetate at 37°C for 30 minutes, followed by washing three times with PBS.

The resultant fluorescence was measured using a fluorescent microplate reader at excitation and emission wavelengths of 485 and 528 nm, respectively.

Amplex Red assay was used to measure hydrogen peroxide levels. Adipocytes were incubated in the Krebs-Ringer phosphate (KRP) (145 mM NaCl, 5.7 mM sodium phosphate, 4.86 mM KCl, 0.54 mM CaCl₂, 1.22 mM MgSO₄, 5.5 mM glucose, pH 7.35) containing the Amplex Red reagent (A22188, Invitrogen). Fluorescence was then measured at excitation 485nm and emission 570nm, respectively.

2.6. RNA-seq Analyses

Total RNA isolated from iWAT of cold-treated SAT1-aKO and control mice was used for RNA-seq analyses. RNA-seq was performed at the UCI Genomic High-Throughput Facility. The Chipster software was used for quantification and statistics of the RNA-seq data [27].

Functional clustering was performed using DAVID analysis, Panther pathway and GO biological process analysis.

2.7. Immunohistochemistry

Tissues collected from cold-challenged SAT1-aKO and control mice were fixed in paraffin. Embedded sections were routinely deparaffinized and endogenous peroxidase was quenched with 3% H₂O₂ in 1×PBS. Then they were stained with antibody against F4/80 (Bio-Rad) to detect macrophages. Images were captured using a Zeiss Detector Imaging System.

2.8. Statistical Analysis

All data were expressed as mean ± SEM. Statistical differences were assessed using unpaired Student's two tailed t tests for two groups and a one-way ANOVA for three groups or more. Statistical significance was assumed at P < 0.05.

3. Results

3.1 Adipose SAT1 is induced by thermogenic stimuli

Cold and β 3-AR agonist are the classic stimuli to induce beige adipocyte biogenesis and BAT activity. Challenging wild type mice with cold for 72 hours strongly stimulated UCP1 and PGC-1 α expression in iWAT and BAT (Supplementary Fig. 1A–1B). β 3-AR agonist CL-316,243 treatment for 4 days also induced UCP1 expression in iWAT and BAT (Supplementary Fig. 1C–1D). Both Cold and CL-316,243 increased the expression of SAT1 in iWAT and BAT (Fig. 1A–1D). The polyamine synthetic enzyme ODC was elevated in iWAT and tended to be increased in BAT of mice exposed to cold. The acetyl polyamine oxidation enzyme APAOX was stimulated in iWAT, but not in BAT by cold and CL-316,243 (Fig. 1A–1D). Consistent with the increased SAT1 mRNA expression, SAT1 enzymatic activity was enhanced in iWAT and BAT by the stimulation of cold and CL-316,243 (Fig. 1E–H). It has to be noted that the baseline SAT1 activity at the room temperature in BAT was approximately 5 times higher than that in iWAT (Fig. 1E–1H). This is partially due to the higher SAT1 mRNA expression in BAT (Supplementary Fig. 1E). These results indicate that SAT1 expression and activity are linked to adipose thermogenesis.

To test whether these effects were cell-autonomous, we isolated preadipocytes from iWAT, differentiated the cells to adipocytes, and treated the cells with CL-316,243. SAT1 expression was rapidly induced by CL-316,243 in 1 hour. The maximum effects were at 4-hour time point. UCP1 expression, on the other hand, did not increase after 1 and 2 hours but elevated at later time points (4 and 8-hour) with CL-316,243 treatment (Fig. 1I). The rapid SAT1 induction could be recapitulated in vivo as acute cold exposure for 4 hours increased SAT1 in BAT and for 12 hours increased SAT1 in both BAT and iWAT (Fig. 1J–1K). The increased SAT1 levels were associated with elevated UCP1 expression in adipose tissue of acute cold-challenged mice (Supplementary Fig. 2A–2B). In addition to cold and adrenergic stimulation, high fat diet (HFD) feeding initially promotes the adipose UCP1 expression [28, 29] (Supplementary Fig. 3A–3B). Interestingly, one-week HFD feeding also elevated adipose SAT1 expression (Supplementary Fig. 3A–3B). The SAT1 induction in adipocytes was partially caused by protein kinase A (PKA) activation because the CL-316,243-induced

SAT1 elevation was attenuated by H-89, a PKA inhibitor (Fig. 1L). In addition, CL-316,243 and cold also increased the expression of adipose nuclear factor (erythroid-derived 2)-like 2 (Nrf2), the major transcription factor regulating SAT1 expression (Supplementary Fig. 4A–4B). These results indicate that thermogenic stimuli enhance polyamine catabolism in adipose tissue.

3.2 Adipose-specific SAT1 knockdown mice develop late-onset obesity

To determine the effects of adipose SAT1 on systemic energy metabolism, we generated adipose-specific SAT1 knockout mice (SAT1-aKO) by crossing adiponectin-Cre to SAT1-flox/flox mice [22]. SAT1 expression was reduced by 87% in BAT, but only 49% and 46% in iWAT and perigonadal WAT (pgWAT), with no changes in liver (Fig. 2A). The relatively modest reduction of SAT1 in WAT was likely caused by abundant SAT1 expression in stromal vascular fraction since isolated adipocytes from iWAT and pgWAT of SAT1-aKO mice showed 80–95% SAT1 reduction (Fig. 2A). The SAT1-aKO mice developed late-onset obesity after 18 weeks of high fat diet (HFD)-feeding (Fig. 2B). The increased adiposity in SAT1-aKO mice was confirmed by Echo-MRI studies showing a 20% increase in relative fat mass and a 9% decrease in relative lean mass (Fig. 2C). Consistently, the weight of iWAT and pgWAT of SAT1-aKO mice was higher than that of controls (Fig. 2D). Metabolic cage studies showed a reduction of VO₂ and VCO₂ in SAT1-aKO mice compared with controls when exposed to cold (Fig. 2E–2G). Glucose tolerance tended to be impaired in SAT1-aKO mice compared with controls before the body weight started to diverge (Fig. 2H–II). However, insulin tolerance test (not shown), insulin, glucose and insulin-glucose products (Supplementary Fig. 5A–5C) were not different between SAT1-aKO and control mice fed a HFD for 22–24 weeks when the body weight was separated. These results indicate that adipose SAT1 plays a major role in regulating systemic energy metabolism.

3.3 Adipose SAT1 regulates beige adipocyte thermogenesis

Since SAT1 was induced by cold and β 3-AR agonist in BAT and iWAT and SAT1-aKO mice showed reduction in cold-induced oxygen consumption, we measured the expression of thermogenic gene markers in BAT and iWAT. Upon cold challenge, the expression of beige adipocyte markers uncoupling protein 1 (UCP1), Cidea, peroxisome proliferator-activated receptor gamma coactivator 1-alpha (PGC-1 α), cytochrome c oxidase polypeptide 7A1 (Cox7a1), acyl-CoA Oxidase 1 (ACOX1), carnitine palmitoyltransferase 1 (Cpt1), cluster of differentiation 36 (CD36) were reduced in iWAT of SAT1-aKO mice compared with controls (Fig. 3A). Peroxisome proliferator-activated receptor gamma (PPAR γ) expression was not altered, suggesting SAT1 knockdown did not affect adipocyte differentiation. Interestingly, BAT thermogenic activity was not significantly affected by SAT1 knockdown (Fig. 3B). The only reduced gene in BAT of SAT1-aKO was PGC-1 α while other thermogenic genes including UCP1 were not altered (Fig. 3B). Therefore, adipose SAT1 is necessary for cold-induced beige adipocyte biogenesis.

We next investigated whether the SAT1 directly activates beige adipocyte biogenesis. N1, N11-diethylnorspermine (DENSp_m) is a polyamine analog that increases SAT1 expression and activity [2, 9]. Treating cultured adipocytes with DENSp_m promoted the expression of beige adipocyte biogenesis markers and increased oxygen consumption (Fig. 3C–3D).

Similarly, overexpression of SAT1 in adipocytes also stimulated beige adipocyte biogenesis (Fig. 3E, Supplementary Fig. 6). SAT1 activation did not affect adipocyte differentiation as evidenced by unaltered PPAR γ expression. These results indicate that adipose SAT1 is necessary and sufficient to promote beige adipocyte thermogenesis.

3.4 Adipose SAT1 knockdown reduces immunity-related genes

To investigate the mechanisms for SAT1-regulated beige adipocyte thermogenesis, we performed RNA-seq and subsequent DAVID functional annotation clustering analysis as well as Panther pathway and GO biological process analysis using iWAT from cold-challenged SAT1-aKO and control mice. Both analysis tools consistently showed over-represented signatures for immunity/inflammation and mitochondrial function/TCA cycle among the downregulated genes (Fig. 4A–4B and Supplementary Table 2–3). Among the upregulated genes, the top three clusters were membrane, glycoprotein and endoplasmic reticulum (Supplementary Table 4). The immunity-related genes included cluster of differentiation (CD) markers, chemokines (C-C and C-X-C motif) and cytokines (Supplementary Table 5). These genes represent signatures of immune cells especially T-lymphocytes (CD3, CD4, Foxp3), dendritic cells (CD83), and macrophages (IL-1R), although adipocytes also express low levels of some of these genes [17, 18, 30]. Quantity RT-PCR of selected immunity-related genes confirmed the reduced expression of cluster of differentiation 14 (Cd14), tumor necrosis factor alpha (Tnf- α), chemokine (C-C motif) ligand 8 (Ccl8), forkhead box P3 (Foxp3), C-X-C chemokine receptor type 6 (Cxcr6), and C-X-C chemokine receptor type 3 (Cxcr3) identified from the RNA-seq analysis (Fig. 4C). In addition, IL-1 β and IL-10 were also decreased and IL-6 and MCP-1 tended to be lower in iWAT of SAT1-aKO mice (Fig. 4C). Consistently, immunohistochemistry staining of iWAT showed a reduction of F4/80 positive macrophage infiltration in iWAT of cold-challenged SAT1-aKO mice compared with controls (Fig. 4D, Supplementary Fig. 7A). Some of the cytokines such as CD14 and IL6 were also reduced in the perigonadal WAT of SAT1-aKO mice compared with controls (Supplementary Fig. 7B). Therefore, adipose SAT1 plays a dual role in regulating beige adipocyte thermogenesis as well as immune cell infiltration.

3.5 SAT1-induced reactive oxygen species regulate beige adipocyte thermogenesis and inflammation

SAT1-activated polyamine flux generates H₂O₂ from oxidation of acetylpolyamines [2, 4]. Reactive oxygen species (ROS) has been shown to mediate beige adipocyte biogenesis and UCP1 expression/activation. ROS also induces cellular inflammation in adipocytes [31, 32]. Consistently, treatment of cultured adipocytes with DENSp_m increased H₂O₂ and total ROS levels, both were neutralized by antioxidant agent N-acetylcysteine (NAC) (Fig. 5A). The elevated H₂O₂/ROS was necessary for the UCP1 expression induced by SAT1 activation since NAC abolished the elevated UCP1 induced by DENSp_m (Fig. 5B). SAT1 activation by DENSp_m also increased Cd14, Tnf- α , Mcp-1 and IL-6 expression in adipocytes (Fig. 5C). NAC reversed the elevated expression of CD14, and tended to reduce Tnf- α and IL-6 elevation (Fig. 5C). DENSp_m-induced expression of UCP1, Tnf- α , Mcp-1 and IL-6 was attenuated by SAT1 siRNA knockdown, indicating the specificity of DENSp_m-activated SAT1 in stimulating beige adipocyte biogenesis and inflammation (Supplementary Fig. 8A–

8B). These data indicate that SAT1-induced ROS production is involved in beige adipocyte thermogenesis and inflammation.

4. Discussion

Adipose SAT1 plays a major role in systemic energy metabolism. However, it is not clear how SAT1 is regulated in adipose tissue. We found that thermogenic stimuli such as cold and β 3-AR agonist induce SAT1 expression and activity in adipose tissue. This appears to be partially mediated by PKA/CREB signal pathway since PKA inhibitor H-89 reduces SAT1 elevation stimulated by β 3-AR agonist (Fig. 1J). Consistently, there is a consensus CREB binding site in the SAT1 promoter [2]. Cold and β 3-AR agonist also stimulate the expression of Nrf2 (supplementary Fig. 1E–1F), a major transcription factor that binds to the polyamine-responsive element of the SAT1 promoter [2]. The elevated Nrf2 likely represents a response to intracellular ROS stimulated by cold and β 3-AR in adipose tissue [33, 34]. Interestingly, SAT1 activation also generates ROS. Therefore, Nrf2 and SAT1 may form a regulatory loop in adipose tissue upon cold or β 3-AR stimulation.

Four SAT1-related mouse models have been reported. Whole body SAT1 knockout mice are obese, while systemic SAT1 overexpression causes leanness [7–9, 35, 36]. Mouse aP2 promoter driven SAT1 transgenic and knockout show similar adiposity phenotype as that in systemic transgenic and knockout mice [8, 35]. However, aP2 promoter activity is not adipose-specific and drives target gene expression in non-adipose tissues/cells such as macrophages [37]. In the current studies, we used adiponectin-Cre to generate truly adipocyte-specific SAT1 knockout mice. The fact that SAT1-aKO developed late-onset obesity provided a conclusive evidence that SAT1 in adipocytes plays a critical role in energy metabolism and adiposity. However, the weight gain of our SAT1-aKO mice appears to be milder than that of whole body and aP2-Cre driven SAT1 knockout mice. It therefore remains possible that SAT1 in other tissues/cells might also play some roles in regulating energy expenditure and adiposity.

One major contribution of our study is the elucidation of the novel role of SAT1 in beige adipocyte biogenesis. SAT1 is sufficient and necessary for beige adipocyte biogenesis. Since SAT1 expression and activity are induced by cold and β 3-AR agonist, SAT1 activation is a critical downstream mediator of cold and β 3-AR stimulation for beige adipocyte biogenesis. This is partially mediated by the direct effects of SAT1 on adipocytes as SAT1 activation increased UCP1 expression and oxygen consumption (Fig. 3C–3E). SAT1 activation generates H₂O₂ and ROS through APAOX and spermine oxidase (SMOX) [1–3]. This appears to be a mechanism for the direct effects of SAT1 on beige adipocyte biogenesis since neutralizing H₂O₂ with NAC abolishes UCP1 elevation induced by SAT1 activation. Although pathologically high ROS are detrimental to cellular function, physiological ROS are important second messengers for multiple biological processes including beige adipocyte biogenesis and activity [31]. For example, overexpression of Sestrin2, an antioxidant protein, suppresses ROS and inhibits UCP1 expression in adipose tissue [32]. A more recent study show that ROS may induce UCP1 cysteine-253 sulfenylation, thereby promote adipocyte thermogenesis [34]. Therefore, SAT1-activated ROS production is an important mechanism for beige adipocyte biogenesis.

RNA-seq analyses surprisingly reveal highly enriched immunity-related genes in addition to mitochondrial function/beige adipocyte markers among the reduced genes in adipose tissue of cold-challenged SAT1-aKO compared with control mice (Fig. 4). The gene markers of T-lymphocytes, dendritic cells, and macrophages are reduced, suggesting decreased immune cell infiltration into adipose tissue of SAT1-aKO mice. Interestingly, knocking out SAT1 from renal proximal tubule also decreases immune cell infiltration and inflammatory response to ischemia/reperfusion injury [38]. The mechanism for the reduced immune cell infiltration is not very clear. SAT1 activation induces ROS and cytokine production in adipocytes. This may trigger immune cell infiltration as a compensatory mechanism. Alternatively, SAT1 activation may decrease intracellular polyamines which have anti-inflammatory properties [1]. Regardless of mechanisms, cold and β 3-AR stimulation increase SAT1 activity and promote immune cell infiltration and cytokine expression in adipose tissue [14]. Extensive adipose inflammation is generally considered to be detrimental to insulin sensitivity, BAT activity and beige adipocyte biogenesis. However, low level inflammation is necessary for adipose remodeling and beige adipocyte activation [19]. SAT1 activation during cold and β 3-AR stimulation may contribute to the low-grade inflammation in adipose tissue, which indirectly promotes beige adipocyte biogenesis.

In summary, SAT1-mediated polyamine catabolism is sufficient and necessary for beige adipocyte biogenesis. This is mediated by the direct effects of SAT1-induced ROS production in adipocytes. SAT1-induced inflammation and immune cell infiltration may also contribute to the beige adipocyte biogenesis indirectly. Thus, the current study identifies SAT1-mediate polyamine catabolism as a novel mechanistic node for beige adipocyte biogenesis during cold and β 3-AR stimulation.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Highlights:

- Polyamine catabolic enzyme spermidine-spermine acetyltransferase (SAT1) is induced in adipose tissue upon cold and β 3-adrenergic stimulation.
- Adipose-specific SAT1 knockout mice develop late-onset obesity with impaired cold-induced beige adipocyte biogenesis.
- SAT1 activation promotes beige adipocyte biogenesis.
- Reactive oxygen species and inflammation induced by SAT1 activation contribute to beige adipocyte biogenesis.
- SAT1 activation is a critical mechanistic node for beige adipocyte biogenesis induced by cold and 3-adrenergic stimulation.

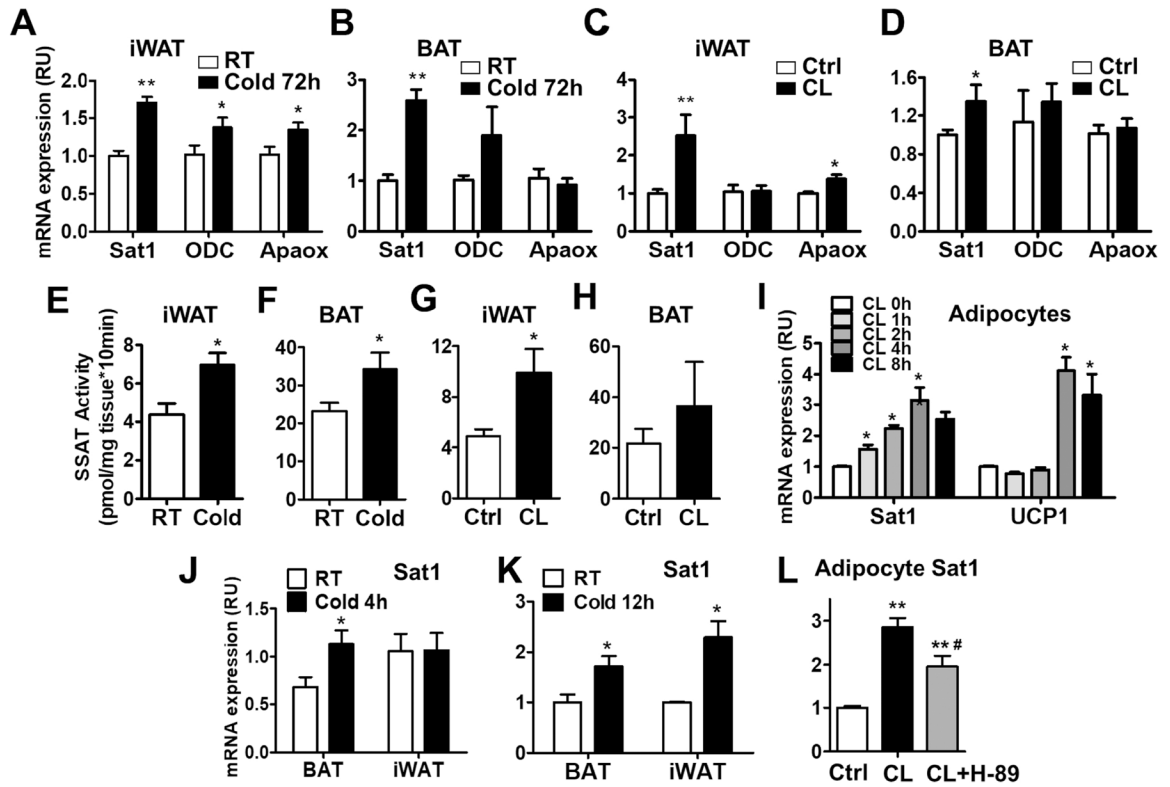


Fig. 1- Cold and β 3-adrenergic receptor agonist stimulate adipose SAT1 expression and activity. (A-B) SAT1, ornithine decarboxylase (ODC) and polyamine oxidase (APAOX) mRNA expression in inguinal (iWAT) and brown (BAT) adipose tissue of cold-challenged mice (4°C for 72 hours). (C-D) SAT1, ODC and APAOX mRNA levels in iWAT and BAT of mice treated with β 3-adrenergic receptor agonist CL 316,243 (CL). (E-H) SAT1 enzymatic activity in iWAT and BAT of mice treated with cold or CL. (I) SAT1 and UCP1 expression in primary adipocytes treated with CL for the indicated times. (J-K) SAT1 expression in iWAT and BAT of mice challenged with cold for 4 hours (J) and 12 hours (K). (L) SAT1 mRNA levels in differentiated adipocytes treated with the PKA inhibitor H-89, followed by stimulation with CL. n=6–10 per group, * $p < 0.05$, ** $p < 0.01$ vs controls (Ctrl). # $p < 0.05$ vs CL.

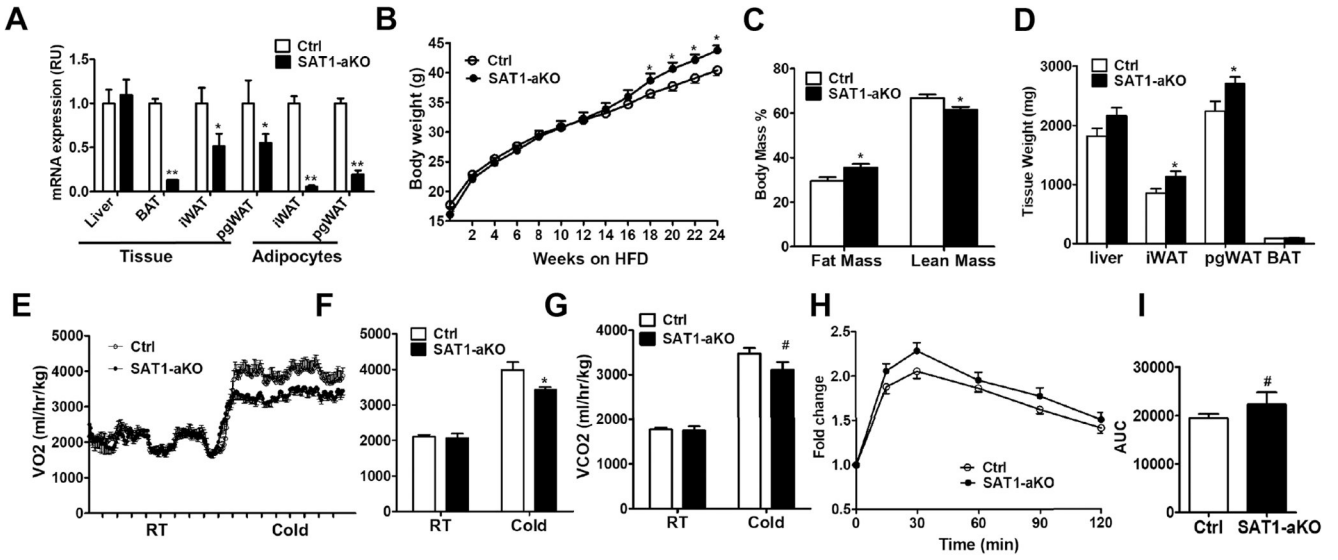


Fig. 2- Adipose-specific SAT1 knockout mice develop late-onset obesity.

(A) SAT1 expression in adipose tissue and adipocytes, (B) body weight, (C) percent fat and lean mass, (D) metabolic organ weight in control (Ctrl) and adipose-specific knockout (SAT1-aKO) mice fed on high fat diet (HFD). (E-G) Oxygen Consumption (E-F) and CO₂ production (G) in SAT1-aKO and control mice under room temperature (RT) and cold (4°C). (H) Glucose tolerance test (GTT) and (I) area under the curve (AUC) in control and SAT1-aKO mice fed a HFD for 12 weeks. n=9–15 per group, except for adipocytes in (A) with n=3, *p<0.05, **p<0.01, #p=0.2 vs Controls (Ctrl).

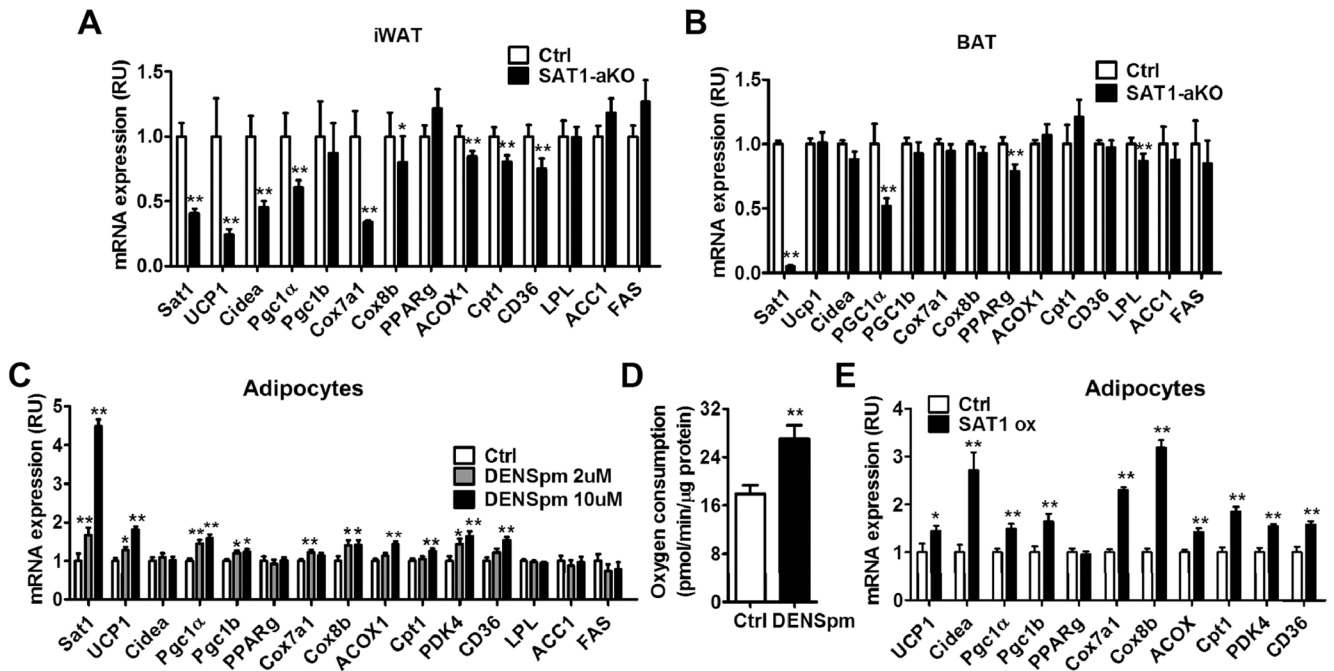


Fig. 3 - Adipose SAT1 regulates beige adipocyte thermogenesis.

(A-B) Browning markers (UCP1, Cidea, Pgc1a, Pgc1b, PPARg, Cox7a1, Cox8b, ACOX1, Cpt1, CD36) and fatty acid synthesis gene expression (ACC, LPL and FAS) in inguinal white adipose tissue (iWAT) (A) and brown adipose tissue (BAT) (B) of cold-challenged SAT1-aKO and control mice. (C-D) Browning gene expression (C) and oxygen consumption in adipocytes treated with SAT1 agonist N1, N11-diethyl norspermine (DENSpm). (E) Browning markers in adipocytes with lentiviral SAT1 overexpression. n=3-8 per group, *p<0.05, ** p<0.01 vs Controls (Ctrl).

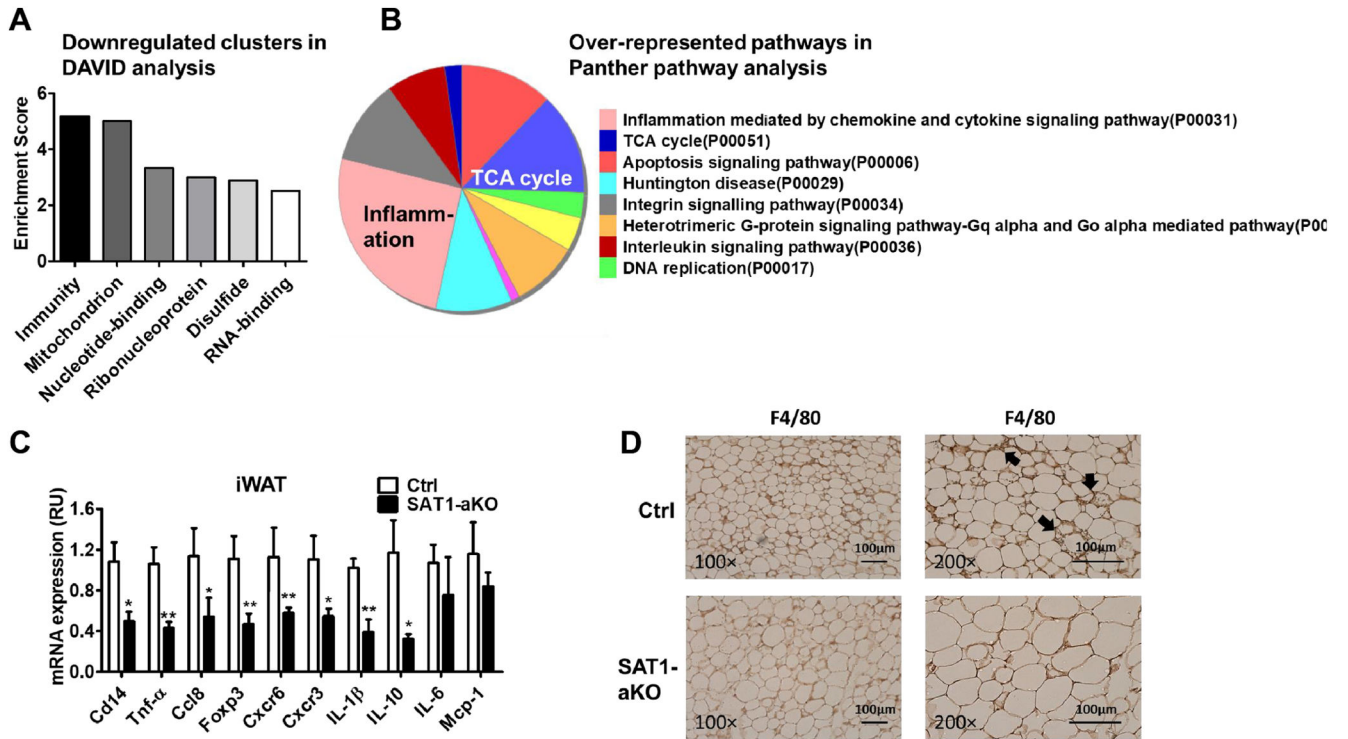


Fig. 4 - SAT1 knockdown reduces immunity-related genes in adipose tissue.

(A-B) DAVID analysis (A) and Panther pathway analysis (B) of RNA-seq results from iWAT of SAT1-aKO and control mice exposed to cold. (C) qPCR measurements of immunity-related gene expression. (D) F4/80 staining of iWAT from cold-challenged SAT1-aKO and control mice. n=6 per group, *p<0.05, ** p<0.01 vs Controls (Ctrl).

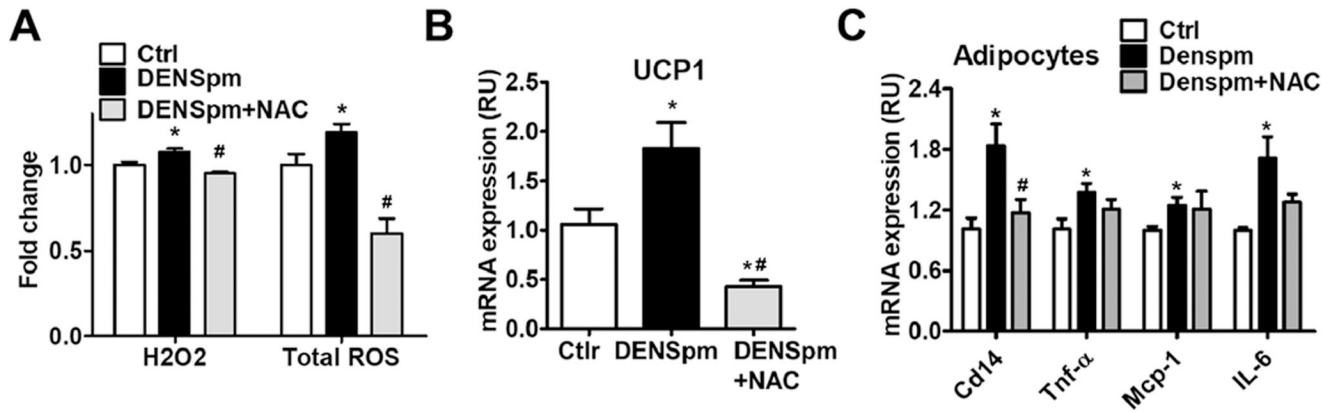


Fig. 5 - SAT1-induced reactive oxygen species regulate beige adipocyte thermogenesis and inflammation.

(A) H₂O₂ levels and total ROS production, (B) UCP1 expression and (C) inflammation-related gene expression in adipocytes treated with SAT1 agonist DENSpm with or without antioxidant N-acetylcysteine (NAC). n=4–8 per group, * p<0.05 vs, ** p<0.01 vs controls (Ctrl), #p<0.05 vs DENSpm.