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dTET3 is necessary for full adipogenesis in vivo

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Reduced adipogenesis by the loss of TET3 in APCs leads to enhanced whole-body metabolism

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**TET3 plays a critical role in white adipose development and diet-induced remodeling**

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**SUMMARY**

Maintaining healthy adipose tissue is crucial for metabolic health, requiring a deeper understanding of adipocyte development and response to high-calorie diets. This study highlights the importance of TET3 during white adipose tissue (WAT) development and expansion. Selective depletion of Tet3 in adipose precursor cells (APCs) reduces adipogenesis, protects against diet-induced adipose expansion, and enhances whole-body metabolism. Transcriptomic analysis of wild-type and Tet3 knockout (KO) APCs unveiled TET3 target genes, including Pparg and several genes linked to the extracellular matrix, pivotal for adipogenesis and remodeling. DNA methylation profiling and functional studies underscore the importance of DNA demethylation in gene regulation. Remarkably, targeted DNA demethylation at the Pparg promoter restored its transcription. In conclusion, TET3 significantly governs adipogenesis and diet-induced adipose expansion by regulating key target genes in APCs.

**INTRODUCTION**

White adipose tissue (WAT) is essential for regulation of energy homeostasis and whole-body metabolism.1 In particular, adipocytes are responsible for storing fat that is released during energy shortage and for secretion of a number of adipokines.1 Both excessive and insufficient amounts of WAT are detrimental to the health of an organism, as both states cause ectopic lipid deposition in non-adipose tissues that results in metabolic insults, including insulin resistance and adipose fibro-inflammation.2,3 Maintaining a moderate amount of healthy WAT is therefore essential for metabolic health, and it is critical to understand mechanisms that control adipocyte developmental and remodeling processes.

Epigenetic changes are modifications to DNA that regulate whether genes are turned on or off. Accumulating evidence indicates that epigenetic factors play important roles in the regulation of adipocyte development and plasticity and that they mediate the interplay between genes and environment.5,6 DNA methylation is a type of reversible epigenetic modification mediated by DNA methyltransferases (DNMTs) and tends to occur at the 5' carbon of the cytosine ring (5C), forming 5-methylcytosine (5mC).10 DNA methylation has long been considered a static epigenetic modification, but emerging evidence suggests that it undergoes dynamic and reversible remodeling.11 TET proteins (TET1, -2, and -3) are a group of enzymes involved in active DNA demethylation. They act as a 5mC dioxygenase, converting 5mC to 5hmC as the first step of DNA demethylation. TET enzymes can also oxidize 5mC or 5hmC to generate 5-formylcytosine (5fC) and/or 5-carboxylcytosine (5caC).11

Our group and others have demonstrated that DNA (de) methylation is involved in multiple aspects of adipose biology.8,9,12–16 Several studies have reported that key adipocyte genes undergo demethylation during adipogenesis, correlating with increased gene expression. For example, Pparg promoter regions have been shown to undergo gradual demethylation during 3T3-L1 adipogenesis.17 This coincides with other epigenetic changes, such as the loss of repressive histone marks (H3K9me3) and the gain of active marks (e.g., H3K27ac and H3K4me3).18 Similarly, reduced DNA methylation has been observed at promoters of other key adipocyte genes (e.g., Cebpa16, Lep,19 Slc2a4,20 and Plin1115) during adipogenesis. Global profiling studies showed that 5hmC is enriched at adipocyte-specific enhancers and in PPARγ+ nuclei of human adipocytes21 and that it co-localizes with PPARγ at enhancers in 3T3-L1 adipocytes.22

A previous study reported that TET1 and TET2 are necessary for 3T3-L1 adipocyte differentiation.17 Other in vitro studies have found that TET1 and TET2 facilitate expression of key adipocyte genes, such as Plin115 and Adipoq15, respectively, by physically interacting with PPARγ. It has therefore been suggested that TET1 and/or TET2 are also critical mediators of DNA
demethylation of adipocyte genes during in vivo adipogenesis. In contrast, we have found that expression of Tet3, rather than Tet1 or Tet2, increases significantly during adipogenesis in naive adipose precursor cells (APCs), as well as in various preadipocyte cell lines.

Here, we explored the role of TET3 in adipogenesis in vivo and the metabolic consequences of Tet3 deficiency. We show that TET3 cell autonomously facilitates adipogenesis. APC-selective Tet3 knockout (KO) mice on either a chow or high-fat diet displayed a decreased adipose mass with reduced adipogenesis while not manifesting a lipodystrophy phenotype. Strikingly, these KO mice were resistant to diet-induced obesity without metabolic impairment. This metabolic advantage was accompanied by reduced fibro-inflammation in WAT. Our genome-wide profiling reveals that TET3 regulates a set of previously unsuspected target genes in APCs, concurrent with modifications of their DNA methylation profile, which are critical for adipogenesis and adipose remodeling. Finally, we validate the importance of TET3-dependent changes of DNA methylation at TET3 target genes as gene regulatory elements by DNA methylation-dependent reporter assays and site-specific modification of DNA demethylation. Taken together, our studies demonstrate that TET3 is a critical epigenetic regulator of homeostatic control of WAT development and diet-induced adipose expansion.

RESULTS

TET3 regulates adipogenesis cell autonomously

Results from our previous research indicate that Tet2 and Tet3 levels are significantly increased during adipogenesis in 3T3-L1 (considered as white), beige, and brown adipocyte cell lines. To determine whether TETs regulate in vivo adipogenesis, if so, which of them might do so, we investigated TET expression patterns during adipogenesis of primary APCs. APCs were defined by expression of PDGFRα, a commonly chosen APC surface marker in stromal vascular fraction (SVF) cells and by the absence of hematopoietic markers (Lin; CD31, CD45, Ter119) (Figure S1A). Upon isolation of APCs using fluorescence-activated cell sorting (FACS) from lean wild-type (WT) mice, we compared Tet expression before and after adipogenesis. The results showed that Tet3 transcription increased by ~3-fold during APC differentiation; in contrast, Tet2 transcript levels did not change, whereas Tet1 transcript rather decreased (Figure S1B). Assessment of a published single-cell RNA sequencing (scRNA-seq) database profiled from mouse WAT revealed an overall positive correlation between Tet3 and Pdgfra transcript expression within Pdgfra+ APCs (Figure S1C). Additionally, we have conducted the tissue distribution analysis of TET3 expression and found that TET3 expression is detected in various tissues, but it is relatively higher in white adipose depots compared with brown adipose tissue (BAT), liver, and skeletal muscle (Figure S1D). This relationship prompted us to investigate the roles of TET3 in adipogenesis in vivo. We generated mouse models of APC-selective Tet3-deficient mice by crossing Tet3-floxed mice with Pdgfra-Cre (Figure 1A), which has been used to target a broad range of APC populations. We first confirmed a highly efficient Tet3 knockout (KD) of >90% in FACS-purified APCs from inguinal WAT (iWAT) and epididymal WAT (eWAT) of KO and WT mice (Figure 1B). To determine the cell-autonomous role of TET3 in adipogenesis, we then FACS-purified Lin−/PDGFRα− iWAT APCs from 8-week-old WT and KO mice on a chow diet and differentiated them in culture by using the standard differentiation inducers (i.e., dexamethasone, 3-isobutyl-1-methylxanthine (IBMX), insulin). KO APCs showed reduced adipogenic competency, as assessed by oil red O (ORO) staining and gene expression analysis of key adipocyte markers (e.g., Pparg, Fabp4, Scd1, and Lipe) (Figures 1C and 1D). To further determine cell-autonomous role of TET3 in adipogenesis, we assessed adipogenic capacity of 3T3-L1 cells upon TET3 overexpression (OE) and Tet3 KD. First, we confirmed that OE and KD increased and decreased 5hmC levels, respectively (Figures 1E and 1I). When assessed by lipid accumulation and gene expression analysis of adipocyte markers, TET3 OE promoted adipogenesis, while Tet3 KD impaired differentiation (Figures 1F–1H and 1J–1L). Together, these data indicate a cell-autonomous role for TET3 in facilitating adipogenesis.

TET3 is necessary for WAT development under chow-fed conditions

We next used Pdgfra-Tet3 KO mice to investigate roles of TET3 in WAT development. When animals were fed a chow diet, no significant difference was found in body weight (BW) between genotypes (Figure 2A). However, Pdgfra-Tet3 KO mice exhibit significant reductions of 22% and 40% in fat and WAT mass, respectively, in comparison to their littermate controls. This was determined by EchoMRI and measurement of tissue weights (Figures 2B and 2C). Histological examination was then carried out using H&E staining (Figures 2D and 2G) and adipocyte size distribution analysis to assess adipocyte formation and morphology. Overall, we found that adipocyte size did not differ between genotypes (Figures 2E and 2H), but total adipocyte number was overall reduced in iWAT and eWAT of KO mice compared with controls (Figures 2F and 2I). Our gene expression analysis of KO mice showed reduced expression of key adipocyte genes, including Pparg, Scd1, Lipe, and Lep, in both iWAT and eWAT depots, and this decrease was particularly pronounced in iWAT (Figures 2J–2M). Importantly, although KO mice had reduced WAT, no differences were observed in their energy expenditure, physical activity, food intake, or respiratory exchange rate (RER) (Figures S2A–S2D). In line with this, no significant change was observed in expression of Ucp1 in various adipose depots, including iWAT and BAT (Figure S2E). We also did not find discernable differences in BAT histology (Figure S2F). These data suggest that reduced adiposity is not due to increased energy expenditure. Together, these data suggest that TET3 is required for the full complement of adipocytes under chow-fed conditions.

Tet3 deficiency in APCs confers protection from diet-induced unhealthy adipose expansion and remodeling

High-fat feeding induces adipose expansion by promoting de novo adipogenesis and hypertrophy. To determine the roles of Tet3 in diet-induced adipose tissue expansion, a cohort of male mice were fed a high-fat diet (HFD). Remarkably, KO mice were generally found to gain less weight, weighing significantly less than control mice after 10 weeks of HFD feeding.
This difference was primarily due to reduced fat mass in KO mice (Figures 3C and 3D). Tissue measurement confirmed that weight of various WAT depots was significantly reduced in KO mice on an HFD compared with control WT littermates (Figure 3E). Analyses of adipocyte size distribution indicated that iWAT and eWAT of KO mice were reduced in size (Figures 3F, 3G, 3I, and 3J), as was total adipocyte number (Figures 3H and 3K). Several key adipocyte genes showed reduced expression in both iWAT and eWAT. The reduction was more pronounced in iWAT, similar to that seen in chow-fed mice (Figures 3L–3N). In agreement with reduced adiposity, serum leptin levels were lower in KO mice than in WT mice (Figure 3O). We also determined whether altered energy homeostasis contributed to reduced mass. Compared with control WT mice, KO male mice on an HFD showed higher energy expenditure and had increased resting whole-body energy expenditure when differences in body mass were accounted for by analysis of covariance (ANCOVA) (Figure S3A), while there were no changes in food intake, physical activity, or RER (Figures S3B–S2D). We therefore investigated whether KO mice have increased adipose thermogenesis. We found that Pdgfra-Tet3 KO mice had increased expression of Ucp1 in iWAT, but not in eWAT or BAT, compared with controls (Figure S3E). Notably, we observed a significant reduction in BAT weight (Figure 3E).
and lipid accumulation in KO mice (Figures S3E and S3F). These data suggest that increased energy expenditure contributes to reduced adiposity in KO animals fed an HFD.

As we observed a reduction in total adipocyte number in KO mice, we then determined whether this was due to altered APC proliferation and/or increased apoptotic rates. To evaluate the levels of apoptosis, we performed terminal deoxynucleotidyl transferase 2’-deoxyuridine-5’-triphosphate (dUTP) nick end labeling (TUNEL) assays, a method based on detection of 5-Bromo-dUTP (Br-dUTP), which are incorporated into fragmented DNA. We observed that the apoptotic rate was generally lower in iWAT KO APCs compared with WT controls, while it remained unchanged in eWAT KO APCs (Figure S4A). These results suggest that the lower adipocyte number was not due to loss of APC pools. Next, APC proliferation was measured by using BrdU, a base analog of thymidine that is incorporated into DNA in place of thymidine, providing a measurement of DNA synthesis and cell proliferation. To examine diet-induced APC proliferation, we fed a cohort of chow-fed mice BrdU-containing water (0.8 mg/kg) for 1 week at 8 weeks of age and put them on an HFD. Following BrdU administration, fractionated SVF cells were stained with surface antibodies for Lin markers and PDGFRα. Cells were then fixed and permeabilized for staining with BrdU antibodies. Our results identified no discernable changes in APC proliferation between genotypes (Figure S4D), suggesting that the reduced de novo adipogenesis observed in KO mice was not due to compromised APC proliferation.

Diet-induced adipose remodeling involves increased inflammation and fibrosis, signifying adipose tissue dysfunction.2,28,29 Our experiments showed reduced macrophage infiltration in KO WAT, as indicated by a lower number of crown-like structures (Figures S4F and S4I) and fewer cells stained with Mac2 (a surface macrophage marker) (Figure S3P). In addition, we noted that KO WAT demonstrated reduced expression of inflammatory cytokines and fibrosis markers (Figures 3L and 3M) and had lower

Figure 2. *Pdgfra-Tet3* KO male mice show reduced adiposity on a chow diet
(A–C) BW (A), body composition (B), and tissue weight (C) of 8-week-old WT and *Pdgfra-Tet3* KO male mice on a chow diet (n = 11 for A and B; n = 7 for C). Quad, quadriceps muscle.
(D–I) H&E staining (D and G), adipocyte size distribution (E and H), and total adipocyte number (F and I) of iWAT and eWAT from WT and *Pdgfra-Tet3* KO mice on a chow diet (n = 3 for E, F, H, and I). The scale bar represents 100 μm.
(J and K) Gene expression analysis in WAT from WT and *Pdgfra-Tet3* KO mice on a chow diet (n = 5).
(L and M) Immunoblotting analysis showing PPARγ protein levels in WAT from WT and *Pdgfra-Tet3* KO mice on a chow (n = 3).
Data are presented as mean ± SEM. *p < 0.05, **p < 0.01, ***p < 0.001 using two-tailed Student’s t test (A–C, E, F, H–K, and M).
quantities of hydroxyproline (a major component of fibrillar collagen) (Figure 3Q).

In both humans and mice, 50%–70% of the bone marrow cavity is filled with bone marrow adipocytes, which likely play a role in local bone homeostasis. Since PDGFRα is expressed in mesenchymal cells in the bone marrow, we sought to investigate whether the Tet3 deficiency in mice PDGFRα+ cells has an impact on bone parameters. To do that, we performed microcomputed tomography (μCT) imaging of the tibiae bones of WT and Pdgfra-Tet3 KO mice and detected no discernible changes in tibiae bone parameters between genotypes in both genders on either diet (Figure S5). Taken together, our data indicate that Tet3 deficiency in APCs results in protection from unhealthy diet-induced adipose expansion and remodeling.

**APC-selective Tet3 deficiency improves metabolic health in mice upon chronic high-fat feeding**

We investigated the metabolic consequences of reduced adipogenesis caused by APC-selective Tet3 deficiency. No significant differences in metabolic profile were observed between WT and

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**Figure 3. Pdgfra-Tet3 KO male mice on an HFD are protected from unhealthy adipose expansion and remodeling**

(A–E) BW (A), whole-body and WAT photographs (B and C), body composition (D), and adipose tissue weight (E) of WT and Pdgfra-Tet3 KO male mice on an HFD. pWAT, perirenal WAT; mWAT, mesenteric WAT; GA, gastrocnemius; EDL, extensor digitorum longus (n = 11 for A and D; n = 10 for adipose tissues and liver; quad, spleen, GA, EDL, and soleus n = 4 [WT] or 3 [KO] in E).

(F–K) H&E staining (F and I), frequency distribution of adipocyte size (G and J), and total adipocyte number (H and K) of WATs from WT and Pdgfra-Tet3 KO mice on an HFD (n = 6).

(L–O) mRNA expression analysis (L and M) and PPARγ protein levels and quantification (N) in WAT of WT and Pdgfra-Tet3 KO mice on an HFD (n = 5 for L and M; n = 6 for N).

(P–Q) Serum leptin levels (O), immunofluorescence staining of Mac2 (P), and quantification of hydroxyproline (Q) in WAT from WT and KO mice on an HFD (n = 10 for O; n = 3 for Q). The scale bar represents 100 μm.

Data are presented as mean ± SEM. *p < 0.05, **p < 0.01, ***p < 0.001 using two-tailed Student’s t test (A, D, E, G, H, J–O, and Q).
Pdgfra-Tet3 KO male mice on a chow diet, including their glucose tolerance and insulin sensitivity, as assessed by glucose and insulin tolerance tests (GTTs and ITTs, respectively) (Figures 4A–4D). We then fed a cohort of WT and KO male mice an HFD starting at 8 weeks of age to determine whether this would affect the metabolic profile. Remarkably, after 6–7 weeks of HFD feeding, KO male mice exhibited improved glucose tolerance and demonstrated hypoglycemia during fasting as well as during GTTs and ITTs (Figures 4E, 4F, and 4G). KO male mice also showed a reduced level of homeostatic model assessment for insulin resistance (HOMA-IR), but there was no significant difference observed in the kinetics of glucose clearance following insulin injection (Figures 4G–4I). Sex-specific differences have been reported in several aspects of adipose biology, including development and responses to metabolic adaptation. Therefore, we examined a cohort of female mice to determine whether TET3 acts in the same manner as a male cohort in this biological context. Our results showed that KO female mice had generally reduced fat mass (Figures S6A–S6C) and improved glucose tolerance following prolonged feeding on an HFD (Figures S6D–S6G), although the extent of these effects was somewhat less pronounced than in the male cohort. We conclude that it is likely that the role of TET3 is largely conserved in both male and female mice.

Figure 4. Pdgfra-Tet3 KO male mice display improved whole-body metabolism on an HFD
(A–I) GTT (A, B, E, and F), ITT (C, D, G, and H), and HOMA-IR (I) from Pdgfra-Tet3 KO and WT male mice reared on a chow at 8 weeks of age after being fed an HFD for 2 months (n = 5 mice for A–D; n = 8 mice for E, F, and I; n = 12 mice for G and H). AOC, area over the curve. Data are presented as mean ± SEM. *p < 0.05, using two-tailed Student’s t test (A–G).
(J) H&E staining of liver sections of Pdgfra-Tet3 KO and WT male mice on a chow and fed an HFD. The scale bar represents 50 μm.
(K) Quantification of triglyceride (TG) in Pdgfra-Tet3 KO and WT male mice fed an HFD (n = 6).
Data are presented as mean ± SEM. *p < 0.05, **p < 0.01, ***p < 0.001 using two-tailed Student’s t test (A–I).
TET3 regulates adipogenesis through key target genes in APCs, particularly extracellular matrix (ECM)-related genes

We and others have demonstrated that TETs control many biological processes by regulating sets of specific target genes. To reveal the underlying molecular mechanisms by which TET3 regulates adipogenesis, we performed genome-wide RNA-seq and DNA methylation profiling studies, comparing WT and KO APCs from mice fed an HFD (Figure 5A). By transcriptomic analysis, we identified 176 and 153 genes that were up- and downregulated in KO APCs, respectively (Figure 5B; Table S1). As TET3 mediates DNA demethylation, we hypothesized that loss of TET3 would lead to DNA hypermethylation of target genes, resulting in gene repression. We therefore focused our attention on the downregulated genes, which were most likely to be the direct targets of TET3. Pathway analysis indicated that several biological pathways involved in ECM remodeling were highly enriched in the downregulated gene sets (e.g., Cilp, C1qtnf7, Col8a1, Mgp) (Figures 5C and 5D). Although it was just under the threshold of significance in the RNA-seq analysis, our qPCR analysis identified Pparg as a downregulated gene in Tet3 KO APCs (Figure 5D). While we observed pathways enriched in upregulated gene sets, we have not found them to be relevant in the Tet3 KD phenotypes we have observed thus far (Figure S7). In addition, these enriched pathways were less statistically significant when compared with the enriched pathways found in the downregulated gene sets.

Excessive ECM production has been linked to unhealthy adipose formation and insulin resistance, though mechanisms underlying these processes are not well understood. Of the ECM-related TET3 target genes, we chose to prioritize to study Cilp, as it demonstrated a cell-autonomous downregulation with Tet3 KD in 3T3-L1 preadipocytes (Figure 5E), increased expression during adipogenesis (Figure 5F), and elevated expression in...
obese WAT (Figure 5G), with hypermethylated CpGs at the gene locus (Figure 6F). Clip encodes the cartilage intermediate layer protein (CILP), a matricellular protein produced by chondrocytes in articular cartilage, and CILP has been implicated in common musculoskeletal disorders, including osteoarthritis and lumbar disc degeneration. It has also been associated with cardiac ECM remodeling in mouse hearts undergoing pathological ventricular remodeling, such as during myocardial infarction and transverse aortic constriction. Notably, recent scRNA-seq studies identified Clip as one of the extracellular markers enriched in specific subpopulations of APCs. Despite its relevant implications, the role of CILP in regulating adipogenesis and adipocyte remodeling has not yet been investigated. Thus, we sought to determine the functional importance of CILP in TET3-mediated adipogenesis by gain-of-function studies. Remarkably, 3T3-L1 cells treated with recombinant CILP had enhanced adipogenic capacity, as assessed by ORO and gene expression analysis (Figures 5H and 5I). Our data suggest that CILP may be a critical molecular target for TET3 in promoting adipogenesis.

TET3-dependent modification of DNA methylation in APCs is critical for target gene regulation and adipogenesis

TETs have been shown to mediate biological processes through DNA-demethylase-dependent and -independent regulation of target genes, depending on the biological context. We therefore investigated whether the catalytic activity of TET3 is essential for regulating adipogenesis. We first performed gain-of-function studies using various TET3 alleles in 3T3-L1 cells: TET3-WT (full-length WT), TET3-CD (catalytic domain only), and TET3-CDM (full length with catalytic-dead mutation) (Figure 6A). When these alleles were overexpressed to a similar degree, OE of TET3-WT and TET3-CD enhanced adipogenesis, whereas OE of TET3-CDM did not (Figures 6B and 6C). Additionally, we conducted a rescue experiment by reintroducing an equivalent amount of TET3-WT vs. TET3-CDM allele in Tet3 KD cells (Figures 6D and 6E). Our results showed that only the WT allele significantly restored diminished adipogenesis in Tet3 KD cells (Figures S8E and S8F). Together, these data demonstrate that DNA demethylase activity of TET3 is required for its effect on adipogenesis.

Next, to determine the epigenetic basis of TET3 regulation of target genes, we profiled DNA methylomes of naive WT and KO APCs from mice fed an HFD using whole-genome bisulfite sequencing (WGBS), a gold-standard method of measuring DNA methylation profiles at bp resolution. We defined differentially methylated regions (DMRs) as at least 5 CpGs in 200 bp bins, with minimal differences in methylation level (>20%, $p < 0.01$). Consistent with the theory that TET3 mediates DNA demethylation, genome-wide Tet3 KO APCs showed a much higher number of hypermethylated DMRs (hyper-DMRs; n = 1811) than hypomethylated DMRs (hypo-DMRs; n = 9) (Figures 6A and 6B). We found that 32.7% (50/153) of downregulated genes in KO APCs were associated with hyper-DMRs within a 50 kb distance of transcription start sites (Figure 6C). To identify relevant biological pathways associated with methylation changes resulting from Tet3 deficiency, we performed pathway analysis of genes linked to hyper-DMRs in KO APCs. Remarkably, several Kyoto Encyclopedia of Genes and Genomes (KEGG) pathways involved in cell remodeling were found to be enriched (Figure 6D), which were consistent with genes downregulated in KO APCs (Figures S5B–S5D). Hypermethylation was observed at the Pparg (Figure 6E) and ECM-related genes, such as Clip and C1qtnf7 (Figures 6F and 6G), and we therefore sought to determine the importance of DNA methylation of hyper-DMRs in gene regulation. We subcloned the target hyper-DMRs near TET3 target genes (i.e., Pparg, Clip, and C1qtnf7) to a DNA-methylation-dependent reporter plasmid (pCpGL-Basic) that lacked backbone CpG dinucleotides. Our results showed that methylase treatment significantly reduced reporter activities of the pCpGL-containing hyper-DMRs of Pparg, Clip, and C1qtnf7 by ~40%–70%, but no effect was found on the pCpGL itself or the vector containing the CpG-free gene activation element (pCpGL-CMV/EF1) (Figure 6H). Furthermore, our TET3 chromatin immunoprecipitation (ChIP) analysis confirmed that TET3 binds to these hypermethylated regions in Tet3 KO APCs (Figures 6I–6K). Together, these data indicate that TET3 regulates expression of target genes by modifying their DNA methylation profiles.

TETs are known to be recruited to DNA by sequence-specific transcription factors in different biological contexts. We therefore queried the DNA sequence of hyper-DMRs in KO APCs to predict the candidate factor(s) that confers TET3 target specificity. Most notably, motif analysis of hyper-DMRs showed high enrichment for the transcription factors known to be important in adipogenesis (Figure 6L). For example, the nuclear factor I (NFI) transcription factors have previously been shown to be pro-adipogenic in multiple adipogenesis models. The CCAAT-enhancer-binding protein (CEBP) family of transcription factors are also well recognized for their critical role in initiating and enhancing adipogenesis. Of note, C/EBPβ and ε have been shown to orchestrate chromatin remodeling in 3T3-L1 adipogenesis. It was interesting to note that the CEBPε motif ranked highly as an enriched at hyper-DMRs in KO, although it was not the most highly expressed CEBP transcription factor in adipocytes. This could be due to the inaccuracy of motif prediction in distinguishing different gene family members.

CRISPR-dCas9-mediated DNA hydroxymethylation at the Pparg promoter is sufficient to increase its expression

PPARγ is a master transcription factor of adipogenesis. It has two main isoforms—γ1 and γ2—that are generated from separate transcriptional start sites, resulting in 30 additional amino acids at the PPARγ2 N terminus. We found that expression of both isoforms of PPARγ was reduced in KO WATs. In particular, our data indicate that TET3 regulates expression of Pparg2 by modifying methylation levels of the promoter region (Figures 6E, 6H, and 6L). Thus, we attempted to introduce DNA hydroxymethylation at the Pparg2 promoter region to determine whether this would be sufficient to alter Pparg2 gene expression. To do that, we used CRISPR-dCas9-based approaches with endogenous cis-regulatory elements that allow target specificity. A chimeric DNA demethylase tool was developed by fusing the TET1 catalytic domain to the nuclease-dead double Cas9 in a
Figure 6. TET3 modifies the DNA methylation profile of key target genes in APCs

(A) Correlation map depicting relative CpG density and methylation levels between WT and Tet3 KO APCs.
(B) Pie chart of hyper- and hypo-DMR counts in Tet3 KO APCs.
(C) Pie chart of downregulated gene counts linked with hyper-DMRs in Tet3 KO APCs.
(D) KEGG pathway most enriched in genes linked to hyper-DMRs in Tet3 KO APCs. ECM-related pathways are depicted in bold.
(E–G) Snapshots of WGBS track showing hypermethylated regions (boxed region) near the TET3 target genes. The height of the gray and red bars represents the degree of CpG methylation in WT and Tet3 KO APCs, respectively.
(H) HEK-293T cells were transfected with methylated (mCpG) and unmethylated (CpG) versions of pCpGL reporter plasmids containing target regions with differential methylation or control sequences (basic and CMV/EF1 were used as a negative and positive controls, respectively). These cells were then assayed for luciferase reporter activity. The C1qtnf7 DMR appeared on the left side of the track that was tested (n = 3).
(I–K) TET3 ChIP-qPCR analysis with WT iWAT at hypermethylated regions in Tet3 KO APCs (n = 3).
(L) The overrepresented transcription factor motifs predicted by motif analysis using HOMER (v.4.11) on the hyper-DMRs in Tet3 KO APCs.
(M and N) Pparg2 mRNA expression (M) and 5mC enrichment at the Pparg2 promoter (N) in 3T3-L1 preadipocytes expressing pINDUCER-dCas9-Tet1CD (dCas9-Tet1CD) or pINDUCER-dCas9-Tet1CDM (dCas9-Tet1CDM) compared with Pparg gRNAs (n = 3 for M and N).

Data are presented as mean ± SEM. **p < 0.01, ***p < 0.001 using two-tailed Student’s t test (H–K, M, and N).
doxycycline-inducible lentivirus (pINDUCER-dCas9-TET1CD).\textsuperscript{57} A catalytic mutant version of this (pINDUCER-dCas9-TET1CDM)\textsuperscript{57} was used as a control. To target the Pparg hyper-DMR, we tested 3 gRNAs subcloned into a lentiviral gRNA expression vector (pLN-U6-gRNA). 3T3-L1 cells underwent lentiviral transduction, producing pINDUCER-dCas9-TET1CD or pINDUCER-dCas9-TET1CDM. These cells reliably expressed the fusion proteins and were then transduced with pLN-U6-gPparg and control gRNAs. We found that pINDUCER-dCas9-TET1CD cells co-expressing Pparg gRNA (particularly #2 gPparg2) increased Pparg expression by approximately 7-fold following doxycycline administration, while there was a marginal increase in pINDUCER-dCas9-TET1CD cells (Figure 6M). Our methylated DNA immunoprecipitation (MeDIP) analysis confirmed that DNA methylation was successfully reduced at the targeted region in pINDUCER-dCas9-TET1CD cells (Figure 6N). Next, to test whether targeted DNA demethylation is sufficient to alter adipogenic capacity, we transduced control and shTet3 KD 3T3-L1 cells with gPparg (#2) and pINDUCER-dCas9-TET1CD. We found that gPparg-expressing cells showed a significant increase in adipogenic gene expression in both control and shTet3 KD cells (Figure 6O). These results indicate that the DNA methylation level of the Pparg2 promoter is critical for modulation of Pparg2 expression and adipogenesis.

**DISCUSSION**

Adipocyte formation and diet-induced adipose expansion involve dramatic changes in epigenetic modifications. Here, we demonstrate that TET3 is a critical epigenetic regulator of *in vivo* adipogenesis by employing Pdgfra-Tet3 KO mice with APC-selective targeting. Our data also show that, rather than causing lipodystrophy, the reduced adipogenesis observed in KO mice is associated with protection from diet-induced obesity and improves whole-body metabolism when mice are challenged with an HFD. Genome-wide transcriptomic and epigenomic profiling studies (followed by functional validation studies) on naive WT and Tet3 KO APCs revealed a set of previously unsuspected TET3 molecular and genomic targets. Furthermore, we successfully induced DNA demethylation at the Pparg promoter region using a programmable CRISPR-dCas9-based demethylase tool and were thus able to alter the expression of Pparg.

While preparing our manuscript, we learned of a highly relevant paper recently published by Park et al.\textsuperscript{53} These authors reported the DNA methylation level of the Pparg2 promoter region using a programmable CRISPR-dCas9-based demethylase tool and were thus able to alter the expression of Pparg.

For example, studies have shown that mice lacking a single copy of Pparg are protected from diet-induced obesity and glucose intolerance,\textsuperscript{62} suggesting that Pparg produces a dose-dependent effect. Most recently, a study generating Pparg isoform-specific KO mice showed that both Pparg1 and g2 KO mice demonstrate resistance to diet-induced obesity and have higher glucose tolerance.\textsuperscript{62} Likewise, our results showing improved organismal metabolism in Pdgfra-Tet3 KO mice fed an HFD are consistent with the reduced Pparg expression levels observed in several previous mouse genetic models.\textsuperscript{62,63} Taken together, the findings of these studies suggest that reducing PPARγ expression or activity results in protection from diet-induced obesity and metabolic dysregulation rather than lipodystrophy. As such, the identification of Pparg as a direct target gene of TET3 and the potential to alter Pparg expression with site-specific DNA demethylation are highly notable findings in the context of developing therapeutic interventions for metabolic diseases.

WAT ECM remodeling is essential for various biological activities in adipose tissue, including effective adipogenesis,\textsuperscript{64} tissue architecture,\textsuperscript{65} and intercellular interactions in the tissue niche.\textsuperscript{66} Studies have shown that excessive accumulation of ECM components and subsequent interstitial deposition of fibrotic material observed in obesity worsens fibrosis, inflammation, and metabolic impairment.\textsuperscript{30,67–69} Collagens are major constituents of adipose tissue ECM. They are primarily produced by adipocytes but also other cell types, including preadipocytes and stem cells.\textsuperscript{70}
Accumulation of various collagens has been observed in obesity; for example, studies in humans reported that collagen VI expression is positively correlated with body mass index, insulin resistance, and hyperglycemia and that it is associated with increased expression of inflammatory genes. Collagen VI KO has also been shown to improve adipocyte survival and enhance whole-body metabolism. Metalloproteinases (MMPs) are proteases that remodel ECM proteins through cleavage. MMP14, for example, has been identified as required for proper adipogenesis in vitro and in vivo. Some MMPs (MMP3, -11, -12, -13, -14) have been shown to be upregulated in obesity, while others (MMP7, -9, -16, and -24) are downregulated. In our investigations, we observed that transcripts of several collagen (e.g., Col4a1, Col4a2, Col5a2, Col6a1, Col6a3, Col6a5, Col15a1, Col8a1) and MMP (Mmp3 and Mmp11) genes were downregulated in KO APCs. We therefore suggest that reduced ECM deposition could be a contributing factor in the reduced fibro-inflammation and metabolic improvement observed in Pdgfra-Tet3 KO mice. Of the ECM-related Tet3 target genes, Cilp was of particular interest to us, as it has been shown to be involved in pathogenic ECM remodeling in common musculoskeletal disorders and several cardiac diseases. Interestingly, we also found that Cilp is highly induced during adipogenesis and in obesity. We therefore investigated the role of Cilp in adipogenesis and found that CILP gain of function promotes adipogenesis in vitro. Addressing the function of CILP in vivo is essential, as adipose tissue remodeling involves intricate interactions with other cell types in the niche. Given its role as a secretary protein, it will be interesting to determine whether CILP affects the metabolic function of adipocytes in a paracrine manner. Adipose remodeling involves interactions between several different cell types in multiple biological contexts, including angiogenesis, inflammation, and fibrosis. Single-cell resolution studies of entire WAT depots from WT and KO tissues will therefore be helpful to better understand the role of Tet3 in adipose tissue remodeling.

In summary, our current study demonstrates a physiological role for Tet3 as a critical epigenetic regulator of adipogenesis and diet-induced adipose remodeling. Our genome-wide transcriptional and epigenetic profiling studies, combined with follow-up functional validation studies, have revealed underlying molecular and epigenetic mechanisms by which Tet3 regulates these processes.

Limitations of the study
We have observed that approximately two-thirds of hyper-DMRs do not exhibit a correlation with downregulated genes in Tet3 APCs. This suggests that these specific DMRs, lacking any associations with genes, might have emerged as secondary consequences of the loss of Tet3. To discern the direct targets of Tet3, further investigations involving Tet3 ChIP-seq studies in APCs will be required.

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SUPPLEMENTAL INFORMATION
Supplemental information can be found online at https://doi.org/10.1016/j.celrep.2023.113196.

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The authors thank Dr. Chao Wang (University of Chinese Academy of Sciences) for sharing the Tet3-Cd mutant construct. We thank Dr. Andreas Stahl (University of California, Berkeley) for shearing the plN-U6-gRNA plasmid. We thank Dr. Michael Rehli (University Hospital Regensburg) for shearing the pCOpGL3-Basic plasmid. This work was funded by R01 DK116008 (NIDDK) and P4F P30DK116074 (Stanford Diabetes Research Center) to S.K.

AUTHOR CONTRIBUTIONS

DECLARATION OF INTERESTS
The authors declare no competing interests.

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REFERENCES


67. Henegar, C., Tordjman, J., Achard, V., Lacasa, D., Cremer, I., Guerre-...


## STAR METHODS

### KEY RESOURCES TABLE

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 RESOURCE AVAILABILITY

Lead contact
Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Sona Kang (kangs@berkeley.edu).

Materials availability
Unique reagents generated in this study are available from the lead contact, Sona Kang (kangs@berkeley.edu).

Data and code availability
- The sequencing data reported here has been deposited in NCBI Gene Expression Omnibus and is accessible through GEO: GSE214483.
This study does not report original code. Any additional information required to reanalyze the data reported in this paper is available from the lead contact upon request.

**EXPERIMENTAL MODEL AND SUBJECT DETAILS**

**Animal studies**

Animal Care Mice were maintained under a 12-h light/12-h dark cycle at constant temperature (23°C) with free access to food and water. All animal work was approved by UC Berkeley ACUC.

Tet3 fl/fl mice were obtained from Dr. Anjana Rao and Pdgfra-Cre was purchased from Jackson Laboratory (#013148). These animals were maintained on a C57BL/6J background. High fat diet (Research Diets, D12331I) was started at 8 week old for 8–12 weeks. For ITT, mice were fasted for 6 h and were i.p. injected with insulin (0.5 U/kg for a chow diet and 0.8 U/kg for HFD cohort). Blood glucose was measured at 0, 15, 30, 60, 90, and 120 min. For GTT, mice were fasted for 6 h and i.p. injected with glucose (1 g/kg for chow and 1.5 g/kg for HFD cohort). Blood glucose levels were measured at 0, 15, 30, 60, and 120 min, using Contour NEXT glucometer and glucose strips.

To calculate the area of the curve (AOC), we subtract the area under the baseline value for a more accurate measure of glucose tolerance and insulin sensitivity, independent of differences in baseline glucose levels. If post-baseline values are lower than the baseline value, the area above the baseline value is considered as the AOC. Metabolic rate was measured by indirect calorimetry in open-circuit Oxymax chambers, a component of the Comprehensive Laboratory Animal Monitoring System (CLAMS; Columbus Instruments). Mice were housed individually and maintained at 25°C under a 12 h light/12 h dark cycle. Activity was monitored in 1 min intervals of infrared beam breaks on the X, Y and z axis and found not to be significantly different for any of the groups. Food and water were available ad libitum.

**Cell culture**

3T3-L1 preadipocytes were obtained from ATCC and these cells were authenticated by the ability to differentiate and confirmed to be mycoplasma negative. The cells were maintained in DMEM supplemented with 10% fetal calf serum and 100 U/ml penicillin, and 100 μg/mL streptomycin. For differentiation, cells at post 2 day confluence were switched to DMEM with 10% fetal bovine serum and exposed to a full (250 μM IBMX, 5 μg/mL insulin, and 1 μM dexamethasone) or sub-optimal induction cocktail (0.4x strength of the full cocktail) from day 0–2, insulin from day 2–4, and switch to DMEM with FBS only. For the ex vivo culture studies, WAT pads were excised, minced, and digested in PBS containing 2 mg/mL Collagenase type II, 10 mM CaCl₂ and 3% bovine serum albumin for 30 min in a shaking water bath at 37°C. Digested fat pads were filtered through 100 μm strainer, followed by centrifugation at 300g for 5 min. Pellet is acquired and incubated with RBC lysis buffer for 5 min at RT. Then, cells are filtered through 40 μm strainer and centrifuged at 500g for 5 min. After washing with 3% BSA/PBS solution, acquired SVFs were stained with surface antibodies for Lin markers (CD31, CD45, and TER119) and PDGFRα. FACS-purified Lin−/PDGFRα− APCs were cultured in DMEM with 10% FBS and differentiated as described above. To generate lentivirus particles, lentiviral constructs were co-transfected with pMD2.G- and psPAX2-expressing plasmids into 293T cells. After 48 h, virus-containing supernatant was collected, filtered through 0.45 μm filters, and added to mature 3T3-L1 adipocytes for 24 h along with 8 μg/mL Polybrene. Transduction efficiency was determined by comparing to cells transduced in parallel with a GFP-expressing lentivirus.

**METHOD DETAILS**

**Protein analysis**

Cell or tissue culture samples were lysed in RIPA lysis buffer containing 50 mMol/L Tris-HCl (pH 7.5), 150 mMol/L NaCl, 1% Triton X-100, 0.1% SDS, and protease inhibitor cocktail (Complete Mini-EDTA free, 1183617001, Roche) for 30 min on ice with rotating samples. Protein concentration was quantified by Bradford analysis. 40–60 μg proteins were used for the assays. Proteins were size fractionated by SDS-PAGE and then transferred to polyvinylidene difluoride or nitrocellulose membrane. After blocking with 5% nonfat dried milk in TBS-Tween (0.25%), the membranes were incubated overnight with the appropriate primary antibodies, followed by secondary antibody incubation for 1 h at room temperature. After washing with TBS-Tween for 30 min, signals were detected using the ECL Assay Kit (NEL104001EA; PerkinElmer) using an iBright CL1500 Imaging System (Lnvitrogen), and quantified with ImageJ.

**Reagents**

Insulin, dexamethasone, isobutylmethylxanthine (IBMX), puromycin, and Oil Red O were purchased from Sigma. Methylene blue was purchased from Ricca Chemical. Collagenase type 2 was purchased from Fisher scientific. Recombinant CILP protein was purchased from Abcam (182802). Antibodies were purchased from Cell Signaling (HSP-90, 4874; GAPDH, 2118; Mouse (G3A1) mAb IgG1, 5415), Sigma (anti-FLAG, F3165), BD Biosciences (anti-CD45-APC, 561018; CD31-APC, 561814; TER119-APC, 561033), Abcam (5mc, ab10805), Active Motif (5hmC, 39769), Santa Cruz (PPARγ, sc-7273), Cedarlane Labs (MAC-2, CL8942AP) and Millipore (TET3, abe383).
RNA extraction and qPCR
Total RNA was extracted from tissues using TRIzol reagent according to the manufacturer’s instructions. cDNA was reverse transcribed from 1 μg of RNA using the cDNA High-Capacity cDNA Reverse Transcription Kit (Applied Biosystems). Quantitative PCR (qPCR) was performed with SYBR green qPCR master mix (AccuPower 2X, Bioneer) using QuantStudio 5 (Applied Biosystems). The relative amount of mRNA normalized to cyclophilin B was calculated using the delta–delta method. Primer sequences are listed in Table S2.

Plasmids
Lentiviral overexpression vectors for TET3-WT, TET3-CD, and TET3-CDM were subcloned into pCDH-CMV-MCS-EF1 vector, and hairpins targeting Tett3 were subcloned at AgeI/EcoRI to pLKO.1 vector. qRNA to target Pparg promoter was cloned at Aarl site to pLN-U6-grRNA. Hairpin and gRNA sequences are shown in Table S2 pMD2.G and psPAX2 were gifts from Didier Trono (Addgene plasmid # 12259, #12260; http://n2t.net/addgene:12259, 12260; RRID: Addgene_12259, 12260). pLKO.1 - TRC cloning vector was a gift from David Root (Addgene plasmid # 10878; http://n2t.net/addgene:10878; RRID: Addgene_10878)

ELISA
For the leptin ELISA, serum leptin was analyzed using mouse leptin ELISA Kit (catalog #90030; Crystal Chem). For the insulin ELISA, mice were fasted for 6 h. Plasma insulin was analyzed using an ultrasensitive mouse Insulin ELISA kit (#90080; Crystal Chem). The endpoint calorimetric assays were performed using a Plate Reader SpectraMax i3. HOMA-IR was calculated by using homeostasis model assessment-2 (HOMA2) index using an online-based calculator.

Methylation-dependent reporter assay
The CpG-free luciferase reporter vector, pCpGL3-Basic was a kindly gift from Dr. Michael Rehli of University Hospital Regensburg. Target genomic sites were amplified using genomic DNA from C57BL/6J mice and inserted to pCpGL3-Basic vector precut with HindIII/BglII for Pparg and BamHI/NcoI for Clip and C1qtnf7. Plasmid methylation was carried out using CpG methyltransferase, M.SsI (New England Biolabs) following to the manufacturer’s protocol. Briefly, 4 μg plasmids were co-incubated with 160 μM S-sadenosylmethionine (SAM), Sss I (12 U) for 8 h at 37°C. Additional 160 μM SAM was added, and incubated for 16 h. The unmethylated plasmids were prepared as described above, but without adding SAM in the reaction. Purified plasmids were quantified and transiently co-transfected in HEK-293T with Renilla using Lipopectamine 3000 reagent (Invitrogen), following the manufacturer’s protocol. After 24 h after transfection, cells were lysed to measure Firefly and Renilla luciferase activity using the Dual-Luciferase Reporter Assay System (Promega). SpectraMax i3 plate reader (Molecular Devices, San Jose, USA) was used to analyze luminescence, and Firefly activity was normalized to Renilla activity.

Dot blotting, ChIP-qPCR, and MeDIP assays
For dot blotting, genomic DNA was extracted using the DNeasy Blood & Tissue kit (Qiagen, 69504) following the manufacturer’s protocol. DNA was denatured at 95°C for 15 min in 0.1 M NaOH. It was neutralized with 1 M NH4OAc on ice and diluted to 300 mg in DNase free water. 200 ng DNA was applied to a positively charged nylon membrane under vacuum with a Dot Blot Microfiltration Apparatus (Bio-Rad). The membranes were briefly washed in 2× SSC buffer (0.3 M NaCl, 30 mM sodium citrate) for 5 min, then baked at 80°C for 5 min, and then cross-linked using a UV Stratalinker 1800. Membranes were blocked with 5% nonfat dried milk in TBS-Tween (0.25%). The membranes were incubated with the 5mC primary antibodies. For a loading control, membranes were stained with methylene blue. Immunoblots were quantified by the ImageJ program. For ChIP-qPCR, cells were cross-linked with 1% formaldehyde for 10 min at room temperature. Cross-linked chromatin was sonicated using an S220 Ultrasonicator (Covaris) to generate DNA fragments of ~200–500 bp. Inputs were taken from cleared lysates, and the rest were rotated 0/N at 4°C with TET3 and IgG antibodies for immunoprecipitation. An aliquot of 20 μL of pre-washed Dynabeads Protein G was added per IP and rotated 1 h at 4°C. Beads were successively washed in low-salt RIPA buffer (20 mM Tris-HCl [pH 8.0], 1 mM EDTA, 1% Triton X-100, 0.1% SDS, 140 mM NaCl, 0.1% Na deoxycholate), high-salt RIPA buffer (20 mM Tris-HCl [pH 8.0], 1 mM EDTA, 1% Triton X-100, 0.1% SDS, 500 mM NaCl, 0.1% Na deoxycholate), LiCl buffer (250 mM LiCl, 0.5% NP40, 0.5% Na deoxycholate, 1 mM EDTA, 10 mM Tris-HCl [pH 8.0]) and TE buffer (10 mM Tris-HCl [pH 8.0] and 1 mM EDTA). Each reaction was then incubated in digestion buffer (50 mM Tris-HCl [pH 8.0], 1 mM EDTA, 100 mM NaCl, 0.5% SDS, proteinase K) for a minimum of 4 h at 65°C to reverse cross-links. DNA was recovered using a phenol-chloroform extraction. Real-time qPCR primers are listed in Table S2. All data were normalized to input. For MeDIP, genomic DNA was sheared using a Covaris S220 to an average of 200–800 bp. Two micrograms of denatured DNA were incubated with 2 μg of anti-IgG or 5mC antibody in IP buffer (10 mM Na-Phosphate pH 7.0, 0.14 M NaCl, 0.05% Triton X-100) for 2 h at 4°C. Antibody-bound DNA was collected with 20 μL of Dynabeads anti-mouse IgG (Invitrogen Dynal AS) for 1 h at 4°C on a rotating wheel and successively washed five times with washing buffer (0.1% SDS, 1 Triton X-100, 2 mM EDTA, 20 mM Tris-HCl pH 8.1, 150 mM NaCl) and twice with TE (10 mM TrisCl, 1 mM EDTA pH 8.0). DNA was recovered in 125 μL of digestion buffer (50 mM Tris pH 8.0, 10 mM EDTA, 0.5% SDS, 35 mg proteinase K) and incubated for 3 h at 65°C. Recovered DNA was used for qPCR analysis. Primers for MeDIP-qPCR studies are listed in Table S2. All data were normalized to input.
TUNEL assay
Apoptosis rate was measured by using the APO-BrdU Fragmentation Assay Kit (BioVision, K401), which detects the DNA fragmentation of apoptotic cells by exploiting the fact that the DNA breaks expose a large number of 3’-hydroxyl ends. These hydroxyl groups can then serve as starting points for terminal deoxynucleotidyl transferase (TdT), which adds deoxyribonucleotides in a template-independent fashion. Addition of the deoxythymidine analog 5-bromo-2’-deoxyuridine 5’-triphosphate (BrdUTP) to the TdT reaction serves to label the break sites, which can be detected by BrdU antibody. In brief, the SVFs were stained with surface antibodies for Lin markers (CD31, CD45, and TER119) and PDGFRα, followed by fixation with 3.6% Formaldehyde (v/v) for 15 min and ice-cold 70% (v/v) ethanol for 30 min on ice. The fixed cells were incubated with DNA labeling solutions, stained with anti-BrdU-FITC, and incubated with DAPI for 30 min at RT. After washing, resuspended cells were analyzed for BrdU positive cells by flow cytometry (BD LSR Fortessa). Cells that were not incubated with antibody were used as a negative control to determine background fluorescence levels and BrdU positive area was set based on the signals from positive control cells provided in the kit.

Cell proliferation assay
To assess cell proliferation in vivo, BrdU was administered in the drinking water at 0.8 mg/mL to incorporate into newly synthesized DNA. Feeds were switched from normal feed to an HFD at the same day of BrdU administration to boost the proliferation of adipose precursors within adipose depots. After 1 week of administration, mice are sacrificed and SVFs are isolated, and stained with surface antibodies for Lin markers and PDGFRα. After fixation with successive incubation with 3.6% Formaldehyde (v/v) and ice-cold 70% (v/v) ethanol, cells are stained with anti-BrdU-FITC antibody, followed by the incubation of DAPI for 30 min at RT. The processed samples are analyzed with flow cytometry (BD LSR Fortessa). Cells that were not incubated with antibody were used as a negative control to determine background fluorescence levels and BrdU positive area was set based on the signals from positive control cells provided in the kit.

Hydroxyproline assays
Hydroxyproline content in adipose was assayed using the Hydroxyproline Colorimetric Assay kit (BioVision, K555) according to the manufacturer’s protocol. Briefly, 100–150 mg of frozen adipose tissues were homogenized and incubated in 12 N HCL for 3 h at 120 °C. Samples were reacted with Chloramine T reagent for 5 min at RT. OD560 measurements were made on a plate reader (SpectraMax i3 Plate reader) at 25°C, and the data were calculated using a standard curve.

Adipocyte size distribution and number analysis
The H&E images were captured using an Axiolmager M1 microscope (Zeiss, Germany). Individual adipocyte area (μm²) was measured using an automated software called Adiposoft. To estimate the total number of cells in the fat pads, the fat pad volume (Vfat_pad) was divided by the average fat cell volume (Vfat_cell). The fat pad volume (Vfat_pad) was calculated by dividing the fat pad mass by the adipose density, which was determined to be 0.95 g/cm³. The density value was obtained through Monte Carlo simulations using adipose tissue derived from a mouse model. The volume of the fat cells was calculated using the following formula, where the mean diameter (d) was calculated as 2 × √(mean cell area/π). To account for the variation in fat cell size, (3π² × d + d³) was used instead of d³ to calculate the fat cell volume, where s represents the standard deviation of the diameter.

Immunofluorescence assay
Adipose tissues were fixed with neutral-buffered formalin and embedded in paraffin. Fixed tissues were permeabilized with PBS containing 0.1% Triton X-100 for 10 min and then incubated with anti-MAC2 antibodies in PBS containing 10% (v/v) FBS for 18 h at 4°C. Immunofluorescent staining was analyzed with a confocal LSM710 microscope (Carl Zeiss Microimaging Inc.).

RNA-seq library generation and analysis
RNA samples were extracted using the RNAqueous-Micro Kit (Cat# AM1931), and the quality of total RNA was assessed by the 2100 Bioanalyzer (Agilent). Libraries were prepared using the Poly(A) RNA Selection Kit V1.5. RNA-seq reads were aligned to the UCSC mm10 genome using Hisat2. Gene expression levels were measured from feature counts using Htseq. Differential-expression analysis was done using Deseq2 and differentially expressed genes upon KO were selected by fold-change >1.25 & FDR <0.05. Enrichment analysis was performed using GSEApy to generate enrichment terms and pathways from the KEGG 2019 Mouse. RNA-seq raw data is available in GEO (GSE214483).

WGBS analysis
Raw reads were quality and adapter trimmed using FastQC (v0.11.7) and TrimGalore (v0.6.1) prior to mapping to the in-silico bisulfite converted mouse genome (GRCh39) using Bismark (v0.18.1). CpG methylation calls were extracted after alignment with Bismark as well. Differentially Methylation Region (DMR) was called by DSS (v2.45.0) in R (v4.1.2) environment. Genomic regions with more than 5 CpGs in 200bp bins, and minimal methylation level differences (>20%, p < 0.01) were defined as DMRs. Functional enrichment
analysis of DMRs was done using GREAT (v4.0.4) by finding genes associated to DMRs. For transcription factor motif enrichment, HOMER (v4.11) analysis was used for finding de novo motif enrichments of DMRs. The program findMotifs.pl with default parameters was used to find motifs on the DMRs. WGBS sequencing data is available in GEO (GSE214483).

**Bone μCT analysis**

Murine tibiae were harvested at necropsy, fixed in paraformaldehyde for 48+ hours, washed 3X with PBS, and stored in Sorensen phosphate buffer at 4°C. μCT analysis was performed as previously described. Briefly, tibiae were placed in a 19-mm diameter specimen holder and scanned over the entire length of the tibiae using a μCT system (μCT100 Scanco Medical, Bassersdorf, Switzerland). The following scan settings were used: voxel size 12 μm, 70 kVp, 114 μA, 0.5 mm AL filter, and integration time 500ms. Density measurements were calibrated to the manufacturer’s hydroxyapatite phantom. Analysis was performed using the manufacturer’s evaluation software with a threshold of 180 for trabecular bone and 280 for cortical bone.

**TG measurements in tissues**

Frozen tissues (100 mg) were weighted and transferred into Eppendorf tubes. The tissues were digested with 1mL of alkali digested buffer (6.2% potassium hydroxide in EtOH solution) for O/N at 55°C. Digested tissues are centrifuged at 5,000 rpm for 5 min and supernatants are acquired. Then, 1.1 supernatants volume of 1M MgCl₂ was added. After 10 min of incubation on ice, samples were centrifuged at 5,000 rpm for 5 min and supernatants were acquired. TG levels were measured with a Serum Triglyceride Determination Kit (TR0100, Sigma) following the manufacturer’s protocol. Briefly, 80 μL of free glycerol reagent was dispensed into a 96-well plate. Then, 1 μL of final supernatants were added and incubated for 15 min at RT. Finally, 20 μL of Lipase containing Triglyceride Reagent was added and incubated for 15 min at RT, followed by absorbance measurement at 540 nm.

**QUANTIFICATION AND STATISTICAL ANALYSIS**

Where appropriate, data are expressed as mean ± SEM. The data were statistically analyzed using unpaired Student’s t-test and two-way ANOVA with Tukey’s multiple comparisons test using GraphPad Prism version 8. The p values below 0.05 were considered statistically significant. *p < 0.05, **p < 0.01, ***p < 0.001. Replicate information is indicated in the figures.