MED1 is a lipogenesis coactivator required for postnatal adipose expansion

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MED1 often serves as a surrogate of the general transcription coactivator complex Mediator for identifying active enhancers. MED1 is required for phenotypic conversion of fibroblasts to adipocytes in vitro, but its role in adipose development and expansion in vivo has not been reported. Here, we show that MED1 is not generally required for transcription during adipogenesis in culture and that MED1 is dispensable for adipose development in mice. Instead, MED1 is required for postnatal adipose expansion and the induction of fatty acid and triglyceride synthesis genes after pups switch diet from high-fat maternal milk to carbohydrate-based chow. During adipogenesis, MED1 is dispensable for induction of lineage-determining transcription factors (TFs) PPAR γ and C/EBP α but is required for lipid accumulation in the late phase of differentiation. Mechanistically, MED1 controls the induction of lipogenesis genes by facilitating lipogenic TF ChREBP- and SREBP1 α -dependent recruitment of Mediator to active enhancers. Together, our findings identify a cell- and gene-specific regulatory role of MED1 as a lipogenesis coactivator required for postnatal adipose expansion.

[**Keywords:** MED1; Mediator; adipose expansion; coactivator; de novo lipogenesis]

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Adipose tissue development starts before birth. After birth, adipose tissues, especially white adipose tissue (WAT), undergo marked expansion with tissue mass and lipid contents increasing from newborn pups to adult mice (Cristancho and Lazar 2011; Berry et al. 2013). Adipose tissue development requires the master adipogenic transcription factor (TF) PPAR γ , which is induced in the early phase of adipogenesis, the differentiation of preadipocytes towards adipocytes (Rosen et al. 2002). PPAR γ cooperates with C/EBP α on transcriptional enhancers to induce the expression of thousands of adipocyte genes that also include major lipogenic TFs ChREBP and SREBP1, which together promote de novo lipogenesis (DNL) in the late phase of adipogenesis (Lefterova et al. 2008; Schmidt et al. 2011).

Liver and adipose tissues are the main sites for DNL in mammals. Fatty acid synthesis is low in liver and adipose tissue of mouse pups but increases dramatically after the switch of diet from high-fat maternal milk to carbohydrate-rich standard laboratory chow (Pearce 1983). DNL is a tightly regulated metabolic process that converts ex-

cess carbohydrates to fatty acids before the synthesis of triglyceride for energy storage. Through glycolysis and the tricarboxylic acid cycle, carbohydrates are converted to citrate. Fatty acid synthesis enzymes, including ATP citrate lyase (ACLY), acetyl-CoA carboxylase α/β (ACACA/ACACB, also known as ACC1/ACC2), fatty acid synthase (FASN), and stearoyl-CoA desaturase-1/2/3/4 (SCD1/2/3/4), act sequentially to convert citrate to fatty acids (Strable and Ntambi 2010; Song et al. 2018). Through the subsequent actions of triglyceride synthesis enzymes GPAM (also known as GPAT1), 1-acylglycerol-3-phosphate O-acyltransferase-1/2 (AGPAT1/2), LIPIN1, and DGAT1, fatty acids are esterified with glycerol-3-phosphate and incorporated into triglyceride (Takeuchi and Reue 2009).

DNL enzyme expression is transcriptionally regulated by major lipogenic TFs carbohydrate-responsive element-binding protein (ChREBP, also known as MLXIPL) and sterol regulatory element-binding protein 1 (SREBP1) in adipose tissues. ChREBP is a major driver for adipocyte DNL (Song et al. 2018). It is expressed as two isoforms ChREBP α and ChREBP β , which are encoded by the same gene but transcribed from different promoters. ChREBP β transcription activity is ~20-fold higher than that of

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ChREBP α (Herman et al. 2012). ChREBP α is constitutively expressed but is sequestered in the cytosol and transcriptionally inactive under low glucose conditions. ChREBP α is activated by high glucose and, in turn, stimulates ChREBP β expression (Abdul-Wahed et al. 2017). Overexpression of the constitutively active form of ChREBP (CA-ChREBP) in mouse WAT induces expression of DNL enzymes such as ACLY, ACACA, FASN, and SCD1 (Nuotio-Antar et al. 2015). ChIP-seq analysis in mouse WAT showed genomic binding of ChREBP on *Acaca*, *Acacb*, *Fasn*, and *Chrebp* genes, suggesting that ChREBP directly regulates transcription of DNL enzymes as well as itself (Poungvarin et al. 2015). SREBP1 also promotes DNL gene expression in adipocytes. Overexpression of the SREBP1 α isoform in adipose tissue increases the expression of fatty acid synthesis genes, leading to increased lipid accumulation in adipocytes (Horton et al. 2003).

In eukaryotes, all transcription is mediated by RNA polymerase II (Pol II) for protein-coding genes. Transcription by Pol II requires transcriptional coregulators including chromatin remodelers, histone modifiers, and the Mediator coactivator complex (Malik and Roeder 2010). The Mediator complex comprises approximately 30 subunits, and it generally serves as a functional bridge between TFs and basal transcriptional machinery including Pol II (Allen and Taatjes 2015). Mediator is enriched at transcriptional promoters and enhancers. One of the Mediator subunits, MED1 (also known as TRAP220, PBP), is often used as a surrogate for the Mediator to identify active enhancers and superenhancers (Whyte et al. 2013). Whole-body *Med1* KO in mice leads to embryonic lethality around E11.5 (Ito et al. 2000; Zhu et al. 2000).

We previously reported that *Med1* KO mouse embryonic fibroblasts show defects in PPAR γ - and C/EBP β -stimulated phenotypic conversion into adipocytes (Ge et al. 2002, 2008). However, it has remained unclear whether MED1 is required for adipose tissue development and expansion in vivo. Using *Adipoq*-Cre- or *Myf5*-Cre-mediated deletion of *Med1* in adipocytes or precursor cells in mice, we show that MED1 is required for postnatal adipose expansion after switching diet from high-fat maternal milk to carbohydrate-based standard chow and that MED1 is largely dispensable for adipose tissue development. By RNA-seq and ChIP-seq analyses, we found that MED1 does not control the expression of early adipogenesis marker *Pparg* but directly controls the transcription of lipogenesis genes that encode fatty acid synthesis enzymes ACLY, ACACA, SCD1, and FASN and triglyceride synthesis enzymes GPAM and AGPAT2 in adipocytes. Mechanistically, MED1 facilitates Mediator binding on active enhancers of ChREBP- and SREBP1 α -regulated lipogenesis genes in adipocytes.

Results

Med1^{f/f};Adipoq-Cre mice show lipodystrophy and fail to expand adipose tissue under high-fat diet

To investigate the functional role of MED1 in adipose tissues, we used conditional knockout (KO) mice targeting

exons 8–10 of *Med1* gene (*Med1^{fl}*, hereafter referred to as *f/f*) (Supplemental Fig. S1A; Jia et al. 2004). Adipocyte-specific *Med1* KO (*Med1^{fl};Adipoq-Cre* [A-KO]) mice were generated by crossing *f/f* mice with *Adipoq-Cre* mice expressing *Adiponectin* promoter-driven Cre (Eguchi et al. 2011). A-KO mice showed similar appearance and total body weight compared with *f/f* mice but had significantly reduced fat mass and increased lean mass at 8 wk (Fig. 1A,B). Adipose tissues including brown adipose tissue (BAT), inguinal WAT (iWAT), and epididymal WAT (eWAT) decreased in A-KO mice, while liver mass increased (Fig. 1C,D). Similar results were obtained from female mice at 8 wk (Supplemental Fig. S1B–E). Histological analysis revealed reduced lipid droplets in adipose tissues of A-KO mice (Fig. 1E). Deletion of *Med1* was verified in adipose tissues (Fig. 1F). These results indicate that deletion of *Med1* in adipocytes leads to lipodystrophy in adult mice.

Blood glucose and insulin levels increased in the serum of A-KO mice, while leptin levels decreased. Free fatty acid, triglyceride, and cholesterol levels were similar between A-KO and *f/f* mice (Fig. 1G). A glucose tolerance test (GTT) and insulin tolerance test (ITT) showed that A-KO mice were insulin-resistant (Fig. 1H–I). Deletion of *Med1* in adipocytes did not affect food intake and total activity but caused a significant increase in total energy expenditure, which was likely due to the increased lean mass (Supplemental Fig. S1F). However, in contrast to *f/f* mice, A-KO mice failed to increase energy expenditure and showed reduced lipolysis upon stimulation with CL316,243, a β 3 adrenergic receptor agonist (Fig. 1J,K). A-KO mice maintained core body temperature when housed at room temperature ($\sim 22^{\circ}\text{C}$), but their body temperature significantly dropped to $\sim 31^{\circ}\text{C}$ after 3 h exposure to an ambient temperature of 6°C (Fig. 1L). Expression of thermogenesis genes *Ucp1* and *Elovl3* decreased in BAT of A-KO mice at room temperature and failed to induce after 3-h cold exposure, indicating thermogenesis defects in A-KO mice (Fig. 1M). These data indicate that adipocyte-selective deletion of *Med1* impairs glucose homeostasis and cold-induced thermogenesis in mice.

Under a high-fat diet (HFD), A-KO mice gained significantly less body weight and fat mass compared with *f/f* mice, but more lean mass (Fig. 2A–C; Supplemental Fig. S2A,B). *Med1* KO had little effect on food intake and energy expenditure calculated by energy balance technique (Fig. 2D–E). After 8 wk of HFD, A-KO mice exhibited severe lipodystrophy phenotypes including dramatically reduced iWAT and eWAT mass, fatty liver, insulin resistance, hyperlipidemia, and reduced serum leptin levels (Fig. 2F–L; Supplemental Fig. S2C–E). These data indicate that mice with adipocyte-selective deletion of *Med1* fail to expand adipose tissue under a HFD.

MED1 is dispensable for embryonic development of BAT

To find out whether MED1 is required for adipose tissue development, we used *Myf5-Cre* mice to delete *Med1* in precursor cells of BAT and skeletal muscle (Tallquist et al. 2000). *Med1^{fl};Myf5-Cre* (M-KO) embryos were

Figure 1. Deletion of *Med1* in adipocytes leads to lipodystrophy. (A–F) All data were from 8-wk male mice fed with a regular diet. (A) Representative morphology of adult mice. (B) Body composition measured by MRI ($n=8$ per group). (C) Representative pictures of BAT, iWAT, eWAT, and liver. iWAT from A-KO mice failed to float in PBS. (D) Average tissue weights are presented as percentage of body weight ($n=6$ per group). (E) H&E staining shows reduced lipid droplet in A-KO adipose tissues. Scale bar, 100 μ m. (F) qRT-PCR analysis of *Med1* mRNA levels ($n=5$ per group). (G) Levels of serum metabolites ($n=8$ per group). (H) Glucose tolerance test (GTT) (left panel) and plasma insulin levels (right panel) ($n=8$ per group). (I) Insulin tolerance test (ITT). Absolute blood glucose levels are shown. (J) Total energy expenditure (EE) after saline or CL316,243 (CL) administration. (K) Lipolysis analysis. Serum levels of free fatty acid or glycerol were measured after saline or CL administration. (L, M) Cold tolerance test. (L) Body temperatures ($n=6$ per group). (M) qRT-PCR of *Ucp1* and *Elov13* in BAT ($n=4$ per group). All quantitative data for mice are presented as means $\pm$ SEM. Statistical comparison between groups was performed using Student's *t*-test. (*) $P < 0.05$, (**) $P < 0.01$.

Figure 2. Mice with adipocyte-selective deletion of *Med1* fail to expand adipose tissue under high-fat diet. Male *f/f* and A-KO mice ($n=8$ per group) were fed with a high-fat diet (HFD) from the eighth week of age. Data for female mice are shown in Supplemental Figure S2. (A–C) Total body weight (A), fat mass (B), and lean mass (C) were measured by MRI during HFD feeding. (D,E) Food intake (D) and total energy expenditure (E). (F) Representative morphology of HFD-fed mice. (G) Representative pictures of the interscapular area (upper area) and abdominal area (lower panel) after 8 wk of HFD feeding. (H) Representative pictures of each fat depot and liver. (I) H&E staining of each fat depot and liver. Scale bar, 100 μ m. (J) Levels of serum metabolites. (K) GTT (left panel) and plasma insulin levels (right panel). (L) ITT. Absolute blood glucose levels are shown. Statistical comparison between groups was performed using Student's *t*-test. (*) $P < 0.05$, (**) $P < 0.01$.

obtained at the expected ratio without showing morphological differences compared with *f/f* embryos (Fig. 3A, B). M-KO embryos did not show observable differences in BAT mass compared with *f/f* embryos (Fig. 3C,D). In addition, *Myf5*-Cre-mediated deletion of *Med1* had little effect on the expression of adipogenesis markers *Pparg*, *Cebpa*, and *Fabp4* but significantly reduced brown adipocyte marker *Ucp1* expression (Fig. 3E). Decreased *Ucp1* expression in *Med1* KO BAT is consistent with previous

observations that MED1 is required for *Ucp1* expression in brown adipocytes in culture (Harms et al. 2015; Iida et al. 2015). Although M-KO mice had largely intact BAT and muscle before birth, they became runts and showed severe growth retardation and reduced survival rates around 3–4 wk after birth, possibly due to muscle development defects (Fig. 3A; Supplemental Fig. S3). These data suggest that MED1 is largely dispensable for embryonic development of BAT.

Figure 3. MED1 is dispensable for embryonic development of BAT. (A–E) *Med1^{fl/f}; Myf5-Cre* (M-KO) mice show largely intact BAT before birth. *Med1^{fl/f}* (f/f) mice were crossed with *Myf5-Cre* to generate M-KO mice. (A) Genotypes of mice at E18.5 and 3–4 wk. The expected ratio of the four genotypes is 1:1:1:1. Numbers of M-KO mice at 3–4 wk were reduced as indicated by an asterisk. Data from 3- to 4-wk M-KO mice are shown in Supplemental Figure S3. (B) Representative morphology of E18.5 embryos. (C) H&E staining of E18.5 embryos. Sagittal sections of the cervical/thoracic area are shown. Scale bar, 80 μ m. (D) H&E staining of BAT at E18.5. (E) qRT-PCR of *Med1*, *Pparg*, *Cebpa*, *Fabp4*, *Ucp1*, and *Scd1* expression in BAT of E18.5 embryos ($n=5$ per group). Statistical comparison between groups was performed using Student's *t*-test. (**) $P < 0.01$.

MED1 is required for postnatal adipose expansion and lipogenesis gene expression

Since the requirement for MED1 to maintain adiposity was observed only after birth, we focused on postnatal adipose expansion in mice, which is stimulated by changes from breastfeeding to carbohydrate-rich diet (Fig. 4A; Pearce 1983). We sought to profile cell type-specific transcriptomes in adiponectin-positive (*Adipoq*⁺) adipocytes during postnatal adipose expansion (Fig. 4B,C). For this purpose, we first crossed *TRAP* (Translating Ribosome Affinity Purification) mice (Zhou et al. 2013) with *Adipoq-Cre* mice to label ribosomes in *Adipoq*⁺ adipocytes. GFP-tagged ribosomes were isolated from interscapular BAT and iWAT of newborn and adult mice for RNA-seq analyses (Fig. 4D). Using a fivefold cutoff for differential gene expression, we identified genes that are induced (1.6%, 214/13,285) or reduced (1.8%, 240/13,285) in *Adipoq*⁺ brown adipocytes from postnatal day 0.5 (P0.5) to 12 wk (Fig. 4E). Induced genes were functionally associated with lipogenesis but not adipogenesis (Fig. 4F). Lipogenesis genes including *Acly*, *Acaca*, *Fasn*, and *Scd1*, which encode four key fatty acid synthesis enzymes, were markedly induced in brown adipocytes from newborn to adult mice (Fig. 4G). Similarly, we identified induced (4.0%, 552/13,799) or reduced (5.4%, 745/13,799) genes in *Adipoq*⁺ white adipocytes from P2.5 to 12 wk (Fig. 4H). Four key lipogenesis enzyme genes, *Acly*, *Acaca*, *Fasn*, and *Scd1*, were induced over fivefold in white adipocytes from P2.5 to 12 wk (Fig. 4G–I). Western blot analyses confirmed

that protein levels of SCD1 and FASN were highly induced during postnatal expansion of BAT and iWAT but not those of adipogenesis markers and master regulators PPAR γ and C/EBP α (Fig. 4J). These results indicate that postnatal adipose expansion is associated with marked induction of lipogenesis enzymes in adipocytes from newborn to adult mice.

To investigate the role of MED1 in postnatal adipose expansion, we isolated fat depots from f/f and A-KO mice at breastfeeding stage (P10) and carbohydrate-rich diet feeding stages (4 wk and 8 wk). We did not observe any discernible differences in total body weight and interscapular WAT between f/f and A-KO mice at P10 (Fig. 5A; Supplemental Fig. S4). However, A-KO mice exhibited nearly complete loss of interscapular WAT as well as reduced BAT and iWAT mass at 4 wk and 8 wk (Fig. 5B,C; Supplemental Fig. S4). These results suggest that MED1 is required for adipose expansion during carbohydrate-rich diet feeding stages but not during breastfeeding. RNA-seq analysis showed that *Adipoq-Cre*-mediated deletion of *Med1* did not affect the expression of adipogenesis marker *Pparg* but impaired the induction of lipogenesis genes including fatty acid synthesis enzyme genes *Acly*, *Acaca*, *Acacb*, *Fasn*, and *Scd1*, triglyceride synthesis enzyme genes *Gpam* and *Agpat2*, and lipid uptake genes *Fatty acid transport 1* (*Fatp1*) and *Fatp2* from P10 to 4 wk and 8 wk in both BAT and iWAT (Fig. 5D–F). Together, these data indicate that MED1 is required for postnatal adipose expansion and lipogenesis gene expression when mice switch from breastfeeding to a carbohydrate-rich diet.

Figure 4. Marked induction of lipogenesis enzymes in adipocytes during postnatal adipose expansion. (A–C) Increased adipose tissue size and lipid accumulation from newborn to adult mice. (A) Schematics of postnatal development stages from newborn to adult mice. (B) Representative pictures of BAT and iWAT of mice at postnatal day 0.5 (P0.5), P2.5, and 8 wk. Adipose tissues from two mice are shown. (C) H&E staining of BAT and iWAT. Scale bar, 80 μ m. (D–J) Lipogenesis enzyme expression is highly induced during postnatal expansion of BAT and iWAT. (D) Schematics of experimental design. *TRAP* mice were crossed with *Adipoq-Cre* to delete the STOP signal and generate mice expressing GFP-fused L10a, an integral component of the 60S ribosomal subunit, in *Adipoq*⁺ adipocytes. GFP-L10a-tagged ribosomes were immunoprecipitated with GFP antibody, and mRNA was purified for RNA-seq. (E,H) RNA-seq analysis of *Adipoq*⁺ adipocytes isolated from BAT (E) or iWAT (H) of *TRAP;Adipoq-Cre* mice at P0.5, P2.5, and 12 wk. The cutoff for induced or reduced genes from P0.5 or P2.5 to 12 wk is fivefold. (F,I) Gene ontology (GO) analysis of gene groups defined in E and H. (G) Expression levels of representative genes are shown in RPKM values of RNA-seq data from BAT and iWAT of *TRAP;Adipoq-Cre* mice. *Gpm* and *Fatp1* are also known as *Gpat1* and *Slc27a1*, respectively. (J) Western blot analysis of FASN, SCD1, C/EBP α , and PPAR γ using whole-cell lysates from BAT (left panel) or iWAT (right panel). β -Actin was used as loading control.

Figure 6. MED1 is required for lipogenesis but not early adipogenesis in culture. Immortalized *Med1^{flf};Cre-ER* brown preadipocytes were treated with 4OHT to delete *Med1*, followed by adipogenesis assay, RNA-seq, or nascent RNA-seq. (A–C) MED1 is required for lipid accumulation and lipogenesis gene expression in culture. (A, left panel) Oil Red O staining at day 7 (D7) of adipogenesis is shown. (Right panel) Lipid accumulation was quantified by extracting Oil Red O using isopropanol and reading absorbance at 510 nm. $n = 3$ biological replicates. Data are presented as means $\pm$ SD. Statistical comparison between groups was performed using Student's *t*-test. (*) $P < 0.05$. (B) RNA-seq analysis at D7 of adipogenesis. The cutoff for differential expression is 2.5-fold. (C) GO analysis of gene groups defined in B. (D–F) MED1 is required for transcription of lipogenesis genes. (D) Expression levels of representative genes are shown in RPKM values. (E) Nascent RNA-seq analysis at D7 of adipogenesis. The cutoff for differential expression is twofold. (F) GO analysis of gene groups defined in E. (G,H) Decreased *Chrebp β* expression in *Med1* KO brown adipocytes. (G) Profiles of nascent RNA-seq data around *Chrebp* locus. (H) qRT-PCR of *Chrebp α* and *Chrebp β* expression at D7 of adipogenesis. All qPCR data in cells are presented as means $\pm$ SD. Two technical replicates from a single experiment were used.

MED1 is required for lipogenesis but not early adipogenesis in culture

To understand how MED1 regulates lipogenesis, we established inducible *Med1* KO (*Med1*^{fl/fl}; *Cre-ER*) brown preadipocytes. Cells were treated with 4-hydroxytamoxifen (4OHT) to delete *Med1*, followed by the induction of adipogenesis. Consistent with the A-KO mouse phenotype, *Med1* KO adipocytes showed a >50% decrease in lipid content at day 7 (D7) of adipogenesis (Fig. 6A). RNA-seq analysis confirmed deletion of exons 8–10 of the *Med1* gene (Supplemental Fig. S5A). MED1 is a subunit of general transcription coactivator complex Mediator. However, only about 1% of expressed genes were down-regulated in *Med1* KO adipocytes at D7 (Fig. 6B). GO analysis showed that down-regulated genes were strongly functionally associated with fatty acid biosynthesis (Fig. 6C). Consistent with data obtained from mouse adipose tissues, deletion of *Med1* led to decreases in expression of lipogenesis genes *Acly*, *Acaca*, *Acacb*, *Fasn*, *Scd1*, *Gpm*, and *Agpat2* but not adipogenesis marker *Pparg* in immortalized and primary adipocytes differentiated in culture (Fig. 6D; Supplemental Figs. S5B, S6).

To better assess the role of MED1 in regulating lipogenesis gene transcription, we performed nascent RNA-seq analysis. Using a twofold cutoff, we found that only ~2% of expressed genes were down-regulated transcriptionally in *Med1* KO adipocytes at D7 (Fig. 6E). Consistent with steady-state RNA-seq data, nascent RNA-seq analysis indicated that deletion of *Med1* reduces transcription levels of lipogenesis genes *Acly*, *Acaca*, *Acacb*, *Fasn*, *Scd1*, and *Gpm* but not adipogenesis markers *Pparg* and *Cebpa* (Fig. 6D–F). Interestingly, while *Med1* KO did not affect the transcription of lipogenic TFs LXR α , LXR β , and SREBP1, it specifically reduced transcription of the *Chrebp* β isoform, which encodes a key lipogenic TF in adipocytes (Fig. 6G,H; Herman et al. 2012; Vijayakumar et al. 2017), suggesting that ChREBP may play a role in MED1-dependent lipogenesis. Together, these results suggest that MED1 is required for lipogenesis but not early adipogenesis in culture.

MED1 is required for Pol II binding on lipogenesis genes in adipocytes

To investigate whether MED1 affects chromatin accessibility and enhancer activation, we performed ATAC-seq (assay for transposase accessible chromatin with high-throughput sequencing) and ChIP-seq analyses of active enhancer mark H3K27ac, and RNA polymerase II (Pol II) at D7 of adipogenesis. On *Med1*⁺ promoters or enhancers in adipocytes, deletion of *Med1* did not affect chromatin accessibility or H3K27ac levels (Fig. 7A,B). However, *Med1* KO led to decreased binding of serine 5 phosphorylated (S5P) initiating Pol II, while having limited effects on serine 2 phosphorylated (S2P) elongating Pol II (Fig. 7A,B; Komarnitsky et al. 2000). GO analysis revealed that genes associated with MED1-dependent S5P-Pol II were strongly functionally associated with lipid metabolism including fatty acid biosynthesis (Fig. 7C). Consistent with nascent

RNA-seq data, deletion of *Med1* clearly reduced both S5P-Pol II and S2P-Pol II binding on *Scd1*, *Fasn*, *Agpat2*, *Acaca*, and *Gpm* loci but not the *Pparg* locus (Fig. 7D,E; Supplemental Fig. S7). These data indicate that, while MED1 is dispensable for chromatin opening and enhancer activation, it is required for Pol II binding on lipogenesis genes in adipocytes.

MED1 facilitates Mediator binding on ChREBP⁺ SREBP1a⁺ lipogenic enhancers in adipocytes

We hypothesized that MED1 regulates Pol II binding on lipogenesis genes by facilitating Mediator binding on lipogenic enhancers. To test this hypothesis, we first identified lipogenic enhancers that are bound by ChREBP or SREBP1a. *Med1*^{fl/fl}; *Cre-ER* brown preadipocytes were infected with retroviruses expressing the triple T7 tagged constitutive active form of ChREBP (T7-CA-ChREBP) or the nuclear form of SREBP1a (T7-nSREBP1a) (Supplemental Fig. S8A). Cells were treated with 4OHT to delete *Med1*, followed by the induction of adipogenesis. We found that both ectopic ChREBP and SREBP1a failed to rescue the reduced lipogenesis in *Med1* KO adipocytes (Fig. 8A,B). Next, we performed ChIP-seq using antibodies against T7 and MED12, which is a representative Mediator subunit (Kagey et al. 2010), at D7 of adipogenesis. Motif analysis of the T7-CA-ChREBP and T7-nSREBP1a binding regions identified the ChREBP and SREBP motifs as top motifs, respectively (Supplemental Fig. S8B). We identified active enhancers that are bound by ChREBP (2627) or SREBP1a (2183) (Supplemental Fig. S9A,E). The majority of these enhancers were bound by both ChREBP and SREBP1a (ChREBP⁺ SREBP1a⁺, 1732). A subset of these lipogenic enhancers (ChREBP⁺, 934/2627; SREBP1a⁺, 915/2183; and ChREBP⁺ SREBP1a⁺, 819/1732) showed reduced MED12 binding in *Med1* KO adipocytes (Fig. 8C; Supplemental Figs. S8C, S9B,C,F,G, J). Notably, lipid metabolism was a top GO term associated with genes that showed MED1-dependent MED12 binding (Fig. 8D; Supplemental Fig. S9D,H). Moreover, ChREBP- and SREBP1a-associated genes that show MED1-dependent MED12 binding exhibited reduced expression in *Med1* KO cells (Fig. 8E), indicating that MED1 facilitates MED12 binding on ChREBP⁺ SREBP1a⁺ lipogenic enhancers to regulate lipogenesis gene expression in adipocytes. Interestingly, ChREBP and SREBP1a directly bind to triglyceride synthesis genes *Agpat2* and *Gpm* in addition to fatty acid synthesis enzyme genes *Scd1*, *Fasn*, *Acaca*, and *Acly* (Fig. 8F; Supplemental Fig. S10). Deletion of *Med1* clearly reduced MED12 binding to lipogenesis genes *Scd1*, *Fasn*, *Acaca*, *Acly*, *Agpat2*, and *Gpm* loci but not the *Pparg* locus in adipocytes. Together with previous data, these results suggest that MED1 is a lipogenesis coactivator required for Mediator binding on ChREBP⁺ SREBP1a⁺ lipogenic enhancers in adipocytes (Fig. 8G).

Discussion

By crossing *Med1*^{fl/fl} with *Adipoq-Cre* or *Myf5-Cre* mice, we selectively deleted *Med1* in adipocytes or precursor

Figure 7. MED1 is required for Pol II binding on lipogenesis genes in adipocytes. Adipogenesis was done as in Figure 6. Cells were collected at D7 of adipogenesis for ATAC-seq and ChIP-seq of H3K27ac, S5P-Pol II, and S2P-Pol II. (A,B) *Med1* KO does not affect chromatin accessibility or enhancer activation but reduces S5P-Pol II binding on MED1⁺ promoters and enhancers in adipocytes. Heat maps (A) and average profiles (B) were aligned around the center of MED1 binding sites on MED1⁺ promoters (4117) and enhancers (8377). Published MED1 ChIP-seq data were used (GSE74189) (Lai et al. 2017). (C) GO analysis of genes associated with MED1-dependent S5P-Pol II. (D,E) MED1 is required for Pol II binding on lipogenesis genes. Profiles of ATAC-seq, H3K27ac enrichment, S5P-Pol II, S2P-Pol II binding, and nascent RNA-seq data around *Pparg*, *Scd1*, *Fasn*, and *Agpat2* gene loci are shown. Profiles around additional lipogenesis genes are shown in Supplemental Figure S7.

Figure 8. MED1 facilitates Mediator binding on ChREBP⁺ SREBP1a⁺ lipogenic enhancers in adipocytes. (A,B) Ectopic CA-ChREBP or nSREBP1a fails to rescue reduced lipogenesis in *Med1* KO. (A) Oil Red O staining at D7 of adipogenesis. (B) qRT-PCR of *Scd1*, *Fasn*, and *Pparg* expression. Two biological replicates were used. (C,D) MED1 is required for MED12 binding on a subset of ChREBP⁺ SREBP1a⁺ active enhancers. (C) Average profiles around the center of T7-CA-ChREBP⁺ active enhancers. (D) GO analysis of genes that are associated with either MED1-dependent or MED1-independent MED12 binding on ChREBP⁺ SREBP1a⁺ active enhancers. (E) MED1 facilitates MED12 binding to promote ChREBP- and SREBP1a-associated gene expression. Fold changes of RPKM values between control (Mock) and *Med1* KO (4OHT) cells are shown in box plots. Statistical significance levels are indicated (Wilcoxon signed rank test, two-sided). (F) MED1 is required for MED12 binding on lipogenesis genes. Profiles of T7-CA-ChREBP, T7-nSREBP1a, and MED12 binding on *Scd1*, *Fasn*, and *Agpat2* gene loci are shown. (G) Proposed model.

cells in vivo. We found, surprisingly, that MED1 is required for postnatal adipose expansion but is largely dispensable for the development of adipose tissues. Consistently, MED1 is required for lipid accumulation but not early differentiation during adipogenesis in culture. Transcriptome analysis in mice and in culture indicates that only 2%–3% of expressed genes are down-regulated by *Med1* KO in adipocytes. Down-regulated genes include key lipogenesis enzymes such as SCD1, FASN, and AGPAT2, but not early adipogenesis markers such as PPAR γ . Nascent RNA-seq in adipocytes reveals that *Med1* KO specifically reduces transcription of the ChREBP β isoform but not other lipogenic TFs. We further demonstrate that MED1 regulates Pol II binding on lipogenesis genes by promoting Mediator binding on ChREBP $^{+}$ SREBP1a $^{+}$ lipogenic enhancers in adipocytes. Together, our findings suggest that the MED1 subunit of the Mediator coactivator complex regulates postnatal adipose expansion by promoting lipogenesis gene expression.

By crossing *Med1*^{fl} with *Myf5-Cre* mice, we observed that *Med1* KO had little effect on the size of embryonic BAT and the expression of adipogenesis markers such as PPAR γ and C/EBP α . A-KO mice showed similar BAT and iWAT tissue masses and adipogenesis marker expression as the control mice at P10. In culture, deletion of *Med1* in preadipocytes does not affect PPAR γ or C/EBP α expression during differentiation of white and brown adipocytes. Together, these results indicate that MED1 is largely dispensable for the general development of adipose tissues. The requirement of MED1 for *Ucp1* induction in BAT and iWAT is consistent with previous findings in cell culture that PRDM16 physically binds to and recruits MED1 to active enhancers of BAT-selective genes and that both PRDM16 and MED1 are required for *Ucp1* induction in brown adipocytes (Harms et al. 2015; Iida et al. 2015). We also show that MED1 is largely dispensable for embryonic development of muscle. However, M-KO mice show severe growth retardation and mortality around weaning age, suggesting that MED1 may play a role in muscle development and function after birth. These observations suggest that MED1 plays functional roles in late stages of adipogenesis and myogenesis rather than early stages of cell differentiation and tissue development.

Using adipocyte-specific transcriptome analysis, we show that postnatal adipose expansion is associated with marked induction of fatty acid synthesis enzymes (Fig. 4D–J). Transcriptome analyses at P10, 4 wk, and 8 wk also reveal marked induction of fatty acid synthesis enzymes as well as lipogenic TF *Chrebp* in BAT and iWAT when mice switch from maternal milk to a carbohydrate-based diet (Fig. 5F). These results explain previous observations that lipogenic enzyme activity and fatty acid synthesis in adipose tissues increase markedly after mice switch from breastfeeding to a carbohydrate-based diet (Pearce 1983). A-KO mice exhibit increasing lipodystrophy after weaning but not during the breastfeeding stage. A-KO mice also show severely impaired induction of fatty acid synthesis genes in BAT and iWAT when switching from high-fat maternal milk to a carbohydrate-based diet.

These data suggest that MED1 is required for carbohydrate-rich diet-induced postnatal adipose expansion and that lipodystrophy in A-KO mice is at least in part due to impaired induction of fatty acid synthesis genes. Consistently, *Med1* KO reduces lipid accumulation and impairs the induction of fatty acid synthesis genes in adipocytes in culture.

We also observed impaired induction of *Agpat2*, *Perilipin 1* (*Plin1*), *Fatp1*, and *Fatp2* in adipose tissues of A-KO mice (Fig. 5F), which likely contributes to the observed lipodystrophy phenotype. AGPAT2 is a critical enzyme for triglyceride synthesis and is highly expressed in adipose tissues. Mutations of AGPAT2 lead to congenital generalized lipodystrophy in humans (Agarwal et al. 2002). Whole-body *Agpat2* KO mice show a severe lipodystrophy phenotype similar to A-KO mice (Cortes et al. 2009). PLIN1 is an adipocyte-specific lipid-droplet coat protein that regulates fat storage and breakdown. Whole-body *Plin1* KO mice show reduced adipose tissue mass (Martinez-Botas et al. 2000). Mutations of *PLIN1* are associated with familial partial lipodystrophy in humans (Gandotra et al. 2011). A-KO mice show impaired induction of *Agpat2* and *Plin1* in BAT and iWAT from P10 to 4 wk (Fig. 5F). *Med1* KO reduces *Agpat2* expression in primary white and brown adipocytes in culture (Supplemental Fig. S6G). These results suggest that the impaired induction of *Agpat2* and/or *Plin1* likely contributes to the lipodystrophy in A-KO mice. Whole-body *Fatp1* KO mice show reduced eWAT mass due to a significant reduction in adipocyte sizes (Wu et al. 2006). We cannot exclude the possibility that decreased expression of *Fatp1/2* contributes to lipodystrophy in A-KO mice.

In *Med1* KO adipocytes, only 1%–2% of the ~16,000 expressed genes are down-regulated, indicating that MED1 is not generally required for transcription. Down-regulated genes are preferentially associated with lipogenesis, in particular fatty acid and triglyceride synthesis, such as *Acaca*, *Elovl6*, *Fasn*, *Scd1*, *Agpat2*, *Gpm*, and *Chrebp* β . ChREBP is a major transcriptional regulator of fatty acid synthesis in adipocytes. Vijayakumar et al. reported that Adipoq-Cre-mediated KO of *Chrebp* reduces the expression of lipogenesis genes including *Acaca*, *Elovl6*, *Fasn*, and *Scd1* in BAT and iWAT and significantly decreases perigonadal WAT mass (Vijayakumar et al. 2017). *Chrebp* KO led to a compensatory increase in *Srebp1* expression in adipocytes, which could explain the mild lipodystrophy phenotype in these mice. SREBP1 is another critical transcriptional regulator for fatty acid synthesis in adipocytes (Horton et al. 2003). However, the whole-body KO of SREBP1 does not affect adiposity or DNL enzyme expression in WAT (Shimano et al. 1997). The fact that KO of either ChREBP or SREBP1 has mild or little effect on adipose tissue mass in mice suggests that ChREBP and SREBP1 have redundant functions in regulating lipogenesis gene expression in adipocytes. We show that MED1 is required for Mediator binding on ChREBP $^{+}$ and/or SREBP1a $^{+}$ lipogenic enhancers. ChREBP and SREBP1a directly bind to not only fatty acid synthesis genes *Scd1*, *Fasn*, *Acaca*, and *Acl* but also triglyceride synthesis genes *Agpat2* and *Gpm* (Fig. 8F; Supplemental

Fig. S10). Deletion of *Med1* leads to reduced MED12 binding on a broad range of lipogenesis genes. Our findings suggest that MED1 is a master lipogenesis coactivator that cooperates with lipogenic TFs ChREBP and SREBP1a in adipocytes (Fig. 8G). Future work will be needed to simultaneously inactivate ChREBP and SREBP1 to determine whether these two TFs represent the major pathways by which MED1 regulates postnatal adipose expansion. Our current analysis uses a candidate approach and involves ChREBP and SREBP1. We cannot exclude the possibility that additional TFs require MED1 for activating lipogenesis genes in adipocytes.

Previous studies have implicated other Mediator subunits in SREBP1-dependent lipogenesis gene expression. A GST-fused SREBP1a activation domain can pull down the Mediator complex from cell nuclear extracts. SREBP1a directly interacts with MED15 (also called ARC105) *in vitro*. Depletion of MED15 down-regulates SREBP1a-dependent *FASN* expression in human cells (Yang et al. 2006). CDK8 and Cyclin C, which associate with the transcriptionally inactive Mediator complex, have been reported as negative regulators of *de novo* lipogenesis in *Drosophila* and mice (Zhao et al. 2012). CDK8 and Cyclin C are dissociated from the Mediator complex upon refeeding with a carbohydrate-rich diet after fasting (Youn et al. 2019). Our finding that MED1 is a lipogenesis coactivator in adipocytes is consistent with these previous reports. Detailed molecular mechanisms by which Mediator regulates lipogenesis through its subunits including MED1, MED15, CDK8, and Cyclin C remain to be fully understood.

Materials and methods

Plasmids, antibodies, and chemicals

A triple T7 (3×T7) double StrepII tag was N-terminally subcloned into retroviral vector pMSCVhyg to generate pMSCVhyg-3xT7 using a pTAG1-hygroTK-N-term 3xT7 double StrepII tag (Brown et al. 2017) as a template. Constitutive active form of mouse ChREBP or nuclear form of mouse SREBP1a was PCR-amplified using full-length clones as templates (Addgene 39235 and 32017) and was subcloned into pMSCVhyg-3xT7 to generate pMSCVhyg-3xT7-CA-ChREBP or pMSCVhyg-3xT7-nSREBP1a. All plasmids were confirmed by DNA sequencing. Anti-MED1 (A300-793A) and anti-MED12 (A300-774A) were from Bethyl Laboratories. Anti-FASN (no. 3810), anti-SCD1 (no. 2794), and anti-T7 (no. 132469) were from Cell Signaling Technology. Anti-C/EBPα (sc-61X) and anti-PPARγ (sc-7196X) were from Santa Cruz Biotechnology. Anti-H3K27ac (ab4729), anti-GFP (ab290), anti-S5P Pol II (ab5131), and anti-S2P Pol II (ab5095) were from Abcam. Anti-β-Actin (A1978) and (Z)-4-hydroxytamoxifen (4OHT) (H7904) were from Sigma.

Generation of mouse strains

Med1^{fl} mice (Jia et al. 2004) were crossed with *Myf5-Cre* (Jackson 007893), *Adipoq-Cre* (Jackson 028020), or *Cre-ER* (Jackson 008463) to generate *Med1*^{fl}/*Myf5-Cre*, *Med1*^{fl}/*Adipoq-Cre*, or *Med1*^{fl}/*Cre-ER*. For genotyping of *Med1* alleles, PCR was performed using the following primers: 5'-TCTCCCCGCTAA TATTCATA-3' and 5'-AAGGAACAAGCCAGCAAGC-3'. PCR

amplified 839 bp from the wild-type and 575 bp from the floxed allele. *Rosa26*^{lsTRAP} (*TRAP*) mice (Jackson 022367) (Zhou et al. 2013) were crossed with *Adipoq-Cre* (Jackson 028020) to generate *TRAP;Adipoq-Cre* mice. For genotyping of *TRAP* alleles, PCR was performed using the following primers: common forward, 5'-AAGGGAGCTGCAGTGGAGTA-3'; wild type reverse, 5'-CCGAAAATCTGTGGGAAGTC-3'; mutant reverse, 5'-CG GGCCATTACCGTAAGTTAT-3'. PCR amplified 197 bp from the wild-type and 284 bp from the *TRAP* allele. All mouse experiments were performed in accordance with the NIH Guide for the Care and Use of Laboratory Animals and approved by the Animal Care and Use Committee of the National Institute of Diabetes and Digestive and Kidney Diseases, National Institutes of Health.

Translating ribosome affinity purification

Brown and inguinal white adipose tissues from newborn (P0.5 for BAT or P2.5 for WAT) and adult (12 wk) *TRAP;Adipoq-Cre* mice were minced in homogenization buffer (50 mM Tris at pH 7.5, 12 mM MgCl₂, 100 mM KCl, 1% NP-40, 100 mg/mL cycloheximide, 1 mg/mL sodium heparin, 2 mM DTT, 0.2 U/mL RNasin, protease inhibitors) and homogenized by Dounce (Type B, 2 mL). After incubation for 10 min on ice, lysates were centrifuged at 13,000g for 10 min at 4°C and the supernatant was transferred to new tubes after removing the top lipid layer. The supernatant was incubated with anti-GFP antibody (5 μg/mL) prebound to Dynabeads Protein A (Invitrogen) for 2 h with end-over-end rotation to capture the GFP-fused L10a subunit of the 60S ribosome. Beads were collected on a magnetic rack and washed twice with low-salt wash buffer (50 mM Tris at pH 7.5, 12 mM MgCl₂, 100 mM KCl, 1% NP-40, 100 mg/mL cycloheximide, 2 mM DTT) and then three times with high-salt wash buffer (50 mM Tris at pH 7.5, 12 mM MgCl₂, 300 mM KCl, 1% NP-40, 100 mg/mL cycloheximide, 2 mM DTT). Immunoprecipitated ribosomes were immediately placed in the RLT buffer to extract RNA using the Qiagen Micro RNeasy kit (Qiagen) following the manufacturer's protocol.

Metabolic studies

For determination of serum metabolites, blood was collected from mice fed with standard laboratory mouse chow (15% calories from fat, 56% calories from carbohydrate, 29% calories from protein; Envigo 7022 NIH-07) or high-fat diet (60% calories from fat, 20% calories from carbohydrate, 20% calories from protein; Research Diets D12492). Serum was obtained by centrifuging blood samples at 12,000g for 4 min at room temperature. Serum insulin and leptin concentrations were measured by an ELISA kit from CrystalChem and R&D Systems, respectively. Serum free fatty acid, triglyceride, and cholesterol concentrations were measured with reagents from Roche Diagnostics GmbH; Pointe Scientific, Inc.; and Thermo Scientific, respectively. For GTTs, mice were fasted overnight for 16 h. For ITTs, mice were fasted for 4 h. Mice received glucose (1 g/kg i.p. [intraperitoneally]), or human insulin (0.75 U/kg i.p.; Humulin, Eli Lilly), respectively, and blood was collected from the tail vein at specific time points. For both tests, blood glucose levels were determined using a portable glucometer (Contour Glucometer, Bayer). Serum concentrations of free fatty acid and glycerol were analyzed with reagents from Roche and Sigma, respectively. Food intake, O₂ consumption, CO₂ production, and locomotor activity were measured over 24-h periods at 22°C and 29.5°C in a comprehensive laboratory animal monitor system (CLAMS) system (2.5-L chambers with plastic floors, using 0.6 L/min flow rate, one mouse per chamber; Columbus Instruments, Inc.) after a 48-h adaptation

period. For the cold tolerance test, mice were individually housed at room temperature (22°C) and then in a cold room (6°C) for 6 h ($n = 6$ per group). Core body temperature was measured using a rectal thermometer (TH-5; Braintree Scientific) before and hourly after cold exposure. Body composition was measured with the EchoMRI 3-in-1 analyzer (Echo Medical Systems). CL316,243 (0.1 mg kg⁻¹; Sigma-Aldrich) or saline were administered intraperitoneally and O₂ consumption was measured at 30°C from 1 to 4 h after injection. For *in vivo* lipolysis, blood was collected 20 min after the injection of CL316,243 or saline. In mice fed with HFD, energy expenditure was calculated by the energy balance technique (Ravussin et al. 2013; Zhuang et al. 2018).

Primary preadipocytes culture, immortalization, and adipogenesis

Primary brown preadipocytes were isolated from interscapular BAT of newborn *Med1*^{fl}; *Cre-ER* pups and immortalized by infecting retroviruses expressing SV40T. Immortalized cells were further infected with retroviruses expressing 3xT7-CA-ChREBP or the 3xT7 vector. Primary white adipocytes were from inguinal WAT of *Med1*^{fl}; *Adipoq-Cre* or *Med1*^{fl} adult mice. For the adipogenesis assay, preadipocytes were plated in growth medium (DMEM plus 10% FBS) 3–4 d before the induction and were induced with 0.02 μM insulin, 1 nM T3, 0.5 mM IBMX, 2 μg mL⁻¹ DEX, and 0.125 mM indomethacin for 2 d. For adipogenesis of primary white preadipocytes, 1 μM rosiglitazone was included throughout the differentiation (Park and Ge 2017; Park et al. 2017).

Western blot and qRT-PCR

Western blot of nuclear extracts or whole-tissue lysates was done as described (Jang et al. 2019). Total RNA was extracted using TRIzol (Invitrogen) and reverse-transcribed using a ProtoScript II first strand cDNA synthesis kit (NEB), following the manufacturers' protocols. qRT-PCR of *Med1* exon 8 was performed using SYBR green primers: forward 5'-CCTGTTTGATGGGATGTCCA-3', and reverse, 5'-GCAGAGATATGCAGATTGCC-3'. *Chrepa* and *Chrebpβ* qRT-PCR primers were described previously (Herman et al. 2012). SYBR green primers for other genes were described previously (Jin et al. 2011; Park and Ge 2017).

RNA-seq library preparation

Total RNA (1 μg) or TRAP-isolated RNA (200 ng) was subjected to the NEBNext poly(A) mRNA magnetic isolation module (NEB) to isolate mRNA and proceeded directly to double-stranded cDNA synthesis. Library construction was done using the NEBNext Ultra II RNA library preparation kit for Illumina (NEB) following the manufacturer's protocol.

Nascent RNA-seq library preparation

Cells (10⁶) were labeled with 0.5 mM ethylene uridine for 30 min at 37°C. After RNA extraction, 1 μg of total RNA was depleted of rRNA using the NEBNext rRNA depletion kit (NEB). To capture nascent RNA, the sample was biotinylated by the Click-iT nascent RNA capture kit (Thermo Fisher C10365) according to the manufacturer's instructions. Double-stranded cDNAs were synthesized using a SuperScript double-stranded cDNA synthesis kit (Invitrogen). Library construction was done using a NEBNext Ultra II DNA library preparation kit for Illumina (NEB) according to the manufacturer's instructions. Sequencing libraries were analyzed with Qubit and pooled and sequenced on a HiSeq3000.

ChIP-seq and ATAC-seq library preparation

ChIP-seq was performed as described in detail previously (Lee et al. 2013, 2017; Lai et al. 2017; Zhuang et al. 2018). ChIP-seq of T7 and MED12 were done in the presence of *Drosophila* spike-in chromatin and antibody following the manufacturers' protocol (Active Motif). ChIP-seq library construction was done using a NEBNext Ultra II DNA library preparation kit for Illumina (NEB) following the manufacturer's protocol. For ATAC-seq, 50,000 cells were washed with PBS and collected in a cold lysis buffer (10 mM Tris-HCl at pH 7.4, 10 mM NaCl, 3 mM MgCl₂, 0.1% Igepal CA-630). After lysis, the nuclear pellet was resuspended in the transposase reaction mix (Illumina, Nextera Tn5 transposase kit) and incubated for 30 min at 37°C. Immediately following transposition, the sample was purified by a Qiagen MinElute kit (Qiagen). Eluted DNA was amplified with PCR using Nextera i7 and i5 index primers (Illumina) and purified with AMPure XP magnetic beads (Beckman Coulter). Sequencing libraries were analyzed with Qubit and pooled and sequenced on a HiSeq2500 or HiSeq3000.

Computational analysis

RNA-seq data analysis Raw sequencing data were aligned to the mouse mm9 genome using STAR software (Dobin et al. 2013). Reads on exonic regions or gene bodies (nascent RNA-seq) were collected to calculate reads per kilobase per million (RPKM) as a measure of gene expression level. Only genes with RPKM > 1 were considered expressed. Gene ontology (GO) analysis of differentially expressed genes was carried out using DAVID bioinformatics resources (<https://david.ncifcrf.gov>).

ChIP-seq peak calling and GO analysis of genomic regions Raw sequencing data were aligned to the mouse mm9 genome using bowtie2. To identify ChIP-enriched regions, we used the "SICER" method (Zang et al. 2009). For H3K27ac enrichment, the window size of 200 bp and the estimated false discovery rate (FDR) threshold of 10⁻¹⁰ were used. For ChIP-seq of S5P-Pol II, S2P-Pol II, T7, and MED12, the window size of 50 bp and the FDR threshold of 10⁻³ were used. Previously published MED1 ChIP-seq data were used (GSE74189). GO analysis of genomic regions was done using GREAT (<http://great.stanford.edu/public/html>).

Motif analysis For motif analysis of 3xT7-CA-ChREBP or 3xT7-nSREBP1a binding sites, we used the SeqPos motif tool in Galaxy Cistrome (<http://cistrome.org/ap/root/>) with default parameters. We selected the top 3000 binding regions to screen enriched TF motifs based on the FDR value provided by SICER.

Normalization of ChIP-seq data For ChIP-seq spike-in normalization, sequences were aligned to the *Drosophila* genome dm6. Normalization factors were determined by counting *Drosophila* tags and applying to mouse tags to generate normalized heat maps and profiles (Fig. 8; Supplemental Fig. S8).

Heat maps and box plots Heat map matrices were generated using in-house scripts with 50-bp resolution and visualized in R using gplots package. The ratio of RPKM values in *Med1* KO and control cells in base 2 logarithm was plotted using box plot, with outliers not shown (Fig. 8). A Wilcoxon signed rank test (two-sided) was used to determine statistical differences.

ATAC-seq data analysis Raw sequencing reads were processed using Kundaje laboratory's atac pipelines (https://github.com/kundajelab/atac_dnase_pipelines). For downstream analysis, we

used filtered reads that remained after removing unmapped reads, duplicates, and mitochondrial reads.

Data availability

RNA-seq, ATAC-seq, and ChIP-seq data sets generated in the paper have been deposited in NCBI Gene Expression Omnibus under accession number GSE160605.

Competing interest statement

The authors declare no competing interests.

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Author contributions: Y.J. and K.G. conceived the study. Y.J., Y.-K.P., J.-E.L., and O.G. performed the methodology. Y.J., Y.-K.P., J.-E.L., D.W., O.G., N.T., and K.G. performed the investigation. J.-E.L., Y.J., and Y.-K.P. were responsible for the software and performed the formal analysis and data curation. Y.J., J.-E.L., and K.G. wrote the original draft of the manuscript. Y.J., J.-E.L., Y.-K.P., N.T., and K.G. reviewed and edited the manuscript. K.G. administered the project and acquired the funding.

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