miR-26 suppresses adipocyte progenitor differentiation and fat production by targeting *Fbxl19*

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Fat storage in adult mammals is a highly regulated process that involves the mobilization of adipocyte progenitor cells (APCs) that differentiate to produce new adipocytes. Here we report a role for the broadly conserved miR-26 family of microRNAs (miR-26a-1, miR-26a-2, and miR-26b) as major regulators of APC differentiation and adipose tissue mass. Deletion of all miR-26-encoding loci in mice resulted in a dramatic expansion of adipose tissue in adult animals fed normal chow. Conversely, transgenic overexpression of miR-26a protected mice from high-fat diet-induced obesity. These effects were attributable to a cell-autonomous function of miR-26 as a potent inhibitor of APC differentiation. miR-26 blocks adipogenesis, at least in part, by repressing expression of *Fbxl19*, a conserved miR-26 target without a previously known role in adipocyte biology that encodes a component of SCF-type E3 ubiquitin ligase complexes. These findings have therefore revealed a novel pathway that plays a critical role in regulating adipose tissue formation in vivo and suggest new potential therapeutic targets for obesity and related disorders.

[**Keywords:** adipocyte progenitor cell; adipogenesis; *Fbxl19*; obesity; white adipose tissue; miR-26; microRNA]

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MicroRNAs (miRNAs) comprise a class of diverse ~21- to 23-nt noncoding RNAs that were first discovered in *C. elegans* and subsequently found to regulate the expression of a large fraction of metazoan mRNAs (Krol et al. 2010; Bartel 2018). Encoded either within the introns of protein-coding genes or transcribed under the control of their own promoters, miRNAs are produced through a defined processing pathway involving sequential endonuclease-mediated cleavages by the RNase III enzymes DROSHA and DICER (Ha and Kim 2014). After maturation, miRNAs are loaded into a member of the Argonaute (AGO) family of proteins, which they guide to sites of partial complementarity predominantly within the 3' UTRs of target mRNAs, resulting in accelerated deadenylation, increased turnover, and reduced translation (Jonas and Izaurralde 2015; Gebert and MacRae 2019). Through the

regulation of extensive target repertoires, miRNAs broadly impact mammalian development and physiology, while miRNA dysfunction has been causatively linked to several human diseases including cancer and cardiovascular disease (Bartel 2009; Mendell and Olson 2012).

Obesity, and associated metabolic abnormalities such as Type 2 diabetes represent a growing public health crisis in Western nations due to the association with cardiovascular disease and all-cause mortality (Martin et al. 2015). Several studies have uncovered important roles for miRNAs in regulating obesity-associated metabolism due to their ability to modulate cellular responses to dietary and metabolic cues such as cholesterol, glucose, and insulin (Najafi-Shoushtari et al. 2010; Rayner et al. 2010;

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Dooley et al. 2016). A fundamental characteristic of obesity is the uncontrolled expansion and accumulation of white adipose tissue (WAT) (Berry et al. 2013). In addition to storing energy during times of caloric surplus, WAT functions as a critical node in the regulation of systemic metabolism by secreting signaling proteins and lipids that influence key physiologic responses such as appetite, glucose homeostasis, and immune responses (Deng and Scherer 2010; Blüher 2014). Under obesogenic conditions, pre-existing adipocytes expand to accommodate excess nutrients in the form of triglycerides. Additionally, new adipocytes are generated from a proliferative population of adipocyte progenitor cells (APCs) that localize to the WAT vasculature where they resemble mural cells. APCs can be distinguished from other nonadipogenic mural cells due to their expression of the master regulator of adipogenesis, peroxisome proliferator-activated receptor γ (PPAR γ 2) (Tontonoz et al. 1994; Wu et al. 1999; Rosen and MacDougald 2006; Tontonoz and Spiegelman 2008). Genome-wide profiling has revealed that PPAR γ 2 occupies the promoters of numerous adipogenic genes to drive adipogenesis and terminal differentiation (Lefterova et al. 2008; Nielsen et al. 2008). While the transcriptional hierarchy that drives APC mobilization and adipogenesis has been extensively characterized, the importance of miRNA-mediated regulation in this process remains unclear. Further insight into the functions of miRNAs within this lineage would illuminate our understanding of adipocyte development and function, and might reveal new therapeutic approaches to counteract obesity.

The miR-26 family (miR-26a-1, miR-26a-2, and miR-26b in mammals) represents a highly conserved family of vertebrate miRNAs that have been implicated in a broad array of developmental and physiological functions, including neuronal differentiation (Dill et al. 2012), muscle development (Dey et al. 2012; Icli et al. 2013), and hepatic glucose and lipid metabolism (Fu et al. 2015). In addition, miR-26 family members have been implicated as tumor suppressors in multiple types of cancer, including hepatocellular carcinoma (Ji et al. 2009; Kota et al. 2009) and colorectal cancer (Zeitels et al. 2014). Despite the extensive prior investigation of the functions of these miRNAs, however, our understanding of their importance in mammalian development and physiology has been limited by the difficulty in studying the consequences of miR-26 loss-of-function in vivo. The three ubiquitously expressed and likely redundant mammalian miR-26 family members are produced from unlinked loci, which has deterred previous efforts to produce complete miR-26 knockout mice.

In this study, we report the application of CRISPR/Cas9-mediated genome editing to generate mice lacking all miR-26 family members. Although analysis of these animals showed that these miRNAs are dispensable for overtly normal development, a critical role for miR-26 as a physiologic suppressor of adult adipogenesis was unexpectedly revealed. miR-26 triple knockout (TKO) mice exhibit excessive fat production due to precocious differentiation of APCs. Furthermore, we provide evidence that regulation of APC differentiation by miR-26 is medi-

ated, at least in part, by repression of *Fbxl19*, a novel miR-26 target that encodes a component of SCF family E3 ubiquitin ligase complexes. Together, these results reveal a new pathway that regulates adipogenesis and uncover potential therapeutic targets for obesity.

Results

Global loss of miR-26 results in excess body fat and dyslipidemia

To investigate the role of miR-26 in mammalian physiology in vivo, CRISPR/Cas9-mediated genome editing was used to generate mice lacking all miR-26 family members. In both mouse and human genomes, all three miR-26 family members reside in the introns of genes coding for Carboxy-terminal domain RNA polymerase II polypeptide A small phosphatase (CTDSP) proteins (Supplemental Fig. S1A), an organization conserved to zebrafish (Dill et al. 2012). Due to the nearly identical sequences of miR-26a and miR-26b, as well as their broad expression in mouse tissues (Fig. 1A; Supplemental Fig. S1B,C), these miRNAs most likely function redundantly. We therefore designed single guide RNAs (sgRNAs) targeting each miRNA, taking advantage of a protospacer adjacent motif (PAM) near the seed sequence (Fig. 1A). Despite a high degree of sequence similarity, the individual sgRNAs did not exhibit detectable cross-targeting of the other miR-26 loci when introduced into embryonic stem (ES) cells (Supplemental Fig. S2A).

miR-26-targeting sgRNAs, together with Cas9 mRNA, were co-injected into fertilized C57BL/6 oocytes, resulting in the generation of insertion/deletion mutations at all three genomic loci (miR-26a-1, miR-26a-2, and miR-26b). Alleles with small deletions that removed the 5' end of each miRNA, including the seed sequences (Fig. 1A), were propagated and combined by standard breeding to yield the total knockout, hereafter referred to as miR-26 triple-knockout (TKO) mice. Initial analyses revealed that miR-26-TKO mice were born at the expected Mendelian ratio, were fertile, and displayed no overt phenotype at weaning (Supplemental Fig. S2B,C; data not shown). Quantitative RT-PCR (qRT-PCR) confirmed the loss of miR-26a and miR-26b expression in multiple tissues (Supplemental Fig. S2D,E). Additionally, the deletion mutations did not affect expression of the miR-26 host transcripts (Supplemental Fig. S2F–H).

A careful examination of young adult mice revealed that while most tissues appeared indistinguishable between wild-type and miR-26-TKO mice, both male and female knockout animals exhibited an approximately twofold to threefold increase in white adipose tissue (WAT) (Fig. 1B,C; Supplemental Fig. S3A). An intermediate phenotype was observed in mice lacking five out of six miR-26 alleles (hereafter referred to as 5/6-KO mice, which harbor only a single intact allele of miR-26a-2). Histologic analysis of WAT demonstrated an absence of adipocyte hypertrophy in TKO mice (Fig. 1D, top panels), suggesting that the expansion of fat depots was a consequence of excessive production of adipocytes.

Figure 1. miR-26-TKO mice have excess body fat and dyslipidemia. (A) Sequences of mouse miR-26a-1, miR-26a-2, and miR-26b genomic loci and deletion alleles generated using CRISPR/Cas9. Protospacer adjacent motif (boxed rectangle), miRNA seed region (red) and sgRNA sequence used for targeting (Cas9 targeting site) are shown. (B) Relative weights of isolated fat depots and major organs from 3-mo-old male mice of the indicated genotypes fed standard chow, normalized to wild type ($n = 6$ mice per genotype). 5/6-KO denotes mice with five of the six miR-26 alleles knocked out with only one wild-type copy of miR-26a-2 intact. (TKO) Triple knockout mice. Fat depots: (IGW) Iguinal; (ISCW) interscapular; (PGW) perigonadal; (RPW) retroperitoneal; (MWAT) mesenteric; (BAT) brown adipose tissue. (C) Representative whole-mount images of subcutaneous and visceral white fat depots of mice from B. (D) Representative H&E stained sections of IGW (top panels) and liver (bottom panels) from mice of the indicated genotypes. Scale bars, 100 μ M. (E) Relative weights of isolated fat depots and major organs from postnatal day 10 male mice ($n = 4$ –6 mice per genotype). No significant differences were noted at this time point. (F) Fat content of male mice at indicated ages measured by whole body NMR ($n = 6$ mice per genotype, per timepoint). (G,H) Serum triglycerides (G) and cholesterol (H) in 3-mo-old male mice ($n = 6$ mice per genotype). For panels B and E–H, data are represented as mean $\pm$ SD. (*) $P < 0.05$; (**) $P < 0.01$; (***) $P < 0.001$, calculated using two-tailed t -test.

Adipocytes are generated from progenitor populations through distinct developmental and adult phases of adipogenesis. While developmental progenitors are primarily responsible for producing a defined number of adipocytes during embryogenesis, adult adipocyte progenitors respond to a variety of environmental and dietary cues such as high-fat diet to produce new adipocytes when increased energy storage is required (Jiang et al. 2014; Wang et al. 2015). In order to determine which phase of adipogenesis is responsible for the increase in WAT in miR-26-TKOs, we first analyzed mice at postnatal day 10 (P10), a time-point at which WAT is predominantly

derived from developmental progenitors. No difference in WAT mass between wild-type and TKO mice was detectable at this age (Fig. 1E; Supplemental Fig. S3B). In contrast, young adult TKO mice, starting as early as 5 wk of age, exhibited a clear and progressive increase in whole body fat content (Fig. 1F; Supplemental Fig. S3C), suggesting excessive activation of the adult adipogenesis program despite a diet of normal mouse chow. Moreover, serum analysis indicated that TKO mice were dyslipidemic, with elevated levels of serum triglycerides (Fig. 1G; Supplemental Fig. S3D) and cholesterol (Fig. 1H; Supplemental Fig. S3E). Despite producing excess WAT, TKO

mice did not exhibit liver steatosis (Fig. 1D, bottom panels), glucose intolerance (Supplemental Fig. S3F–I), or an increase in overall body weight (Supplemental Fig. S3J, K). Furthermore, metabolic cage studies demonstrated that wild-type and TKO mice had similar food intake, physical activity, and energy consumption, as indicated by oxygen consumption and carbon dioxide production (Supplemental Fig. S4A–D). Thus, these data support a previously unrecognized role for miR-26 as a physiologic suppressor of adult adipogenesis.

miR-26a overexpression protects mice from diet-induced obesity

Given the significant expansion of WAT in miR-26-TKO mice, we hypothesized that this miRNA family may function as a rheostat to control fat production in adult animals. To further test this model, we took advantage of a previously described transgenic mouse line with doxycycline (dox)-inducible miR-26a expression (Fig. 2A, *M2rtTA*; *eGFP.miR-26a* mice) (Zeitels et al. 2014) to determine whether enforced expression of this miRNA

could suppress WAT expansion in adult mice. High fat diet (HFD) is a potent stimulant of APC differentiation and resultant de novo adipocyte production (Joe et al. 2009; Wang et al. 2013b; Jeffery et al. 2015; Vishvanath et al. 2016). As expected, control mice (*M2rtTA* only) fed a HFD for 12 wk exhibited significant weight gain and an overall total body fat content close to 40% (Fig. 2B,C). In contrast, mice with global overexpression of miR-26a were strongly resistant to weight gain and HFD-induced WAT expansion (Fig. 2B–E). Notably, despite a substantial reduction in total fat mass, hepatic lipid storage was not altered in HFD-fed miR-26-overexpressing mice (Fig. 2D; data not shown). We also note that an apparent decrease in pancreas weight in miR-26a overexpressing animals (Fig. 2D) was most likely due to a reduction in contaminating visceral adipose tissue in dissected pancreata from these mice. In addition to lower overall body fat, HFD-fed miR-26a transgenic mice exhibited lower serum glucose and lipid levels compared with controls (Fig. 2F–I). Thus, enforced miR-26a expression protected against the adipogenic and metabolic effects of HFD.

Figure 2. Global overexpression of miR-26a protects mice from HFD-induced obesity. (A) Schematic of the dox-inducible miR-26a transgene (*eGFP.miR-26a*) (Zeitels et al. 2014). TRE, Tetracycline-response element; pA, poly(A) signal. (B) Body weight of double transgenic (*M2rtTA*; *eGFP.miR-26a*, abbreviated *M2rtTA-26a*) and control (*M2rtTA* only, abbreviated *M2rtTA*) male mice ($n = 6-8$ per genotype) on dox and HFD for 12 wk. *M2rtTA-26a* and *M2rtTA* male transgenic mice were switched to dox-containing water (2 mg/mL) at 6 wk of age. Two weeks later, HFD was initiated and body weight was recorded weekly. (C) Whole body fat content of mice in B after 12 wk on HFD analyzed by NMR. (D) Relative weights of isolated fat depots and major organs from mice from B after 12 wk on HFD. (E) Representative images of isolated subcutaneous (IGW) and visceral (PGW and RPW) fat depots from mice of the indicated genotypes from B. (F–I) Blood glucose (F), serum triglycerides (G), cholesterol (H), and nonesterified free fatty acids (I) in mice from B after 12 wk on HFD. Data presented in B–I are part of the same experiment and representative of three independent experiments. Data represented as mean $\pm$ SD. (*) $P < 0.05$; (**) $P < 0.01$; (***) $P < 0.001$, calculated using two-tailed t -test.

To test whether miR-26a overexpression could also protect mice from weight gain on normal chow, control and transgenic mice were analyzed after 1 yr of dox administration. Similar to the results obtained after feeding a HFD, miR-26a transgenic mice on normal chow gained less weight, exhibited less total body fat, and displayed dramatically smaller fat depots than control animals (Supplemental Fig. S5A–E). Together with the loss-of-function studies, these results demonstrate that miR-26 levels dictate the extent of fat production in adult mice.

miR-26 regulates adipogenic differentiation in vitro

These initial studies used animal models with global miR-26 loss- and gain-of-function. In order to investigate whether this miRNA family functions intrinsically within the adipocyte lineage to control adipogenesis, multiple experimental strategies were pursued. First, we confirmed that miR-26 is expressed within APCs. The stromal vascular fraction (SVF) of WAT, obtained by enzymatic disso-

ciation of adipose depots that have been depleted of mature adipocytes, constitutes a heterogeneous mixture of mural cells, endothelial cells, immune cells, and APCs. Fluorescence-activated cell sorting (FACS) was used to isolate APCs from the SVF of *Adipotrak* mice (*Pparg-tTA; TRE-H2B-GFP*), in which *Pparg*-expressing APCs are labeled with GFP (Tang et al. 2008). miR-26a and miR-26b expression was readily detectable in APCs and modestly enriched relative to the GFP-negative SVF population (Supplemental Fig. S6A,B).

We next examined the consequences of modulation of miR-26 activity on adipogenesis using an in vitro SVF differentiation assay (Fig. 3A). To avoid systemic effects of miR-26 loss-of-function from influencing APC numbers or behavior prior to in vitro culture, SVF cultures were established from wild-type mice and miR-26 was inhibited after SVF isolation using locked nucleic acid (LNA) antisense oligonucleotides (Fig. 3B). Acute miR-26 inhibition resulted in a robust induction of adipogenic gene expression (Fig. 3C) and adipocyte differentiation (Fig. 3D) in

Figure 3. miR-26 regulates adipogenesis in vitro. (A) Schematic of isolation and in vitro differentiation of stromal vascular fraction (SVF) from pooled fat depots of adult mice. (B) qRT-PCR analysis of mature miR-26a, miR-26b, and miR-19b (an unrelated control miRNA) in wild-type SVF cultures transfected with miR-26 LNA antisense inhibitors (26-LNA) or scrambled control LNA oligonucleotides (Sc-LNA). $n = 3$ biological replicates per condition. (C) qRT-PCR analysis of adipogenic gene expression following in vitro differentiation of miR-26 LNA-transfected SVF cultures from B, relative to scrambled control transfected SVFs. (D) Representative images of Oil Red O-stained SVF cultures from B. (E) qRT-PCR analysis of miR-26a or miR-19b levels in SVFs derived from *M2rtTA; eGFP.miR-26a* transgenic mice, cultured with or without dox (1 µg/mL) for 4 d prior to induction of differentiation. $n = 3$ biological replicates per condition. (F) qRT-PCR analysis of adipogenic gene expression following in vitro differentiation of dox-treated SVF cultures from E, relative to untreated cultures. (G) Representative images of Oil Red O-stained SVF cultures from E. Data represented as mean $\pm$ SD.

this system. Increased adipogenesis was also observed in SVF cultures derived from young miR-26-TKO mice relative to cultures derived from age- and sex-matched wild-type animals (Supplemental Fig. S6C–E). Conversely, doxycycline-induced overexpression of miR-26a in SVF cultures from *M2rtTA; eGFP.miR-26a* mice (Fig. 2A) strongly inhibited adipogenesis (Fig. 3E–G), while doxycycline treatment of SVFs obtained from *M2rtTA* control mice had no antiadipogenic effect (Supplemental Fig. S6F–H). Together, these data support a direct role for miR-26 as a suppressor of adipogenesis via its activity within the adipocyte lineage.

miR-26a expression in adult SMA⁺ APCs confers resistance to HFD-induced obesity

To further assess whether miR-26 expression within adipocyte progenitors is sufficient to suppress adult adipogenesis in vivo, we utilized a miR-26a transgene that is activated by both dox treatment and Cre expression (Fig. 4A, *M2rtTA; LSL.eGFP.miR-26a* mice) (Zeitels et al. 2014). Adult APCs reside in the perivascular niche of fat depots and resemble pericytes or mural cells (Rodeheffer et al. 2008; Tang et al. 2008; Gupta et al. 2012; Berry and Rodeheffer 2013). A smooth muscle actin (SMA)-

Figure 4. Overexpression of miR-26a in adult APCs prevents HFD-induced obesity. (A) Schematic of the dox- and Cre-inducible miR-26a transgene (*LSL.eGFP.miR-26a*) crossed to tamoxifen-inducible *SMA-Cre^{ERT2/+}* for overexpression of miR-26a in adult APCs. (B) Experimental time course. Tamoxifen (TM) was administered to 6-wk-old *M2rtTA; LSL.eGFP.miR-26a; SMA-Cre^{ERT2/+}* (*M2/26a/Cre*) and control *M2rtTA; SMA-Cre^{ERT2/+}* (*M2/Cre*) mice, followed by administration of dox to activate miR-26 expression in the APC lineage. Mice were then fed a HFD for 12 wk. (C) Body weight of control (*M2/Cre*) and transgenic (*M2/Cre/26a*) mice on HFD ($n = 6$ –8 mice per genotype) recorded weekly. (D) Whole body fat content of mice from C measured by NMR after 12 wk on HFD. (E) Relative weights of isolated fat depots and major organs of mice from C after 12 wk on HFD. (F) Representative whole-mount images of subcutaneous IGW and visceral PGW and RPW fat depots from mice from C after 12 wk on HFD. (G–J) Blood glucose (G), serum cholesterol (H), triglycerides (I), and non-esterified fatty acids (J) from mice from C after 12 wk on HFD. Data represented as mean $\pm$ SD. (*) $P < 0.05$; (**) $P < 0.01$; (***) $P < 0.001$ calculated using two-tailed t -test.

Cre^{ERT2/+} allele, which is expressed in adult APCs but not mature adipocytes (Wendling et al. 2009; Jiang et al. 2014), was used to activate the miR-26 transgene within this lineage. Transgenic mice (*M2rtTA*; *LSL.eGFP.miR-26a*; *SMA-Cre^{ERT2/+}*) were administered tamoxifen to activate Cre-recombinase, then treated with dox to turn on miR-26a expression in APCs (Fig. 4B). Mice were subsequently fed a HFD for 12 wk. Overexpression of miR-26a in the SMA⁺ APC lineage conferred strong protection against HFD-induced weight gain (Fig. 4C). Relative to control mice, miR-26a overexpressing mice had lower total body fat, smaller fat depots, and reduced serum glucose and lipids (Fig. 4D–J). These data further support a cell-autonomous role for miR-26 in suppressing adult adipogenesis, although it remains possible that additional effects of miR-26 overexpression on food intake, nutrient absorption, or energy expenditure also contribute to protection from weight gain.

Precocious differentiation of APCs in miR-26-TKO mice

To definitively assess whether loss of miR-26 triggers adipogenic differentiation in vivo, we performed a lineage tracing experiment to monitor the differentiation of APCs to mature adipocytes. To indelibly label adult APCs and their descendants, we utilized *SMA-Cre^{ERT2/+}* in combination with a reporter allele that produces tdTomato when coexpressed with Cre (*R26R^{tdT/+}*) (Fig. 5A; Wendling et al. 2009). Cre recombinase was activated at P25 by tamoxifen administration and the fate of labeled APCs was followed over time to determine the extent to which they gave rise to mature adipocytes (Fig. 5B). As expected, analysis of adipose depots 2 d after Cre activation resulted in robust labeling of the vasculature, most likely representing smooth muscle and perivascular cells including APCs (Fig. 5C, top panels). Importantly, at this time point, no reporter expression was observed in mature adipocytes. Similarly, very few labeled adipocytes were present >3-wk later at P50 in wild-type mice (Fig. 5C, bottom panels; Supplemental Fig. S7A). In contrast, extensive labeling of mature adipocytes was apparent at P50 in miR-26-TKO mice. To determine whether this increase in APC-derived adipocytes could be the result of an expansion of the APC population in miR-26-TKO mice, we quantified the numbers of tdT⁺; Sca-1⁺ cells (Sca-1 distinguishes APCs from smooth muscle) in fat depots from tamoxifen-treated animals using flow cytometry. These experiments confirmed that loss of miR-26 does not result in an overt increase in the overall number of APCs (Supplemental Fig. S7B). Together, these data strongly support the conclusion that miR-26 is necessary to suppress precocious differentiation of APCs into mature adipocytes.

To determine the physiologic consequences of precocious APC differentiation in miR-26-deficient animals, we next examined cohorts of wild-type and miR-26-TKO mice fed a HFD. Young (6-wk-old) wild-type and miR-26-TKO mice exhibited comparable weight gain over a 12-wk period on a HFD (Fig. 5D,E). Surprisingly, however, older miR-26-TKO mice gained significantly less weight and accumulated significantly less body fat than age-

matched controls (Fig. 5F–I). These data suggest that excessive differentiation of APCs in young miR-26-deficient mice results in a depletion of the APC pool, ultimately resulting in defective production of new adipocytes in response to caloric excess later in life.

Fbxl19 is a novel miR-26 target that promotes adipogenesis

To investigate the mechanisms through which miR-26 suppresses adult adipogenesis, we performed RNA-seq on primary SVF cultures from wild-type or miR-26-TKO mice, or *M2rtTA*; *eGFP.miR-26a* mice with or without dox treatment (Fig. 6A). From these data, we sought genes that (1) were up-regulated at least 1.5-fold in miR-26-TKO cells, (2) were down-regulated at least 1.5-fold in dox-treated *M2rtTA*; *eGFP.miR-26a* cells, and (3) had multiple conserved predicted miR-26 binding sites according to Targetscan7.2 (Agarwal et al. 2015). Two genes, which to our knowledge have not previously been implicated in the regulation of adipogenesis, fulfilled these criteria: *F-box and leucine rich repeat protein 19* (*Fbxl19*) and *STE20 related adaptor β* (*Stradb*). FBXL19 is a component of Skp1–Cullin–F-box E3 ubiquitin ligase complexes and has been shown to trigger proteasome-mediated degradation of protein targets involved in inflammation, apoptosis, and cell migration (Zhao et al. 2012, 2013; Dong et al. 2014). More recently, a role for FBXL19 in the epigenetic activation of lineage-specific genes during embryonic stem cell differentiation was uncovered (Lee et al. 2017; Dimitrova et al. 2018). STRADB is a pseudokinase that regulates activity of the LKB1 kinase (Boudeau et al. 2003). Up-regulation of *Fbxl19* and *Stradb* in miR-26-TKO SVF cultures or down-regulation in dox-treated *M2rtTA*; *eGFP.miR-26a* cultures, was confirmed by qRT-PCR (Fig. 6B,C; Supplemental Fig. S7C,D). An increase in FBXL19 protein in miR-26-TKO cultures was further documented by Western blotting (Fig. 6D).

To directly assess the role of *Fbxl19* and *Stradb* in adipogenesis, siRNA was used to knockdown these genes in primary SVF cultures (Supplemental Fig. S7E). As expected, knockdown of *Pparg2*, an essential regulator of adipogenesis, strongly inhibited adipogenic differentiation (Supplemental Fig. S7F). While *Stradb* knockdown had no effect, reduced *Fbxl19* expression dramatically impaired adipogenesis in wild-type SVFs (Fig. 6E; Supplemental Fig. S7F). Accordingly, adipogenic gene expression was strongly reduced in *Fbxl19*, but not *Stradb*, knockdown cultures (Fig. 6F; Supplemental Fig. S7G). Moreover, knockdown of *Fbxl19* in miR-26-TKO cultures restored adipogenesis to wild-type levels (Fig. 6E,F). Conversely, overexpression of FBXL19 robustly stimulated adipogenesis to an extent that was comparable to the effect of PPARγ2 overexpression (Fig. 6G,H) and strongly induced adipogenic gene expression (Fig. 6I).

Finally, we tested the ability of miR-26 to directly regulate *Fbxl19* expression. The *Fbxl19* 3' UTR contains two closely spaced highly conserved 8mer binding sites for miR-26 (Fig. 6J; Agarwal et al. 2015). Placement of these sites within the 3' UTR of a luciferase reporter transcript

Figure 5. Loss of miR-26 results in precocious differentiation and depletion of APCs. (A) Schematic of transgenes used for lineage tracing of adult APCs in wild-type and miR-26-TKO mice. (B) Experimental time course. Adult wild-type and miR-26-TKO mice were pulsed with tamoxifen (TM) at postnatal day 25 (P25) to label the APCs with tdTomato using SMA-Cre^{ERT2/+} and R26R^{tdT/+}. One-half of the animals from both genotypes were sacrificed on P27 to ensure efficient labeling. Reporter expression in the remaining mice was analyzed at P50 to assess adipocyte labeling. (C) Representative stained sections of IGW fat depots from mice of the indicated genotypes, induced with tamoxifen at P25 and analyzed at P27 (Pulse, top half) or P50 (Chase, bottom half). R26R^{tdT/+} expression was detected by antibody staining for tdTomato (red) on FFPE sections. Nuclei are marked with DAPI (blue) and Perilipin (Plin) marks mature adipocytes (green). These data are representative of two independent experimental cohorts of animals. (D) Body weight of mice of indicated genotypes fed HFD starting at 6 wk of age ($n = 10$ – 12 mice per genotype) recorded weekly. (E) Whole-body fat content of mice in D measured by NMR after 12 wk on HFD. (F) Body weight of mice of indicated genotypes fed HFD starting at 14 wk of age ($n = 10$ – 12 per genotype) recorded weekly. (G) Whole-body fat content of mice in F measured by NMR after 12 wk on HFD. (H) Representative whole-mount images of subcutaneous (IGW) and visceral (PGW and RPW) fat depots from mice in F after 12 wk on HFD. (I) Relative weights of isolated fat depots and major organs of mice in F after 12 wk on HFD. Data represented as mean $\pm$ SD. (*) $P < 0.05$; (**) $P < 0.01$, calculated using two-tailed t -test.

Figure 6. *Fbx19* is a conserved miR-26 target that promotes adipogenesis. (A) Experimental design to identify candidate miR-26 targets that regulate adipogenesis. RNA-seq was used to identify (1) genes up-regulated in miR-26-TKO vs. wild-type SVF cultures and (2) genes down-regulated in dox-treated versus untreated SVF cultures from miR-26a transgenic mice (*M2rtTA; miR-26a*). Genes were further filtered for Targetscan7.2-predicted miR-26 targets (<http://www.targetscan.org>). (B,C) qRT-PCR analysis of *Fbx19* expression in wild-type and miR-26-TKO SVFs (B) and miR-26a transgenic SVFs with or without dox treatment (C). *n* = 3 biological replicates. (D) Western blot of FBXL19 protein levels in wild-type and miR-26-TKO SVFs. Data are representative of three independent experiments. α-tubulin (TUB) represents a loading control. (E) Representative images of Oil Red O-stained in vitro differentiated wild-type or miR-26-TKO SVF cultures following siRNA knockdown of *Fbx19*. siNT, nontargeting control siRNA. (F) qRT-PCR analysis of adipogenic gene expression in SVF cultures in **E**. (G) Western blot of Flag-tagged mouse FBXL19 or PPARγ2 in transfected wild-type SVF cultures. (H) Representative images of Oil Red O-stained SVF cultures after overexpression of the indicated proteins. (I) qRT-PCR analysis of adipogenic gene expression in SVF cultures with GFP or FBXL19 overexpression. (J) Schematic representation of conserved miR-26 binding sites in the 3' UTR of *Fbx19*. Mutations introduced to disrupt miR-26-binding are shown below the alignment and highlighted in red. (K) Relative firefly luciferase activity of reporter constructs containing wild-type or mutant miR-26-binding sites from the *Fbx19* 3' UTR following cotransfection with control or miR-26a mimics. Data represent the average of three independent experiments, each performed with technical triplicates. Data represented as mean ± SD (*). ** P < 0.01, calculated using two-tailed t-test.

was sufficient to confer miR-26-mediated repression, which was abolished by mutations that disrupted interaction with the miR-26 seed sequence (Fig. 6K). Together, these data validate *Fbxl19* as a direct miR-26 target and support its role as a regulator of adipogenesis that functions downstream from the miR-26 family within the APC lineage.

Discussion

Obesity, or excess body fat, is a global health problem that represents a major risk factor for several chronic diseases, including type 2 diabetes, nonalcoholic fatty liver disease, cardiovascular disease, stroke, and cancer. The molecular mechanisms that regulate adipose tissue homeostasis are key to understanding the pathogenesis of obesity and metabolic dysfunction. Nevertheless, the genes and pathways that influence adipocyte size and number in vivo remain incompletely understood. Here we describe a new pathway that plays a critical role in the control of adipogenesis in vitro and in vivo. Through the characterization of mice lacking the highly conserved miR-26 miRNA family, we unexpectedly discovered that loss of these miRNAs results in the dramatic expansion of adipose tissue early in adult life. Several lines of evidence, including in vitro differentiation assays, lineage-specific miRNA overexpression experiments, and fate mapping of adult APCs in knockout animals, established that this phenotype is due to a cell-autonomous function of miR-26 as a suppressor of adipogenic differentiation. Furthermore, *Fbxl19*, which encodes a component of Skp–Cullin–F-box-containing (SCF) E3 ubiquitin ligase complexes and epigenetic regulator, was validated as a novel miR-26 target that functions downstream from this miRNA family to drive adipogenesis. These studies therefore establish a previously uncharacterized miR-26-*Fbxl19* axis as a major regulator of APC mobilization and adipose tissue expansion in mammals (Fig. 7).

Given their ability to proliferate and produce new adipocytes, a population of perivascular APCs are now appreciated to function as crucial regulators of WAT maintenance and expansion (Berry et al. 2016; Hepler et al. 2017). Genetic lineage tracing methodologies have been instrumental in establishing the importance of this cell population for adult adipose tissue homeostasis. Fate mapping studies have provided direct evidence that

a subpopulation of mural cells characterized by distinct marker gene expression (SMA^+ ; $PDGFR\beta^+$; $PDGFR\alpha^+$; $CD34^+$; $Sca1^+$; $Zfp423^+$; $Ppar\gamma2^+$; $CD31^-$; $CD45^-$) function as adipocyte precursors that contribute to the maintenance and high-fat diet-induced expansion of adipose depots in adult mice (Gupta et al. 2012; Jiang et al. 2014; Vishvanath et al. 2016). Our studies of the miR-26 family have revealed a new pathway that robustly regulates the behavior of these cells in vivo. Loss of miR-26 triggers precocious APC differentiation, resulting in adipocyte hyperplasia and WAT expansion, in a manner that resembles the physiologic response to caloric excess. Over time, excessive APC differentiation leads to a depletion of this progenitor compartment, such that older miR-26-TKO mice exhibit impaired weight gain and adipogenesis when fed a HFD. On the other hand, enforced miR-26 expression strongly suppresses activation of APCs, robustly suppressing WAT expansion. These findings have therefore uncovered a previously unanticipated miRNA-mediated checkpoint that plays a critical role in controlling APC mobilization.

The discovery of a role for miR-26 as a regulator of adult adipogenesis expands our understanding of the functions of this miRNA family in mammalian metabolic homeostasis. Previously, miR-26 was found to be down-regulated in the livers of obese humans and mice (Fu et al. 2015; Kim et al. 2019). Hepatic overexpression of miR-26a was demonstrated to improve, whereas miR-26 inhibition was shown to decrease insulin sensitivity (Fu et al. 2015). Our studies have revealed the adipose tissue, and the APC population specifically, as another major site of miR-26 action in the regulation of systemic metabolism. Enforced miR-26 expression in the APC lineage was sufficient to block HFD-associated weight gain, adipose tissue expansion, hyperglycemia, and hyperlipidemia. Interestingly, it has recently been suggested that exosome-mediated miRNA transfer from adipose tissue-derived immune cell populations to other tissues, such as liver, muscle, and pancreas, modulates insulin sensitivity (Ying et al. 2017; Guay et al. 2019). In light of these findings, and the potent effects of miR-26 expression on APC differentiation, future studies are warranted to examine whether transfer of miR-26 from immune cells to APCs provides a mechanism to regulate the APC population and, consequently, fat production.

In addition to revealing a critical role for miR-26 in controlling adult adipogenesis, these studies have also uncovered a new role for FBXL19 as a major regulator of this process. FBXL19 is a component of SCF E3 ubiquitin ligase complexes that mediate targeted ubiquitylation of proteins destined for proteasomal degradation (Jin et al. 2004). Of the >70 F-box proteins that have been identified in humans, the most well-studied facilitate degradation of cell cycle regulators (Nakayama and Nakayama 2006). Previous work has identified a limited number of ubiquitylated substrates of FBXL19, including members of the Ras and Rho family of small GTPases and a receptor for IL-33, implicating the protein in diverse cellular functions including proliferation, migration, apoptosis, and immune responses (Zhao et al. 2012, 2013; Dong et al.

Figure 7. miR-26 suppresses adipocyte progenitor cell differentiation. miR-26 functions within the APC lineage to inhibit mobilization and subsequent adipocyte production by down-regulating FBXL19, a novel driver of adipogenesis.

2014). More recent work has highlighted a nuclear role for FBXL19 as a CpG island-binding protein via its N-terminal CxxC-Zn finger domain (Lee et al. 2017; Dimitrova et al. 2018). Upon binding CpG islands upstream of genes involved in lineage specification, FBXL19 activates gene expression by promoting RNF20-mediated histone H2B mono-ubiquitylation and/or by directly recruiting the mediator complex, thereby promoting embryonic stem cell differentiation. In principle, both the targeted proteasomal degradation of FBXL19 ubiquitylation substrates as well as the activation of lineage-specific gene expression could contribute to the adipogenic properties of this protein. Elucidation of the downstream pathways controlled by FBXL19 in adipocyte progenitors therefore represents an important priority for future work and is likely to reveal new mechanisms that impact adipose tissue homeostasis.

In sum, the data reported here highlight the potentially beneficial effects of increasing miR-26 activity, or inhibiting its downstream targets, in the setting of obesity. This concept is further supported by the observation that sustained global overexpression of miR-26 does not result in any overt toxicities in mice (Zeitels et al. 2014). Further investigation of the miR-26-*Fbx19* pathway as well as the identification of additional miR-26 targets that mediate the beneficial metabolic activities of these miRNAs promises to reveal new insights into the biology of obesity and potentially new therapeutic strategies.

Materials and methods

Mouse strains

miR-26-TKO mice were generated and maintained on a pure C57Bl/6 background. Mice were housed in a 12-h light-dark cycle with normal chow (Harlan Teklad, TD.2916) provided ad libitum. Wild-type C57Bl/6 mice (Jackson Laboratory, 000664) were obtained from the mouse-breeding core at the University of Texas Southwestern Medical Center and maintained in the same colony. *SMA-Cre^{ERT2}/+* mice were generously provided by Dr. Pierre Chambon (IGCMB, Strasbourg, France). Activation of *Cre* recombinase in these mice was achieved by tamoxifen administration (0.1 mg/g body weight) by oral gavage. Generation of *LSL:eGFP-miR-26a* transgenic mice was described previously (Zeitels et al. 2014). To induce miR-26a expression, dox (2 mg/mL in 1% sucrose) was provided in drinking water starting at 6 wk of age. Unless otherwise specified, for HFD studies, male mice were fed a high-fat high-sucrose diet (Research Diets, D12331) for a total duration of 12 wk, starting at 6–8 wk of age. All animal studies were approved by the University of Texas Southwestern Medical Center Animal Care and Use Committee in accordance with NIH guidelines (Animal protocols 2017-102001 and 2015-101187).

T7 endonuclease assay

Plasmid pX330 expressing Cas9 and sgRNA (Cong et al. 2013) was linearized with *Bbs1*. A pair of oligonucleotides (sequences provided in Supplemental Table S1) corresponding to each of the individual miR-26 family members was annealed and ligated to the linearized vector. E14 mouse embryonic stem (mES) cells were maintained in Dulbecco's knockout media supplemented with 15% fetal bovine serum (Sigma), 2 mM GlutaMAX (Life Technol-

ogies), nonessential amino acids (1×), Sodium pyruvate (1 mM), LIF (Millipore), β-ME (Sigma), 100 U/mL penicillin, and 100 μg/mL streptomycin. Cells were transfected with plasmid DNA using X-Fect mESC Transfection Reagent (Clontech, PT5004-2). After 48 h, genomic DNA was extracted using DNeasy Blood and Tissue kit (Qiagen) following manufacturer's protocol. Genomic DNA surrounding the CRISPR target site for each of the miR-26 genes was PCR amplified (primer sequences in Supplemental Table S1) and the products were purified using QiaQuick spin columns (Qiagen). A total of 200 ng of the purified PCR product was subjected to denaturation followed by gradual reannealing to enable heteroduplex formation (for 10 min at 95°C, 95°C to 85°C ramping at 2°C/sec, 85°C to 25°C ramping at 0.25°C/sec, and hold for 1 min at 25°C). Reannealed PCR products were digested with T7 endonuclease (New England Biolabs), and the digested products were analyzed on a 1% agarose gel.

CRISPR/Cas9-mediated genome editing of miR-26 loci

For editing of miR-26 genomic loci, Cas9 mRNA and sgRNAs were coinjected into the pronucleus of fertilized C57BL/6 one-cell eggs following in vitro transcription (Wang et al. 2013a). Briefly, T7-Cas9 or T7-sgRNA PCR products (primer sequences in Supplemental Table S1) were gel purified and used as the template for in vitro transcription (IVT) using the mMESSAGE mMACHINE T7 ULTRA kit (Life Technologies) or the MEGAshortscript T7 kit (Life Technologies), respectively. Both Cas9 mRNA and sgRNAs were purified using the MEGAclear kit (Life Technologies) and eluted in RNase-free water. Founders with disrupted miRNA seed sequences were identified by TA cloning and Sanger sequencing. Founders with deletions of the seed sequence were crossed with C57BL/6 mice for six generations followed by intercrossing to generate the global miR-26 triple knockout. Since the probability of obtaining a TKO and wild-type in the same cross is very low (1/64 in a triple heterozygote cross), knockouts were compared to age- and sex-matched wild-type C57BL/6 mice bred under the same conditions. Additionally, as a littermate control, 5/6-KO mice (deleted for five of the six miR-26 alleles, bearing only one functional allele of miR-26a-2) were also included in most analyses.

SVF culture and differentiation

All in vitro differentiation assays were performed as previously described (Tang et al. 2008) using primary stromal vascular fraction (SVF) cultures derived from 6- to 8-wk-old adult mice. Briefly, subcutaneous and visceral white fat depots were pooled, minced, and digested by slow shaking in adipocyte isolation buffer (100 mM HEPES at pH 7.4, 120 mM NaCl, 50 mM KCl, 5 mM glucose, 1 mM CaCl₂, 1.5% BSA, 1 mg/mL collagenase type I) for 2 h at 37°C. The suspension was then centrifuged at 800 rpm for 10 min to yield the floating adipocyte layer and the crude stromal vascular (SV) pellet. The pellet was first suspended in erythrocyte lysis buffer, filtered through a 100-μm mesh, and spun at 1300 rpm for 5 min. The resulting pellet was suspended in 1× PBS, passed through a 40-μm mesh, and spun at 1300 rpm for 5 min to obtain the SVF. Cells from the SVF were cultured in DMEM supplemented with 10% FBS. Dox was added to induce miR-26a overexpression in SVF cultures derived from transgenic mice (*M2rtTA; eGFP-miR-26a*) at a concentration of 1 μg/mL. LNA inhibitors for miR-26a and miR-26b (Exiqon, Inc.) and siRNA SMARTpools (Dharmacon) targeting mouse *Fbx19*, *StradB*, and *Pparγ2* were reverse-transfected into primary SVFs using RNAiMax (Thermo Fisher Scientific). Knockdown efficiency was monitored 48 h after transfection by qRT-PCR. Overexpression of FBXL19 and

PPAR γ 2 in primary SVF cultures was achieved by nucleofection using the SE cell line 4D-Nucleofector X-kit (Lonza). Flag-tagged ORF clones for FBXL19 (Origene, MR209945) and PPAR γ 2 (Origene, MR208132) were transfected into wild-type SVFs ($\sim 3 \times 10^5$ cells) using the CM137 program. Differentiation was induced 2 d after the cells had reached confluency using an adipogenic cocktail containing 5 μ M dexamethasone, 0.5 μ g/mL insulin, and 0.5 mM isobutylmethylxanthine in culture medium. After 72 h, medium was replaced with DMEM containing 0.5 μ g/mL insulin. Oil Red O staining was performed as previously described (Tang et al. 2008). Briefly, differentiated adipocytes were fixed in 4% paraformaldehyde for 30 min, washed with deionized water, and incubated in 60% isopropanol for 10 min. Adipocytes were then stained in Oil Red O solution for 30 min, rinsed with tap water, and imaged using a Nikon Eclipse TS100 inverted microscope.

Metabolic phenotyping assays

Blood glucose was measured in tail blood using a Bayer Contour blood glucose monitoring system. For serum measurements, blood was collected by cardiac puncture, incubated at room temp for 30 min, and centrifuged at 8000 rpm for 10 min to obtain the serum. Serum triglyceride, cholesterol, and NEFA were measured in adult mice in the UT Southwestern Metabolic Phenotyping Core. GTT was performed as previously described (Zeve et al. 2012). Total body fat mass of mice was measured by NMR spectroscopy using a Minispec mq spectrometer (Bruker).

Metabolic cage studies

Wild-type, 5/6-KO, and miR-26-TKO mice on normal chow were monitored in metabolic chambers. Mice were housed individually and acclimatized to the metabolic chambers for 5 d before data collection was initiated. For the subsequent 5 d, food intake, VO $_2$, VCO $_2$, and physical activity were monitored over a 12-h light/dark cycle with food provided ad libitum.

RNA extraction, quantitative RT-PCR, and RNA-seq

Total RNA was isolated using the miRNeasy Mini Kit (Qiagen) according to the manufacturer's instructions. Pre-designed TaqMan primers and probes (Life Technologies) were used to measure the levels of mature miR-26a, miR-26b, miR-19b, and U6 snRNA according to the manufacturer's protocol. All other mRNAs were reverse transcribed using PrimeScript reverse transcription master mix (Takara) and transcript abundance was assayed using Power SYBR Green master mix (Life Technologies). All qRT-PCR values were normalized using 18S rRNA expression as a loading control. Primer sequences are provided in Supplemental Table S1.

RNA-seq experiments were performed on triplicate samples from SVFs from (1) wild-type C57BL/6 and miR-26-TKO mice, and (2) *M2rtTA*; *eGFP.miR-26a* mice with or without 1 μ g/mL dox treatment for 4 d. Library prep and RNA-seq were performed by the UTSW McDermott Center Next-Generation Sequencing Core. Sequencing libraries were prepared with the TruSeq Stranded mRNA Library Prep Kit (Illumina) and sequenced using the 75-bp single-read protocol on a NextSeq 500 (Illumina). Reads were mapped to the mouse reference genome (GRCm38) using HISAT2 (version 2.1.0) (Pertea et al. 2016). Only reads uniquely mapped to the genome were kept for future analysis. Aligned reads were counted per gene ID using feature counts (v.1.6.0) (Liao et al. 2014) based on the annotation of GENCODE M16

(Mudge and Harrow 2015). Differential expression analysis was performed using edgeR (3.24.0) (McCarthy et al. 2012).

Histology, immunohistochemistry, and Western blotting

Hematoxylin and eosin (H&E) staining and immunocytochemistry were performed on paraffin sections as described previously (Tang et al. 2008). Isolated tissues were fixed in 10% formalin, embedded in paraffin, and sectioned with a HM 325 microtome. tdTomato reporter expression in paraffin sections was visualized by immunostaining with a mouse monoclonal antibody against DsRed (Takara, 632392) used at a 1:200 dilution. Adipocytes were identified by immunostaining with anti-Perilipin-1 (Abcam, ab61682) used at 1:1000. For Western blotting, whole-cell extracts of cultured SVFs were immunoblotted with anti-FBXL19 (1:1000; Abcam, ab172961), anti- α -Tubulin (1:2000; Sigma, T6199), and anti-Flag epitope (1:1000; Cell Signaling, 2368).

Flow cytometry

For FACS analysis of tdTomato $^{+}$ cells, freshly isolated SVFs from 6-wk-old control (*SMA-Cre^{ERT2/+}*; *R26R^{tdT/+}*) or knockout (*SMA-Cre^{ERT2/+}*; *R26R^{tdT/+}*; *miR-26-TKO*) mice were stained for stem cell antigen Sca-1 (1:200 antimouse Ly-6A/E antibody, BD biosciences, 560653) for 30 min on ice followed by two washes with staining buffer (PBS with 3% FBS and 2 mM EDTA, pH 8.0). Cells were analyzed with a BD FACS Aria at the UT Southwestern Flow Cytometry Core. Data analysis was performed using BD FACS Diva software.

Luciferase reporter assays

The wild-type reporter construct was created by cloning a 372-bp fragment of the mouse *Fbxl19* 3' UTR encompassing the two conserved miR-26 binding sites into the XbaI site of pGL3-control (Promega). Mutagenesis was performed by amplifying the plasmid using overlapping oligonucleotides that contained mutations in both miR-26-binding sites (oligonucleotide sequences in Supplemental Table S1). HCT116 cells (1×10^5 per well) were plated in a 24-well plate 16 h prior to transfecting 100 ng of the indicated reporter construct, along with 5 ng of phRL-SV40 (Promega) and 25 nM of control or miR-26a mimic (Dharmacon) using Lipofectamine 2000 (Thermo Fisher Scientific). Firefly and renilla luciferase activity were assayed 48 h after transfection using the Dual Luciferase Reporter Assay System 4 (Promega) according to the manufacturer's protocol.

Data access

RNA-seq data have been deposited in GEO under accession code GSE134736.

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