Dnmt3a is an epigenetic mediator of adipose insulin resistance

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Abstract Insulin resistance results from an intricate interaction between genetic make-up and environment, and thus may be orchestrated by epigenetic mechanisms like DNA methylation. Here, we demonstrate that DNA methyltransferase 3a (Dnmt3a) is both necessary and sufficient to mediate insulin resistance in cultured mouse and human adipocytes. Furthermore, adipose-specific Dnmt3a knock-out mice are protected from diet-induced insulin resistance and glucose intolerance without accompanying changes in adiposity. Unbiased gene profiling studies revealed Fgf21 as a key negatively regulated Dnmt3a target gene in adipocytes with concordant changes in DNA methylation at the Fgf21 promoter region. Consistent with this, Fgf21 can rescue Dnmt3a-mediated insulin resistance, and DNA methylation at the Fgf21 locus was elevated in human subjects with diabetes and correlated negatively with expression of FGF21 in human adipose tissue. Taken together, our data demonstrate that adipose Dnmt3a is a novel epigenetic mediator of insulin resistance in vitro and in vivo.

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Introduction Insulin resistance (IR) is generally defined as a reduced ability of insulin to provoke metabolic responses, such as glucose uptake. Several mechanistic drivers of IR have been proposed, including defects in the insulin signal transduction pathway, dysfunctional mitochondrial respiration, and ER stress (Boucher et al., 2014; Gregor and Hotamisligil, 2011; Houlistis et al., 2006; Ozcan et al., 2004; Samuel and Shulman, 2012). The data supporting the involvement of these non-mutually exclusive pathways are convincing, but additional mechanisms are also likely to participate in the development of IR. Nuclear pathways in particular (i.e. transcriptional and epigenomic mechanisms) are likely to play an important role, given that IR often develops over a time course that is inconsistent with rapid signal transduction events (Kang et al., 2015; Tan et al., 2015). Furthermore, there is a large body of data from humans and rodents indicating that the propensity for IR can be passed transgenerationally; such ‘fetal programming’ studies strongly imply an epigenetic basis for IR (Carone et al., 2010; Heijmans et al., 2008; Li et al., 2010; Tobi et al., 2014). Finally, thiazolidinedione drugs, which activate the transcription factor PPARγ, provide one of the only pharmacological therapies for IR (Soccio et al., 2014; Step et al., 2014; Sugii et al., 2009). Taken together, these
DNA methylation is a reversible epigenetic mark involving the covalent transfer of a methyl group to the C-5 position of a cytosine residue, usually in the context of a CpG doublet, by DNA methyltransferases (Dnmts) (Robertson, 2005). Long thought to be a static epigenetic mark, emerging evidence suggests that methylated DNA undergoes dynamic and reversible remodeling in somatic cells during developmental, physiological, and pathogenic pathophysiological processes (Barres et al., 2013; 2009; Benner et al., 2015; Kirchner et al., 2016; Multhaup et al., 2015; Siersbaek et al., 2017). As an example, a bout of intense exercise can induce DNA hypomethylation at the PPARC1a locus in human skeletal muscle within hours, followed by increased gene expression; these effects were reversed after resting (Barrés et al., 2012). Mounting evidence points to a role for DNA methylation in the pathogenesis of metabolic disorders. For example, the obesity-prone agouti mouse has reduced DNA methylation at the regulatory region for the Agouti gene, leading to enhanced expression and subsequent inhibition of the satiety-inducing MC4 receptor (Xie et al., 2013). Multiple studies have shown that changes in DNA methylation at key metabolic genes, such as Cox7a1 (Rönn et al., 2008), IGF-2 (Heijmans et al., 2008), and Pomc (Ehrlich et al., 2010), associate with various metabolic insults including aging, obesity, anorexia, and prenatal exposure to famine. Moreover, recent genome-wide profiling studies have identified distinct global DNA methylation patterns that associate with obesity and diabetes in humans (Dayeh et al., 2014; Feinberg et al., 2010; Kirchner et al., 2016; Multhaup et al., 2015; Nilsson et al., 2015; Volkmar et al., 2012; Volkov et al., 2017). Altered DNA methylation is linked to the transgenerational passage of metabolic disorders (Ng et al., 2010; Wei et al., 2014). Despite numerous studies showing a link between DNA methylation and metabolic dysregulation, the cause-and-effect relationship between the two remains largely unknown.

In mammals, five Dnmt family members have been identified: Dmnt1, Dmnt2 (also known as Trdm1), Dmnt3a, Dmnt3b and Dmnt3l, yet only three (Dmnt1, 3a, and 3b) possess DNA methyltransferase activity (Robertson, 2005). Dmnt1 (the maintenance DNA methyltransferase) has a preference to methylate hemimethylated DNA whereas Dmnt3a and 3b prefer unmethylated DNA as substrate, and thus act as de novo DNA methyltransferases (Robertson, 2005). Dmnt3l is homologous to the other Dnmt3s but lacks catalytic activity (Robertson, 2005) and Dmnt2 has sequence homology to all Dnmts but methylates cytoplasmic tRNA, and not DNA (Robertson, 2005). Administration of a small molecule Dmnt inhibitor has been shown to improve insulin sensitivity in the setting of obesity, in part by demethylating the Adipoq promoter, and a role for Dmnt1 in this process was suggested by genetic knockdown studies in cultured adipocytes (Kim et al., 2015).

Here, we identify Dmnt3a as a key epigenetic determinant of obesity-associated IR in adipose tissue. Dmnt3a plays a causal role in the development of cell autonomous IR in mouse and human adipocytes in a manner that is dependent on its catalytic activity. Consistent with results from adipocytes in vitro, we find that adipose-specific Dmnt3a deficiency in vivo confers protection from diet-induced IR and glucose tolerance without accompanying changes in body weight or adiposity. In addition, unbiased gene expression profiling suggests that Dmnt3a acts as primarily as a gene repressor, and reveals a set of adipocyte-specific target genes, including Fgf21. Further, we demonstrate that Dmnt3a mediates IR, at least in part, by methylating specific cis-regulatory elements in the Fgf21 gene and thus suppressing its expression. We also provide evidence that similar mechanisms might be at play in human adipose tissue. These studies identify Dmnt3a as a novel epigenetic mediator of IR in adipocytes.

**Results**

**Adipose expression of Dnmts inversely correlates with insulin sensitivity**

To identify a potential role for Dnmts in the pathogenesis of insulin resistance, we compared their expression levels in insulin-sensitive metabolic tissues (i.e. liver, muscle, and adipose depots) in various metabolic settings. In a diet-induced obesity (DIO) mouse model, Dmnt1 and Dmnt3a mRNA expression was elevated in adipose tissue, especially in epididymal and mesenteric WAT (Figure 1A, Figure 1—figure supplement 1A). Dmnt3a expression was significantly induced by HFD in skeletal...
muscle, but to a lesser degree than in adipose tissue. No effect of HFD was observed in liver (Figure 1—figure supplement 1B–C). In genetically obese ob/ob mice, mRNA expression of all three functional Dnmts was elevated in adipose tissue (Figure 1B). Importantly, expression of these genes was significantly reduced after treatment with the insulin sensitizer rosiglitazone (Rosi) (Figure 1B). Increased adipose expression of Dnmt3a was confirmed by immunoblotting in DIO mice (Figure 1C, D). Relative expression of Dnmt3a was higher in eWAT compared to skeletal muscle and liver compared in normal chow-fed mice (Figure 1—figure supplement 1D–E). Dnmt2 and Dnmt3l mRNA displayed minimal expression in adipose tissue (not shown). Together, this led us to hypothesize that Dnmts may have a direct functional role in the pathogenesis of adipose IR.
Knock-down of Dnmt3a confers protection from drug-mediated IR

Given the inverse correlation of adipose Dnmt expression with insulin sensitivity, we sought to test the cell autonomous role of Dnmts in the development of IR. To that end, individual Dnmts were knocked-down in mature 3T3-L1 adipocytes using two distinct short hairpin RNAs (shRNAs) (Figure 2—figure supplement 1A–F). To promote IR, cells were treated with dexamethasone (Dex) and TNF-α (TNF) (Houstis et al., 2006; Kang et al., 2015) for an additional four days. Knock-down of individual Dnmts did not have a major impact on the basal level of insulin sensitivity when assessed by insulin-stimulated glucose uptake (Figure 2—figure supplement 2A–E). However, knock-down of Dnmt3a, but not Dnmt1 or Dnmt3b, protected adipocytes from Dex- and TNF-mediated IR as assessed by insulin-stimulated glucose uptake and signal transduction (Figure 2A,B, Figure 2—figure supplement 2A–E). Given the elevated expression of adipose Dnmts in obesity, we were prompted to test whether overexpression of Dnmts per se is sufficient to drive IR in adipocytes. Individual Dnmts were expressed in mature 3T3-L1 adipocytes by lentiviral delivery. Interestingly, over-expression of Dnmt3a and 3b, but not Dnmt1, potently inhibited insulin-stimulated glucose uptake (Figure 2C). Additionally, we tested whether the causal role of Dnmt3a in the development of insulin resistance is conserved in human adipocytes. Overexpression of Dnmt3a powerfully inhibited insulin-stimulated glucose uptake in in vitro differentiated adipocytes from human adipose derived stem cells (Figure 2D). Together, our gain- and loss-of-function studies in vitro show that Dnmt3a is both necessary and sufficient to mediate IR in adipocytes in a cell autonomous manner.

DNA methyltransferase activity is critical for Dnmt3a to mediate IR

Studies have shown that Dnmt3a can act as a gene repressor using both catalytic-dependent and independent mechanisms (Bachman et al., 2001; Fuks et al., 2001). Thus, we sought to determine whether DNA methyltransferase activity is required to mediate IR. We initially pursued a pharmacological approach, using two synthetic Dnmt inhibitors, 5-azacytidine (5-Aza) (Stresemann and Lyko, 2008), a nucleoside analogue serving as a pseudo-substrate for Dnmts, and RG-108 (Brueckner et al., 2005), a non-nucleoside-based drug. Remarkably, both Dnmt inhibitors blocked Dex- and TNF-mediated IR (Figure 3A,B, Figure 3—figure supplement 1A,B). In addition to these synthetic inhibitors, the naturally occurring Dnmt inhibitors (Rajavelu et al., 2011) epigallocatechin gallate (ECCG) and theaflavin-3,3’-digallate (TF-3) also showed insulin-sensitizing activity in the setting of Dex- and TNF-mediated IR (Figure 3—figure supplement 1C–F). These data suggest that DNA methyltransferase activity is necessary for Dnmt3a to cause IR. To further probe this hypothesis, we overexpressed a catalytically inactive allele of Dnmt3a (C706S) (Figure 3C). Together, the pharmacological and genetic evidence indicates that DNA methyltransferase activity is essential for Dnmt3a to mediate IR.

Adipose-specific Dnmt3a deficiency confers protection from diet-induced insulin resistance and glucose intolerance

We next sought to determine whether loss of Dnmt3a in adipocytes could confer protection from diet-induced adipose tissue and systemic IR in vivo. To address this, we generated adipose-specific Dnmt3a knock-out mice (Dnmt3a<sup>AdiKO</sup>) by crossing Dnmt3a floxed mice (Dnmt3a<sup>f/f</sup>) (Nguyen et al., 2007) to adiponectin-Cre mice (Eguchi et al., 2011) (Figure 4—figure supplement 1A,B). On chow diet, male WT and Dnmt3a<sup>AdiKO</sup> animals did not show any obvious change in their body weight, insulin sensitivity, glucose tolerance, or insulin levels (Figure 4—figure supplement 1A–D). On HFD (45% fat), Dnmt3a<sup>AdiKO</sup> mice still showed no significant differences in body weight or composition relative to littermate controls (Figure 4A,B). Moreover, the effect of long-term fasting (16 hr) on body weight or body composition did not differ between genotypes (Figure 4C), suggesting that gross energy homeostasis is not altered in the knock-out animals. Despite no obvious change in energy balance, high fat fed male Dnmt3a<sup>AdiKO</sup> mice exhibited improved glucose tolerance (Figure 4D,E) and insulin sensitivity (Figure 4F,G) with reduced serum insulin in the fed state (Figure 4H). Consistent with their metabolic improvement, male Dnmt3a<sup>AdiKO</sup> mice displayed increased insulin signal transduction in adipose tissue, as demonstrated by increased insulin-stimulated phosphorylation of IRS and AKT (Figure 4I,J). Similarly, female Dnmt3a<sup>AdiKO</sup> mice display improved insulin sensitivity and glucose tolerance on HFD without a change in body weight (Figure 4—figure
Figure 2. Dnmt3a is sufficient and necessary to mediate cell autonomous IR in vitro. (A) Two independent hairpins against Dnmt3a (versus scrambled control shRNA, shScr) were delivered to mature 3T3-L1 adipocytes (Day 8) via lentiviral transduction. Cells were then treated with Dex and TNF for 4 days and assessed for insulin-stimulated glucose uptake ([3H]-2-DG assay, n = 6, p<0.05). Shown is the % of insulin-stimulated glucose uptake rescued by Dnmt3a knockdown (n = 6, p<0.05). (B) Cells in (A) with #2 shDnmt3a hairpin and control cells were withdrawn from serum for 6 hr, stimulated with 20 nM insulin for 5 min and subjected to immunoblotting with antibodies against total and phospho-AKT and –IRS-1 with and without insulin. (C) Basal and insulin-stimulated glucose uptake in mature 3T3-L1 and (D) in in vitro-differentiated primary human adipocytes transduced with lentivirus expressing Dnmts or the empty plasmid vector, pCDH (n = 6, p<0.05, Student’s t-test, mean ±s.e.m.).

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Figure supplement 1. Knock-down efficiency of Dnmt hairpins.
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Figure supplement 1—source data 1. Figure 2—figure supplement 1—source data.
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Figure supplement 2. Dnmt3a is necessary for Dex- and TNF-mediated IR.
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Figure supplement 2—source data 1. Figure 2—figure supplement 2—source data.
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We next assessed whether insulin sensitization could be due to either enhanced browning or reduced inflammation of WAT. With regard to the former, we found no consistent change in the expression of major thermogenic genes such as \textit{Ucp1} and \textit{Ppargc1a} in either iWAT and BAT (Figure 4—figure supplement 4A,B). Additionally, gene expression analysis showed a non-significant trend toward reduced expression of proinflammatory genes such as \textit{Tnfa} and \textit{Ccl2} (which encodes MCP-1) in the eWAT of \textit{Dnmt3a}\textsuperscript{AdiKO} mice (Figure 4—figure supplement 5A). We also saw no obvious change in the density of crown-like structures between WT and KO tissues (Figure 4—figure supplement 5B). Together, these results demonstrate that adipose-specific deletion of \textit{Dnmt3a} confers protection from diet-induced IR in a manner that is independent of adiposity, inflammation, or browning of white adipose tissue.

Unbiased gene profiling studies reveal that \textit{Fgf21} is a key metabolic target gene of \textit{Dnmt3a} in adipocytes

\textit{Dnmt3a} affects developmental processes by regulating non-overlapping sets of cell type-specific target genes (Challen et al., 2011; Nguyen et al., 2007; Nishikawa et al., 2015). To gain mechanistic insight into \textit{Dnmt3a}-mediated IR at the molecular level, we performed RNA-seq in 3T3-L1 adipocytes, in both gain- and loss-of-function contexts. We have successfully used this approach to enhance our ability to identify \textit{bona fide} targets (Kang et al., 2012). We found that 97 genes were repressed after shRNA-mediated \textit{Dnmt3a} knock-down, while only eight genes were up-regulated in the presence of excess \textit{Dnmt3a}; the single gene showing concordant regulation was \textit{Dnmt3a} itself (Figure 5A). Conversely, 290 genes were induced by \textit{Dnmt3a} knock-down, and 36 genes showed diminished expression after \textit{Dnmt3a} overexpression; seven genes were concordantly regulated by both manipulations (Figure 5B and C and Supplementary file 1). These data are consistent with the expectation that \textit{Dnmt3a} works primarily as a repressor of gene expression. Among the concordantly-regulated negative target genes of \textit{Dnmt3a}, we were immediately struck by the presence of \textit{Fgf21}. \textit{FGF21} is known to facilitate glucose uptake in adipocytes (Ge et al., 2011; Lee et al., 2014; Minard et al., 2016), and is a positive target gene of PPAR\(\gamma\) (Muise et al., 2008). Furthermore, mice lacking \textit{FGF21} do not support Rosi-mediated insulin sensitization in vivo (Dutchak et al., 2012). We confirmed by Q-PCR that \textit{Fgf21} expression is indeed reduced in cultured adipocytes.
Figure 4. Adipose-specific Dnmt3a KO mice display improved whole body insulin sensitivity and glucose tolerance on HFD. (A) Body weight of Dnmt3a AdiKO and control mice on HFD (n = 7, 9, p<0.05, Student’s t-test, mean ± s.e.m.). (B) Body composition after 10 weeks of HFD (n = 7, 9). (C) Effect of long-term fasting (16 hr) on body weight or body composition by EchoMRI. Shown is % loss after fasting (n = 14, 14, p<0.05, Student’s t-test, mean ± s.e.m.). (D) Glucose tolerance test after 9 weeks on HFD (n = 7, 9, p<0.05, Student’s t-test, mean ± s.e.m.). (E) The area under curve (AUC) in (D). (F) Insulin tolerance test after 11 weeks on HFD (n = 7, 9, p<0.05, Student’s t-test, mean ± s.e.m.). (G) AUC in (F). (H) Fed and fasted insulin levels measured by ELISA (n = 7, 9 p<0.05, Student’s t-test, mean ± s.e.m.) (I) Immunoblot of total and phospho-AKT and IRS-1 in eWAT after IP insulin. (J) Quantification of western blot in (I).
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Figure supplement 1. Adipose-specific deletion of Dnmt3a.
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Figure supplement 1—source data 1. Figure 4—figure supplement 1—source data.
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Figure supplement 2. Metabolic features of Dnmt3a AdiKO mice on chow.
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Figure supplement 2—source data 1. Figure 4—figure supplement 2—source data.
DOI: https://doi.org/10.7554/eLife.30766.020
Figure supplement 3. Metabolic features of female Dnmt3a AdiKO on HFD.
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Figure supplement 3—source data 1. Figure 4—figure supplement 3—source data.
DOI: https://doi.org/10.7554/eLife.30766.022
Figure supplement 4. Gene expression profile of thermogenic genes in Dnmt3a AdiKO mice on HFD.
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Figure supplement 4—source data 1. Figure 4—figure supplement 4—source data.
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Figure 4 continued on next page
overexpressing Dnmt3a, is increased in cells expressing shDnmt3a, and most importantly, is up-regulated in the adipose tissue of Dnmt3a AdiKO mice (Figure 5D). Unlike adipose tissue levels, serum levels of Fgf21 were reduced in HFD-fed Dnmt3a AdiKO mice compared to controls (Figure 5—figure supplement 5–). To determine whether restoration of FGF21 levels could ameliorate Dnmt3a-mediated IR, we treated 3T3-L1 adipocytes that overexpress Dnmt3a with recombinant FGF21, which restored insulin sensitivity in a dose-dependent manner (Figure 5E).

Dnmt3a-mediated changes in Fgf21 gene expression involve changes in DNA methylation

In general, the level of DNA methylation at gene promoters is inversely correlated with mRNA expression (Jones, 2012). Given the functional significance of FGF21 in Dnmt3a-mediated IR, we further examined whether Dnmt3a directly alters methylation of the 5 kb upstream promoter regions of the Fgf21 gene (Figure 6A). Using methylated DNA precipitation PCR (MeDIP-qPCR), we detected genomic DNA hypermethylation at Fgf21 promoter regions in Dnmt3a overexpressor 3T3-L1 cells compared to controls (Figure 6B). Conversely, DNA hypomethylation of these same regions was seen in Dnmt3a knock-down cells (Figure 6B). In addition to the in vitro models, we conducted MeDIP-qPCR analysis using genomic DNA from fractionated adipocytes of Dnmt3a AdiKO and control animals, finding reduced DNA methylation at the Fgf21 promoter in KO cells (Figure 6C). Consistent with DNA methylation pattern at the Fgf21 promoter region, co-transfection of Dnmt3a potently inhibited promoter activity of Fgf21 (Figure 6D). Together, these data suggest that Dnmt3a-mediated suppression of Fgf21 involves changes in DNA methylation profile at key promoter regions.

DNA methylation of FGF21 is increased in human adipose tissue of subjects with type 2 diabetes

To translate some of our findings to humans, we investigated DNA methylation and gene expression of FGF21 in human adipose tissue collected from diabetic vs. non-diabetic human subjects. In line with our mouse data, we found hypermethylation of four CpG sites annotated to FGF21 in adipose tissue of subjects with type 2 diabetes compared with non-diabetic controls (Figure 7A and Supplementary file 2). Further, we found a negative correlation between DNA methylation and mRNA expression of FGF21 in human adipose tissue (Figure 7B–E). Together, these data support a role for increased DNA methylation of FGF21 in the regulation of adipose FGF21 expression in association with metabolic disease.

Discussion

While genetics plays an important role in obesity and T2D, genetic differences cannot fully explain many features of these conditions, such as discordance in monozygotic twins, and the close relationship with lifestyle factors (McCarthy, 2010; Morris et al., 2012; Voight et al., 2010; Zeggini et al., 2008). Of note, the vast majority of GWAS-proven allelic variants associated with T2D correspond much better to altered insulin secretion and islet function rather than to IR per se (Voight et al., 2010). Hence, forms of non-genetic variation, such as epigenetic alterations, must also be considered. This notion has been borne out by a recent epigenome-wide association study (EWAS) linking alterations in DNA methylation to whole-body insulin sensitivity (Zhao et al., 2012). Importantly, even genetic variants may cause disease in a DNA methylation-dependent manner, as has recently been shown for T2D and obesity (Dayeh et al., 2013; Elliott et al., 2017; Nilsson et al., 2014; Olsson et al., 2014; Volkov et al., 2016). We have also shown that adipose tissue from patients with T2D exhibit numerous methylation and expression differences relative to unrelated healthy controls, with many of the same changes also seen in twin pairs discordant for T2D (Nilsson et al.,...
In this study, we identified increased expression of an enzymatic effector of DNA methylation, Dnmt3a, as a key epigenetic driver of IR in vitro and in vivo.

DNA methylation, like other forms of epigenomic modification, has been an attractive therapeutic target because of their plasticity and because they offer an opportunity to reprogram cells into a more healthy state. For example, administration of pan-inhibitors of histone deacetylase exerts beneficial metabolic effects in both mice and humans, such as increased energy expenditure, insulin sensitivity and secretion (Christensen et al., 2011; Daneshpajooh et al., 2017; Sharma and Taliyan, 2016; Ye, 2013). DNA methyltransferase inhibitors have also been shown to have an insulin-
sensitizing effect in vivo and in vitro (Kim et al., 2015). In line with pharmacological studies, mutant mouse models carrying loss-of function histone modifiers (e.g. Mll2, Ehmt1, Jmhd2a Lsd1) were shown to have profound impact on whole body metabolism (Lee et al., 2006; Ohno et al., 2013; Sambeat et al., 2016; Tateishi et al., 2009; Zeng et al., 2016). It should be noted that metabolic consequences in mouse models carrying mutations of histone modifiers appear to be mainly attributable to changes in body weight. Unlike these examples, adipose-specific Dnmt3a KO mice are relatively unique in that they display improved insulin sensitivity without a change in adiposity or body weight. Similarly, the effect on insulin action is not dependent on increased beige or brown adipocytes, or on reducing intra-adipose inflammation.

Our in vitro study indicates that overexpression of Dnmt3a is sufficient to induce cell autonomous IR, however, it does not seem to be the case in vivo. Kamei et al showed that transgenic expression of Dnmt3a per se was not sufficient to render whole body IR and glucose intolerance (Kamei et al., 2010) on chow, HFD, or high methyl diet with chow diet. Authors observed only a mild increase in the gene expression of proinflammatory cytokines (i.e. Mcp1 and Tnfa) when animals were put on HFD. It should be noted that the major caveat of this study is the use of aP2 promoter to drive the transgene expression (Jeffery et al., 2014; Kang et al., 2014; Lee et al., 2013); Potentially, non-adipose expression and earlier expression during development might have a compounding factor in this study. Despite these notions, it will be highly interesting to address how the combination of

Figure 6. Dnmt3a-mediated methylation of the Fgf21 gene promoter. (A) Schematic showing the tested regions for MeDIP-qPCR at the Fgf21 locus. Target regions were chosen using MethFinder (http://www.urogene.org/methprimer) on regions proximal (5 kb) and distal (50 kb) to the Fgf21 transcription start site. (B) MeDIP-qPCR was performed using genomic DNA extracted from 3T3-L1 adipocytes transduced with GFP vs. Dnmt3a or shScr vs. shDnmt3a. Shown is relative fold enrichment of 5mC of Dnmt3a overexpressor and KD cells over control cells. (C) MeDIP-qPCR was performed using genomic DNA from Dnmt3a KO and Flox mice on HFD. Shown is relative fold enrichment of 5mC of KO over WT samples. (D) Fgf21-promoter reporter (1497/+5) assay performed by co-transfection of the indicated amounts of Dnmt3a expression vector. (n = 3 mice, p<0.05, Student’s t-test, mean ±s.e.m.).

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high methyl donor diet and HFD would affect the metabolic profile of these transgenic animals compared to controls.

Our unbiased gene expression profiling studies led us to identify Fgf21 as a key target gene in adipocytes, and the data from diabetic humans suggests that this could be a conserved mechanism of IR in both mice and humans. Fgf21 is better known as a hepatokine, and the liver is the main source of circulating Fgf21 (Markan et al., 2014). Adipose Fgf21 probably does not circulate, but rather acts in an autocrine/paracrine manner (Dutchak et al., 2012; Markan et al., 2014). In fact, levels of Fgf21 in serum were moderately reduced in Dnmt3a KO animals compared to controls, in contrast to significantly elevated adipose tissue levels. Studies have shown a positive correlation between insulin and circulating Fgf21 levels (Chavez et al., 2009; Chen et al., 2008; Zhang et al., 2008), suggesting that insulin may be a key determinant of Fgf21 in serum. Thus, it is possible that the reduced insulin levels in Dnmt3a KO mice on HFD may have led to reduction of Fgf21 production in liver. This notion is in line with the finding that Fgf21 serum levels are often elevated in obesity-associated IR (Fisher and Maratos-Flier, 2016; Morrice et al., 2017), possibly reflecting resistance to Fgf21 (Fisher et al., 2010).

We propose that the local activity of Fgf21 within the fat pad is important for the improved insulin sensitivity in the knock-out mice, although formal proof of this idea would require generating adipose tissue-specific Dnmt3a and Fgf21 double knockout mice. In line with this view, there are previous studies indicating that adipose Fgf21 is insulin-sensitizing. For example, three independent studies demonstrated that mutant mice devoid of Fgf21 signaling in adipose tissue secondary to ablation of FGFR1 or bKlotho, were refractory to the insulin-sensitizing effect of Fgf21 administration.

Figure 7. FGF21 DNA methylation in adipose tissue from subjects with T2D compared with controls. (A) Four CpG sites annotated to FGF21 with significant difference in adipose tissue DNA methylation between subjects with type 2 diabetes and controls. (p<0.01, paired Wilcoxon statistics, mean ±s.d.). (B–D) Correlations between DNA methylation and mRNA expression for CpG sites significantly associated with T2D in human adipose tissue. (r = Pearson correlation coefficient.).

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Some of these studies (Adams et al., 2013; Ding et al., 2012) suggested that adiponectin is an essential mediator of Fgf21 metabolic action; the other study (BonDurant et al., 2017) suggested that adiponectin is dispensable. In our studies, albeit it did not reach statistical significance, there was a trend toward increased mRNA expression of Adipoq in both L1 and primary Dnmt3a KO adipocytes (not shown).

Control of Fgf21 expression appears to be highly tissue-specific. For example, PPARγ is an important driver of Fgf21 in adipose tissue, while PPARα serves that role in liver (Dutchak et al., 2012). Other studies have also demonstrated that Fgf21 is positively regulated by PPARγ in fat (Muise et al., 2008; Wang et al., 2008). In addition to focused studies on Fgf21 promoter regions, global DNA methylation profiling will be necessary to identify differentially methylated regions in the setting of Dnmt3a gain- and loss-of function.

A recent study demonstrated that Dnmt1 may contribute to obesity-associated inflammation and insulin resistance by promoting hypermethylation at the Adipoq locus (Kim et al., 2015). Concordant with this study, we also see elevated Dnmt1 levels in the adipose tissue of obese animals. However, knockdown of Dnmt1 did not rescue insulin sensitivity in the setting of Dex or TNF, and overexpression of Dnmt1 did not reduce insulin-stimulated glucose uptake in cultured adipocytes. Based on those criteria, we chose to focus on Dnmt3a, and showed that adipose-specific deletion of this gene promoted insulin sensitivity even in weight-matched obese animals. Of note, we did not identify Adipoq as a regulated target after either deletion or overexpression of Dnmt3a.

Despite some effort, we have not yet identified the mechanism by which Dnmt3a expression is increased in obesity. We found that Dnmt3a is not induced by lipids (e.g. palmitate), pro-inflammatory signals (e.g. IL-6, LPS), high doses of insulin, or high glucose in cultured adipocytes (not shown). In our study, in vivo administration of thiazolidinediones normalized the expression of Dnmt3a in ob/ob mice but not in lean control animals. The mechanism by which PPARγ represses gene expression is not entirely clear. Trans-repression through binding of inflammatory transcription factors like NF-κB and AP-1 has been proposed as a general mechanism for this type of effect (Glass and Saijo, 2010) although we note that there are PPARγ binding sites near the Dnmt3a TSS (not shown). It is thus unclear if direct or indirect (i.e. trans-repression) mechanisms account for the actions of rosiglitazone on Dnmt3a expression in obesity.

In summary, we have identified a novel role for Dnmt3a as an epigenetic mediator of adipose IR in vitro and in vivo. Future studies will be required to fully elucidate the full range of its target genomic regions and metabolic effects.

Materials and methods

Cell culture

3T3-L1 preadipocytes were obtained from ATCC and maintained and differentiated as described (Kang et al., 2012). These cells were authenticated by the ability to differentiate and confirmed to be mycoplasma negative. To generate lentivirus particles, lentiviral constructs were co-transfected with pM2DG- and psPAX-expressing plasmids into 293T cells. After 48 hr, virus-containing supernatant was collected, filtered through 0.45 μm filters, and added to mature 3T3-L1 adipocytes for 24 hr along with 8 μg/ml Polybrene. Transduction efficiency was determined by comparison with cells transduced in parallel with a GFP-expressing lentivirus. For the ex vivo system, subcutaneous adipose tissue from wild-type C57Bl/6 mice was fractionated with digestion buffer (10 mg/ml collagenase D, 2.4 units of dispase II, 10 mM CaCl₂ in PBS). Cells from the stromal-vascular fraction (SVF) were plated in culture and differentiated as described previously (Qureshi et al., 2014). To differentiate hASCs, Rosi (5 μM) was added in addition to IBMX (0.5 mM), insulin (5 μg/ml), and dexamethasone (1 μM), from day 0–2.

3H-2-DG assay

Mature 3T3-L1 adipocytes were incubated in serum-free DMEM for 4–6 hr. Cells were then washed three times with KRH buffer (12 mM HEPES, pH 7.4, 121 mM NaCl, 5 mM KCl, 0.33 mM CaCl₂, and 1.2 mM MgSO₄) and incubated for 20 min in KRH buffer in the absence or presence of 50 nM insulin. Cells were treated with 2-deoxy-d-[2,6-3H]-glucose (0.33 μCi/ml) for another 10 min. Glucose uptake was stopped quickly by three rapid washes with KRH buffer containing 200 mM glucose and 10 μM...
cytochalasin B on ice. Cells were solubilized in 0.1% SDS for 30 min, and radioactivity was measured by liquid scintillation counting.

Reagents
2-Deoxy-d-[2,6-3H]-glucose was purchased from PerkinElmer NEN radiochemicals. Insulin, dexamethasone, isobutylmethylxanthine (IBMX), cytochalasin B, glucose, 2-deoxyglucose, 5-Aza, ECGG, and TF-3 were purchased from Sigma-Aldrich. Recombinant TNF-α and Fgf21 were purchased from Millipore. RG-108 was purchased from Stemgent.

Antibodies
Antibodies were purchased from Cell Signaling (Dnmt3a, 3598S; Akt, 9272; pAkt [S473], 3787; IRS-1, 2382), and from Thermo Fisher (β-actin, MA5-14739, pIRS1[pY612], 44–816G).

Animals
For rosiglitazone studies, female mice ob/ob and ob/+ were treated starting at 8 weeks of age with rosiglitazone by gavage (10 mg/kg body weight) in 0.5% carboxymethylcellulose versus 0.5% carboxymethylcellulose vehicle control daily for 6 weeks. For chow and high-fat feeding studies, male C57Bl/6J mice were put on diet beginning at 8 weeks of age and continued for three months (n = 8 per dietary condition). Samples were collected from the perigonadal fat pad.

For histology, adipose tissues were fixed with neutral-buffered formalin and embedded in paraffin and sections were stained with H&E.

For in vivo insulin signaling assay, after overnight fast, insulin (10 U/kg, 5 min) or saline IP were given to WT and KO mice on HFD. After 10 min various tissues were harvested and stored at –80°C until use. Tissue samples were homogenized in cell signaling lysis buffer containing protease inhibitors (Roche) and phosphatase inhibitors (Sigma-Aldrich) and subjected to western blotting. All animal work was approved by the BIDMC IACUC and/or the UC Berkeley ACUC.

RNA extraction and quantitative PCR
Total RNA was extracted from cells or tissues using TRIzol reagent according to the manufacturer’s instructions. cDNA was reverse-transcribed from 1 μg of RNA using the RETROscript first strand synthesis kit (Ambion). Quantitative PCR (qPCR) was performed with SYBR Green qPCR Master Mix (Applied Biosystems) using a 7900HT Fast Real-Time PCR System (Applied Biosystems) and CFX96 Touch (Bio Rad). Primer sequences are listed in Supplementary file 3. The relative amount of mRNA normalized to cyclophilin B was calculated using the delta–delta method (Livak and Schmittgen, 2001).

Plasmids
Hairpins against Dnmt1, Dnmt3a, and Dnmt3b were purchased from Sigma. Lentiviral overexpression vectors for Dnmt1, 3a, and 3b were subcloned into pcDH using various multicloning sites (XbaI/NotI for Dnmt1, EcoRI/NotI for Dnmt1, Dnmt3a, Dnmt3a-CM) and NotI sites and hairpins targeting Dnmts were subcloned at AgeI/EcoRI or purchased from Open Biosystems. Hairpin sequences are shown in Supplementary file 3. Fgf21-luc (1497/+5) was generously provided by Dr. Steven Kliewer (UT Southwestern).

MeDIP-qPCR
Genomic DNA was sheared by sonication to an average of 200–800 bp size. Two microgram of denatured DNA was incubated with 2 μg of anti-5-methylcytidine antibody (Epigentek) in IP buffer (10 mM Na-Phosphate pH 7.0, 0.14 M NaCl, 0.05% Triton X-100) for 2 hr at 4°C. Antibody-bound DNA was collected with 20 μl of Dynabeads anti-mouse IgG (Invitrogen Dynal, Oslo, Norway) for 1 hr at 4°C on a rotating wheel and successively washed five times with washing buffer (0.1% SDS, 1% Triton X-100, 2 mM EDTA, 20 mM Tris-HCl pH 8.1, 150 mM NaCl), and twice with TE (10 mM Tris-Cl, 1 mM EDTA pH 8.0). DNA was recovered in digestion buffer 125 μl buffer (50 mM Tris pH 8.0, 10 mM EDTA, 0.5% SDS, 35 μg proteinase K) and incubated for 3 hr at 65°C. Recovered DNA was used for qPCR analysis. Primers for MeDIP-qPCR studies are listed in Supplementary file 3. All data are normalized to input.
RNA-seq studies

mRNA was purified from 100 ng of total RNA using the Ribo-Zero Magnetic Gold Kit (catalog MRZG126, Illumina). Libraries were prepared using the TruSeq RNA Library Preparation Kit v2 (catalog RS-122–2001, Illumina) according to the manufacturer’s protocol starting with the EPF step. Sequencing was performed on the Illumina HiSeq2500. RNA-Seq data are aligned using TopHat2 (Kim et al., 2013). Reads are assigned to transcripts using featureCounts and an mm9 genome modified to minimize overlapping transcripts (Liao et al., 2014). Differential expression analysis of the data is performed using EdgeR (Robinson et al., 2010).

DNA methylation and mRNA expression in human adipose tissue

DNA methylation was analyzed in adipose tissue from 28 subjects with type 2 diabetes and 28 controls as well as in adipose tissue of 14 monozygotic twin pairs (n = 28) using Infinium HumanMethylation450 BeadChips (Illumina) according to our previous study where we also present the characteristics of these subjects (Nilsson et al., 2014). Their characteristics are presented in Supplementary file 4. mRNA was analyzed in adipose tissue from the 12 monozygotic twin pairs (n = 24) using GeneChip Human Gene 1.0 ST arrays (Affymetrix, Santa Clara, CA) according to the manufacturer’s recommendations. In the present study, we studied mRNA expression and DNA methylation of CpG sites annotated to FGF21. DNA and RNA were extracted from human adipose tissue as presented (Nilsson et al., 2014).

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Author contributions

Dongjoo You, Formal analysis, Validation, Investigation, Carried out most of biological experiments and assisted with preparing the manuscript; Emma Nilsson, Data curation, Formal analysis, Investigation, Collected human adipose tissue and analyzed DNA methylation and gene expression with human sample; Danielle E Tenen, Investigation, Methodology, Generated RNA-Seq library; Anna Lyubetskaya, Formal analysis, Generated and analyzed RNA-Seq data; James C Lo, Resources, Helped with metabolic characterization of knock-out mouse models; Rencong Jiang, Investigation, Carried out insulin signaling assay; Jasmine Deng, Investigation, Carried out gene expression analysis of several biological experiments; Brian A Dawes, Formal analysis, Helped with RNA-Seq data analysis, Generated figures, Submitted GEO data; Allan Vaag, Investigation, Collected human adipose tissue and analyzed DNA methylation and gene expression with human sample; Charlotte Ling, Resources, Investigation, Collected human adipose tissue and analyzed DNA methylation and gene expression data, Supervised overall human data set; Evan D Rosen, Supervision, Funding acquisition.
Writing—original draft, Writing—review and editing, Supervised experiments and edited the manuscript and revision for the revision; Sona Kang, Supervision, Investigation, Writing—original draft, Writing—review and editing, Designed most of biological experiments, Supervised experiments, Wrote draft and revision for the revision.

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Ethics
Animal experimentation: This study was performed in strict accordance with the recommendations in the Guide for the Care and Use of Laboratory Animals of the National Institutes of Health. All animal work was approved by the BIDMC IACUC (056-2017) and/or the UC Berkeley ACUC (AUP-2015-08-7887).

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Additional files

Supplementary files

• Supplementary file 1. Genes that are up-regulated (UP) and down-regulated (DOWN) Dnmt3a vs. GFP and shDnmt3a/shScr.
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• Supplementary file 2. DNA methylation of four CpG sites annotated to Fgf21 showing differential DNA methylation in adipose tissue of a case control study for type 2 diabetes.
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• Supplementary file 3. Oligonucleotide sequences used in this manuscript.
  DOI: https://doi.org/10.7554/eLife.30766.037

• Supplementary file 4. Clinical characteristics of study subjects included in the discordant twin cohort and the case control cohort.
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