The Secreted Enzyme PM20D1 Regulates Lipidated Amino Acid Uncouplers of Mitochondria

Graphical Abstract

Highlights
- PM20D1 is a secreted enzyme that regulates N-acyl amino acids in vivo
- N-acyl amino acids are endogenous metabolites that uncouple mitochondria
- Increased PM20D1 or N-acyl amino acid administration augments energy expenditure

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In Brief
Brown and beige fat cells secrete an enzyme that tacks lipids on to amino acids. These N-acyl amino acids directly activate mitochondria for thermogenesis.

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The Secreted Enzyme PM20D1 Regulates Lipidated Amino Acid Uncouplers of Mitochondria


SUMMARY

Brown and beige adipocytes are specialized cells that express uncoupling protein 1 (UCP1) and dissipate chemical energy as heat. These cells likely possess alternative UCP1-independent thermogenic mechanisms. Here, we identify a secreted enzyme, peptidase M20 domain containing 1 (PM20D1), that is enriched in UCP1⁺ versus UCP1⁻ adipocytes. We demonstrate that PM20D1 is a bidirectional enzyme in vitro, catalyzing both the condensation of fatty acids and amino acids to generate N-acyl amino acids and also the reverse hydrolytic reaction. N-acyl amino acids directly bind mitochondria and function as endogenous uncouplers of UCP1-independent respiration. Mice with increased circulating PM20D1 have augmented respiration and increased N-acyl amino acids in blood. Lastly, administration of N-acyl amino acids to mice improves glucose homeostasis and increases energy expenditure. These data identify an enzymatic node and a family of metabolites that regulate energy homeostasis. This pathway might be useful for treating obesity and associated disorders.

INTRODUCTION

Adaptive thermogenesis has gained increasing attention as a process to fight the epidemic of obesity and type 2 diabetes. Mammals have brown and beige fat, two tissues that express uncoupling protein 1 (UCP1) and that are specialized to dissipate stored chemical energy in the form of heat (Cannon and Nedergaard, 2004; Harms and Seale, 2013). Either ablation of brown or beige cells (Cohen et al., 2014; Lowell et al., 1993) or knockout (KO) of the Ucp1 gene (Feldmann et al., 2009) predisposes mice to obesity and diabetes. Conversely, increasing the number or activity of brown and beige cells is protective against weight gain and metabolic disease (Seale et al., 2011).

Most studies of adaptive thermogenesis and thermogenic fat have centered on the expression and function of UCP1. This protein catalyzes a “proton leak” whereby protons that are pumped out of the mitochondrial matrix in the electron transport chain (ETC) are transported back across the inner mitochondrial membrane (Nicholls et al., 1978; Rouxset et al., 2004). This results in oxidative metabolism with no production of ATP, a process referred to as uncoupled respiration. While UCP1 is a very important part of adaptive thermogenesis, in principle, any biochemical process that requires energy and is not linked to energy storage or work can function as a thermogenic event. Indeed, data have emerged indicating that UCP1 is not the only mediator of this process (Kazak et al., 2015; Ukopec et al., 2006). Moreover, other carriers of the mitochondrial SLC25 family, of which UCP1 is only one member (SLC25A7), also have the ability catalyze a proton leak across the inner mitochondrial membrane (Brand et al., 2005).

In addition to storing chemical energy, adipose cells are now recognized to be important sensors of energy balance and secrete many bioactive proteins, including adipin, leptin, and adiponectin (Kershaw and Flier, 2004). Proteins secreted by brown and beige fat cells in particular have not been extensively or systematically studied (Svensson et al., 2016; Wang et al., 2014). We recently developed the UCP1-TRAP mouse to identify the gene expression signature of brown and beige cells in vivo, regardless of their anatomical localization (Long et al., 2014). These initial experiments elucidated a smooth muscle-like origin for beige adipocytes, while also providing a comprehensive molecular inventory of the thermogenic fat cell in vivo.

Here, by combining the UCP1-TRAP molecular inventory with other datasets, we have compiled a core thermogenesis gene set that is co-expressed in vivo with Ucp1. We hypothesized that such a gene set might be used to identify secreted proteins that play a significant role in adaptive thermogenesis. This analysis led to the identification of a previously unstudied secreted enzyme, peptidase M20 domain containing 1 (PM20D1). We demonstrate that PM20D1 is a biosynthetic enzyme for a class of N-lipidated amino acids in vivo, and these metabolites function as endogenous uncouplers of mitochondrial respiration in a UCP1-independent manner.
RESULTS

PM20D1 Is Expressed in UCP1+ Adipocytes and Promotes Energy Expenditure In Vivo

To identify secreted factors from brown and beige adipocytes, we first generated a list of “core thermogenesis” genes that are co-expressed with Ucp1 in vivo. To this end, we identified the overlapping gene set from the following publicly available microarray/RNA sequencing (RNA-seq) datasets: (1) enrichment in the classical brown adipose tissue (BAT) versus the epididymal white fat (eWAT); (2) equivalent expression in both brown and beige cells in vivo using the TRAP method; and (3) induction in the cold in the subcutaneous inguinal white fat (iWAT) following 1 or 5 weeks cold exposure (Long et al., 2014; Seale et al., 2007; Xue et al., 2009) (Figure 1A). Thirty-two genes passed these filters (Table S1). Consistent with our original search strategy, cross-referencing of these 32 genes with the Universal Protein Resource (UniProt) demonstrated that half of these candidates (16 out of 32) were mitochondrial in subcellular localization (Table S1). Of the remainder, only one, PM20D1, contained a signal peptide without any transmembrane domains, two features characteristic of a classically secreted protein.

We validated the original microarray/RNA-seq datasets in a new cohort of mice and found that Pm20d1 mRNA was higher in BAT versus the other fat depots and cold-inducible in the iWAT depot (Figures S1A and S1B). Further supporting Pm20d1 co-expression with Ucp1 in vivo, Pm20d1 and Ucp1 mRNA were coordinately upregulated in the iWAT following treatment of mice with the β-adrenergic receptor agonist CL-316,243 (Figure S1C) and coordinately downregulated in the eWAT following high fat diet (Figure S1D). Shotgun proteomics with tandem mass tag (TMT) labeling confirmed the presence of PM20D1 in blood, though its circulating levels were unchanged following 10 days of cold exposure (Figures S1E and S1F).

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S1F; Table S2). This is presumably because, in addition to UCP1+ fat cells, liver and kidney also highly express PM20D1 (Figure S1A). To confirm that PM20D1 can be secreted from intact cells, we generated a C-terminal flag-tagged PM20D1 cDNA construct and transfected this plasmid into HEK293A cells. PM20D1 was detected both in cells and in conditioned media, whereas flag-tagged GFP was found exclusively in the cellular fraction (Figure S1G). These data demonstrate that PM20D1 is a bona fide secreted factor enriched in UCP1+ versus UCP1− adipocytes and induced in adipose tissues by cold exposure.

To assess the functions of PM20D1 in vivo, mice were tail vein-injected with adeno-associated viral vectors (serotype AAV8) expressing PM20D1 or GFP. These vectors are primarily taken up and expressed by the liver, although other tissues may also express them (Zincarelli et al., 2008). One week following the injections, the mice were placed on high fat diet (HFD). Increased circulating PM20D1 was observed by western blots of the plasma at 40 days post-injection (Figure 1B). At room temperature (22°C), mice with augmented circulating PM20D1 showed blunted weight gain (final weight means ± SEM: GFP, 44.2 ± 1.0 g; PM20D1, 39.8 ± 1.5 g; 10% weight difference) compared to the control animals (Figure 1C). A similar blunting of weight gain was observed at thermoneutrality (30°C) (final weight means ± SEM: GFP, 35.0 ± 0.8 g; PM20D1, 31.8 ± 0.7 g; 9% weight difference; Figure 1D), where the sympathetic nervous input to adipose tissues is decreased. Body composition analysis by MRI at the end of the experiment at thermoneutrality revealed that the weight difference was due exclusively to a 30% reduction in fat mass in those animals receiving PM20D1 compared to GFP; there were no effects on lean mass (Figures 1E and 1F). In a separate cohort of mice treated with AAV-PM20D1 or AAV-GFP at room temperature, we performed whole body indirect calorimetry measurements at a time point prior to the divergence in weights (4 weeks HFD; Figure S2A). These analyses revealed significantly augmented VO2 (Figure 1G) and VCO2 (Figure S2B), indicative of increased energy expenditure. Most importantly, this occurred with no changes in movement (Figure 1H) or food intake (Figure S2C).

**PM20D1 Regulates N-Lipidated Amino Acids In Vivo**

Changes in energy expenditure with no change in physical movement are usually indicative of activation of a thermogenic gene program in the classical brown fat, subcutaneous iWAT (termed browning), or both. In addition, recent work has indicated the presence of an additional thermogenic pathway based on a futile cycle of creatine phosphorylation (Kazak et al., 2015). Surprisingly, the increased metabolic rates observed here were not accompanied by any broad molecular change corresponding to these processes, such as an induction of UCP1 (Figures 2A and 2B), or changes in gene expression related to creatine metabolism (Figure 2A). These data suggest that PM20D1 increases whole body energy expenditure through an alternative mechanism.

To further investigate how PM20D1 augmented energy expenditure, we examined the detailed molecular consequences of PM20D1 actions in the blood. PM20D1 is one of five members of the mammalian M20 peptidase family, but remains entirely uncharacterized with respect to its endogenous substrates and products. Despite their annotation as metallopeptidases, the other four other mammalian M20 family members possess peptidase activity on a variety of small molecule substrates (Teufel et al., 2003; Van Coster et al., 2005; Veiga-da-Cunha et al., 2014). We therefore performed untargeted polar small molecule liquid chromatography-mass spectrometry (LC-MS) profiling of plasma from mice injected with AAV-GFP or AAV-PM20D1 viral vectors (Smith et al., 2006). Manual inspection of differential peaks revealed the most robust difference occurred in a metabolite with an m/z = 428, which was increased in mice injected with AAV-PM20D1 versus AAV-GFP (Figure 2C; Table S3). We identified this m/z = 428 peak as the [M-H]− ion of N-oleoyl phenylalanine (C18:1-Phe, chemical formula C_{27}H_{42}O_{2}N, expected m/z = 428) (Figure 2D). This identification was confirmed by tandem mass spectrometry (MS/MS) experiments, which revealed a predominant product ion of m/z = 164 corresponding to the phenylalanine anion (Figure 2E; chemical formula C_{16}H_{29}O_{2}N, expected m/z = 164). Further supporting this identification, both synthetic C18:1-Phe and the endogenous m/z = 428 peak eluted at similar retention times by LC (Figure 2F).

We then developed a targeted multiple reaction monitoring (MRM) program to assess the scope of N-acyl amino acids regulated by PM20D1 in plasma in vivo. By absolute quantitation using a C15-Phe internal standard, the basal plasma concentrations of specific members of the N-acyl amino acids in AAV-GFP injected mice were found to be in the 1–100 nM range (Table S4). PM20D1 predominantly elevates medium and long N-acyl Phe (C14–C18; Figures 3A and 3B) and also N-oleoyl amino acid metabolites with large and hydrophobic head groups (C18:1-Phe and C18:1-Leu/Ile; Figures 3C and 3D). Similar changes in N-acyl amino acids were observed for mice injected with AAV-PM20D1 at room temperature (Figure S3). Because Pm20d1 mRNA is induced in the iWAT following cold exposure (Figures 1A and 1A), we also assessed plasma levels of the N-oleoyl amino acids following cold exposure by targeted MS. Six hours or 2 days cold exposure increased specific members of this class (C18:1-Leu/Ile and C18:1-Val; Figure 3E). Long-term cold exposure (16 days) significantly elevated the levels of most N-oleoyl amino acids that were measured (Figure 3E). Therefore, both PM20D1 and its N-acyl amino acid products are physiologically co-regulated by cold exposure.

**PM20D1 Is a Bidirectional N-Acyl Amino Acid Synthase and Hydrolase In Vitro**

Although the endogenous presence of N-acyl amino acids has been previously described (Tan et al., 2010), their biosynthesis has remained a long-standing mystery. The increase of these metabolites in AAV-PM20D1 mice suggested that PM20D1 might be an enzyme responsible for N-acyl amino acid biosynthesis in vivo. To investigate this process in vitro, we generated purified, mammalian recombinant mouse PM20D1 for enzymatic assays. As expected, purified mouse PM20D1 migrated as an ~60 kDa band by Coomassie staining (Figure S4A). Significant formation of C18:1-Phe was observed by LC-MS when recombinant PM20D1 was incubated in the reaction mixture with physiologically relevant concentrations of free oleate and Phe (300 μM and 100 μM, respectively; Figures 4A and 4B). Among different amine head groups, Phe was the amino acid most efficiently phosphorylated (Kazak et al., 2015). Surprisingly, the increased metabolic rates observed here were not accompanied by any broad molecular change corresponding to these processes, such as an induction of UCP1 (Figures 2A and 2B), or changes in gene expression related to creatine metabolism (Figure 2A). These data suggest that PM20D1 increases whole body energy expenditure through an alternative mechanism.
Figure 2. Lack of Classical Browning and Identification of Increased N-Oleoyl Phenylalanine in Mice Injected with AAV-PM20D1

(A and B) mRNA expression of the indicated genes in BAT, iWAT, and eWAT (A) and western blot of UCP1 and mitochondrial proteins (B) from male C57BL/6 mice at thermoneutrality after tail vein injections of AAV-GFP or AAV-PM20D1. Mice were 7 weeks old at the time of injection, high fat diet (HFD) was started 7 days post injection of AAV-GFP or AAV-PM20D1. Mice were 7 weeks old and high fat diet (HFD) was started 7 days post injection. Mice were maintained at room temperature for the duration of the experiment. (A and B) Data are from 47 days post injection. (A) n = 8/group, mean ± SEM, *p < 0.05. (B) n = 4–5/group.

(C) Chromatogram at m/z = 428 from plasma of male C57BL/6 mice after tail vein injection of AAV-GFP or AAV-PM20D1. Mice were 7 weeks old at the time of injection, high fat diet (HFD) was started 7 days post injection and mice were maintained at room temperature for the duration of the experiment. The comparative metabolomics was performed on plasma harvested 54 days post injection. n = 4/group, *p < 0.05.

(D) N-oleoyl phenylalanine (C18:1-Phe).

(E and F) MS/MS spectra (E) and retention time (F) of endogenous (top) or synthetic (bottom) C18:1-Phe.

See also Table S3.

Based on sequence homology with the other members of the mammalian M20 family and Uniprot annotations, we generated three point mutations in PM20D1 that we anticipated would disrupt catalytic activity. Due to differences in protein stability and expression, following flag-immunoaffinity purification, each construct was titrated such that approximately equivalent protein amounts were used in the subsequent enzymatic assays (Figure 4G). Under synthase activity reaction conditions using oleate and Phe as substrates, or hydrolase activity reaction conditions using C18:1-Phe, only wild-type (WT) PM20D1 possessed enzymatic activity, whereas no activity was observed for any of the mutants (Figures 4H and 4I). In these in vitro assays, we observe 1.2% ± 0.1% and 94.0% ± 0.8% conversion (means ± SEM, n = 3) in the synthase and hydrolase direction, respectively for the wild-type enzyme. Therefore, the synthase and hydrolase activities are embodied within the PM20D1 polypeptide and not a co-purifying protein. PM20D1 appears to require residues predicted to coordinate divalent cations, and both synthase and hydrolase activities are coordinately disrupted by point mutations in the cation binding sites.

Figure 4C). PM20D1 could also condense other amino acids with oleate, although less efficiently than Phe (Figure 4C). The negatively charged amino acid glutamate, as well as ethanolamine (EA), were not substrates for the synthase reaction (Figure 4C). PM20D1 was also capable of using arachidonate as a fatty acid donor (Figure 4D) and showed strong preference for free oleate over oleoyl-coenzyme A (C18:1-CoA; Figure 4E). Thus, free fatty acids and free amino acids are substrates for PM20D1 and its N-acyl amino acid synthase activity shows selectivity for specific amino groups and acyl donors.

We also observed that incubation of C18:1-Phe with PM20D1 liberated free oleate, indicating that PM20D1 can also act as an N-acyl amino acid hydrolase (Figures 4A and 4F). The hydrolase activity was apparently more promiscuous than the synthase activity since PM20D1 hydrolyzed all N-oleoyl amino acids tested (Figure 4F). In contrast, N-oleoyl ethanolamine (C18:1-EA), a well-characterized substrate for fatty acid amide hydrolase (FAAH) (Cravatt et al., 1996; Saghatelian et al., 2004), was not a PM20D1 substrate under these conditions (Figure 4F).
Finally, we generated purified, recombinant human PM20D1, which shares 71% identity and 86% similarity with the mouse enzyme; the human protein also shows complete conservation of the H125, D127, and H465 residues required for catalysis (Figures 4J and 4K). Human PM20D1 also possessed N-acyl amino acid synthase and hydrolase activities (Figures 4J and 4K), demonstrating the conservation of this enzymatic activity in the various species of mammalian protein from these two mammalian species.

Taken together, these studies indicate that PM20D1 is a bidirectional enzyme that can generate N-acyl amino acids from amino acids and fatty acids and can also hydrolyze N-acyl amino acids into amino acids and free fatty acids. The changes in the various species of N-acyl amino acids in plasma from AAV-PM20D1 injected mice are therefore likely to reflect a balance of the relative synthase and hydrolase activities on, as well as the relative concentrations of, the particular substrates and products. Differences between N-acyl amino acids regulated by PM20D1 overexpression and cold exposure (Figure 3) likely reflect different levels of fatty acid or amino acid substrates under various physiologic conditions. However, we cannot exclude other, as of yet unknown enzymes that might perform similar chemical reactions.

**N-Acyl Amino Acids Are Endogenous Uncouplers of Respiration in Cells**

To directly test the effect of N-acyl amino acids on respiration, we acutely treated differentiated primary BAT adipocytes first with the ATP synthase inhibitor oligomycin to block coupled respiration and then with C18:1-Phe (50 μM). Cellular oxygen consumption was significantly augmented by C18:1-Phe treatment (maximal oxygen consumption rates [OCR] of 198% versus oligomycin-treated basal respiration; and 156% versus DMSO treated basal respiration; and 285% and 214%, respectively, versus oligomycin-treated basal respiration; and 285% and 214%, respectively, versus DMSO). Similarly, in differentiated primary iWAT adipocytes, both C20:4-Gly and C20:4-Phe augmented respiration in the presence of oligomycin (maximal OCR of 235% and 243%, respectively). As a control, arachidonate itself also increased the oxygen consumption rate very modestly, but not nearly as much in magnitude as the N-acyl amino acids (Figure 5B).

The major biological mechanism currently understood for inducing uncoupled respiration in adipocytes is through the action of UCP1 (Rouset et al., 2004). However, C20:4-Phe...
Figure 4. Enzymatic Activity of PM20D1 In Vitro
(A) Schematic of synthase and hydrolase reaction of free fatty acid and free amino acid to N-acyl amino acid.
(B) Relative levels of C18:1-Phe generated in vitro from Phe (100 μM), oleate (0.03–1.5 mM), and purified mouse PM20D1-flag.
(C–E) Relative levels of C18:1-amino acid generated in vitro from the indicated head group (100 μM) and purified mouse PM20D1-flag using either oleate (1.5 mM, C), arachidonate (1.5 mM, D), or oleoyl-coenzyme A (C18:1-CoA, 0.7 mM, E). EA, ethanolamine.
(F) Relative levels of oleate generated in vitro from the indicated N-acyl amide substrates (100 μM) and purified mouse PM20D1-flag. C18:1-EA, N-oleoyl ethanolamine.
(G) Anti-flag western blot of immunoaffinity purified mouse PM20D1-flag or the indicated point mutants.
(H) Relative levels of C18:1-Phe generated in vitro from Phe (100 μM), oleate (1.5 mM), and the indicated wild-type (WT) or mutant PM20D1-flag protein.
(I) Relative levels of oleate generated in vitro from C18:1-Phe (100 μM) and the indicated wild-type (WT) or mutant PM20D1-flag protein.
(J) Relative levels of C18:1-amino acid generated in vitro from the indicated head group (100 μM), oleate (1.5 mM), and purified human PM20D1-flag.
(K) Relative levels of oleate generated in vitro from the indicated N-acyl amide substrate (100 μM) and purified human PM20D1-flag.
For (B)–(F) and (H)–(K), enzymatic assays were carried out in PBS at 37°C for 1.5 hr, n = 3/group, mean ± SEM, *p < 0.05, **p < 0.01, for reaction with PM20D1 versus reaction omitting PM20D1, or reaction with PM20D1 versus reaction with heat-denatured PM20D1. Y-axes indicates relative ion intensity normalized to 1 nmol of a D3,15N-serine internal standard that was doped in during the extraction process prior to MS analysis.
See also Figure S4.
The following non-standard abbreviations are used: C18:1-Phe-NH2 (N-oleoyl phenylalanine amide), C18:1-Phe-OCH3 (N-oleoyl phenylalanine methyl ester), C18:1-Leu (20:1, 1–12, July 14, 2016), 1–12, July 14, 2016

produced virtually identical increases in uncoupled respiration in both UCP1-WT and KO cells (maximal OCR of 221% and 214%, in UCP1-WT and UCP1-KO cells, respectively, versus the oligomycin-treated basal respiration, \( p > 0.05 \) between genotypes; Figure 5C), establishing that UCP1 is not required for this effect. Consistent with these observations, N-acyl amino acids were

Figure 5. Effects of N-Acyl Amino Acids on Respiration in Cells

(A–C) Oxygen consumption rates (OCRs) of differentiated primary BAT cells (A), differentiated primary iWAT cells (B), and differentiated primary BAT cells from UCP1-WT or UCP1-KO mice (C), treated with the indicated compounds for the indicated time. Adipocytes were differentiated and analyzed on day 5.

(D–H) OCRs of C2C12 cells (D–F and H) or U2OS cells (G) treated with the indicated compounds for the indicated time.

(E) Time courses for representative experiments.

(F) uATP values for representative experiments.

(G) The following non-standard abbreviations are used: C18:1-Phe-NH2 (N-oleoyl phenylalanine amide), C18:1-Phe-OCH3 (N-oleoyl phenylalanine methyl ester). (H) The following non-standard abbreviations are used: C20:4-Phe, C20:4-Dopamine, C20:4-GABA, C20:4-Ser, C20:4-Gly, C20:4-NAT (N-arachidonoyl taurine), C20:4-NAT (N-arachidonoyl taurine), C20:4-GABA (N-arachidonoyl gamma-amino butyric acid).

For (A)–(H), n = 3–6/group, mean ± SEM, *p < 0.05, **p < 0.01, ***p < 0.001 for treatment versus DMSO at the same time point.
also competent inducers of uncoupled respiration in cell types that completely lack UCP1, including the mouse myoblast cell line C2C12 (Figures 5D–5F) and the human osteosarcoma cell line U2OS (Figure 5G). These data show that N-acetyl amino acids are endogenous chemical uncouplers of mitochondrial respiration in a UCP1-independent manner.

We next explored the structural requirements for N-acetyl amino acids to exert their effects on uncoupled respiration. A direct comparison of C18:1-Leu and oleate demonstrated that the N-acetyl amino acid conjugate was significantly more potent than the free fatty acid alone (Figure 5D). Modification of the free amino acid carboxylate of C18:1-Phe, either by methyl esterification (C18:1-Phe-OCH3), or by primary amidation (C18:1-Phe-NH2), completely abrogated uncoupling activity (Figure 5E), indicating that the amino acid carboxylate moiety is required for activity. We observed some amino acid head group selectivity to the respiration effects, as C18:1-Gln, but not C18:1-Lys, possessed the uncoupling activity (Figure 5F). Modification of the lipid chain revealed that only C16-, C18-, and C18:1-Phe possessed uncoupling activity, whereas saturated acyl chains that were too short (C12:0-Phe) or too long (C20:0-Phe) completely lacked uncoupling activity (Figure 5G). Finally, testing multiple commercially available fatty acid amides revealed that most did not stimulate uncoupled respiration, unless they possessed these general structural features outlined above (e.g., C20:4-GABA; Figure 5H).

**N-Acetyl Amino Acids Directly Uncouple Mitochondria and Interact with Mitochondrial Proteins**

To directly assess the ability of N-acetyl amino acids to uncouple mitochondria, we employed two approaches. First, we isolated mitochondria from the BAT tissues and treated these mitochondria with increasing concentrations of C18:1-Phe (10–100 μM). Isolated BAT mitochondria increased respiration in a dose-dependent manner following C18:1-Phe treatment (Figure 6A), indicating that N-acetyl amino acids do not require any other cellular components or organelles for their uncoupling effects. Second, we used tetramethylrhodamine methyl ester (TMRM) fluorescence to directly measure the mitochondrial membrane potential in live cells. As expected, treatment of C2C12 cells with oligomycin increased the membrane potential (Figure 6B). Co-treatment of oligomycin with C18:1-Phe (50 μM) decreased the TMRM fluorescence by ∼45% (Figure 6B). As a positive control, the well-known chemical uncoupler FCCP (0.4 μM) also reduced TMRM fluorescence by an even larger magnitude (∼70% reduction). Therefore, N-acetyl amino acids can directly augment uncoupled respiration in isolated mitochondria, resulting in a decreased mitochondrial membrane potential.

The structure activity relationships (SAR) observed with N-acetyl amino acids-induced uncoupling (Figure 5), and the direct effects of N-acetyl amino acids on isolated mitochondria (Figure 6A) are consistent with specific binding interactions between these metabolites and mitochondrial proteins (Niphakis et al., 2015). To identify the proteins that may be mediating the uncoupling by N-acetyl amino acid, we synthesized a photo-crosslinkable version (Figure 6C). This molecule, which we term “photo-probe,” contains a modified Met amino acid with a photo-cross-linking diazarine side chain and a fatty acid-alkyne for downstream click chemistry applications. Photo-probe (50 μM) was a competent inducer of uncoupled respiration in C2C12 cells, demonstrating that the alkyne and diazarine modifications did not affect the bioactivity (Figure 6D). Moreover, photo-probe demonstrated robust, UV-dependent crosslinking as assessed by in-gel TAMRA fluorescence (Figure 6E).

C2C12 cells were selected for LC-MS/MS analysis of photo-probe targets since they demonstrate robust N-acetyl amino acid induced uncoupling. To this end, C2C12 cells were incubated with photo-probe (20 μM, “probe only” samples), or co-incubated with both photo-probe (20 μM) and a C20:4-Phe competitor at 5-fold excess (100 μM, “probe + competitor” samples). Cells were then UV irradiated on ice and lysed by sonication. Probe-labeled proteins were conjugated to biotin-N3 by click chemistry, streptavidin-enriched, and subject to LC-MS/MS analysis with spectral counting. In total, 149 proteins were identified that showed >50% competition by C20:4-Phe (Figure 6F; Table S5). Of these, 31 proteins (21%) are localized to the mitochondria by Uniprot annotation, including six members of the SLC25 carrier family (Figure 6F; Table S5). Notably, the two most abundantly detected proteins in the entire dataset were the mitochondrial SLC25A4 and SLC25A5 (also known as ANT1 and 2). In addition to their ADP/ATP symport activity, these transporters have previously been demonstrated to translocate protons across the inner membrane (Brand et al., 2005). Taken together, these data are consistent with a model where N-acetyl amino acids increase uncoupled respiration by liganding SLC25 family members, including SLC25A4 and SLC25A5, and increasing SLC25-mediated proton flux into the matrix.

**Administration of N-Acyl Amino Acids to Mice Increases Energy Expenditure and Improves Glucose Homeostasis**

We next sought to determine the physiologic effects of N-lipidated amino acid administration to mice. Diet-induced obese (DIO) mice were treated daily with vehicle, oleate, or C18:1-Leu (25 mg/kg, intraperitoneally [i.p.]). After 8 days treatment, mice receiving C18:1-Leu lost 4.1 ± 0.3 g, whereas mice treated with oleate or vehicle only lost 0.3 ± 0.2 g and 0.6 ± 0.1 g, respectively (means ± SEM; Figure 7A). Over this time course, food intake was slightly but significantly reduced in mice receiving C18:1-Leu (17% reduction versus vehicle-treated mice; Figures 7B and 7C). Body composition analysis at the end of this experiment showed the weight loss induced by C18:1-Leu was entirely accounted for by a difference in fat mass (Figure 7D). Lastly, C18:1-Leu-treated mice also showed improvements in GTT versus either vehicle- or oleate-treated mice (Figure 7E).

To assess the effects of C18:1-Leu on whole body energy expenditure, we performed indirect calorimetry measurements in a separate cohort of mice treated daily with vehicle or C18:1-Leu (25 mg/kg/day, i.p.). After eight daily injections, mice were placed into metabolic cages and injected for an additional 2 days (Figure S5). Mice treated with C18:1-Leu showed significantly augmented VO2 (Figure S7F) and VCO2 (Figure S5F) compared with vehicle-treated mice and also slightly reduced movement (Figure 7G). In this cohort, C18:1-Leu-treated mice also showed a reduced food intake over the final 2 days of the experiment (Figure S5H), but not during the earlier time period (days 0–7; Figures S5C and S5D). Finally, the respiratory
Membrane potential

C20:4-Gly reduced food intake, these compounds nevertheless dose. While chronic treatment of mice with C18:1-Phe or post-injection, respectively, following a single 30 mg/kg i.p. m

0.3 (15 mg/kg/day, i.p.). Blood levels of C18:1-Phe were 3.0 daily with C18:1-Phe (30 or 50 mg/kg/day, i.p.) or C20:4-Gly in vivo, we analyzed additional cohorts of DIO mice treated

analyzed by MS (see the Experimental Procedures). Proteins satisfying the following filtering criteria are shown: >50% reduction in peptide counts with competitor present versus without competitor, and detection of at least one peptide in all three probe only samples. Comparisons in which no peptides were

See also Table S5.

Photo-probe with 100 (F) Proteins in C2C12 cells that showed C20:4-Phe competed photo-probe labeling. C2C12 cells were incubated with 20 μM photo-probe (50 μM), or in combination with TAMRA-N3. Control cells that were not UV-irradiated were kept under ambient light (on ice, 10 min).

OCR, which is normalized to 100%. n = 3–4/group, mean ± SEM, n = 3) at 2 hr and 6 hr –UV +UV fluorescence In-gel TAMRA

maximal increase in OCR (as % of oligomycin basal) (average spectral counts)

control cells and cat-

exchange ratio (RER) was significantly lower in mice receiving C18:1-Leu, indicating a switch to fats as a metabolic fuel type (Figure S5G).

The uncoupling activity of N-acyl amino acids appears to be a property of this entire class of metabolites, at least in cells. To explore the generality of N-acyl amino acid bioactivity in vivo, we analyzed additional cohorts of DIO mice treated daily with C18:1-Phe (30 or 50 mg/kg/day, i.p.) or C20:4-Gly (15 mg/kg/day, i.p.). Blood levels of C18:1-Phe were 3.0 ± 0.3 μM and 0.4 ± 0.1 μM (means ± SEM, n = 3) at 2 hr and 6 hr post-injection, respectively, following a single 30 mg/kg i.p. dose. While chronic treatment of mice with C18:1-Phe or C20:4-Gly reduced food intake, these compounds nevertheless significantly augmented VO2, with no effects on movement (Figures S6 and S7). Measurements of plasma aspartate aminotransferase (AST) and alanine aminotransferase (ALT), as well as plasma cytokines, revealed no significant elevations in mice treated with C20:4-Gly versus vehicle (Figures S7G–S7J). Taken together, these data demonstrate that N-acyl amino acids can directly augment whole body energy expenditure, reduce fat mass, and improve glucose clearance in mice.

DISCUSSION

We have characterized a previously unstudied enzyme, PM20D1, that is enriched in UCP1+ versus UCP1− cells and catalyzes the condensation of fatty acids and certain amino acids to form N-acyl amino acids. While these lipidated metabolites had previously been detected in biological tissues, little was known about their biosynthesis or physiologic functions. Here, we show that N-acyl amino acids function as endogenous uncouplers of mitochondrial respiration, even in cells lacking UCP1.

Figure 6. Effects of N-Acyl Amino Acids in Mitochondria and Identification of N-Acyl Amino Acid-Interacting Proteins
(A) Oxygen consumption rates (OCRs) of freshly isolated BAT mitochondria treated with the indicated compounds for the indicated times. Respiration was measured with 10 mM pyruvate and 5 mM malate as substrates, and FCCP and rotenone were used at 2 μM and 3 μM, respectively. n = 4–5/group, **p < 0.01. (B) Tetramethyl rhodamine methyl ester (TMRM) fluorescence in C2C12 cells following 20 min treatment with oligomycin alone (1 μM), or in combination with C18:1-Phe (10 or 50 μM) or FCCP (0.4 μM), n = 3/group, mean ± SEM. "p < 0.05, **p < 0.01. (C) Chemical structure of the N-acyl amino acid photo-crosslinkable probe ("photo-probe"). (D) OCR of C2C12 cells treated with DMSO or the photo-probe (50 μM), Data is shown as the maximal increase in OCR as a percentage of the oligomycin basal OCR, which is normalized to 100%. n = 3–4/group, mean ± SEM, **p < 0.01. (E) TAMRA in-gel fluorescence of C2C12 cells treated with the photo-probe (50 μM, 20 min), followed by UV irradiation (on ice, 10 min), cell lysis, and click chemistry with TAMRA-N3. Control cells that were not UV-irradiated were kept under ambient light (on ice, 10 min). (F) Proteins in C2C12 cells that showed C20:4-Phe competed photo-probe labeling. C2C12 cells were incubated with 20 μM photo-probe ("probe only") or 20 μM photo-probe with 100 μM C20:4-Phe competitor ("probe + competitor"). Cells were then UV-irradiated, lysed, subjected to click chemistry with biotin-N3, and analyzed by MS (see the Experimental Procedures). Proteins satisfying the following filtering criteria are shown: >50% reduction in peptide counts with competitor present versus without competitor, and detection of at least one peptide in all three probe only samples. Comparisons in which no peptides were detected in “probe + competitor” samples were assigned a fold change of 15. See also Table S5.
PM20D1 in adipose tissues. Knockout mice will definitely establish the contribution of various knockout mice will definitely establish the contribution of various

tissue-specific knockout mice will definitely establish the contribution of various tissues to circulating PM20D1 levels and the physiologic roles of PM20D1 in adipose tissues.

These data suggest that either PM20D1, or N-acyl amino acids themselves, might be used therapeutically for the treatment of obesity and other obesity-associated disorders. The therapeutic use of synthetic chemical uncouplers has been limited by untoward and even fatal side effects (Grundlingh et al., 2011). Whether a wider therapeutic window for N-acyl amino acids exists remains to be determined. Although administration of several distinct N-acyl amino acids to mice augments energy expenditure and promotes weight loss, we also observe a small but significant reduction in food intake, at least under our chronic dosing regimen. The total weight loss observed in the DIO mice treated with N-acyl amino acids is likely due to a combination of both reduced food intake and augmented energy expenditure. Further exploration of other naturally occurring lipidated amino acids or chemical modifications of such molecules might identify compounds that can dissociate the beneficial from any potential undesired effects. Alternatively, PM20D1 protein injections might also be used to augment N-acyl amino acid levels in vivo.

Besides potential therapeutic applications, these studies on the enzymology of PM20D1 address long-standing questions regarding biosynthetic pathways for N-acyl amino acids. Historically, an enzymatic activity involving the condensation of fatty acids or chemical modifications of such molecules might identify compounds that can dissociate the beneficial from any potential undesired effects. Alternatively, PM20D1 protein injections might also be used to augment N-acyl amino acid levels in vivo.
acids and amino acids has been previously reported in tissues (Fukui and Axelrod, 1961), but its molecular identity has remained unclear. The data here provide strong evidence that PM20D1 is an enzyme responsible for these activities. From an chemical equilibrium point of view, we calculate a thermodynamic equilibrium of ~1% conversion to N-acyl amino acid, based on the in vitro synthesize reactant concentrations (1.5 mM oleate and 100 μM Phe) and the equilibrium constant for a related amide condensation and hydrolysis reaction (Katayma et al., 1999). Our experimentally observed N-acyl amino acid generation in the synthase direction (1.2% ± 0.1% conversion; mean ± SEM) are consistent with these calculations. Moreover, such concentrations of fatty acid and amino acid reactants are within the physiologic range (Stegink et al., 1991). The energetic driving force of the synthase reaction in vivo is likely to arise from a disequilibrium of the fatty acid and amino acid reactants and their N-acyl amino acid products, analogous to the proton-motive force that drives ATP production by ATP synthase.

An outstanding issue is the molecular target(s) responsible for the uncoupling activity of N-acyl amino acids. The photo-cross-linking experiments provide direct evidence that N-acyl amino acids engage members of the SLC25 family of inner mitochondrial carriers, including ANT1 and ANT2. Notably, the function of this photo-probe reagent requires both diazirine and alkyne “ends” of the molecule. Consequently, it must be the intact photo-probe, and not a hydrolyzed product, that is interacting with proteins. The proton conductance activity of the ANTs, or other SLC carriers, might be directly or allosterically activated by N-acyl amino acid binding. Loss-of-function studies will ultimately be required to definitely establish such a role for these carriers.

In summary, these data identify a new enzymatic node and a class of lipidated metabolites that might be used for the treatment of human obesity and diabetes and to modulate thermogenesis more generally.

EXPERIMENTAL PROCEDURES

Plasmids and Viruses
Full-length mouse Pm20d1 cDNA (GE Dharmaco) was subcloned with an in-frame C-terminal 6xHis and Flag tag into pENTR/D-TOPO (Thermo Fisher Scientific) according to the manufacturer’s instructions. This construct was subsequently cloned into the appropriate expression vectors as described in the Supplemental Experimental Procedures. Full-length C-terminal Flag tagged human Pm20d1 cDNA (Origene) was used directly for transient transfection experiments.

Chemicals
The full inventory of commercially available compounds, purchased from Sigma, Cayman Chemical Company, or Santa Cruz Biotechnology, as well as the synthesis of N-acyl amino acids, is described in the Supplemental Experimental Procedures.

General Animal Information
Animal experiments were performed according to procedures approved by the Beth Israel Deaconess Medical Center IACUC. Mice were maintained in 12 hr light:dark cycles at 22°C and fed a standard irradiated rodent chow diet. All experiments on wild-type mice were performed with male C57BL/6 mice purchased from Jackson Laboratories, except for the AAV experiments, which were performed with male C57BL/6 mice from Charles River. UCP1-KO (stock #017478) and 16- to 20-week DIO mice (stock #380050) were obtained from Jackson. Details of the mouse injections and treatments can be found in the Supplemental Experimental Procedures.

Indirect Calorimetry and Body Composition Measurements
Energy expenditure, O2 consumption, CO2 production, respiratory exchange ratio, total locomotor activity, and food intake measurements were made with a 16-cage Columbus Instruments Oxymax Comprehensive Lab Animal Monitoring System at ambient room temperature (21°C–23°C). Mice were acclimated for 1 day in metabolic cages prior to data collection. Whole-body composition was assessed with an EchoMRI 3-in-1 on conscious mice.

In Vitro Enzymatic Assays
Flag-tagged mouse and human PM20D1 were immunopurified using magnetic Flag-M2 beads (Sigma Aldrich) as detailed in the Supplemental Experimental Procedures. In vitro PM20D1-catalyzed synthesis of N-acyl amino acids was measured by incubating purified PM20D1 protein with oleate (1.5 mM) and phenylalanine (0.1 mM) in PBS (to 100 μl final volume) at 37°C for 1.5 hr. In vitro PM20D1-catalyzed N-acyl amino acid hydrolysis was measured by incubating purified PM20D1 protein with the indicated N-acyl amino acid (0.1 mM) in PBS (to 100 μl final volume) at 37°C for 1.5 hr. Reactions were terminated by placing the vials at −80°C and analyzed as described below by LC-MS.

Measurements of Metabolites In Vivo and Enzyme Activities In Vitro by LC-MS
Frozen serum (30 μl) for polar metabolomic analyses were extracted in 160 μl of 1:1 acetonitrile/methanol with inclusion of internal standard D2,15N-serine (1 nmol). Activity assays (100 μl) were extracted in 400 μl 1:1 acetonitrile/methanol. Following 30 s of thorough vortexing and 1 min of bath sonication, the polar metabolite fraction (supernatant) was isolated by centrifugation at 13,000 x g for 10 min. This supernatant (10 μl) was analyzed by SRM-based targeted LC-MS/MS, or untargeted LC-MS. The Supplemental Experimental Procedures contains details about MS parameters used for these analyses.

Cellular Respiration Measurements
Cellular oxygen consumption rates were determined using an XF24 Extracellular Flux Analyzer (Seahorse Biosciences) as detailed in the Supplemental Experimental Procedures.

Statistics
The Student’s t test was used for pair-wise comparisons, and ANOVA was used for indirect calorimetry experiments. Unless otherwise specified, statistical significance was set at p < 0.05.

SUPPLEMENTAL INFORMATION
Supplemental Information includes Supplemental Experimental Procedures, seven figures, and five tables and can be found with this article online at http://dx.doi.org/10.1016/j.cell.2016.05.071.

AUTHOR CONTRIBUTIONS

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