Adipocyte-specific Nos2 deletion improves insulin resistance and dyslipidemia through brown fat activation in diet-induced obese mice

Vanessa Rodrigues Vilela, Nolwenn Samson, Renato Nachbar, Lia Rossi Perazza, Gabriel Lachance, Volatiana Rokotarivelo, Carolina Centano-Baez, Patricia Zancan, Mauro Sola-Penna, Kerstin Bellmann, Vincenzo Di Marzo, Mathieu Laplante, André Marette

ABSTRACT

Objective: Inducible nitric oxide (NO) synthase (Nos2) is a well-documented inflammatory mediator of insulin resistance in obesity. Nos2 expression is induced in both adipocytes and macrophages within adipose tissue during high-fat (HF)-induced obesity.

Methods: Eight-week-old male mice with adipocyte-selective deletion of the Nos2 gene (Nos2fl/fl) and their wildtype littermates (Nos2fl/+) were subjected to chow or high-fat high-sucrose (HFHS) diet for 10 weeks followed by metabolic phenotyping and determination of brown adipose tissue (BAT) thermogenesis. The direct impact of NO on BAT mitochondrial respiration was also assessed in brown adipocytes.

Results: HFHS-fed Nos2fl/fl mice had improved insulin sensitivity as compared to Nos2fl/+ littermates. Nos2fl/fl mice were also protected from HF-induced dyslipidemia and exhibited increased energy expenditure compared with Nos2fl/+ mice. This was linked to the activation of BAT in HFHS-fed Nos2fl/fl mice as shown by increased Ucp1 and Ucp2 gene expression and augmented respiratory capacity of BAT mitochondria. Furthermore, mitochondrial respiration was inhibited by NO, or upon cytokine-induced Nos2 activation, but improved by Nos2 inhibition in brown adipocytes.

Conclusions: These results demonstrate the key role of adipocyte Nos2 in the development of obesity-linked insulin resistance and dyslipidemia, partly through NO-dependent inhibition of BAT mitochondrial bioenergetics.

Keywords Nitric oxide synthase; Dyslipidemia; Obesity; Insulin resistance; brown adipose tissue; Mitochondrial respiration

1. INTRODUCTION

Nitric oxide (NO) is synthesized by nitric oxide synthase enzymes (NOS). The inducible member of the NOS family, NOS2 (formerly iNOS), is activated in response to bacterial endotoxins, cytokines, and nutrient overload [1]. NOS2 can produce large amounts of NO over prolonged periods of time and plays a key role in host defense through its bacteriostatic action [2]. However, Nos2 is also a key mediator of inflammation and insulin resistance, and chronic NOS2 induction in obesity interferes with insulin signaling to phosphatidylinositol 3-kinase PI3K (PI3K)/Akt in metabolic tissues [3,4]. NOS2 expression is induced in both adipocytes and macrophages within adipose tissue during high-fat (HF)-induced obesity [5,6]. Genetic invalidation of Nos2 protects mice from insulin resistance mediated by obesity, lipid infusion, and lipopolysaccharide (LPS) challenge [4,7–9]. Nos2 is induced in all metabolic tissues during obesity [7,8,10,11] and mostly found in macrophages in adipose tissue [12]. However, myeloid Nos2 deletion did not protect mice from insulin resistance, whereas pharmacological Nos2 inhibition improved insulin action in HF-fed mice [13]. These results suggest that Nos2 expression in metabolic cells is key to disrupt metabolic homeostasis in obesity. To directly test this hypothesis, we generated mice with adipose-specific Nos2 deletion using Cre/loxP recombination. Interestingly, these mice showed improved insulin sensitivity and lipid profile upon HFHS feeding, which were associated with brown adipose tissue (BAT) activation. We further showed using cultured BAT cells that Nos2 activation can directly inhibit mitochondrial oxidative metabolism.

2. METHODS

2.1. Animals and genotyping

Mice containing flox sites spanning Exon 2 of Nos2 were created (Ingenious, Stony Brook, NY, USA), crossed with B6.Cg-Tg (ACT FLPe)

1 Quebec Heart & Lung Institute, Université Laval, 2725 Ch Ste-Foy, Québec, QC, G1V 4G5, Canada 2 Institute of Nutrition and Functional Foods, Centre NUTRISS, Université Laval, 2440 Boulevard Hôtel-Dieu-Saint-Sacrement 1710, Québec, QC, G1V 0A6, Canada 3 Canada Excellence Research Chair Microbiome-Endocannabinoidome Axis in Metabolic Health (CERC-MEND), Canada

4 Vanessa Rodrigues Vilela and Nolwenn Samson contributed equally to this work.

5 Present address: Department of Pharmaceutical Biotechnology, Federal University of Rio de Janeiro, Rio de Janeiro, RJ, 21941-902, Brazil.

*Corresponding author. The Quebec Heart and Lung Institute, Hôpital Laval, Pavillon Marguerite d’Youville Bureau Y4340, Ste-Foy, Québec, G1V 4G5, Canada. E-mail: andre.marette@criucpq.ulaval.ca (A. Marette).

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9205DyN/j mice (Jackson Laboratory, Bar Harbor, ME, USA) to remove the neoeycin cassette and then backcrossed for at least 12 generations on C57BL/6J background (stock No. 000664, Jackson Laboratories). Adipocyte-specific (Nos2fl/−/−, Nos2 knockent mice were generated by crossing floxed Nos2 (Nos2fl/+) mice with Adipoq-Cre (kindly provided by E. Rosen and backcrossed for at least 10 generations on C57BL/6J background). Genomic DNA was extracted from samples using the DNA REDExtract-N-Amp PCR kit (Sigma), and genotyping was performed for floxed Nos2 and Adipoq-Cre. Primers used for floxed Nos2 were 5′-ggC ACC TAA gAg AgA gTCT ggT CT CC-3′ and 5′-CTG AGG ATA GTA GAG GGC CA TC-3′. Adipoq-Cre-specific recombination of Nos2 exon 2 was detected by PCR using primers binding to intron 1 (5′-CTTG6CTCAGGATGCTGAT-3′ (forward)) and intron 2 (5′-TTGACACCTCCAGTTGCTA-3′ (reverse)) genomic DNA outside of the floxed Nos2 sites. Genomic DNA was isolated from mouse tissues. Mapping of the recomined alleles was done by directly cloning PCR products into pGEM-T-easy vector (Promega #A1360) followed by T7p DNA sequencing (Genotyping and Sequencing platform, CHU de Québec-Université Laval Research Center). To determine Nos2 ablation in the adipocyte compartment of white (eWAT, iWAT) and brown (BAT) adipose tissues, mice were injected with LPS (3 mg/kg; i.p.) or saline for 6 h to acutely induce Nos2 expression. Adipose depots were rapidly removed, cleaned of non-esterified fatty-acids (NEFA) (Wako Chemicals, Richmond, VA), cholesteral (Randox Laboratories, Crumlin, UK) and free glycerol levels (Sigma–Alrich) were enzymatically determined according to the manufacturer’s instructions.

2.4. Blood metabolite measurements

Blood glucose was measured by glucometer (Accu-Chek). Plasma insulin was measured using ultrasensitive ELISA kit (Alpco, Salem, USA); C-peptide was determined using ELISA kit (Crystal Chem, Elk Grove Village, USA). Plasma triglyceride (TG) (Randox Laboratories, Crumlin, UK), non-esterified fatty-acids (NEFA) (Wako Chemicals, Richmond, VA), and free glycerol levels (Sigma–Alrich) were enzymatically determined according to the manufacturer’s instructions.

2.5. Measurement of plasma and tissue nitrate and nitrate

Nitrite and nitrate levels in plasma, epidymidal and inguinal adipose tissue were measured by fluorometric spectrophotometry after reducing nitrate to nitrite using nitrite reductase and the NAHD regeneration system [14]. Briefly, blood was collected in tubes containing EDTA and centrifuged for 10 min at 3200 × g to obtain plasma. Tissues were grounded in liquid nitrogen using a pestle and mortar. The tissue powder was resuspended in 5 volumes of Tris buffer (20 mM Tris pH 7.5). Tissue lysates were centrifuged at 2800 × g for 20 min and protein concentration of the supernatant was measured by the BCA™ protein assay (Thermo Scientific, Illinois, USA). Plasma and tissue lysates were then centrifuged at 5000 × g (4 °C) in a Ulntrafree-Microcentrifuge 10000 NMWL filter unit. Fluorescence was measured at λex 360 nm and λem 450 nm.

2.6. Protein extraction and western blotting

Fifty milligram of brown adipose tissue was pulverized with a pestle and mortar in liquid nitrogen, and the powder was homogenized in 4 vol of homogenization buffer (20 mM Tris—HCl pH 7.5, 150 mM NaCl, 1 mM CaCl2, 1 mM MgCl2, 10% glycerol, 1% Igepal CA-630, 10 mM NaF, 2 mM Na3VO4, 1 mM PMSF, and protease inhibitors). For western blot analysis, crude protein lysates were solubilized in sample buffer and loaded onto a 6% acrylamide gel and subjected to SDS-PAGE, transferred to nitrocellulose membrane. Non-specific binding sites were blocked with 3% (w/v) dry nonfat milk in TBS-T (100 mM Tris, 1.5 mM NaCl, pH 8.0 and 0.5% Tween 20) and the membrane was incubated overnight with antibodies against mitochondrial complexes I to V (Abcam, #110413, dilution 1:1000). Secondary antibodies were purchased from Jackson Immuno Research Laboratories and diluted 1:1000. The substrate (WBKLS0500, Millipore) was used to image the blots. Bands were detected and recorded using a Bio–Rad Gel Doc 2000 gel documentation system (Bio–Rad, Hercules, CA, USA). Densitometric analysis was conducted using ImageJ software from the National Institutes of Health.

2.7. RNA analysis

Total RNA was isolated using TRIzol reagent (Thermo Fisher Scientific) and total RNA purification was performed using a RNeasy mini kit (Qiagen). Total RNA was used for reverse transcription using High-Capacity cDNA Reverse Transcription Kit (Thermo Fisher Scientific) and cDNA levels were measured by quantitative PCR as described before (27) or by qPCR-based TaqMan® open array (ThermoFisher). Primers are listed in Supplementary Tables 3 and 4.

Insulin tolerance was determined in 6-h-fasted mice after 8 weeks on diet. Insulin (0.65 U/kg BW) was injected intraperitoneally before blood collection via the saphenous vein at 0, 5, 10, 15, 20, 30, and 60 min. For glucose tolerance tests (performed 10 weeks post-diet treatment), 12-h-fasted mice were given glucose (1 g of glucose/kg body weight) by gavage, and blood was drawn from the saphenous vein at 0, 15, 30, 60, 90, and 120 min. Additionally, blood samples (~ 30 μL) were collected at each time point during oGTT for the determination of insulin concentration.
2.8. Cell culture
T37i cells were cultured in DMEM/Ham’s F12 medium supplemented with glutamine (2 mM), Hepes (15 mM) and FBS (10%). Confluent cells were differentiated for 7 days in the presence of insulin (20 nM) and triiodothyronine (2 nM).

2.9. Oxygen consumption
Differentiated T37i cells were seeded in Seahorse 24 well plate at a density of 40000/well and treated for 24 h with a cytokine cocktail (625 U/ml IFNγ, 2.5 ng IL-1β, 0.625 ng/ml TNF-α) with or without 1400W (100 μM). After washing the cells thrice with Seahorse XF Medium (DMEM w/o phenol red supplemented with pyruvate (1 mM), glutamine (2 mM) and glucose (10 mM)), the Seahorse mitostress protocol was run. Oxygen consumption was monitored by sequential addition of Oligomycin (1.5 μM), FCCP (1.5 μM), Rotenone (1 μM) and Antimycin A (1 μM). Oligomycin inhibits the activity of the ATP synthase (complex V) and provides information about the ATP-linked cellular respiration. FCCP triggers the disruption of the mitochondrial membrane potential. Under this condition, the cells can reach their maximal oxygen consumption through complex IV. Finally, the addition of Rotenone and Antimycin A, which inhibit the complex I and III, respectively, shuts down mitochondrial respiration to determine non-mitochondrial consumption.

Figure 1: Determination of selective disruption of Nos2 in adipose tissue. Mice containing flox sites before and after Exon 2 of Nos2 were created and backcrossed for at least 10 generations on a C57BL/6J background. Adipocyte-specific (Nos2lox/−) Nos2 knockout mice were generated on a C57BL/6J background by crossing mice homozygous for floxed Nos2 (Nos2fl/k0) with Adipoq-Cre mice (A). Recombined alleles from genomic DNA isolation (B). *: unspecific band. Δ: deletion. Nos2 expression in adipocytes isolated from epididymal (C), inguinal (D) and brown (E) adipose tissue of saline- and LPS-injected mice; N = 2—3.
Figure 2: Nos2−/− KO prevents HFHS diet increase in brown adipose tissue mass irrespectively of body weight gain. Nos2−/− KO mice and wild-type (Nos2+/+) mice were fed either a chow or a high-fat/high sucrose (HFHS) diet for 10 weeks. Total weight gain (A), total energy intake during 10 weeks (B), weight of brown adipose tissue (interscapular) (C). Nos2 mRNA expression in epididymal (D), inguinal (E) and brown (F) adipose tissue. Nitrite/nitrate content in epididymal (C), inguinal (D) adipose tissue and plasma (E). n = 11 – 12 mice/group. Two-way ANOVA with a Student-Newman-Keuls post hoc test was applied to calculate the significance of the differences between groups. Data are expressed as the mean ± SEM. ***p < 0.001 for diet effect and #p < 0.05 for genotype difference.
Figure 3: Adipocyte-specific knockout of Nos2 increases insulin sensitivity and alleviates diet-induced hyperinsulinaemia in HFHS-fed mice. 12 h fasting glycemia (A), insulinemia (B) and C-peptide (C). Mice 10 wk on HFHS diet were fasted overnight (12 h) and an oral glucose tolerance test (D) was performed after gavage with glucose (1 g/kg body weight). Blood samples were collected at each time point during OGTT for insulinaemia and C-peptide determination (E, F). Mice 8 wk on HFHS diet were 6 h fasted and an insulin tolerance test (E) was performed after an intraperitoneal injection of insulin (0.65 UI/kg body weight). n = 11–12 (A, D, E); n = 8–11 (B); n = 7–11 (G), n = 3–6 (C, F).

Two-way ANOVA with a Student-Newman-Keuls post hoc test was applied to calculate the significance of the differences between groups. Data are expressed as the mean ± SEM.

* p < 0.05, ** p < 0.01 and *** p < 0.001 for diet effect; # p < 0.05 and ## p < 0.01 for genotype difference. HFHS, high fat/high sucrose; oGTT, oral glucose tolerance test; ipITT, intraperitoneal insulin tolerance test.
respiration. The OCR measurements were normalized to total DNA content.

2.10. Statistical analysis
All data are presented as means ± SEM. Two-way ANOVA with a Student-Newman-Keuls post hoc test was used to assign significance to the comparisons between groups (GraphPad Software 7, La Jolla, CA, USA). All results were considered statistically significant at \( P < 0.05 \).

3. RESULTS

3.1. Adipocyte-specific ablation of Nos2 reduces HFHS diet-induced brown adipose tissue mass
To determine the specific role of Nos2 in adipose tissue, we generated adipocyte-specific (Nos2ADKO) Nos2 knockout mice by crossing homozygous Nos2ADKO mice containing a floxed exon2 with adiponectin-Cre (Adipoq-Cre) mice, respectively (Figure 1A). Nos2fl/fl and Nos2ADKO animals were born at the expected Mendelian ratio with no morphological differences. To assess the specificity of the adiponectin-Cre driven recombination of the Nos2 floxed alleles, genomic DNA was isolated from different tissues. The site of recombination on the Nos2 gene was mapped, and adiponectin-Cre driven excision of exon2 was confirmed (Figure 1A). Finally, we tested the specificity of the deletion by comparing the rate of Nos2 exon 2 recombination in different adipose tissues, liver and skeletal muscle. We observed the deleted Nos2 (ΔNos2) alleles in all white and brown adipose tissue depots of Nos2ADKO animals but not in muscle and liver (Figure 1B) demonstrating that Nos2 deletion is efficiently working and specific for adipose tissue. Isolation of adipose cells from eWAT, iWAT, and BAT of saline- and LPS-injected mice were also performed to demonstrate a marked reduction of LPS-induced Nos2 expression in white and brown adipocytes from Nos2ADKO mice (Figure 1C–E).

Adipocyte-selective Nos2 knockout did not impact normal adipose tissue function. Indeed, we found no change in the plasma levels of adiponectin and leptin, two key adipokines. No genotypic differences were observed for body composition, iWAT, eWAT and BAT histology in mice that were fed regular chow before the start of the dietary intervention (Supplementary Fig. 1).

Nos2ADKO mice were not protected from HFHS-diet induced obesity, gained the same amount of weight, and had similar energy intake as compared to their littermate controls (Figure 2A,B). The weight of white fat masses as well as other tissues were not different between Nos2ADKO and Nos2fl/fl mice (Supplementary Table 1). Interestingly, BAT weight was significantly lower in HFHS-fed Nos2ADKO mice as compared to HFHS-fed Nos2fl/fl mice (Figure 2C). Nos2 expression was significantly reduced in BAT of HFHS fed Nos2ADKO mice, but not in visceral and subcutaneous adipose tissue (Figure 2D–F). The residual Nos2 expression in adipose tissue from Nos2ADKO mice may represent its undeleted enzyme expression in macrophages and other immune cells, which are the predominant cellular sites of Nos2 expression in adipose tissue [12]. Accordingly, Nos2 deletion in adipocytes did not affect the typical inflammatory features of epididymal and brown adipose tissues in obese mice, as revealed by the lack of changes in Adgre1 expression (encoding the F4/80 antigen) as well as M1 and M2 activation markers between HFHS-fed Nos2ADKO and Nos2fl/fl mice (Supplementary Figs. 2A and C). On the other hand, we found a tendency for increased Adgre1 expression in iWAT of Nos2ADKO mice on HFHS diet compared to Nos2fl/fl mice (Supplementary Fig. 2B), suggesting a partial compensatory increase in macrophage infiltration in this fat depot in mice lacking adipocyte Nos2.

3.2. HFHS-fed Nos2ADKO mice showed increased insulin sensitivity and reduced plasma triglycerides
We next determined the impact of adipocyte-specific Nos2 disruption on whole body glucose homeostasis. Fasting blood glucose and C-peptide were not different, but fasting insulinaemia was reduced in HFHS-fed Nos2ADKO mice as compared to HFHS Nos2fl/fl littermates (Figure 3A–C). Oral glucose tolerance tests (oGTT) showed similar glucose excursion curves between Nos2ADKO and HFHS Nos2fl/fl mice on either diets (Figure. 3D). However, glucose-stimulated insulin response during the oGTT was significantly improved in HFHS-fed Nos2ADKO mice versus HFHS-fed Nos2fl/fl further suggesting that adipocyte Nos2 disruption improved insulin sensitivity in obese animals (Figure 3E). C-peptide release was slightly different at 15 min between the HFHS fed groups although there was no significant difference in C-peptide secretion from 30 to 120 min when the insulin responses were markedly different between the genotypes (Figure 3E–F). This suggests that the reduced insulin levels in Nos2ADKO mice on HFHS diet demonstrate an improvement in insulin sensitivity in these mice and/or higher insulin clearance. The improvement of insulin sensitivity was more directly confirmed by the finding of improved insulin tolerance in HFHS fed Nos2ADKO mice compared to their HFHS-fed Nos2fl/fl counterparts (Figure 3G).

HFHS-fed Nos2ADKO mice also exhibited an improved lipid profile as compared to HFHS-fed Nos2fl/fl controls as revealed by reduced plasma levels of triglycerides, cholesterol, and non-esterified fatty acids (Figure 4A–C). Plasma glycerol levels were also reduced by adipocyte Nos2 ablation but only reached statistical significance in the chow-fed animals (Figure 4D). TG levels in inguinal adipose tissue were increased in Nos2ADKO mice on both diets compared to their littermate controls (Figure 4E), but no genotypic differences were observed in epididymal adipose tissue (Figure 4F). These results together with lower levels of NEFA and a slight reduction in glycerol levels might suggest reduced lipolysis in iWAT. Importantly, neither insulin sensitivity nor plasma TG levels or tissue weights in HFHS-fed mice were affected by adiponectin-Cre expression per se (see Supplementary Fig. S3 and Supplementary Table 2) validating that the metabolic benefits seen in HFHS-fed Nos2ADKO mice are directly related to selective Nos2 ablation in adipocytes.

3.3. Adipocyte-specific Nos2 ablation increases energy expenditure and BAT activation in HFHS-fed mice
Indirect calorimetry measurements in metabolic cages revealed reduced oxygen consumption in HFHS-fed Nos2fl/fl as compared to the chow-fed counterparts, but this altered phenotype was fully restored in HFHS-fed Nos2ADKO mice (Figure 5A) suggesting that adipose tissue Nos2 is responsible for reduced energy expenditure in obese animals. No differences were observed in CO2 production, ambulatory movements or the respiratory exchange ratio between HFHS-fed Nos2ADKO and Nos2fl/fl mice (Figure 5B–D).

Given the changes in energy expenditure and reduction of BAT weight suggesting the activation of the tissue in HFHS-fed Nos2ADKO mice, we next sought to further investigate the potential role of BAT function in improved insulin sensitivity and lipid homeostasis in this model. Remarkably, we found that adipocyte Nos2 ablation markedly raised uncoupling protein-1 (Ucp1) expression in BAT of HFHS-fed mice (Figure 6A). We also found a similar increased Ucp2 expression in BAT of HFHS-fed Nos2ADKO mice compared with HFHS-fed Nos2fl/fl mice.

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Expression of other genes implicated in energy metabolism in BAT such as peroxisome proliferator-activated receptor alpha (Ppara), thermogenic gene type II iodothyronine deiodinase (Dio2), G-protein coupled bile acid receptor 1 (Gpbar1) and peroxisome proliferative activated receptor, gamma, coactivator 1-alpha (Ppargc1a) were not affected by Nos2 deletion (Figure 6A). Total mitochondrial expression of OXPHOS complexes was higher in BAT of HFHS-fed Nos2\(^{AD-C0}\) KO mice compared to their HFHS-fed Nos2\(^{fl/fl}\) littermates (Figure 6C) and these differences were related to specific increases in the levels of complexes II, III and V (Figure 6B). Interestingly, the gene expression of lipoprotein lipase (Lpl), a key enzyme controlling the hydrolysis of TG-rich particles, increased in Nos2\(^{AD-C0}\) KO animals suggesting a rise in free fatty acid (FFA) delivery to BAT, also consistent with the downregulation of angiopoietin-like 4 (Angptl4) in BAT of HFHS-fed Nos2\(^{AD-KO}\) and HFHS-fed Nos2\(^{fl/fl}\) mice. We also examined whether Nos2 deletion may induce browning of white adipose tissues. While no changes were observed in inguinal fat (Figure 7A), we found that Nos2 deletion in adipocytes reduced Lpl expression in epididymal fat, which reached the level of significance in chow-fed animals (Figure 7B). However, unlike the findings in BAT, no changes were observed in the expression of Angptl4 in visceral fat of HFHS-fed Nos2\(^{AD-KO}\) and HFHS-fed Nos2\(^{fl/fl}\) mice.

3.4. Nitric oxide directly impairs mitochondrial oxygen consumption in brown adipose cells

To confirm that NOS2 controls BAT thermogenesis, we next used T37i brown adipocytes in culture to investigate whether NOS2 induction directly modulates mitochondrial function in these cells using the Seahorse system. NOS2 expression was induced by incubating brown adipocytes with cytokines as previously described [15]. NOS2 activation and NO production was confirmed by accumulation of nitrite during cytokine exposure and was completely blocked when using the selective NOS2 inhibitor 1400W (Figure 8A). Seahorse analysis revealed that oxygen consumption rate (OCR) was strongly decreased by cytokine treatment as compared to untreated cells but that selective inhibition of NOS2 activity with 1400W completely reversed the defective mitochondrial respiration (Figure 8B). Addition of different doses of the NO donor SNP confirmed that exogenous NO...
These results confirm that NOS2 is a cell-autonomous regulator of mitochondrial respiration in BAT, thus providing evidence that NOS2 ablation in BAT is a key mechanism underlying improved energy expenditure in HFHS-fed Nos2AD/C0 KO mice.

4. DISCUSSION

We have previously shown that whole body Nos2 deletion protects mice from high-fat diet-induced insulin resistance [8]. Nevertheless, the target tissue mediating the beneficial effects of Nos2 has not yet been identified. Here, we evaluated the metabolic effects of a specific disruption of Nos2 in adipose tissue of chow- and HFHS-fed mice. Adipocyte Nos2 disruption improved HFHS-induced insulin resistance and dyslipidemia. These effects, which were independent of the changes in body weight gain, were accompanied by a decrease in fat accumulation in BAT and a rise in Ucp1 expression and oxygen consumption in BAT. NOS2 induction in BAT cells impaired oxygen consumption, which was completely restored when NOS2 activity was inhibited, but mimicked by exogenous NO addition.

We and others have demonstrated the effect of whole body Nos2 deletion in obese and lean mice using different mouse models [8,10,13,16–20]. Importantly, improvement in insulin sensitivity was observed in whole body Nos2 KO mice treated for 12–18 weeks with HFD [10,20] or in obese ob/ob mice lacking Nos2 [21]. Whole body Nos2 KO mice are also protected from insulin resistance during lipid infusion as compared to wild-type mice [4]. Pharmacologic inhibition of NOS2 using L-NIL also improved fasting hyperglycemia and improved insulin sensitivity in obese diabetic ob/ob mice [18]. Although induction of NOS2 expression is highest in macrophages, Lu et al. [13] showed that specific Nos2 ablation in myeloid cells did not protect from insulin resistance, suggesting that non-myeloid cells or tissues are responsible for the beneficial effects of Nos2 invalidation on insulin sensitivity.

The present study reveals a key role of adipocyte NOS2 in mediating insulin resistance and some metabolic alterations in obesity since specific ablation of Nos2 in adipocytes is sufficient to ameliorate insulin sensitivity as well as the lipid profile. However, some reported effects of systemic Nos2 deletion were not observed in Nos2AD/C0 KO mice. In addition to modulating insulin sensitivity, we have reported that whole-body invalidation of Nos2 in mice led to hyperphagia [8], which was not observed in the current study, suggesting that Nos2 in the central nervous system or other tissues may control food intake.

One major finding of this study is that NOS2 in BAT plays a key role in metabolic homeostasis. Brown adipose tissue is a unique organ that uses UCP1 to uncouple oxidative phosphorylation from ATP production to produce heat. The discovery of a functional BAT in human has greatly contributed to stimulate the interest in this tissue over the last decade.

Even though most human brown fat deports disappear after infancy, a recent study using PET-scan has shown that these fat deports amount for 1.5% of total body mass corresponding to 4.3% of total fat mass. More importantly, up to 90% of human adult BAT could be activated BAT [22]. Also, the function of UCP1 in mitochondria of BAT in human.

**Figure 5**: Adipocyte-specific knockout of Nos2 increases oxygen consumption in HFHS-fed mice. Oxygen consumption (A), respiratory exchange ratio (B), total ambulatory movement (C) and carbon dioxide production (D) were measured by indirect calorimetry. Two-way ANOVA with a Student-Newman-Keuls post hoc test was applied to calculate the significance of the differences between groups. Data are expressed as the mean ± SEM. n = 9–12 (A,F), n = 5–6 (B,C), n = 10–12 (D,E). *p < 0.05 and **p < 0.001 for diet effect; *p < 0.05 and #p < 0.01 for genotype difference.
Figure 6: Adipocyte-specific knockout of Nos2 increases BAT activation in HFHS-fed mice. BAT mRNA expression of genes related to energy metabolism: uncoupling protein 1 (Ucp1), uncoupling protein 2 (Ucp2), peroxisome proliferator-activated receptor alpha (Ppara), type 2 iodothyronine deiodinase (Dio2), G-coupled protein receptor (Tgr5) and peroxisome proliferator-activated receptor-gamma coactivator (Ppargc1a) (A). Protein expression of total OXPHOS complexes (B) and complexes I-IV (C) in BAT. BAT mRNA expression of genes related to lipid metabolism: Lipoprotein lipase (Lpl) and Fasting-induced adipose factor/Angiopoietin-like 4 (Angptl4) (D). Two-way ANOVA with a Student-Newman-Keuls post hoc test was applied to calculate the significance of the differences between groups. Data are expressed as mean ± SEM. n = 9–12. *p < 0.05, **p < 0.01 and ***p < 0.001 for diet effect; #p < 0.05, ##p < 0.01 and ###p < 0.001 for genotype difference.
Figure 7: Nos2 ablation did not change the expression of genes related to browning effect. Genes expressed in energy metabolism, lipid handling and browning in subcutaneous (A) and visceral adipose tissue (B). Tissues were extracted from Nos2<sup>−/−</sup> and wild-type (Nos2<sup>fl/fl</sup>) mice after 6 h of fasting during sacrifice. Two-way ANOVA with a Student-Newman-Keuls post hoc test was applied to calculate the significance of the differences between groups. Data are expressed as the mean ± SEM. n = 9–12. *p < 0.05 and ***p < 0.001 for diet effect; #p < 0.05 for genotype difference.
Figure 8: Nitric oxide affects oxygen consumption in brown adipose cells. Differentiated T37i cells were exposed to cytokine cocktail with or without 1400W (100 μM) for 24 h. Nitrite was determined in the supernatant (A) and oxygen consumption rate (OCR) was determined by running the Seahorse mitostress protocol (B). T37i cells were exposed to different concentrations of sodium nitroprusside (SNP) for 16 h and OCR was determined as in B (C). Data are expressed as the mean ± SEM, n = 3 (A, B); n = 2 (C).
and mouse is comparable [23], highlighting the importance of studies using mouse models. We found a robust increase in Ucp1 gene expression in BAT of Nos2\(^{-}\text{ko}\) mice. Becerril and colleagues [21] also showed UCP1 activation in whole body Nos2 KO mice although they also observed a reduction in food intake, which is a potential confounding factor. It is known that increased BAT thermogenesis is tightly correlated with increased UCP1 expression in BAT [24]. Interestingly, Nos2 deletion in brown adipocytes also increased Ucp2 gene expression in BAT of Nos2\(^{-}\text{ko}\) mice, which could also be involved in both, the raised energy expenditure and improved lipid profile of these mice. Indeed, UCP2 is also involved in cold-induced BAT thermogenesis [25] whereas the lack of UCP2 also led to higher TG concentration in BAT favoring glucose utilization [25]. It has also been shown that BAT takes up plasma TG via lipolysis [26,27], which is well aligned with the increase in Lpl expression we have observed here and the reduction in plasma TG, which is also in accordance with the effect of adipocyte Nos2 ablation on Angpt14, a well known LPL inhibitor.

Although BAT activation in mice lacking adipocyte Nos2 was associated with increased energy expenditure and improved glucose and lipid metabolism within 7–10 weeks of dietary treatments, we did not observe body weight reduction in Nos2\(^{-}\text{ko}\) mice as compared to their WT flexed littermates. It is possible that longer term changes in BAT thermogenesis and energy metabolism are needed to translate into significant body weight change in mice. This is supported by the previous findings that BAT transplantation improves glucose homeostasis and insulin sensitivity within 8 weeks, but that body weight changes became only significant after 12 weeks in these mice [28]. Further analysis of BAT in our study also points toward the role of Nos2 in regulating mitochondrial capacity of brown adipocytes in Nos2\(^{-}\text{ko}\) mice, as indicated by increased levels of mitochondrial complex II, III, and IV protein expression, which suggests more demand for these complexes in the absence of Nos2 in BAT. This mechanism was confirmed using T37i brown fat cells whereby Nos2 induction was shown to directly impair mitochondrial respiration, and it was completely reversed by inhibiting NOS2 activity. Moreover, direct addition of exogenous NO dose dependently impaired oxygen consumption rate in these cells, demonstrating that Nos2-derived NO is a key role for mitochondrial function and decreases lipoxidation of these mitochondria.

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**Conflict of Interest**

The authors declare no competing financial interests. AM has been the holder of Pfizer/CIHR partnered research chair and received grants from Danone Nutricia Research, Acasti Pharma, Allysta inc. None of these funding sources are relevant to this publication. No other potential dualities of interest are associated with this article.

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**Supplementary data**

Supplementary data to this article can be found online at https://doi.org/10.1016/j.molmet.2022.101437.

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