Increased Sirt1 secreted from visceral white adipose tissue is associated with improved glucose tolerance in obese Nrf2-deficient mice

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ABSTRACT

Obesity is associated with metabolic dysregulation characterized by insulin resistance and glucose intolerance. Nuclear factor E2-related factor (Nrf2) is a critical regulator of the stress response and Nrf2-deficient mice (Nrf2−/−) are protected against high fat diet (HFD)-induced metabolic derangement. We searched for factors that could underline this favorable phenotype and found that Nrf2−/− mice exhibit higher circulating levels of sirtuin 1 (Sirt1), a key player in cellular homeostasis and energy metabolism, compared to wild-type mice. Increased Sirt1 levels in Nrf2−/− mice were found not only in animals under standard diet but also following HFD. Interestingly, we report here that the visceral adipose tissue (eWAT) is the sole source of increased Sirt1 protein in plasma. eWAT and other fat depots displayed enhanced adipocytes lipolysis, increased fatty acid oxidation and glycolysis, suggesting autocrine and endocrine actions of Sirt1 in this model. We further demonstrate that removal of eWAT completely abolishes the increase in circulating Sirt1 and that this procedure suppresses the beneficial effect of Nrf2 deficiency on glucose tolerance, but not insulin sensitivity, following a HFD regime. Thus, in contrast to many other stressful conditions where Nrf2 deficiency exacerbates damage, our study indicates that up-regulation of Sirt1 levels specifically in the visceral adipose tissue of Nrf2−/− mice is a key adaptive mechanism that mitigates glucose intolerance induced by nutritional stress.

1. Introduction

Obesity and associated metabolic dysfunction constitute a great threat for global health and the study of mechanisms implicated in these conditions may help to identify potential therapeutic targets. The white adipose tissue (WAT) is crucial in obesity due its fat storing capacity as well as its endocrine function, promoted by the secretion of several factors and hormones that influence whole body metabolism [1]. For example, WAT releases leptin and adiponectin to control appetite and insulin sensitivity, and produces inflammatory factors such as interleukin-6 (IL-6) and tumor necrosis factor-α (TNF-α) that are involved in the metabolic complications of obesity. These and other observations led to a shift in the perception of WAT from that of an inert tissue to that of a highly plastic organ that can be manipulated to provide beneficial effects in obesity. In line with this, we have recently demonstrated that a switch in adipose tissue metabolism towards glycolysis induced by a carbon monoxide-releasing molecule (CORM-401) reduces body weight gain and improves insulin resistance in HFD-fed mice [2]. In addition, stimulation of brown-like features in WAT, based on the idea that brown fat augments energy expenditure as opposed to the energy storage function of WAT, has been reported. Treatment with fibroblast growth factor 21 (FGF21), irisin or the over-expression of sirtuin 1 (Sirt1), were shown to confer ‘browning’ characteristics to WAT and exert beneficial effects on insulin and glucose tolerance [3]. In particular, Sirt1 is a NAD+-dependent protein deacetylase that plays a crucial role in controlling cellular metabolism as well as the energy status of the organism, and has been shown to protect mice against HFD-induced metabolic dysfunction [4]. Additional key roles of

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Sirt1 include inhibition of adipogenesis and increased free fatty acids utilization in WAT [5], while in the liver Sirt1 enhances gluconeogenesis and inhibits glycosylation via forkhead box protein O1 (FOXO1) [6]. Overexpression of Sirt1 also ameliorates hepatic steatosis and glucose intolerance in obese mice by inducing FGFR21 [7].

Previous studies have highlighted a crosstalk between Sirt1 and nuclear factor erythroid 2-related factor 2 (Nrf2), the transcription factor that activates the response to stress conditions by regulating the expression of a variety of detoxifying and antioxidant genes including NAD(P)H-quione oxidoreductase 1 (NQO1), glutathione S-transferases (GSTs) and heme oxygenase-1 (HO-1) [8–11]. Besides its well-described cytoprotective activities against tissue damage triggered by oxidative stress and inflammation [12–14], the possible implications of Nrf2 in the regulation of metabolism started to emerge only recently [15]. However, whether Nrf2 plays a protective or deleterious role in obesity is unclear as divergent data have been published so far. In fact, pharmacologic induction of Nrf2 by the synthetic triterpenoid CDDO-imidazolide prevents body weight gain, adipogenesis and hepatic lipid accumulation in mice [16]. Activation of Nrf2 also improves leptin regulation of metabolism started to emerge only recently [15]. How
timetolerant in obese mice by inducing FGF21 [7].

Nrf2
generated from the TAAM laboratory (Orlés, France). To take in
consideration that mice were from different suppliers, even if from the
same genetic background, mice from both groups were received at the
age of 4 weeks, put into quarantine for a 4 weeks period in order to
acclimatize them to our animal facility. Thus, the same environment and
nutritional states were applied prior to the beginning of our protocols.
Mice were housed under controlled conditions of temperature (21 ± 1 °C), hygrometry (60 ± 10%) and lighting (12 h per day). Animals were
acclimatized in the laboratory for one week before the start of the
experiments. Mice were fed either a standard diet SD (A04, SAFE Diet,
Augs, France) or a HFD (60 kcal % fat, Brogaarden, DK). All animals
received care according to institutional guidelines, and all experiments
were approved by the Institutional Ethics committee (licence number
16–090). During follow-up, animals underwent body weight, food
intake, metabolic and echocardiography assessments. For tissue collec-
tion, mice were euthanatized 6 h after fasting and organs and blood were
collected and processed for further evaluation.

2. Materials and methods

2.1. Animals

Wild type (WT) C57/BL6J mice (8 weeks old) were purchased from
Janvier Labs (France). Nrf2−/− mice (8 weeks old), which had been
backcrossed with C57/BL6J mice and kept as homozygous, were pur-
chased from the TAAM laboratory (Orlés, France). To take in
consideration that mice were from different suppliers, even if from the
same genetic background, mice from both groups were received at the
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collected and processed for further evaluation.

2.2. Fasting blood glucose, glucose and insulin tolerance tests

Whole-body glucose tolerance and insulin sensitivity were assessed in
all groups at weeks 12th and 13th by intraperitoneal glucose (GTT) and
insulin (ITT) tolerance tests, respectively. First, blood was collected via
tail clip to measure fasting blood glucose (Caresens® N, DinnoSanteTM).
Then, mice received glucose (1.5 g/kg) or insulin (0.3 UI/kg) in solution
by intraperitoneal injection and blood glucose was measured at 15, 30,
60, 90 and 120 min after the injection. The HOMEostasis Model
Assessment of Insulin Resistance (HOMA-IR) adjusted to rodents was
calculated as ((glucose (mg/dl)/18) x [insulin (ng/ml)/0.347])/108.24 as
depicted in [24]. The area under the curve (AUC) for the glucose
excursion was calculated using Graph Prism.

2.3. Body composition

Body composition (fat mass, free water, and total water) was measured
by magnetic resonance imaging (EchoMRI-900TM, Echo Medical System,
Houston, TX) as previously described [25,26]. Conscious mice were
placed in a thin-walled plastic cylinder with a plastic insert that func-
tioned to limit the movement while the mice were in the EchoMRI in-
strument. Body composition analysis was measured in mice exposed to a
low-intensity electromagnetic field and the amount of heat and lean mass
detected was expressed as a percentage of total body weight.

2.4. Removal of visceral adipose tissue (eWATectomy)

A total of 10 male WT C57/BL6J mice (WT-eWAT) and 10 male
Nrf2−/− C57/BL6J mice (Nrf2-eWAT), 10 weeks of age and weighing
approximately 25–30 g, were used for adipose tissue removal performed
by epididymal fat resection (eWATectomy). Before eWATectomy, mice
underwent baseline assessment of metabolic functions (glucose toler-
ance and insulin tolerance test). Mice were anesthetized with inhaled
isoflurane and a 1-cm single abdominal midline incision was made.
Bilateral epididymal fat pads were lifted from the peritoneal cavity onto
a sterilized and humidified surgical drape, dissected with an electronic
scalpel, and removed without damaging the testicular blood supply.
The sham operation was performed in the same manner without fat pads
removal. The abdominal peritoneum was closed with prolyne sutures
and the skin was closed with silk sutures (Angiotech, Canada). After
eWATectomy, mice were subjected to a 8 week HFD regime. GTT and
ITT tests were performed at week 7 and mice sacrificed at the end of the
protocol for tissue analysis.

2.5. Preparation of eWAT-conditioned medium (eCM)

Epididymal white adipose tissue (eWAT, 0.1 g) was collected and
kept at room temperature in a 24-well plate with 1 ml/well of Dulbec-
cos’s modified Eagle’s medium DMEM (Life Technologies, France).
The tissue was minced into ~1 mm2 pieces and incubated for 1 h at 37 °C and
5% CO2 prior to transferring it into a new plate with fresh DMEM me-
dium containing 4.5 g glucose, 2 mM glutamine, 1% free fatty acid
bovine serum albumin, 1% antibiotic and antimycotic solution (Sigma
Aldrich). eWAT-conditioned medium (eCM) was collected 24 h after
incubation and stored at ~80 °C for further analysis.

2.6. Plasma and eCM analysis

Enzyme-linked immunosorbent assay (ELISA) kits were used to measure insulin (ALPCO Diagnostics, Salem, NH), FGFR21 (R&D Systems,
with Bruker AV-III console operating at 600.13 MHz. All spectra were acquired in 3 mm NMR tubes using a Bruker 5 mm QCI cryogenically cooled NMR probe. Plasma samples were prepared and analyzed according to the Bruker In-Vitro Diagnostics research (IVDr) protocol. Sample preparation consisted of combining 50 μl of plasma with 150 μl of buffer supplied by Bruker Biospin specifically for the IVDr protocol. For 1D 1H NMR, data was acquired using the 1D-NOE experiment, which filters NMR signals associated with broad line widths arising from proteins that might be present in plasma samples that adversely affect spectral quality. Experiment conditions included: sample temperature of 310K, 96k data points, 20 ppm sweep width, a recycle delay of 4 s, a mixing time of 150 ms, and 32 scans. Lipoprotein subclass analysis was performed using regression analysis of the NMR data which is done automatically as part of the IVDr platform as previously described [27].

2.7. Analysis of plasma lipids and metabolites

Following an 8 h fast, a blood sample was obtained via the orbital sinus under isoflurane anesthesia for analysis of plasma lipids and metabolites. Nuclear magnetic resonance NMR spectroscopy analysis of plasma samples was performed using a 14.0 T Bruker magnet equipped with a Bruker AV-III console operating at 600.13 MHz. All spectra were acquired in 3 mm NMR tubes using a Bruker 5 mm QCI cryogenically cooled NMR probe. Plasma samples were prepared and analyzed according to the Bruker In-Vitro Diagnostics research (IVDr) protocol. Sample preparation consisted of combining 50 μl of plasma with 150 μl of buffer supplied by Bruker Biospin specifically for the IVDr protocol. For 1D 1H NMR, data was acquired using the 1D-NOE experiment, which filters NMR signals associated with broad line widths arising from proteins that might be present in plasma samples that adversely affect spectral quality. Experiment conditions included: sample temperature of 310K, 96k data points, 20 ppm sweep width, a recycle delay of 4 s, a mixing time of 150 ms, and 32 scans. Lipoprotein subclass analysis was performed using regression analysis of the NMR data which is done automatically as part of the IVDr platform as previously described [27].

2.8. Analysis of mRNA expression

After an initial extraction step by mixing Extract-All (Eurobio, France) and chloroform to samples, total RNA purification was performed with a column extraction Kit (RNeasy Mini®, Qiagen, Germany). Double-strand cDNA was synthesized from total RNA with the High-Capacity cDNA Reverse Transcription Kit (Life Technologies, Carlsbad, CA). Quantitative real-time PCR (qPCR) was performed in a StepOnePlus Real-Time PCR System using commercially available TaqMan primer-probe sets (Life Technologies, Carlsbad, CA). Gene expression was assessed by the comparative CT (ΔΔCT) method with β-actin as the reference gene.

2.9. Western blot analysis

Snap-frozen eWAT samples (200 mg) were lysed in cell lysis buffer (Cell Signaling, Danvers, MA France) supplemented with 1% phenylmethylsulfonyl fluoride (PMSF). Protein samples were resolved on 12% bis-Tris gels followed by transfer to nitrocellulose membrane. Antibodies for AMPK (#2532S), AMPK-P (#2535), Foxo1 (#2880) were from Santa Cruz. Bands were visualized by enhanced chemiluminescence and quantified using ImageJ software.

2.10. Histology and immunohistochemistry

Fresh visceral adipose tissue (eWAT) was fixed in 10% phosphate-buffered formalin overnight. Paraffin wax sections of 5 μm were processed for immunostaining. Hematoxylin-eosin (H&E) stained images from mice were analyzed using Adiposoft software. Perilipin antibodies (#9349) were obtained from Cell Signaling, Sir1 antibodies (GTX17532) were from GeneTex Inc. and CD68 antibodies (ab125512) were from Abcam. Electron microscopy (EM) investigation was performed on eWAT sections fixed with glutaraldehyde and stained with osmium tetroxide. Primary adipocytes from eWAT samples were stained with Mitotracker green (M7514) from Invitrogen. 3D-Z stack of images were acquired using confocal microscope Zeiss LSM510 and analysis was performed using Fiji software.

2.11. Pre-adipocytes isolation and culture

Pre-adipocytes were isolated from eWAT obtained from WT or Nrf2−/− mice. Fat pads were placed in Hanks Balanced Salt Solution (HBSS) medium (Life Technologies) under a tissue culture hood, minced into ~1 mm3 pieces and transferred into fresh digestion medium containing 70% HBSS, 30% Bovine Serum Albumin (Sigma Aldrich) and 1 mg/mL collagenase type II (Life Technologies). Samples were incubated at 37°C for 1 h under shaking (120 rpm). The resulting material was filtered through 100 μm mesh and flushed with stop digestion medium collagenase-free. Adipocytes, pre-adipocytes and stromavascular components were separated by centrifugation at 1500 rpm for 5 min at room temperature. The pre-adipocytes were collected and transferred into a new tube with 1:1 v/v DMEM growth media containing 10% new born calf serum, 1% penicillin/streptomycin and 2.4 nM insulin. Pre-adipocytes were centrifuged at 1500 rpm for 5 min at room temperature, medium was removed and the pre-adipocyte pellet was washed with 10 mL growth medium. This washing step was repeated twice. The pre-adipocytes pellet was resuspended, plated in 12 well plate and cultured for 5-6 days for differentiation at 37°C in 5% CO2.

2.12. Preparation of stromal and adipocyte fractions

Stromal cells and mature adipocytes were isolated from epididymal adipose tissue from WT and Nrf2−/− male mice. Adipose tissue was harvested, weighed, finely minced and digested in HBSS containing 1 mg/mL collagenase type II (Sigma) for approximately 40 min at 37°C with gentle shaking. The digested material was passed through a 100 μm filter (Falcon-352360), washed with HBSS+10%FBS+P/S and spun at 300 g × 5min. The medium was removed, floating mature adipocytes and stromal vascular fraction (SVF) were collected and washed again. Isolation of mRNA was performed by RNeasy Lipid Tissue Mini Kit (Quiagen). Adiponectin (ADIPOQ), a late marker of adipocyte differentiation, was used to confirm the purity of the fractions.

2.13. 3T3-L1 cell culture

3T3-L1 murine pre-adipocytes (reference 088SP-L1-F) were purchased from the ZenBio company (NC, USA) and cultured in an atmosphere of 5% CO2 at 37°C using DMEM supplemented with 10% newborn calf serum. For adipocyte differentiation, cells were stimulated with 3T3-L1 differentiation medium containing IBMX (500 μM), deoxy-methasone (250 nM) and insulin (175 nM) for 2 days after cells reached confluency. The medium was changed to DMEM containing 10% FBS and insulin (175 nM) after 2 days and adipocytes were then kept into DMEM containing only 10% FBS. Prior to the experiments, adipocytes were subjected to serum deprivation for 16 h with DMEM supplemented with 0.5% FBS.

2.14. siRNA treatment

siRNA for Nrf2 (reference L-040766-00-0005) was purchased from Horizon Discovery LTD company (Cambridge, UK) and siRNA treatment was performed according to manufacturer’s instructions.

2.15. Cellular bioenergetic analysis using the Seahorse Bioscience XF analyzer

Bioenergetic profiles of the adipocytes were determined using a Seahorse Bioscience XF42 Analyzer (Billerica, MA, USA) that provides real-time measurements of oxygen consumption rate (OCR), indicative of mitochondrial respiration, and extracellular acidification rate (ECAR), an index of glycolysis as previously described by our group [2].
2.16. Determination of ATP, NAD\(^+\) and NADH

NAD\(^+\) and NADH levels were determined in frozen adipose tissue samples using an enzymatic method (EnzyChrom, BioAssays Systems, Hayward, CA) according to manufacturer’s instructions. Intracellular ATP levels were measured at the end of the experiments with the XF Analyzer using an ATPLiteTM Bioluminescence Assay Kit (PerkinElmer, Courtaboeuf, France) according to manufacturer’s instructions.

2.17. Statistical analysis

Data are expressed as mean values ± standard error of the mean (SEM). Statistical analysis was performed by one or two-way analysis of variance (ANOVA) with Fisher multiple comparison test. The result were considered significant at p-value<0.05.

3. Results

3.1. Nrf2-deficient mice exhibit enhanced Sirt1 levels originating from WAT

A representative photograph of 22 week-old mice reveals that the Nrf2\(^{-/-}\) mouse is slightly smaller than its WT counterpart (Fig. 1A). When mice were fed a standard diet (SD) for 14 weeks, we did not observe a significant difference in the changes of total body weight and food intake between the two groups (Fig. 1B). Nevertheless, Nrf2\(^{-/-}\)
mice remained slightly smaller and displayed significantly higher total lean mass and lower fat mass compared to WT mice (Fig. 1C). Interestingly, the plasma levels of the main adipose tissue cytokines leptin, adiponectin and resistin were also markedly lower in Nrf2−/− mice than in WT mice (Table 1). We did not find any difference in glucose and insulin metabolism assessed by the glucose (GTT) and insulin (ITT) tolerance tests, respectively (Fig. 1D and 1E). Similarly, plasma levels of glucose, insulin and lipids (triglycerides, non-esterified fatty acid, cholesterol, apolipoproteins, low and high-density lipoproteins (LDL and HDL, respectively)) were all comparable between WT and Nrf2−/− mice (Table 1). However, among the various parameters measured, we noticed that plasma Sirt1 levels were strongly enhanced in Nrf2−/− mice compared to their WT counterparts (Fig. 1F). To identify the source of Sirt1, we examined the expression of the Sirt1 gene by qPCR in various organs. Interestingly, and compared to WT mice, we detected a significant up-regulation of Sirt1 mRNA expression only in eWAT of Nrf2−/− mice, while no difference was observed in liver, muscle, heart, subcutaneous white (SubWAT) and brown adipose tissues (BAT) (Fig. 1G). Immunofluorescence staining revealed that the enhanced Sirt1 gene expression in eWAT of Nrf2−/− mice coincided with an increase of Sirt1 protein levels (Fig. 1H). Notably, the marked increase in Sirt1 mRNA expression in eWAT of Nrf2−/− mice was originating from both the adipocyte and stromal vascular fraction (SVF) as indicated by the similar level of Sirt1 in these two type of cellular components (Fig. 1I). To confirm further that an increase in Sirt1 expression is due to the absence of Nrf2, we cultured 3T3-L1 fully differentiated adipocytes and silenced the Nrf2 gene by siRNA transfection. As shown in Fig. 1J, we found that ablatting Nrf2 in adipocytes causes an increased Sirt1 gene expression. These data clearly show that Nrf2 deficiency results in a distinctive adaptation characterized by increased circulating levels of Sirt1, which most likely originates from enhanced secretion of Sirt1 from eWAT.

3.2. Nrf2 deficiency enhances lipolysis and fatty acid oxidation in adipose tissue

Because Sirt1 controls lipid metabolism in adipocytes [5,28], we further investigated the impact of Nrf2 deficiency on lipolysis, fatty acid oxidation and lipogenesis. We first noticed that the amount of eWAT in Nrf2-deficient mice is significantly reduced, with much smaller and denser adipocytes compared to WT mice (Fig. 2A–D). BAT and SubWAT of Nrf2−/− mice were also smaller and exhibited a reduced adipocyte size than in WT (Supplementary Figs. 1 and 2), even in the absence of increased Sirt1 gene levels (Fig. 1G). Secondly, the expression of several genes involved in lipolysis and fatty acid oxidation in eWAT of Nrf2−/− mice was significantly higher compared to WT. These included forkhead box protein O1 (FoxO1), adipose triglyceride lipase (Atgl), fatty acid binding protein 4 (Fabp4) and perilipin (Plin1), peroxisome proliferator-activated receptor gamma coactivator 1-α (PGC1α) and peroxisome proliferator-activated receptor alpha (PPARα) (Fig. 2E). In contrast, lipogenic genes such as peroxisome proliferator-activated receptor gamma (PPARγ), fatty acid synthase (Fasn) and sterol regulatory element-binding transcription factor 1 (Srebp1c) in eWAT of Nrf2−/− were all lower than in WT mice (Fig. 2E). A similar profile for adipocytes lipolytic and lipogenic gene expressions was found in SubWAT and for some genes (i.e. PGC1α) also in BAT (Supplementary Figs. 1 and 2). We also show that increased Fox1 gene expression correlated with up-regulation of FOXO1 protein in eWAT of Nrf2−/− mice in association with enhanced AMPK and phosphorylated-AMPK proteins (Fig. 2F). Since FOXO1, AMPK and phosphorylated-AMPK are downstream targets of Sirt1, these data suggest a Sirt1-driven metabolic adaptation of eWAT resulting from Nrf2-deficiency. Moreover, we found a significant increase in perilipin, a protein known to promote hydrolysis of lipid droplets in adipose tissue (Fig. 2G), and higher levels of non-esterified free fatty acid (NEFA) secreted in the conditioned media from eWAT of Nrf2−/− mice compared to WT (Fig. 2H). Finally, we demonstrated by electron microscopy that eWAT from Nrf2−/− mice contained more lipid droplets than eWAT from WT animals (Fig. 2I). Altogether, these data establish that: 1) Nrf2 deficiency modifies adipose tissue metabolism, driving it towards fatty acids mobilization and elimination; 2) this modification is associated with an increased Sirt1 production, as demonstrated by activation of Sirt1 downstream targets in eWAT and consistent with the recognized role of Sirt1 in fatty acid utilization [28]; 3) in addition to autocrine signaling on eWAT, it is likely that Sirt1 secreted by eWAT and circulating in blood also affects BAT and SubWAT fatty acid metabolism in an endocrine manner.

3.3. Nrf2 deficiency enhances bioenergetic metabolism in adipocytes

Because of the enhanced fatty acid metabolism in eWAT, we next assessed the impact of Nrf2 deficiency on bioenergetics in adipocytes and adipose tissue punches isolated from Nrf2−/− and WT mice. Mitochondrial content, and NAD+/NADH ratio were not significantly different between the two groups, although citrate synthase activity tended to increase (Fig. 3A, 3B and 3C). Using the Seahorse analyzer (Fig. 3D and 3F), we also found that Nrf2−/− adipocytes displayed a slightly higher basal and ATP-linked oxygen consumption rate (OCR), a significant increase in maximal respiration (Fig. 3E) as well as higher glycolysis and glycolytic capacity compared to WT adipocytes (Fig. 3G). Concomitantly, Nrf2−/− adipocytes exhibited higher ATP levels compared to WT adipocytes (Fig. 3H). The increased ATP originates from both oxidative phosphorylation, as evidenced by the experiments conducted in the presence of the glycolysis inhibitor 2-deoxyglucose, and glycolysis, revealed by the experiments performed in the presence of the ATP synthesis inhibitor oligomycin (Fig. 3H). Importantly, a

Table 1

|                  | SD            | HFD           |
|------------------|---------------|---------------|
|                  | Nrf2−/−       | Nrf2−/−       |
|                  | WT            | WT            | HFD            |
| Glucose (mg/dL)  | 139.3 ± 0.4   | 111.3 ± 0.6   | 231.1 ± 0.8   | 149.4 ± 0.9  |
| Insulin (mg/mL)  | 4.0 ± 0.4     | 0.8 ± 0.2     | 0.2 ± 0.1     | 1.2 ± 0.2    |
| HOMA-IR          | 0.8 ± 0.2     | 0.6 ± 0.2     | 2.8 ± 0.6     | 1.3 ± 0.4    |
| Triglycerides    | 143.1 ± 4.8   | 142.6 ± 1.3   | 137.6 ± 1.9   | 147.5 ± 2.1  |
| NEFA (mmol/L)    | 4.8 ± 0.4     | 4.4 ± 0.3     | 4.2 ± 0.3     | 6.3 ± 0.5    |
| Cholesterol (mg/dL) | 51.5 ± 0.2 | 25.8 ± 1.0     | 25.3 ± 1.5     | 43.6 ± 2.7 |
| LDL (mg/dL)      | 12.1 ± 0.8    | 2.5 ± 0.5     | 2.5 ± 0.5     | 2.4 ± 0.4    |
| HDL (mg/dL)      | 13.8 ± 0.4    | 13.8 ± 1.0    | 5.3 ± 0.3     | 43.6 ± 2.4   |
| Ldl/HDL (arbitrary units) | 0.4 ± 0.3 | 0.5 ± 0.1 | 1.0 ± 0.1 | 1.7 ± 0.3 |
| ApoA1 (mg/dL)    | 65.7 ± 1.3    | 65.5 ± 1.9    | 89.6 ± 2.4    | 81.3 ± 2.4   |
| ApoA2 (mg/dL)    | 12.1 ± 0.9    | 12.1 ± 1.1    | 20.8 ± 1.2    | 1.2 ± 0.4    |
| ApoB100 (mg/mL)  | 30.4 ± 0.8    | 31.3 ± 0.8    | 51.5 ± 1.2    | 39.2 ± 1.5   |
| Leptin (ng/mL)   | 2.1 ± 0.7     | 0.6 ± 0.2     | 57.9 ± 1.9    | 18.0 ± 2.5   |
| Adiponectin (µg/mL) | 7.8 ± 0.9 | 5.3 ± 0.2 | 12.3 ± 0.4 | 4.9 ± 0.1 |
| Resistin (ng/mL) | 0.9 ± 0.9     | 0.5 ± 0.5     | 0.5 ± 0.4     | 0.4 ± 0.3    |
| β-hydroxybutyrate (mM/L) | 0.4 ± 0.5 | 0.5 ± 0.4 | 0.5 ± 0.4 | 0.5 ± 0.4 |
| FGF21 (ng/mL)    | 0.64 ± 0.2    | 0.78 ± 0.1    | 1.4 ± 0.9     | 1.98 ± 0.3   |

Apo: apolipoprotein; HDL: high density lipoprotein; HOMA-IR: Homeostastic model assessment of insulin resistance; LDL: low density lipoprotein; NEFA: Non-esterified fatty acid; FGF21: Fibroblast growth factor 21. Results are shown as mean ± SEM. *p < 0.05 vs. SD WT group; #p < 0.05 vs. HFD WT group.
A similar increase in mitochondrial oxygen consumption and glycolytic activity was found in ex vivo adipose tissue punches collected from Nrf2/−/− mice compared to WT eWAT (Fig. 3I and 3J). These results indicate that adipocytes from Nrf2/−/− mice are more metabolically active in vitro and ex vivo. Accordingly, Nrf2/−/− mice displayed increased plasma levels of both citric acid and lactic acid compared with WT mice under a SD regime (Table 2), confirming that Nrf2 deficiency is characterized by an overall increase in energetic metabolism in adipocytes.

3.4. Improved glucose metabolism and lipid profile in Nrf2-deficient mice fed a high fat diet (HFD)

We next asked whether the beneficial changes in glucose and lipid metabolism promoted by Nrf2 deficiency protect against obesity. For this, WT and Nrf2/−/− mice were subjected to a HFD regime for 14 weeks as previously described by our group [2]. As shown in Fig. 4, at the end of this protocol Nrf2/−/− mice displayed a lower body weight despite having a higher food intake (Fig. 4A and 4B) compared to WT mice. Nrf2/−/− mice also had a higher lean mass and a lower fat mass (Fig. 4C) as well as improved glucose metabolism and insulin tolerance compared to their WT counterpart (Fig. 4D and 4E), confirming previously published data [20]. In line with these findings, Nrf2/−/− mice under HFD showed reduced levels of plasma glycemia and insulin after fasting and improved HOMA-IR (Table 1). Although plasma triglyceride and NEFA levels remained unchanged, HDL, LDL, ApoA1, ApoA2 and ApoB100 were all significantly reduced in Nrf2/−/− mice after HFD (Table 1). As previously reported [22], plasma levels and the hepatic gene expression of FGF21 were significantly higher in Nrf2/−/− mice (Table 1 and Supplementary Fig. 3G) together with increased circulating lactic acid and citric acid levels (Table 2). Interestingly, and similarly to mice fed a SD, Nrf2/−/− obese mice maintained significantly higher levels of plasma Sirt1 compared to WT obese mice (Fig. 4F). No difference in the size of adipocytes was observed between the two groups (Fig. 4G and 4H) but the visceral adipose tissue of Nrf2/−/− mice still displayed a similar increase in mitochondrial oxygen consumption and glycolytic activity was found in ex vivo adipose tissue punches collected from Nrf2/−/− mice compared to WT eWAT (Fig. 3I and 3J). These results indicate that adipocytes from Nrf2/−/− mice are more metabolically active in vitro and ex vivo. Accordingly, Nrf2/−/− mice displayed increased plasma levels of both citric acid and lactic acid compared with WT mice under a SD regime (Table 2), confirming that Nrf2 deficiency is characterized by an overall increase in energetic metabolism in adipocytes.

3.4. Improved glucose metabolism and lipid profile in Nrf2-deficient mice fed a high fat diet (HFD)

We next asked whether the beneficial changes in glucose and lipid metabolism promoted by Nrf2 deficiency protect against obesity. For this, WT and Nrf2/−/− mice were subjected to a HFD regime for 14 weeks as previously described by our group [2]. As shown in Fig. 4, at the end of this protocol Nrf2/−/− mice displayed a lower body weight despite having a higher food intake (Fig. 4A and 4B) compared to WT mice. Nrf2/−/− mice also had a higher lean mass and a lower fat mass (Fig. 4C) as well as improved glucose metabolism and insulin tolerance compared to their WT counterpart (Fig. 4D and 4E), confirming previously published data [20]. In line with these findings, Nrf2/−/− mice under HFD showed reduced levels of plasma glycemia and insulin after fasting and improved HOMA-IR (Table 1). Although plasma triglyceride and NEFA levels remained unchanged, HDL, LDL, ApoA1, ApoA2 and ApoB100 were all significantly reduced in Nrf2/−/− mice after HFD (Table 1). As previously reported [22], plasma levels and the hepatic gene expression of FGF21 were significantly higher in Nrf2/−/− mice (Table 1 and Supplementary Fig. 3G) together with increased circulating lactic acid and citric acid levels (Table 2). Interestingly, and similarly to mice fed a SD, Nrf2/−/− obese mice maintained significantly higher levels of plasma Sirt1 compared to WT obese mice (Fig. 4F). No difference in the size of adipocytes was observed between the two groups (Fig. 4G and 4H) but the visceral adipose tissue of Nrf2/−/− mice still displayed a
different profile compared to WT. This included increased perilipin content (Fig. 4G and 4I) and an elevated expression of genes involved in lipolysis and fatty acid oxidation (Fig. 4K). Moreover, a slight, non-significant, decrease in macrophage infiltration, which is known to contribute to adipose tissue low-grade inflammation during obesity, was observed in Nrf2−/− mice (Fig. 4G and 4J). Of note, the metabolic protection afforded by Nrf2 deficiency against HFD was evident not only in the adipose tissue but also in the liver, where our analysis revealed reduced hepatic injury (ALT/AST) and steatosis and decreased expression of certain genes involved in lipogenesis (Supplementary Figs. 3A–C). These data support the concept that Nrf2 deficiency counteracts the metabolic derangements caused by HFD resulting in improved phenotype of the adipose tissue and other organs such as the liver.

3.5. eWAT removal suppresses the increase in Sirt1 and impairs glucose tolerance in Nrf2-deficient obese mice

Since eWAT of Nrf2−/− mice is the only tissue expressing high Sirt1 content (Fig. 4G and 4I) and an elevated expression of genes involved in lipolysis and fatty acid oxidation (Fig. 4K). Moreover, a slight, non-significant, decrease in macrophage infiltration, which is known to contribute to adipose tissue low-grade inflammation during obesity, was observed in Nrf2−/− mice (Fig. 4G and 4J). Of note, the metabolic protection afforded by Nrf2 deficiency against HFD was evident not only in the adipose tissue but also in the liver, where our analysis revealed reduced hepatic injury (ALT/AST) and steatosis and decreased expression of certain genes involved in lipogenesis (Supplementary Figs. 3A–C). These data support the concept that Nrf2 deficiency counteracts the metabolic derangements caused by HFD resulting in improved phenotype of the adipose tissue and other organs such as the liver.

Since eWAT of Nrf2−/− mice is the only tissue expressing high Sirt1 levels, we hypothesized that eWAT-derived Sirt1 is responsible for protecting Nrf2−/− mice against the metabolic disruption caused by HFD. To test this hypothesis, we directly removed eWAT from WT and Nrf2−/− mice (eWATectomy) prior to subjecting them to HFD for 8 weeks (see schematic diagram in Fig. 5A). Plasma levels of Sirt1, body weight and metabolic parameters were assessed at the end of the protocol. Notably, we found that removal of fat completely abolished the increase in circulating Sirt1 characteristic of Nrf2 deficiency to levels comparable to sham-operated WT mice (Fig. 5B), unequivocally demonstrating that eWAT is the main source of Sirt1 in Nrf2−/− mice.
Importantly, Nrf2−/− mice subjected to eWATectomy became glucose intolerant in a manner similar to WT mice under HFD (Fig. 5E) but retained their enhanced insulin sensitivity in both sham and eWATectomy conditions (Fig. 5D). Moreover, eWATectomy did not affect body weight gain induced by HFD since Nrf2−/− mice still exhibited lower body weight gain compared to WT (Fig. 5C). We also found that lactic acid, which was augmented in Nrf2−/− mice both under SD and HFD regimes (Table 2), was normalized by eWATectomy to WT levels (Fig. 5F) while circulating citric acid remained elevated. Circulating lipids were still similar in Nrr2-deficient animals before and after eWATectomy (data not shown). Taken together, these results indicate that Sirt1 produced by eWAT is a critical factor in preserving improved glucose metabolism in Nrf2−/− mice during obesity.

4. Discussion

Our study reveals for the first time that Nrf2 deficiency in lean and obese mice is characterized by increased systemic levels of Sirt1. We establish the visceral adipose tissue as the sole source of high Sirt1. Our results demonstrate that Sirt1 expression is strongly induced by HFD-mediated lipid dysregulation. Our findings on improved insulin-resistance and glucose intolerance in murine Nrf2 deficiency after HFD are in line with previous studies reporting a protective role of Nrf2 deficiency in obese mice through FGF21 overexpression after challenge with HFD [22]. However, in that study and ours, lean Nrf2−/− mice did not display increased FGF21 levels and it was unclear why Nrf2 deficiency caused a different body composition even under standard diet. We therefore looked for other plasma metabolic markers that could explain this difference between the genotypes and discovered that Sirt1 was significantly higher in the circulation of lean Nrf2−/− mice. By evaluating its gene expression in other organs, we demonstrated that in Nrf2−/− mice, the visceral adipose tissue was the only tissue where Sirt1 was elevated compared to WT. We fully validated this finding by showing that this increase was not detected in the liver, muscle, heart, SubWAT and BAT and that circulating Sirt1 returned to WT levels when eWATectomy was carried out in Nrf2−/− mice. Interestingly, Sirt1 levels remained higher also after HFD, supporting its contribution to the beneficial metabolic adaptation of Nrf2−/− mice to an obesogenic regime.

It is intriguing that Sirt1 levels increased exclusively in the adipose tissue in response to whole mouse Nrf2 deficiency and a detailed investigation is required to understand the underlying mechanisms. Nevertheless, we speculate that Nrf2 suppresses Sirt1 expression, as indicated also by our preliminary findings showing that silencing Nrf2 with siRNA in 3T3-L1 pre-adipocytes is directly associated with up-regulation of Sirt1 mRNA. Whether this is due to direct control of Sirt1 gene expression by the Nrf2 transcription factor or because Nrf2 deficiency results in higher oxidative stress that in turn may stimulate Sirt1 [29] is unknown. A very recent report showed that HFD induced marked oxidative stress in adipose tissue with concomitant increase in Nrf2 [30], suggesting that oxidative stress may be a participating factor. These authors also reported that Nrf2 activation promoted lipid accumulation and lipogenesis in adipocytes during obesity [30], supporting our current results on the beneficial role of Nrf2 deficiency against HFD-mediated lipid dysregulation.

Sirt1 is known to be a key metabolic sensor in various metabolic tissues [31]. Sirt1 promotes lipolysis via FOXO1 and adipocyte triglyceride lipase (ATGL)-dependent mechanisms [28] and stimulates fat utilization in white adipose tissue by repressing PPARγ [5]. Consistent with this, our results demonstrated that Nrf2 deficiency led to decreased adipocyte size, increased expression of the lipolysis-related genes Foxo1, Atgl, Fabp4, Perilipin1 and decreased expression of the adipogenesis gene PPARγ in both lean and obese mice. A higher lypolysis activity was corroborated by electronic microscopy imaging of visceral adipose tissue showing more numerous lipid droplets in Nrf2−/− than WT mice, thus indicating a more metabolically active adipose tissue in Nrf2−/− animals. This effect was accompanied by enhanced mitochondrial oxidative and glycolytic activities in primary adipocytes in vitro and in ex vivo adipose tissue. Although a previous study on embryonic fibroblast cells from Nrf2-deficient mice showed an increased oxygen consumption compared to WT cells [32], other reports have demonstrated a decrease in oxygen consumption in cells lacking Nrf2 [33] but these studies were performed on cell types different from adipocytes. Our main findings on gene expression data in different adipose tissue depots, phenotypic and plasma analysis of circulating HDL and LDL as well as apoproteins levels and bioenergetic profile of eWAT strongly support a positive impact of Nrf2 deficiency on systemic metabolism that becomes highly relevant during HFD conditions.

Our work highlights that Sirt1 activation is a crucial adaptive mechanism following the lack of Nrf2, adding to previous investigations which identified increased expression of FGF21 as contributing factor to the improved metabolic phenotype of Nrf2−/− mice under HFD [22]. We also found that plasma levels of FGF21 were elevated in Nrf2−/− mice after HFD and we suggest here that Sirt1 produced by the adipose tissue and FGF21 originating from the liver act in a concerted manner to ameliorate energy homeostasis of Nrf2 deficient animals under HFD. Recent studies report that Sirt1 promotes FGF21 signaling in oxtocin neurons and stimulates oxtocin transcription through Nrf2, contributing to the homeostatic regulation of macronutrient-based diet selection in mice [34,35]. Thus, we cannot exclude that Sirt1 overproduction in eWAT of Nrf2−/− mice may also affect FGF21 levels via an endocrine action on the liver.

Our findings on improved insulin-resistance and glucose intolerance in murine Nrf2 deficiency after HFD are in line with previous studies.
Calorimetry assessments have also shown that Nrf2−/− mice under HFD display a significant increase in energy expenditure compared to WT mice [30,32]. This, combined with our novel data on Sirt1 overexpression derived exclusively from eWAT, prompted us to determine the specific role of Sirt1 on metabolism in Nrf2−/− mice. Because we observed a reduction of Sirt1 to WT levels after eWATectomy in Nrf2-deficient mice, we asked ourselves: what would be the metabolic consequences of subjecting Nrf2−/− mice lacking eWAT to HFD? Our results clearly demonstrate that following the HFD regime, Nrf2−/− mice still exhibit improved insulin sensitivity but lose the protection against glucose intolerance, narrowing down the unique impact of Sirt1 on glucose metabolism in these conditions. Plasma lactic acid also returned to WT levels after eWATectomy, substantiating the role of Sirt1 in glucose metabolism in Nrf2−/− mice and consistent with a study demonstrating the importance of Sirt1 in glucose tolerance but not insulin resistance [4]. In contrast, circulating citric acid remained elevated, indicating that other players, originating from tissues different from eWAT, contribute to metabolic adaptation in Nrf2−/− mice. These additional factors (including FGF21) likely explain why Nrf2 deficient animals are still protected from HFD-induced weight gain and insulin resistance even after eWATectomy.
5. Conclusions

In conclusion, our study advances our understanding on the role Nrf2 plays in modulating energetic metabolism and identifies novel pathways that are controlled by this master regulator of the stress response. This is the first study reporting a link between Nrf2 deficiency and Sirt1 expression in visceral adipose tissue and their beneficial implications in the regulation of glucose metabolism. The use of Nrf2 inhibitors alone or in combination with other drugs could be an effective therapeutic strategy to counteract the metabolic dysfunction and glucose deregulation that characterize diseases such as obesity and type 2 diabetes.

Declaration of competing interest

The authors declare no conflict of interest.

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Abbreviations

ALT alanine transaminase  
AST aspartate transaminase  
DMEM Dulbecco’s Modified Eagle Medium  
DPBS Dulbecco Phosphate Buffer Solution  
ECAR extracellular acidification rate  
eWAT epididymal adipose tissue  
FBS fetal bovine serum  
Nrf2 nuclear factor (erythroid-derived 2)-like 2
glucose tolerance test
glucose tolerance test
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protein extractions, ELISA and immunostaining. G.D. read the manuscript. A.C. acids as well as analysis of plasma lipids. S.M. performed RNA and script. R.M. is the guarantor of this work and, as such, had full access to

SIRT 1 sirtuin-1
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