Elevated UCP1 levels are sufficient to improve glucose uptake in human white adipocytes

D. Tews a,2,*, T. Pula a,2, J.B. Funcke a,1, M. Jastroch b, M. Keuper b, K.M. Debatin c, M. Wabitsch a, P. Fischer-Posovszky a

 a Division of Pediatric Endocrinology and Diabetes, Department of Pediatrics and Adolescent Medicine, Ulm University Medical Center, Ulm, Germany
 b Department of Molecular Biosciences, The Wenner-Gren Institute, Stockholm University, SE-106 91 Stockholm, Sweden
 c Department of Pediatrics and Adolescent Medicine, Ulm University Medical Center, Ulm, Germany

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A B S T R A C T

Brown adipose tissue (BAT) has been considered beneficial for metabolic health by participating in the regulation of glucose homeostasis. The browning factors that improve glucose uptake beyond normal levels are still unknown but glucose uptake is not affected in UCP1 knockout mice. Here, we demonstrate in human white adipocytes that basal/resting glucose uptake is improved by solely elevating UCP1 protein levels. Generating human white Simpson-Golabi-Behmel syndrome (SGBS) adipocytes with a stable knockout and overexpression of UCP1, we discovered that UCP1 overexpressing adipocytes significantly improve glucose uptake by 40%. Mechanistically, this is caused by higher glycolytic flux, seen as increased oxygen consumption, extracellular acidification and lactate secretion rates. The improvements in glucose handling are comparable to white-to-brown transitions, as judged by, for the first time, directly comparing in vitro differentiated mouse brown vs white adipocytes. Although no adipogenic, metabolic and mitochondrial gene expressions were significantly altered in SGBS cells, pharmacological inhibition of GLUT1 completely abrogated differences between UCP1+ and control cells, thereby uncovering GLUT1-mediated uptake as permissive gatekeeper. Collectively, our data demonstrate that elevating UCP1 levels is sufficient to improve human white adipocytes as a glucose sink without adverse cellular effects, thus not requiring the adrenergic controlled, complex network of browning which usually hampers translational efforts.

1. Introduction

Brown adipose tissue (BAT) enables mammals to maintain their body temperature by non-shivering thermogenesis (NST). BAT is highly prevalent in newborns and essential for their survival in absence of other thermogenic means [1]. It is also detectable in adults, where its presence is associated with lower body weight [2]. Besides participating in the regulation of body temperature and energy homeostasis, BAT activity has beneficial effects on systemic metabolism [3,4]. As such, it increases glucose tolerance and insulin sensitivity, and decreases dyslipidemia and body weight [3,4]. Consequently, BAT is currently discussed as a targeted organ to treat obesity and obesity-associated diseases [5].

The key component mediating thermogenesis is uncoupling protein 1 (UCP1). UCP1 is located in the inner mitochondrial membrane, where it dissociates cellular respiration from the generation of ATP. Upon certain stimuli such as cold exposure, sympathetically-derived nor-epinephrine activates β3 adrenergic receptors on brown adipocytes, triggering the release of free fatty acids (FFA) and subsequent activation UCP1 [1]. The persistence of BAT from neonates into adulthood, however, is limited. The induction of “beige” or “brite” adipocytes in human white adipose tissue (WAT) represents another approach to combat excessive nutrient energy. In mice, cold-induced WAT browning improves systemic glucose and lipid homeostasis [6]. Mirabegron, a FDA-approved β3-adrenergic receptor agonist for overactive bladder, also induces in human WAT a complex regulatory network for

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thermogenesis, including UCP1, TMEM26, CIDEA and many other browning genes [7,8].

Data from in vitro studies suggest that glucose uptake and glycolysis are strictly required for adrenergically stimulated oxidative metabolism of murine brown adipocytes [9]. Ectopic UCP1 expression in murine WAT reduces plasma glucose levels [10]. Indeed, glucose uptake has been widely used to detect BAT activity in both mice and humans under basal and cold-exposed conditions with 18F-FDG-PET/CT being the gold standard of BAT detection and activity measurement [11]. This method is based on the observation that BAT is metabolically more active than other tissues [2,12,13]. It thus stands to reason that BAT takes up more glucose than WAT. Interestingly though, knockout of UCP1 in mice does not lead to diminished glucose uptake indicating that thermogenesis and glucose flux are dissociated in BAT and the molecular basis of elevated glucose uptake in BAT still has to be identified [14].

The aim of this study was to delineate whether UCP1 is a main driver of glucose uptake in adipocytes. To this end, we compared the cellular glucose uptake in murine white and brown adipocytes isolated and differentiated from inguinal white and interscapular brown adipose tissue. Furthermore, we knocked out and overexpressed UCP1 in Simpson-Golabi-Behmel (SGBS) cells, a unique human model system for white adipocytes to study adipogenesis [15,16].

2. Materials and methods

2.1. Materials

Standard chemicals were purchased from Merck (Darmstadt, Germany), media and buffers for cell culture were purchased from ThermoFisher Scientific (Waltham, USA).

2.2. Murine cell isolation and culture

6–8 week old C57BL/6J mice were sacrificed by cervical dislocation and inguinal white as well as interscapular brown adipose tissues were dissected. Stromal vascular cells (SVC) were isolated by collagenase digestion and subjected to adipogenic differentiation as described [17]. Briefly, tissues were digested using collagenase and stromal cells were harvested by centrifugation. Cells were seeded in DMEM:F12 containing 10% FCS until reaching subconfluency. For subsequent experiments, cells were trypsinized and reseeded into 12 well plates. Adipogenic differentiation was induced for 12 days DMEM:F12 medium supplemented with 10% FCS, 850 nM insulin, 1 nM T3, 125 μM indomethacin, and 1 μM rosiglitazone. For the first 2 days, 500 μM isobutylmethylxanthine, and 25 nM dexamethasone were added.

2.4. Generation of UCP1-overexpressing and UCP1-deficient cells

For UCP1 overexpression constructs, the UCP1 open reading frame was PCR amplified from SGBS adipocyte cDNA using specific primer pairs harboring attB sites (Supplementary table S2), inserted into pDONR221 using Gateway BP Clonase II (ThermoFisher Scientific), and was verified by Sanger sequencing. UCP1 overexpression cassettes were transferred from pDONR221-UCP1 to pLent6.3/V5-DEST using Gateway cloning-compatible SleepingBeauty transposon plasmid (Addgene #34879, kindly provided by Zsuzsanna Izsvak) at a stable insertion of 9000 bp. UCP1 overexpression was confirmed by qRT-PCR and Western Blot.

For UCP1 knockout constructs, sRNA duplexes targeting the UCP1 gene and non-targeting controls were cloned into the pMuLE ENTR U6 stuffer sRNA scaffold L1-L4 plasmid (Multiple Lentiviral Expression Kit, Addgene #1000000060, kindly provided by Ian Frew) [18]. Correct insertion of the sRNA duplexes was verified by Sanger sequencing. A Gateway cloning-compatible SleepingBeauty transposon plasmid ‘pMuSE eGFP-P2A-PuroR DEST’ [19] was derived from pShuttle-c2 (Addgene #60495, kindly provided by Eric Kowarz) and pLenti3.6/V5-DEST (ThermoFisher Scientific) using routine cloning methods. The specific pMuLE ENTR U6-sRNA plasmids were recombined with pMuLE ENTR CMV-hCas9 R4-R3, pMuLE ENTR MCS L3-L2, and pMuSE eGFP-P2A-PuroR DEST plasmids using LR Clonase II Plus (ThermoFisher Scientific). SGBS preadipocytes were co-transfected with the resulting pMuSE U6-sRNA + CMV-hCas9 + RPBSA-eGFP-P2A-PuroR plasmids and the SleepingBeauty-expressing pCMV(CAT)7-T7-SB100 plasmid (Addgene #34879, kindly provided by Zsuzsanna Izsvak) at a mass ratio of 19:1 using a Neon Transfection System (ThermoFisher Scientific) with 3 × 104 μl pulses of 1400 V. Stable bulk cultures were generated by puromycin selection. UCP1 knockout was confirmed by Western Blot. All essential sequences are listed in Supplemental Table S1.

SGBS cells were cultured and differentiated into adipocytes using a protocol described before [15]. Briefly, cells were seeded into cell culture vessels in DMEM:F12 containing 3.3 mM biotin, 1.7 mM panthotenate, and 10% FCS. Adipogenic differentiation was induced for 14 days in serum-free DMEM:F12 medium supplemented with 10 μg/ml apo-transferrin, 10 nM insulin, 200 pM T3, and 1 μM cortisol. For the first 4 days, 2 μM rosiglitazone, 250 μM isobutylmethylxanthine, and 25 nM dexamethasone were added.

2.5. Gene expression analysis

RNA was isolated using the Direct-Zol RNA Kit (Zymo Research, Irvine, USA). 0.5 μg of total RNA was reverse transcribed using SuperScript II Reverse Transcriptase (ThermoFisher Scientific). Relative expression of target genes was analyzed by quantitative real-time PCR using the ssoAdvanced Universal SYBR Green Supermix on a CFX Connect Real Time PCR Detection System (BioRad, Munich, Germany). Expression values were calculated using the dCt method with hypoxanthine-guanine phosphoribosyltransferase (HPRT) as a reference gene. Sequences of the primers are shown in Supplemental Table S2.

2.6. Western Blot

Cells were washed with ice-cold PBS and total protein was extracted by scraping them in a lysis buffer (10 mM Tris-HCl at pH 7.5, 150 mM sodium chloride, 2 mM EDTA, 1% TX-100, 10% glycerol, 1X Complete Protease Inhibitors (Roche, Mannheim, Germany)). After 30 min incubation on ice, cell debris were pelleted by centrifugation. Protein content in the supernatants was determined using the Bradford Protein Assay (BioRad). For immunodetection, 10–20 μg of protein were separated by SDS-PAGE on Bolt Bis-Tris Plus Gels (ThermoFisher Scientific) and transferred to nitrocellulose membranes by Western Blotting (ThermoFisher Scientific). The following antibodies were used: mouse anti-UCP1 (R&D Systems #536435, Minneapolis, UCA), mouse anti-OPXPHOS protein cocktail (Abcam #ab110411, Cambridge, UK), rabbit anti-TIM23 (Abcam #ab16329), rabbit anti-perilipin A (Abcam #ab3526), rabbit anti-PPARG (Cell Signaling Technology #2443), rabbit anti-adiponectin (GeneTex #GTX121777), mouse anti-GAPDH (HyTest #5G4, clone 6C5, Turku, Finland), mouse anti-β-actin (Merck #A5441).

2.7. Glucose metabolism

In vitro differentiated adipocytes were washed with PBS and incubated in serum-free medium overnight. On the day of measurement, medium was replaced with glucose-free Krebs- Ringer buffer (130 mM NaCl, 10 mM MgSO4, 2.5 mM NaH2PO4, 4.6 mM KCl, 2.5 mM CaCl2,
2.5 mM sodium pyruvate, 5 mM HEPES, pH 7.4). To determine basal glucose uptake, [14C]-2-deoxy-D-glucose (0.2 μCi/well, PerkinElmer, Waltham, USA) was added and the cells were incubated for 15 min at 37 °C. Subsequently, cells were washed with ice-cold PBS and harvested with 100 mM NaOH. Incorporation of [14C]-2-deoxy-D-glucose was measured on a β-counter. In some experiments, the cells were pre-treated for 1 h with 10 μM Cytochalasin B or 100 nM of the selective GLUT1 inhibitor BAY-876 before addition of [14C]-2-deoxy-D-glucose. Data obtained from SGBS cells are given in counts per minute (cpm). As SVC isolated from iWAT and BAT were different in cell size and rate of adipogenic differentiation, we counted the number of adipocytes in the cell culture dishes. Raw cpm values were then normalized to adipocyte numbers per well (cpm/10^5 adipocytes).

For quantification of lactate secretion, the cells were incubated in serum-free medium overnight and then in DMEM medium supplemented with 10 mM glucose, 1 mM pyruvate, 2 mM GlutaMaxx, and 1% bovine serum albumin. Media supernatants were harvested after 16 h and assayed for lactate content using the Lactate Colorimetric Assay Kit (Biovision, Milpitas, USA).

### 2.8. Mitochondrial quantification

Citric acid synthase activity was assayed as a measure for mitochondrial content and activity as described before [20]. In brief, 5 μg total protein were added to reaction buffer (100 mM Tris HCl pH 8.1, 100 μM 5,5′-dithiobis-2-nitrobenzoic acid (DTNB), 300 μM acetyl-CoA, 0.1% Triton X 100) and specific activity was determined by measuring conversion of DNTB to 2-nitro-5-benzoic acid (TNB) at a wavelength of 405 nm after adding 500 μM oxaloacetate as substrate.

DNA was isolated from cells using the DNeasy Blood & Tissue Kit (Qiagen, Hilden, Germany). Content of both nuclear and mitochondrial DNA was quantified by qRT-PCR using specific primers as described before [20]. Mitochondrial/nuclear ratio was calculated as a relative measure of mitochondrial DNA content.

### 2.9. Subcellular fractionation

Cells were scraped into fractionation buffer (20 mM HEPES pH 7.4, 250 mM sucrose, 10 mM KCl, 1.5 mM MgCl2, 1 mM EDTA, 1 mM DTT, Complete Protease Inhibitors (Roche) and homogenized by passing them through a 25 G needle 10 times. Nuclei were separated by centrifugation at 750g for 5 min at 4 °C. The supernatant was centrifuged at 10,000 g for 20 min at 4 °C to obtain the mitochondrial fraction. The supernatant comprising the cytosolic fraction was precipitated by adding four volumes of ice-cold acetone, incubation for 60 min at −20 °C, and centrifugation at g for 30 min at 4 °C. The protein precipitates were dissolved in equal amounts of 1X Bolt LDS Sample Buffer (ThermoFisher Scientific) and subjected to Western Blot analysis.

### 2.10. Cellular respiration and extracellular acidification

Cells were plated in 96-well cell culture microplates (XFe96, Agilent Technologies, Santa Clara, USA) and differentiated into adipocytes for 14 days. One day before measurement, the culture medium was changed to insulin-free medium. On the day of measurement, the cells were incubated for 1 h in bicarbonate-free DMEM medium containing 5 mM HEPES, 10 mM glucose, 1 mM pyruvate, 2 mM GlutaMaxx, and 1% bovine serum albumin. Oxygen consumption and extracellular acidification rates (OCR and ECAR) were measured simultaneously using a Seahorse XFe96 Flux Analyzer (Agilent Technologies). To mimic thermogenic activation, the cells were treated with 0.5 mM dibutyryl cyclic adenosinomonophosphate (cAMP). Uncoupled (proton leak) respiration was profiled by injecting 2μM oligomycin (inhibiting the ATP synthase), and full substrate oxidation capacity was determined by injecting 4μM carbonylcyanide-p-trifluoromethoxyphenylhydrazone (FCCP, a chemical uncoupler). Non-mitochondrial respiration was determined by injecting 1.5 μM antimycin A and 1.5 μM rotenone (inhibiting electron flux through complex I and III). OCRs were determined by machine algorithms and plotted against time. Data were either normalized for lipid content by staining with NileRed or to cell content by staining with JanusGreen [21].

### 2.11. Mitochondrial membrane potential

To estimate mitochondrial membrane potential, the cells were stained with 30 nM MitoTracker Red CMXRos (ThermoFisher Scientific) for 30 min at 37 °C. Subsequently, the cells were washed with PBS, trypsinized, and centrifuged. The cell pellets were resuspended in PBS and the cells were analyzed with the FACSCalibur (BD Biosciences, Heidelberg, Germany).

### 2.12. Statistics

If not stated otherwise, data from three independent experiments are expressed as mean ± standard error of the mean (SEM). For statistical comparison, analysis of variance (ANOVA) or t-tests were used as indicated. A p < 0.05 was considered statistically significant. GraphPad Prism version 7.01 (GraphPad Software, San Diego, USA) was used for all analyses.

### 3. Results

#### 3.1. Elevated basal glucose uptake in mouse brown vs white adipocytes

To determine a difference in the capacity to take up glucose on a cellular level, we isolated stromal-vascular cells from WAT and BAT and subjected them to adipogenic differentiation in vitro. After 10 days of differentiation, the murine cells showed histological signs of adipocytes, i.e. round shape and incorporated lipid droplets (Fig. 1A and B), significantly higher expression of UCP1 and CIDEA in BAT vs WAT cells, and higher expression of HOXC8 in WAT cells (Fig. 1C). To our knowledge, specific glucose uptake rates have never been experimentally compared between isolated BAT vs WAT cells. In line with previous assumptions on heterogeneous bulk tissue in vivo, brown adipocytes in vitro display the expected significantly higher basal glucose uptake rates per cell (cpm/10^5 adipocytes), compared to white adipocytes (Fig. 1D).

#### 3.2. Generation of UCP1-deficient human adipocytes

To investigate whether UCP1 is involved in the regulation of cellular glucose uptake and utilization, we generated adipocytes with either loss or gain of UCP1-functionality. As model system we used human Simpson-Golabi-Behmel syndrome (SGBS) cells, which were derived from subcutaneous WAT of an infant affected by the eponymous syndrome [15]. These cells undergo adipogenic differentiation in the presence of a PPARγ agonist and express low levels of UCP1 at late adipogenesis [20].

We generated UCP1-deficient cells using the CRIPSR-Cas9 system. UCP1 protein was completely absent in bulk cultures of knockout adipocytes as assessed by Western Blot analysis (Fig. 2A), with no difference in adipogenic differentiation, as judged by protein expression of adipogenic markers (Fig. 2A). UCP1-deficient adipocytes displayed a significantly reduced proton leak respiration by 17% compared to control cells (Fig. 2B). As expected from recent mouse data in vivo [14], the basal glucose uptake (Fig. 2C) as well as the extracellular acidification rates (ECARs) were not different between control and UCP1-deficient cells (Fig. 2D). Collectively, the knockout of UCP1 imposed only minor effects on the glucose metabolism of white adipocytes.
3.3. Generation of UCP1-overexpressing human preadipocytes

Next, we aimed to increase the expression of UCP1 in SGBS cells using a lentiviral system. Successful overexpression in the undifferentiated preadipocyte state was demonstrated by Western Blot (Fig. 3A), with no UCP1 detectable in empty vector-transduced control cells. Following subcellular fractionation, UCP1 was found to be localized in mitochondria as expected (Fig. 3B). UCP1 function was verified by assessing mitochondrial respiration by plate-based respirometry using a Seahorse extracellular flux analyzer. UCP1 overexpression had no statistically significant impact on basal respiration and FCCP-induced maximal substrate oxidation capacity (Fig. 3C). Importantly, UCP1-overexpressing cells have markedly increased proton leak respiration by 440% and reduced ATP-linked respiration rates by 57% (Fig. 3D). Taken together, this set of data demonstrates that UCP1 is fully functional, even in preadipocytes.

3.4. UCP1 overexpression does not interfere with adipogenesis or mitochondrial biogenesis

Control and UCP1-overexpressing preadipocytes were subjected to adipogenic differentiation. Both cell types displayed similar morphological features including cell shape, lipid droplet number, and lipid droplet size (Fig. 4A). In line, the rate of adipogenic differentiation (Fig. 4C) and triglyceride levels (Fig. 4D) were comparable. In day 14 adipocytes, UCP1 protein was still overexpressed compared to control cells (Fig. 4B), while the mRNA expression of the adipogenic marker genes PPARG, FASN, and LEP (Fig. 4E) were comparable. In addition, markers of adipocyte browning (e.g. PGC1A, PRDM16 and CIDEA) were not affected by UCP1 overexpression (Fig. 4E).

Assessing mitochondrial activity revealed an increased basal respiration in UCP1-overexpressing adipocytes compared to control cells by 50% (Fig. 4F and G), mainly due to the increase in proton leak respiration, which was 200% higher compared to control cells. Treatment with dibutyryl-cAMP strongly enhanced respiration in UCP1-overexpressing adipocytes by 230%. In contrast, cAMP had only a minor effect (30% increase) on respiration in control cells (Fig. 4F and G). UCP1-overexpressing adipocytes revealed a reduced mitochondrial membrane potential (Fig. 4H), further corroborating UCP1 activity. Intriguingly, the mitochondrial content was not affected by UCP1 overexpression as determined by protein expression of components of the electron transport chain (Fig. 4I), the activity of citrate synthase

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**Fig. 1. Comparison of glucose uptake in murine in vitro differentiated white and brown adipocytes.** (A) Stromal-vascular cells isolated from inguinal WAT and interscapular BAT were differentiated for 10 days into mature adipocytes. (B) mRNA expression of the BAT-specific markers UCP1 and CIDEA and the WAT-specific marker HOXC8 was measured by qRT-PCR. (C) Basal uptake of 14C-deoxy-glucose was determined in adipocytes after serum starvation overnight, cpm values were normalized to adipocyte number. Data (n = 3) are shown as means + SEM, *p < 0.05, **p < 0.01 (Student’s t-test). (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)

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**Fig. 2. CRISPR/Cas9-mediated UCP1 knockout in human adipocytes.** UCP1-deficient cells were generated by co-expressing Cas9 and specific sgRNAs targeting UCP1 in SGBS preadipocytes. (A) The cells were differentiated into mature adipocytes. The knockout of UCP1 and the protein levels of adipogenic markers adiponectin (AdipoQ), peroxisome proliferator-activated receptor gamma (PPARG), and perilipin A (PLIN) were confirmed by Western Blot. NC, non-targeting control. (B) Oxygen consumption rates were determined in a plate-based respirometer. ATP: oligomycin-sensitive OCR; leak: oligomycin resistant OCR. (C) Basal glucose uptake was determined by 14C-deoxy-Glucose incorporation. (D) Extracellular acidification rates were determined as in (B). Data obtained from both negative controls and UCP1 KO cell types were merged. Data of 3–4 independent experiments are shown as mean + SEM. ***, p < 0.01 (Student’s t-test).
14C-deoxy-glucose in basal medium without further stimulation, to drive cellular glucose uptake in adipocytes, cells were treated with 3.5. Elevated UCP1 levels enhance basal glucose uptake and glycolytic flux (Fig. 4J), as well as the relative mitochondrial DNA content (Fig. 4K).

3.5 Elevated UCP1 levels enhance basal glucose uptake and glycolytic flux in adipocytes

To investigate whether elevated levels of UCP1 are solely sufficient to drive cellular glucose uptake in adipocytes, cells were treated with 14C-deoxy-glucose in basal medium without further stimulation, followed by measurements of glucose uptake. Glucose uptake was significantly increased in UCP1-overexpressing compared to control adipocytes by 40% (Fig. 5A), accompanied by enhanced production of lactate (Fig. 5B). These data suggest that UCP1 overexpression is sufficient, without further exogenous stimulation, to induce glucose uptake and to enhance glycolytic flux in adipocytes.

We next tested whether inhibition of glucose metabolism inhibits UCP1 activity. Pretreatment with 2-deoxy-glucose blunted glycolytic rates, as judged by extracellular acidification rates (Fig. 5C). Interestingly, the corresponding OCR values were still significantly higher in UCP1-overexpressing adipocytes upon 2-deoxy-glucose treatment, indicating that these effects persist in the absence of glucose. Pretreatment with 2-deoxy-glucose blunted glycolytic rates, as judged by extracellular acidification rates (Fig. 5C). Interestingly, the corresponding OCR values were still significantly higher in UCP1-overexpressing adipocytes upon 2-deoxy-glucose treatment, indicating that these effects persist in the absence of glucose.

4. Discussion

Since the early 2000s, 18F-FDG-PET/CT has been used for imaging BAT activity, clearly demonstrating that cold-activated BAT takes up glucose in amounts comparable to other high-glucose consuming tissues such as the brain [2,12,13,23]. It is commonly assumed that this high glucose uptake contributes to glycemic control, at least upon cold exposure. This is supported by studies demonstrating in subjects with higher BAT content lower glucose levels [24] and improved insulin sensitivity [4]. Weir et al. have recently shown that the increased glucose uptake by BAT does not necessarily require activation by cold, but that it also occurs at 25°C. Using microdialysis, they convincingly demonstrated that BAT takes up significantly more glucose and releases more lactate than WAT [25], indicating a higher glycolytic activity even under basal conditions. These findings underline the widely accepted notion that BAT acts as a glucose sink. To the best of our knowledge, however, glucose uptake has never been assessed directly by comparing white and brown adipocytes in vitro. In confirmation of the prevalent notion, we exemplify for the first time that brown adipocytes take up significantly more glucose than white adipocytes.

As recently reported, inhibiting glucose uptake has pronounced effects on both basal and isoproterenol-stimulated respiration in brown adipocytes [9]. This suggests that glucose and its metabolic breakdown products are essential for thermogenesis [9]. There are, however, also data from UCP1 knockout mice indicating that glucose uptake into BAT is independent of UCP1-dependent thermogenic activity [14,26]. In contrast, some indirect hints for UCP1-dependency of glucose metabolism were observed in mice with ectopic UCP1 expression in epididymal WAT [10]. Ectopic UCP1 expression significantly reduces blood glucose levels on both normal and high-fat diets. Interestingly, surgical and pharmacological denervation of WAT were ineffective at abrogating the effects of UCP1 overexpression, indicating that these effects persist independent of neuronal innervation [10].

Although mechanistic aspects of UCP1 function have been studied extensively in model systems such as isolated liposomes [27], yeast [28], mouse brown adipocytes [29,30], Chinese hamster ovary (CHO) cells [31], human embryonic kidney (HEK) cells [32], and HeLa cells [33], these studies were never focused on and thus, never revealed whether UCP1 activity per se is sufficient to enhance glycolytic flux, in particular in adipocytes. One study using in 3T3-L1 cells reports increased glucose uptake upon ectopic UCP1 expression, but did not experimentally prove UCP1 function [34].
Here, we introduce human SGBS preadipocytes and adipocytes that stably overexpress UCP1 as a unique, homologous model system to study physiological and pathophysiological relevant aspects of UCP1 function in detail. We demonstrate that UCP1 is functionally active in SGBS preadipocytes and adipocytes under basal as well as lipolytic conditions. This suggests that the activity of ectopically expressed UCP1 in intact white adipocytes is sufficient to increase proton leak, without further induced lipolysis by adrenergic induction. This is not intuitively expected considering that UCP1 per se is not active, at least not in isolated mitochondria in the absence of free fatty acids [30]. Moreover, while it was previously suggested that UCP1 overexpression affects mitochondrial integrity [30], no signs of mitochondrial dysfunction or

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cell viability were observed in our human adipocyte model system. Neither electron transport chain complexes, nor citrate synthase activity, nor relative mitochondrial DNA content were affected, emphasizing that forced UCP1 expression may be without adverse effects. Importantly, cellular proliferation (data not shown) and adipogenic differentiation were not affected in UCP1-overexpressing cells. Thus, we suggest that the constitutive activity of UCP1 in white adipocytes, probably due to free fatty acids from basal lipolysis, may offer the potential to improve glucose metabolism.

In line with previous publications [14], knocking out UCP1 did not alter glucose utilization, but overexpression impacts glucose uptake and lactate production in adipocytes, demonstrating enhanced glycolytic flux. Of note, the relative increase in glucose uptake rates is comparable to murine white vs brown adipocyte differences. Interestingly, increased lactate production supports a model of aerobic glycolysis that should operate independent of UCP1-induced glucose oxidation. Thus, the UCP1-dependent reduction of mitochondrial ATP production is possibly driving glycolytic ATP production to maintain cellular homeostasis. This integrates well with findings from human studies showing increased glycolytic activity in BAT [25]. We want to point out that undifferentiated preadipocytes overexpressing UCP1 may contribute to differences seen in glucose uptake. However, the majority of cells in the cell culture dish were differentiated adipocytes.

Inhibition of glycolysis by 2-deoxy-glucose partially inhibited respiration, but did not abolish the difference in oxygen consumption between UCP1-overexpressing and control adipocytes. This suggests that respiratory activity in our cell model is partially fueled by another substrate, such as lipids and amino acids. It has recently been proposed...
that glucose uptake in brown adipocytes is mediated either by GLUT1 [35] or both GLUT1 and GLUT4 [9]. In our model system of white adipocytes, selective inhibition of GLUT1 completely eradicated the differences in glucose uptake between UCP1-overexpressing and control cells identifying GLUT1 as an important gatekeeper of glycolytic flux. The mRNA expression levels of glucose transporters were not altered by UCP1 overexpression suggesting an alternative level of regulation, e.g. an increased translocation of GLUT1 to the cell surface as described earlier [35].

Taken together, we demonstrate that brown adipocytes take up significantly more glucose under basal conditions than white adipocytes in vitro underlining that brown adipocytes are in general more active metabolically. We show that UCP1 and the insulin-independent glucose transporter GLUT1 are crucially involved in establishing this glucose-consuming phenotype of adipocytes. With this study we furthermore provide a unique cellular model system of UCP1-expressing human adipocytes. It allows detailed studies on human UCP1 function and its impact on cell metabolism, and is suitable to screen for UCP1 modulators, thus facilitating the discovery of new drugs to correct glucose metabolism.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.redox.2019.101286.

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