UCLA

UCLA Previously Published Works

Title

Enhanced hexose-6-phosphate dehydrogenase expression in adipose tissue may contribute to diet-induced visceral adiposity.

Permalink

<https://escholarship.org/uc/item/2df9c372>

Journal

International journal of obesity (2005), 42(12)

ISSN

0307-0565

Authors

Liu, Limei
Wang, Ying
Wang, Jian
et al.

Publication Date

2018-12-01

DOI

10.1038/s41366-018-0041-1

Peer reviewed

Enhanced hexose-6-phosphate dehydrogenase expression in adipose tissue may contribute to diet-induced visceral adiposity

Limei Liu^{1,2} · Ying Wang¹ · Jian Wang^{1,3} · Yunzhou Dong⁴ · Scarlett Chang¹ · Xiwen Liu¹ · Kabirullah Lutfy^{2,5} · Hong Chen⁴ · Theodore C. Friedman¹ · Meisheng Jiang⁶ · Yanjun Liu^{1,3}

Received: 15 August 2017 / Revised: 30 November 2017 / Accepted: 27 December 2017
© Macmillan Publishers Limited, part of Springer Nature 2018

Abstract

Background Visceral fat accumulation increases the risk of developing type 2 diabetes and metabolic syndrome, and is associated with excessive glucocorticoids (GCs). Fat depot-specific GC action is tightly controlled by 11 β -hydroxysteroid dehydrogenase (11 β -HSD1) coupled with the enzyme hexose-6-phosphate dehydrogenase (*H6PDH*). Mice with inactivation or activation of *H6PDH* genes show altered adipose 11 β -HSD1 activity and lipid storage. We hypothesized that adipose tissue *H6PDH* activation is a leading cause for the visceral obesity and insulin resistance. Here, we explored the role and possible mechanism of enhancing adipose *H6PDH* in the development of visceral adiposity in vivo.

Methods We investigated the potential contribution of adipose *H6PDH* activation to the accumulation of visceral fat by characterization of visceral fat obese gene expression profiles, fat distribution, adipocyte metabolic molecules, and abdominal fat-specific GC signaling mechanisms underlying the diet-induced visceral obesity and insulin resistance in *H6PDH* transgenic mice fed a standard of high-fat diet (HFD).

Results Transgenic *H6PDH* mice display increased abdominal fat accumulation, which is paralleled by elevated lipid synthesis associated with induction of lipogenic transcription factor C/EBP α and PPAR γ mRNA levels within adipose tissue. Transgenic *H6PDH* mice fed a high-fat diet (HFD) gained more abdominal visceral fat mass coupled with activation of GSK3 β and induction of XBP1/IRE1 α , but reduced pThr³⁰⁸ Akt/PKB content and browning gene *CD137* and *GLUT4* mRNA levels within the visceral adipose tissue than WT controls. HFD-fed *H6PDH* transgenic mice also had impaired insulin sensitivity and exhibited elevated levels of intra-adipose GCs with induction of adipose 11 β -HSD1.

Conclusion These data provide the first in vivo mechanistic evidence for the adverse metabolic effects of adipose *H6PDH* activation on visceral fat distribution, fat metabolism, and adipocyte function through enhancing 11 β -HSD1-driven intra-adipose GC action.

Introduction

Obesity is a growing global public health problem that affects billion people worldwide. Overweight, especially visceral obesity has been recognized as the most dangerous risk factor for developing type 2 diabetes mellitus (T2DM),

These author contributed equally: L. Liu, Y. Wang and J. Wang

✉ Yanjun Liu
dryanjunliu@hotmail.com

¹ Division of Endocrinology, Metabolism & Molecular Medicine, Charles R. Drew University of Medicine & Sciences, David Geffen School of Medicine, University of California, 1731 E. 120th St, Los Angeles, CA 90059, USA

² Department of Endocrinology & Metabolism, Shanghai Jiaotong University Affiliated Sixth People's Hospital, Shanghai Diabetes Institute, Shanghai, People's Republic of China

³ Department of Neonatology, The First Hospital of Jilin University, Changchun, People's Republic of China

⁴ Vascular Biology Program, Boston Children's Hospital, Harvard Medical School, Boston, MA, USA

⁵ College of Pharmacy, Western University of Health Sciences, Pomona, CA, USA

⁶ Department of Molecular and Medical Pharmacology, David Geffen School of Medicine, University of California, Los Angeles, CA 90095, USA

metabolic syndrome, and cardiovascular diseases [1, 2]. Hormone glucocorticoids (GCs) regulate adipose tissue differentiation, function, distribution, insulin sensitivity, and in excess, will lead to the pathological state. Patients with GC excess (Cushing's syndrome) develop pronounced abdominal visceral fat deposition associated with the progress of metabolic syndrome with dyslipidemia, insulin resistance, T2DM, and hypertension [3]. In target cells, GC action is regulated by 11 β -hydroxysteroid dehydrogenase type 1 (11 β -HSD1) at pre-receptor levels [4]. 11 β -HSD1 is an intracellular endoplasmic reticulum (ER) lumen-resident enzyme, and its major function in vivo is the NADPH-dependent conversion of inert cortisone (11-dehydrocorticosterone in rodents) to active cortisol (corticosterone) and therefore amplifies tissue GC production. Increasing GC production in adipose tissue caused by 11 β -HSD1 induces pronounced visceral fat accumulation and insulin resistance in mice [5]. Clinical observations also have shown that adipose 11 β -HSD1 activity is increased in typical obese humans and patients with metabolic syndrome [6, 7]. In contrast, mice lacking 11 β -HSD1 are resistant to diet-induced obesity and metabolic syndrome [8]. These results implicate the importance of adipose 11 β -HSD1-driven local GC production in the progression of visceral obesity and T2DM.

11 β -HSD1 amplification of intracellular active GC production requires a high [NADPH]/[NADP⁺] ratio, which can be crucially provided by the enzyme hexose-6-phosphate dehydrogenase (*H6PDH*) [9, 10]. *H6PDH* is an enzyme localized to the ER lumen that converts glucose-6-phosphate (G6P) and NADP to produce NADPH, thereby driving 11 β -HSD1 reductase activity and consequently intracellular GC production [11, 12]. Patients with gene mutations in *H6PDH* show defects in 11 β -HSD1 activity and GC production [13]. Consistent with the physiological relevance of *H6PDH* for 11 β -HSD1, *H6PDH* knockout mice are unable to reproduce active cortisol from cortisone by 11 β -HSD1, and have improved lipogenesis and insulin sensitivity [14]. These results demonstrate that *H6PDH* is required for 11 β -HSD1 amplifying local GC signaling related to the progression of obesity and insulin resistance.

In contrast, subcutaneous *H6PDH* mRNA levels are elevated in the worsening metabolic disorders associated with increased local cortisol production in obese patients with the metabolic syndrome [15]. Similarly, there is increased expression of *H6PDH* in abdominal visceral and subcutaneous fat in patients with T2DM as well as in obese diabetic rodents compared to normal controls [16–18]. These observations imply that fat-specific elevation of *H6PDH* expression may represent an emerging mechanism in the pathogenesis of metabolic syndrome and thus agents that reduce endogenous 11 β -HSD1 and consequently can improve or delay the progress of T2DM and obesity. We

recently reported that transgenic mice overexpressing *H6PDH* in adipose tissue (aP2-*H6PDH* Tg mice) exhibit feature of metabolic syndrome with abdominal region fat accumulation, fasting hyperglycemia, dyslipidemia with lipolysis, and elevated intra-adipose GC levels [19], implicating that adipose tissue *H6PDH* is able to couple the regulation of cellular GC metabolism and fat mass linked to the pathogenesis of obesity and insulin resistance. Thus, the aP2-*H6PDH* transgenic mouse may serve as a good model to further explore the potential impact of adipose *H6PDH* activation in the etiology of visceral adiposity and the adverse metabolic consequence relative to visceral obesity/metabolic syndrome in humans. However, it is unclear whether enhanced adipose tissue *H6PDH* expression would render the transgenic mice even more susceptible to obesity and insulin resistance when fed a Western-type diet.

In order to gain the new insight of these important issues, we conducted in vivo studies to investigate the key molecular mechanism underlying the adverse effects of enhanced adipose *H6PDH* on abdominal fat deposition by analysis of lipogenic and adipogenic metabolism profile of visceral adipose tissue in transgenic aP2-*H6PDH* mice. We also investigated at the molecular levels whether adipose tissue *H6PDH* activation would exacerbate visceral adiposity and insulin resistance in obesity by characterization of obese gene expression profiles, adiposity signaling molecules, visceral fat distribution, browning marker gene expression, adipose metabolic parameters, and abdominal visceral fat-specific GC action mechanisms underlying the diet-induced obese syndrome in HFD-fed transgenic aP2-*H6PDH* mice. Our data provide the first in vivo mechanistic evidence for the adverse metabolic effects of enhanced adipose *H6PDH* on fat distribution, adipose function, and homeostasis through enhancing 11 β -HSD1-driven intra-adipose GC action.

Materials and methods

Animals and experimental models

Adult male transgenic mice overexpressing *H6PDH* under the adipocyte fatty acid-binding protein (*aP2*) gene (aP2/*H6PDH* Tg) and their age-matched (10 weeks) wild-type (WT) C57bl/6 control mice were bred and genotyped, as previously described [19]. Eight animals have been used for each experimental group, the numbers of animal used were determined to achieve desired power of 0.8. Animals were housed in standard SPF conditions under a 12:12-h light–dark cycle with water ad libitum. Mice were maintained on either a control diet (Research Diets D12328) or a HFD (45 % energy from Fat, Research Diets D12451; Research Diets Inc., New Brunswick, NJ, USA) for 4 or

14 weeks. For each genotype, mice are divided randomly into the control diet or HFD group. Food intake and body weights were recorded weekly. At the end of the experiments, blood samples were collected, and adipose fat depots were weighed, dissected, and frozen rapidly in liquid nitrogen. Animal protocols were approved by the Institutional Animal Care and Use Committee of the Charles R. Drew University (Los Angeles, California, USA). Blood samples were analyzed for insulin and FFA levels using ELISA kit (Abcam, Cambridge, MA, USA), as previously described [18].

Glucose and insulin tolerance tests

After 20 weeks on a control or a HFD, aP2/H6PDH transgenic and wild-type mice were fasted overnight, and blood samples were taken at the indicated time points following the intraperitoneal injection (i.p) with glucose (1.0 g/kg D-glucose). For the insulin tolerance test, mice were injected with insulin after fasting for 6 h (i.p, 0.75 units/kg; Novolin R; Eli Lilly, Indianapolis, IN), and the glucose levels were measured at the indicated time points.

Glucose uptake in adipose tissue

Glucose uptake was measured as previously reported [20]. Briefly, fresh epididymal fat pads were cut into less than 5-mg pieces and then incubated with 0.5% BSA at 37 °C for 30 min in DMEM. Tissues were washed with 1% BSA in Ringer bicarbonate-HEPES (KRBH) buffer (Sigma K4002) and incubated in KRBH solution with 0.1% BSA in the absence or presence of 100 nM insulin for 5 min at 37 °C. Tissues were further incubated for 30 min in the same solution with adding 10 mmol/L glucose and 2.0 μ Ci 2- 3 H] DG (NEN Life Science, Boston, MA). The uptake reaction was terminated, and radioactivity was measured by using a liquid scintillation counter.

RNA extraction and quantitative real-time RT-PCR

Total RNA was isolated from tissues using the TRIzol reagent (Invitrogen), and the first-strand cDNA was synthesized from 2.0 μ g mRNA using a commercially available kit (Thermo Fisher Scientific). The primers were designed with Primer express software 2.0 and were as follows: mouse 11 β -HSD1 (F: 5'-CCTTGGCCTCATAGACACAGAAAC-3'; R: 5'-GGAGTCAAAGGCGATTTT CA-3'), *H6PDH* (F: 5'-TGGCTACGGGTGTTTTTGAA-3'; R: 5'-TA TACAGGTAC ATCTCCTCTTCCT-3'), *ACC* (F: 5'-TGTAATCTGGCTGCATCCATTAT-3'; R: 5'-TGGTAGACTGCCCGTGTGA-3'), *ACL* (F: 5' ATGCCAAGACCA TCCTCTCACT-3'; R: 5'-TCTACAATGCCCTTGAAGGT-3'), *GR* (F: 5'

-TGCTATGCTTTGCTCCTGAT CTG-3';R: 5'-TGTCAGTTGATAAAACCGCTGC-3'), *FOXO1* (F: 5'-GGACAGCCGC GCAAGACCAG-3'; R: 5'-TTGAATTCTTCCAGCCCCGCCGA-3'), *SIRT1* (F: 5'-AATC CCGGACTTCAGATCCC C-3';R: 5'-CAACATGAAAAAGGGCTTGGG-3'), *TXB 1*(F: 5'-CGCTGTGGGACGAGTTCAAT-3'; R:5'-CGGCCATGGGATCCATT-3'), *CD137* (F: 5'-TTGGGAACATTTAATGACCAGA-3'; R: 5'-TCCCGGTCTTAAGCACAGAC-3'), *IRE α* (F: 5'-CCTACAAGAGTATGTGGAGC-3'; R: 5'-GGTCTCTGTGAA-CAATGTT GAGAG-3'), *XBPI* (F: 5'-GGATTTGGAA-GAAGAGAACCACAA-3'; R: 5'-CCGTGAG TTTTCTCCCGTAAAA-3') and *G6PT* (F: 5'-AGGCCTTG-TAGGAAGCATTGC-3'; R: F: 5' TCACCGTTACTCG-GAAGAGGAA-3'). RT-PCR was performed using SYBR kits in the ABI Prism 7700 System (Thermo Fisher Scientific). Relative mRNA expression levels were calculated using the comparative CT method and were expressed as a relative value after normalization to 18S.

Oil red staining with mounting medium DAPI

Fat tissue was fixed in freshly prepared 4% paraformaldehyde, then imbedded in O.C.T., and frozen sections were cut and stained with Oil Red O (Sigma, St. Louis, MO). Cells were mounted with Vestashield mounting solution with 4', 6'-diamidino-2-phenylindole (Vector Laboratories) and visualized under a fluorescence microscope.

Adipose tissue corticosterone assays

Corticosterone levels were measured in adipose tissue as previously described [21–23]. Briefly, epididymal and mesenteric fat depots were isolated, weighed and homogenized in dPBS buffer (PH 7.4) by a tissue solubilizer. The adipose tissue steroids were extracted twice with ethyl acetate (5 mL/g adipose tissue). The extractions were evaporated and the residues were dissolved in sample analysis buffer. Adipose corticosterone levels were measured using a corticosterone EIA kit (Oxford Biomedical Research). 11 β -HSD1 activity was evaluated by the addition of 1 mM NADPH and 250 nM 11-DHC with [3 H]11-DHC as the tracer to microsomes in KRB solution at 37 °C for 1–2 h. Steroids were separated and the conversion of [3 H]11-DHC to [3 H]CORT was determined by scintillation counting of radioactivity, as previously described [19].

Protein extraction and Western blot analysis

Adipose tissues were homogenized on ice and protein concentrations were determined by Bradford assay as previously described [18]. Adipose proteins were fractionated

Fig. 1 Lipogenic gene ACC and ACL mRNA and protein expression in epididymal fat (Ep fat), mesenteric fat (Mes fat) and subcutaneous fat (Sub fat) of aP2-H6PDH (■) or WT (□) mice. **a, d, g** Relative expression of ACC and ACL mRNA levels were measured by real-

time RT-PCR and normalized to 18S ($n = 8$). **b, c, e, f, h, i** Relative expression levels of t-ACC, t-ACL, pSer⁷⁹ ACC, and pSer⁴⁵⁵ ACL proteins were normalized to GAPDH. Data are means $\pm$ SE of 7–8 mice/group. * $P < 0.01$ vs. WT controls; ** $P < 0.001$ vs. WT controls

on 4–15% acrylamide SDS-PAGE gels (BioRad, Hercules, CA, USA) and transferred onto nitrocellulose membranes for the Western analysis of pSer⁷⁹ ACC, pSer⁴⁵⁵ ACL, p-threonine³⁰⁸ kinase Akt/PKB, IR, IRS-1, *GLUT4*, pSer⁹ GSK3 β , pSer⁷²⁴ IRE α , ACC, ACL, XBP1, p-IRE α , IRE α , and GAPDH (Cell Signaling, Danvers MA). Proteins signals were quantified by Eagle Eye II Quantitation System (Stratagene, CA, USA).

Statistical analyses

All data are expressed as the mean $\pm$ SEM. Data were compared using an unpaired Student's *t*-test. To compare multiple groups, one-way ANOVA was used. When ANOVA revealed significant differences, then group comparisons were performed using the Newman–Keul's post hoc test. A $P < 0.05$ was considered significant.

Fig. 2 Alterations of obese gene expression in epididymal fat (Ep fat), mesenteric fat (Mes fat) and subcutaneous fat (Sub fat) between aP2-H6PDH (■) or WT (□) mice. **a, c, e** Relative expression levels of lipid metabolic gene FAS and PEPCK, and lipogenic gene transcription C/EBP α , PPAR γ , and SREBP mRNA levels in epididymal fat (**a**), mesenteric fat (**c**), and subcutaneous fat (**e**) were normalized to 18 S. **b, d, f** Relative expression of leptin, FOXO1, and SIRT1 mRNA levels in epididymal fat (**b**), mesenteric fat (**d**) and subcutaneous fat (**f**) of aP2-H6PDH Tg (■) or WT (□) mice. Data are mean $\pm$ SE from six to eight mice per group. * P < 0.01 vs. WT controls; ** P < 0.005 vs. WT controls

Results

Transgenic aP2-H6PDH mice have altered lipogenic gene expression in adipose tissue

Adipose *H6PDH* expression is associated with abdominal fat storage [14, 19]. To ascertain whether elevated adipose *H6PDH* expression influences lipogenic profile, we

assessed the expression of lipid synthase acetyl-CoA carboxylase (ACC) and ATP-citrate lyase (ACL), the major markers of lipid content in adipose tissue of aP2/H6PDH Tg mice. As shown in Fig. 1, epididymal fat mRNA levels encoding ACC and ACL in aP2-H6PDH Tg mice were increased compared with that of the WT controls (P < 0.01; Fig. 1a). Furthermore, epididymal fat total ACC and ACL protein levels were increased with elevated (1.9-fold and

Fig. 3 ER stress response and the relative alterations of Akt/PKB and GSK3 β gene expression in epididymal fat (Ep fat) of aP2-H6PDH Tg (■) and their WT controls (□). **a, b** Relative expression of IRE α , XBP1, AKT, and GSK3 β mRNA levels measured by real-time RT-PCR and normalized to 18S. **c, e** Western blotting analyses were performed to compare the expression levels of t-IRE α , XBP1, and pSer⁷²⁴ IRE. **d, f** Relative adipose p-Thr³⁰⁸ Akt and pSer⁹ GSK3 β protein was standardized to GAPDH. * P < 0.01 vs. WT controls; ** P < 0.02 vs. WT controls

2.8-fold respectively) phosphorylated levels of Ser⁷⁹ACC and Ser⁴⁵⁵ ACL in aP2-H6PDH mice compared with WT controls (Fig. 1b, c). Similarly, higher ACC and ACL mRNA levels (Fig. 1d) with elevated ACL, pSer⁷⁹ACC, and pSer⁴⁵⁵ ACL contents were observed in mesenteric visceral fat from the aP2-H6PDH Tg mice, but with unchanged total ACC protein levels compared to that of WT controls (Fig. 1e, f). In addition, subcutaneous adipose ACC and ACL mRNA levels were higher in aP2-H6PDH Tg mice compared with WT controls (Fig. 1g). Moreover, ACC and ACL protein expression levels were markedly elevated with increased levels of pSer⁷⁹ACC and pSer⁴⁵⁵ ACL in aP2-H6PDH mice compared with WT controls (Fig. 1h, i). Consistent with the increased lipid biosynthesis in adipose tissue, fatty acid synthase (FAS) mRNA levels were increased by 2-fold and 2.8-fold, respectively, in epididymal and mesenteric visceral fat in aP2-H6PDH Tg mice compared with WT controls (Fig. 2a, c). Real-time RT-PCR analysis also showed that FFA re-esterification enzyme

PEPCK mRNA for TG synthesis, were markedly elevated in total adipose tissue from aP2-H6PDH Tg (Fig. 2a, c, e; P < 0.01). These data indicate that aP2-H6PDH Tg mice had increased adipose lipid content.

The effects of enhanced adipose tissue H6PDH expression on obese gene expression in fat tissue

To investigate the mechanism underlying the differential changes in lipogenesis in aP2-H6PDH Tg and WT mice, the mRNA encoding lipogenic gene transcripts C/EBP α , PPAR γ , and SREBP were analyzed. C/EBP α mRNA levels were increased by 4-fold in epididymal fat, 3.6-fold in mesenteric fat (Fig. 2a, c), and 3-fold in subcutaneous fat (Fig. 2e) in aP2-H6PDH mice compared with controls. Adipose PPAR γ , a key factor regulating fat distribution mRNA expression were increased by 2.5-fold in epididymal fat, 2-fold in mesenteric fat with a more than 6-fold in subcutaneous fat of the aP2-H6PDH Tg mice compared

Fig. 4 Body weight and visceral fat mass of aP2-H6PDH transgenic and WT mice. **a** Body weight gain of aP2-H6PDH and WT mice fed chow diet (C) or high-fat diet (HFD) (○, non-Tg mice fed chow diet; ■, Tg mice fed chow diet; △, non-Tg mice fed HFD; ●, Tg mice fed HFD, $n = 8-10$ mice/group). **b** Ep fat and Mes fat mass of *H6PDH* mice fed control chow diet or HFD. **c** Lipid droplet accumulation was monitored by fluorescent microscopy using Oil Red O staining with DAPI. **d, e** Relative mesenteric fat obese gene mRNA levels in aP2-H6PDH Tg (▨) and their WT controls (■) and WT mice fed HFD. **f** Relative alterations of Akt, p-Thr³⁰⁸Akt and pSer⁹ GSK3β content in Mes fat of mice fed HFD. * $P < 0.01$ and # $P < 0.02$ vs. WT controls; ** $P < 0.02$ vs. HFD-fed controls; † $P < 0.02$ and ‡ $P < 0.01$ vs. HFD-fed

WT controls. **g** Schematic diagram representing the consequences of enhancing adipose *H6PDH* expression. *H6PDH* activation provides the fuel for the production of 11β-HSD1-driven intra-adipose CORT and results in fat accumulation through: (1) CORT stimulation of IREα/XBP1 pathway activates C/EBPα and PPARγ expression, facilitating lipogenesis and TG synthesis, (2) C/EBPα can activate 11β-HSD1, further enhancing CORT availability and (3) CORT-mediated suppression of p-Akt induces insulin resistance and activation of GSK3β signaling that is required for adipogenesis. Additionally, GSK3β activation can promote CORT to bind/activate GR and further amplifying GC action in adipose tissue

with WT controls ($P < 0.01$; Fig. 2a, c, e). Moreover, SREBPα mRNA expression was increased by 1.9-fold in epididymal fat, 1.7-fold in mesenteric visceral fat, and 5-fold in subcutaneous fat ($P < 0.01$; Fig. 2a, c, e) in aP2-H6PDH Tg mice, respectively. Consistent with the phenotype of fat accumulation, aP2-H6PDH Tg mice also had elevated fat mRNA levels encoding leptin, FOXO1 and SIRT1 in adipose tissue (Fig. 2b, d, c). These results support that aP2-H6PDH Tg mice had upregulation of obese genes associated with fat accumulation.

Transgenic aP2-H6PDH mice exhibit increased ER stress marker expression in fat tissue

Considering that the ER stress proteins have been shown to be crucial for adiposity in fat tissue through mechanism integrating various pathways leading to obesity [24–26], we analyzed the expression levels and the phosphorylation state of XBP1 and IREα, two key transcriptional factors of ER stress in adipose tissue. Real-time RT-PCR analysis revealed that epididymal fat IREα and XBP1 mRNA levels

Fig. 5 Glucose and insulin tolerance tests in aP2-H6PDH Tg mice and WT controls. **a** Glucose tolerance test (GTT; □, non-Tg mice fed chow diet (C); ○, non-Tg mice fed high-fat diet (HFD); ■, Tg mice fed chow diet; ●, Tg mice fed HFD). **b** Insulin tolerance test (ITT; △, non-Tg mice fed chow diet; ○, Tg mice fed chow diet; ▲, non-Tg mice fed HFD; ●, Tg mice fed HFD). **c** Serum FFA levels. **d** Adipose glucose uptake in visceral fat tissue from aP2-H6PDH mice and WT control. Data are means $\pm$ SE of 7–8 mice/group. $^{\dagger}P < 0.02$ vs. HFD-fed WT controls; $^*P < 0.001$ and $^{\#}P < 0.01$ vs. WT controls; $^{**}P < 0.01$ vs. HFD-fed WT controls

were increased by 1.5-fold and 1.45-fold in aP2-H6PDH Tg mice compared with their WT controls (Fig. 3a). Parallel to these observations, IRE α protein level was increased with elevated pSer²⁹⁸ IRE α content by 2.7-fold in epididymal fat of aP2-H6PDH mice ($P < 0.01$; Fig. 3c, e). Consistent with these results, aP2-H6PDH Tg mice also had elevated XBP1 protein expression in epididymal fat compared with WT controls (Fig. 3c, e). These data indicate that activation of IRE α /XBP1 pathway is involved in adipose adiposity of aP2/H6PDH Tg mice.

Transgenic aP2-H6PDH mice activated GSK3 β with suppressed Akt signaling in fat tissue

Since Akt can regulate adipogenesis via inhibition of GSK3 β [27, 28], a master activator of adipogenesis, we next examined if the molecular basis for the induction of visceral fat adiposity in aP2-H6PDH Tg mice was associated with altered Akt and GSK3 β signaling. Consistent with visceral fat obesity, epididymal fat GSK3 β mRNA levels (Fig. 3b; $P < 0.01$) were higher with elevated pSer⁹GSK3 β content by 1.7-fold in aP2-H6PDH Tg mice (Fig. 3d, f). Reciprocally, epididymal fat Thr³⁰⁸Akt/PKB phosphorylation in aP2-H6PDH Tg mice was lower, with no effect of Akt mRNA and total Akt protein levels compared to their respective

WT controls (Fig. 3d, f). These data indicate that activation of GSK3 β with reduction of Akt signaling may be a potential mechanism for adipogenesis in adipose tissue of aP2-H6PDH Tg mice.

Transgenic aP2-H6PDH mice exhibits exacerbation of HFD-induced accumulation of visceral fat

To further investigate the impact of enhanced adipose H6PDH in the development of visceral obesity, we assessed the effect of high-fat feeding on energy homeostasis, body weight, food intake, and abdominal visceral fat pad mass in transgenic aP2-H6PDH mice. The mean body weight gain of mice fed HFD for 14 weeks were significantly increased when compared with regular chow diet feeding (Fig. 4a). However, there were no significant differences in body weight (Fig. 4a) and food intake (data not shown) between aP2/H6PDH mice and WT controls on control diet or HFD. In contrast, HFD-fed aP2/H6PDH mice gained more visceral fat mass (increased by 27% in epididymal fat and 23% in mesenteric visceral fat (Fig. 4b) than that of WT controls fed HFD, reflecting altered fat distribution with increased visceral fat mass. Oil red O staining with mounting media DAPI showed increased lipid droplet content in visceral fat from aP2-H6PDH Tg mice fed on both control diet and

Fig. 6 Adipose target gene expression and corticosterone levels in epididymal fat (Ep fat) and mesenteric fat (Mes fat) of aP2-H6PDH (white bar) and WT mice fed HFD (black bar). **a, b** Quantitative real-time RT-PCR analysis demonstrating the relative alterations of *H6PDH*, *G6PT*, *11β-HSD1*, and *GR* mRNA levels in Ep fat (**a**) and Mes fat (**b**) of aP2-H6PDH and WT mice fed HFD. **c** Western blot analysis of expression of *11β-HSD1* in epididymal fat (Ep fat) and mesenteric fat (Mes fat) of aP2-H6PDH and WT mice fed HFD. **d**

Adipose *11β-HSD1* reductase activity was measured in adipose microsomes using 11-dehydrocorticosterone (DHC) as the substrate in the presence of NADPH. Enzyme activity was expressed as % DHC converted to corticosterone (**b**). **e** Tissue corticosterone concentrations in epididymal fat and mesenteric visceral fat of aP2-H6PDH and WT mice fed HFD. **f** Plasma corticosterone levels in aP2-H6PDH and WT mice fed HFD. Data are means ± SE of 7–8 mice/group. * $P < 0.001$ vs. HFD-fed WT controls; † $P < 0.01$ vs. HFD-fed WT controls

HFD when compared with their respective WT controls (Fig. 4c, d). Consistent with adipose redistribution, mesenteric visceral fat *PPARγ*, *SREBP*, and *C/EBPα* expression were higher with elevated mRNA levels of *IRE* and *XBP1* (Fig. 4d, e), which paralleled to the elevated *IREα* and *XBP1* protein with activation of pSer²⁹⁸ *IREα* in mesenteric fat of aP2/H6PDH mice (Fig. 4f). Likewise, HFD-fed aP2-H6PDH Tg mice also had increased mesenteric fat mRNA levels encoding leptin (Fig. 4d) with upregulation of Ser⁹ *GSK3β* phosphorylation, but reduced p-Thr³⁰⁸ *Akt/PKB* level (Fig. 4f). These data show that elevation of *H6PDH* in adipose tissue promotes the HFD-induced phenotype of abdominal fat accumulation.

In contrast, we found that HFD-fed aP2-H6PDH Tg mice had decreased browning marker gene *CD137* and browning-related gene *GLUT4* mRNA levels in mesenteric fat, but did not significantly change *TBX* mRNA levels (Fig. 4e) in visceral fat, reflecting decreased the browning of white adipose tissue in aP2-H6PDH Tg mice may be involved in the development of HFD-induced abdominal visceral obesity.

Transgenic aP2-H6PDH mice have impaired glucose tolerance and insulin sensitivity

We then determined if elevated adipose *H6PDH* would render the transgenic mice more susceptible to diet-induced insulin resistance. We measured the rate of glucose disposal and insulin sensitivity by the glucose and insulin tolerance tests and analyzed adipose metabolic function. The fasting blood glucose levels of aP2-H6PDH Tg and WT mice fed HFD were higher than those of their respective controls fed chow diet (WT control diet: 158 ± 12.6 mg/dL; WT high-fat diet: 274 ± 17 mg/dL; *H6PDH* Tg control diet: 198 ± 11 mg/dL; *H6PDH* Tg high-fat diet: 337 ± 14 mg/dL) (Fig. 5a). Furthermore, higher plasma glucose levels during GTT (Fig. 5a) were observed in HFD-fed aP2-H6PDH Tg mice compared to that of WT mice fed HFD. During the ITT, aP2-H6PDH Tg mice on HFD also had decreased glucose disposal in response to insulin compared with WT mice fed HFD (Fig. 5b). Moreover, plasma FFA, which is stimulated by GCs [30] and can cause adipose insulin resistance [31], was elevated in HFD-fed aP2-H6PDH Tg mice (Fig. 5c). Additionally, basal and insulin-stimulated glucose uptake in

adipose tissue were lower in aP2-H6PDH mice (Fig. 5d), reflecting the cause of adipose insulin resistance.

Transgenic aP2-H6PDH mice exhibit enhanced 11 β -HSD1 expression and corticosterone level in fat tissues

Adipose *H6PDH* expression have shown to be associated with corticosterone production in adipose tissue [14, 17]. We next determined if the adverse metabolic phenotypes observed in aP2-H6PDH Tg mice fed on HFD are partly due to changes in local GC metabolism in this tissue. We found that HFD-fed aP2-H6PDH Tg mice exhibited elevated *H6PDH* and 11 β -HSD1 mRNA expression in epididymal fat and mesenteric visceral fat (Fig. 6a, b). Moreover, we observed that epididymal fat and mesenteric fat G6PT mRNA, the key driver supplying G6P for H6PDH-mediated NADPH generation within the ER lumen for 11 β -HSD1 [11], were higher in HFD-fed aP2-H6PDH Tg mice ($P < 0.01$; Fig. 6b). Consistent with these observations, epididymal and mesenteric fat 11 β -HSD1 protein expression and activity were higher in HFD-fed aP2-H6PDH mice than their respective HFD-fed controls (Fig. 6c, d). In parallel with the increase in 11 β -HSD1 activity, adipose corticosterone levels were increased by 38% in epididymal fat and by 35% in mesenteric fat in aP2-H6PDH Tg mice compared with WT controls fed HFD (Fig. 6c), indicating increased intra-adipose GC production. Furthermore, adipose GC receptor (GR α) mRNA that is activated by GCs [30], were also elevated in epididymal and mesenteric fat of aP2-H6PDH Tg mice fed on HFD (Fig. 6a, b), reflecting increased GR-mediated local GC action in abdominal adipose tissue. However, there were no significant differences in plasma corticosterone levels between aP2-H6PDH Tg and WT mice fed on HFD, although HFD-fed aP2-H6PDH Tg mice had slightly elevated plasma corticosterone levels (Fig. 6d). These results suggest that elevated intra-adipose GC levels may account, at least in part, for the adverse metabolic phenotype observed in aP2-H6PDH Tg mice fed on HFD, despite normal circulating GC levels.

Discussion

The data presented here suggest that a new fat depot-specific mechanism driving the adverse fat distribution and visceral fat accumulation is the result of enhanced adipose H6PDH-mediated induction of intra-adipose GC production with upregulation of 11 β -HSD1. We found that *H6PDH* transgenic mice induced visceral fat adiposity and the adverse metabolic phenotypes are correlated with elevated 11 β -HSD1-driven local corticosterone level in visceral adipose tissue. We observed that aP2-H6PDH Tg mice

exhibit elevated 11 β -HSD1 and increased epididymal and mesenteric visceral fat mass, hyperglycemia, insulin resistance, hyperlipidemia along with elevated GC-inducible lipogenic enzyme ACC and ACL expression. Increased adipose tissue *H6PDH* could elevate ER luminal NADPH ability to drive 11 β -HSD1 amplification of local GC action and consequently leads to lipogenesis and increased adipose mass linked to visceral fat deposition. This interpretation is further supported by recent report revealing that patients with obesity and metabolic syndrome have increased *H6PDH* mRNA levels and elevated cortisol production with increased 11 β -HSD1 in visceral fat and subcutaneous fat [15–18]. In addition, adipose *H6PDH* activation was correlated with elevated G6PT mRNA level in visceral fat of aP2-H6PDH Tg mice. Elevated G6PT expression ensures ER luminal G6P availability to increase *H6PDH* and may thus provide an additional mechanism to maintain adipose *H6PDH* activation in the aP2-H6PDH Tg mice.

Interestingly, we found that the visceral adiposity and its adverse metabolic phenotype were tightly associated with elevated intra-adipose corticosterone levels in aP2-H6PDH mice, but circulating corticosterone levels were elevated only slightly in response to HFD. Endorsing this hypothesis, is the demonstration of increased adipose GC production leading to visceral obesity and metabolic syndrome, despite the normal plasma GC levels in aP2-11 β -HSD1 mice [5]. Our data suggest that activated *H6PDH* coupled with elevation of 11 β -HSD1-driven intracellular GC production in adipose tissue may be an emerging mechanism to account for the accumulation of visceral fat by increased lipogenesis in aP2-H6PDH mice.

It is well known that tissue-specific action of GCs is mediated by activation of intracellular GR [31]. We found that aP2-H6PDH mice have enhanced 11 β -HSD1 and simultaneously induced fat GR expression with concomitant elevation of adipose GSK3 β phosphorylation that is activated by GCs [31]. Indeed, GSK3 β has been identified as the key ligand-dependent transcription activator and its functional signaling is required for GC-dependent transcriptional activation of GR [32, 33], a ligand-activated transcription factor that is crucially required for GC action. These findings support a concept that activation of GSK3 β signaling itself may be an additional mechanism to ensure adipose H6PDH-coupled 11 β -HSD1-driven local GC action. Induction of GSK3 β signaling could allow GCs to bind/activate intracellular GR and thus promotes GR-mediated intra-adipose corticosterone action leading to visceral fat accumulation and the adverse metabolic phenotype observed in aP2-H6PDH mice. This is consistent with the previous data showing that GSK3 β is required for adipose adipogenesis in insulin-resistant animals [34, 35]. Additionally, we also found that the pThr³⁰⁸Akt content, a marker of adipose insulin sensitivity that is suppressed by

GCs [23], were reduced coupled with decreased glucose uptake in response to elevated local GC levels with concomitant induction of GSK3 β in adipose tissue of aP2-H6PDH Tg mice. Reduction in phosphorylated AKT may indicate failures in insulin signaling and, therefore, a lower *GLUT4* translocation to the cell membrane. This could explain in part the insulin resistance observed in this animal model.

Given that Akt is also a negative regulator of obesity and insulin resistance through inhibition of GSK3 β [35, 36], reduction of p-Thr³⁰⁸Akt could also promote GSK3 β activation and, in turn, visceral fat adiposity and local insulin resistance. These findings support our hypothesis that visceral fat lipid accumulation of aP2-H6PDH mice may be mediated, at least, in part by activation of GSK3 β with suppression of Akt.

Consistent with increased lipogenic gene expression profile of adipose tissue, SREBP expression is elevated in response to increased *H6PDH* coupled with 11 β -HSD1 in visceral fat in aP2-H6PDH mice. SREBP is stimulated by insulin [37] and aP2-H6PDH mice had elevated insulin levels, which is associated with high-SREBP levels and visceral fat accumulation observed in aP2/H6PDH Tg mice. Importantly, we also observed that another two key lipogenic transcriptional activator mRNA encoding PPAR γ and C/EBP α that are positively regulated by GCs [38, 39], were also elevated in response to activation of obese gene mRNA levels encoding SIRT1 and FOXO1 with induction of 11 β -HSD1 in adipose tissue of aP2-H6PDH mice. Moreover, subcutaneous fat PPAR γ and C/EBP α mRNA levels were also increased in aP2-H6PDH mice. Elevated adipose PPAR γ and C/EBP α could thus promote the progress of lipogenesis and adipogenesis linked the development of adipose adiposity in aP2-H6PDH mice. This interpretation is supported by studies reporting that activation of C/EBP α and PPAR γ are necessary and sufficient to initiate adipogenesis and obesity in insulin-resistant rodents [40, 41]. In addition, aP2-H6PDH mice also exhibit reduced expression of visceral fat *CD137* [42, 43], the key gene for begin and browning of white adipocytes that is decreased in adipose tissue in obese animals [42, 43]. Moreover, we also observed that brown fat-related gene encoding *GLUT4* expression that is high in browning of white adipocytes [44], was decreased in fat tissue of aP2-H6PDH mice. These findings suggest that reduced adipose *CD137* and *GLUT4* may also contribute to adipose adiposity in aP2-H6PDH mice.

In addition, the current study also observed that induction of adipose C/EBP α and PPAR γ expression paralleled the elevated expression of adipose IRE α and XBP1, two key ER stress transcription factors activated by GCs [45] in response to elevated adipose GC levels in aP2-H6PDH mice. Indeed, IRE α /XBP1 are required for the

transcriptional activator of C/EBP α during adipogenesis [24] and aP2-H6PDH mice show elevated IRE α expression and its phosphorylation in visceral fat tissues. Consistent with this concept, we also found that the adipose XBP1 levels, a specific substrate for IRE1 α [46, 47], were increased in aP2-H6PDH mice, suggesting that activation of IRE α /XBP1 itself could promote the activation of C/EBP α and lipogenesis in adipose tissue. This is consistent with the role of IRE α /XBP1 in inducing lipogenesis and adipogenesis [24]. Moreover, IRE α /XBP1 is upregulated by insulin [48] and elevated plasma insulin levels may contribute to higher IRE α and XBP1 in visceral fat from aP2-H6PDH mice. Consistent with the earlier finding that PPAR γ is activated by XBP1 [33], here, we observed elevated XBP1 and increased PPAR γ expression in fat tissue of aP2-H6PDH mice. These findings imply that elevation of XBP1/IRE α could contribute to high C/EBP α and PPAR γ levels, which is sufficient to trigger adipogenesis and the progression of fat accumulation in aP2-H6PDH mice. This interpretation is supported by earlier observation that adipose IRE α and XBP1 are increased with obesity and insulin resistance in obese mice [25, 26]. Adipose *H6PDH* activation-mediated induction of IRE α /XBP1 may thus provide an additional mechanism to maintain C/EBP α and PPAR γ linked to increased lipogenesis and fat accumulation in aP2-H6PDH mice.

In summary, our current findings provide the first in vivo mechanistic evidence for the contribution of increased adipose *H6PDH* expression to fat accumulation, insulin resistance, and its relative metabolic disorders. We further demonstrated that the adverse effects of *H6PDH* activation on adipose function and insulin sensitivity may be largely driven by elevated 11 β -HSD1 amplifying local GC action relative to changes in obese gene expression profile and reduction of glucose uptake associated with reduction of insulin signaling molecules. These results imply that fat-specific manipulation of *H6PDH* may have potential as a new therapeutic strategy, and we may expect that selective inhibition of adipose *H6PDH* may represent novel and potential pharmaco-therapeutic approaches to target fat 11 β -HSD1 and treat T2DM and visceral obesity.

Author contributions LL, YW, and JW contributed to the design and performance of experiments. LL, YW, JW, YD, SC, XL, and YL acquired and analyzed the data. MJ generated aP2-*H6PDH* transgenic mouse line. YL drafted and writing of the manuscript and supervised this project. MJ, KL, TCF, and CH reviewed this manuscript. All authors discussed and agreed on the results, and gave final approval.

Funding YL is supported by NIH grant SC1DK104821 and LSI grant F00333003007. TCF is supported by NIH grant 2R24DA017298. LL is supported by grants from the Project of National Natural Science Foundation of China (8147102 and 8127086) and the Shanghai Leading Talent (SLJ15055). YD is supported by AHA grant 12SDG8760-002.

Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interest.

References

- Després JP, Lemieux I, Bergeron J, Pibarot P, Mathieu P, Larose E, et al. Abdominal obesity and the metabolic syndrome: contribution to global cardiometabolic risk. *Arterioscler Thromb Vasc Biol.* 2008;28:1039–49.
- Himabukuro M. Cardiac adiposity and global cardiometabolic risk: new concept and clinical implication. *Circ J.* 2009;73:27–34.
- Bamberger CM, Schulte HM, Chrousos GP. Molecular determinants of glucocorticoid receptor function and tissue sensitivity to glucocorticoids. *Endocr Rev.* 1996;17:245–61.
- Bujalska IJ, Kumar S, Stewart PM. Does central obesity reflect “Cushing’s disease of the omentum. *Lancet.* 1997;349:1210–3.
- Masuzaki H, Paterson J, Shinyama H, Morton NM, Mullins JJ, Seckl JR, et al. A transgenic model of visceral obesity and the metabolic syndrome. *Science.* 2001;294:2166–70.
- Tomlinson JW, Stewart PM. Mechanisms of disease: selective inhibition of 11 β -hydroxysteroid dehydrogenase type 1 as a novel treatment for the metabolic syndrome. *Nat Clin Pract Endocrinol Metab.* 2005;1:92–9.
- Morton NM, Seckl JR. 11 β -hydroxysteroid dehydrogenase type 1 and obesity. *Front Horm Res.* 2008;36:146–64.
- Kotelevtsev Y, Holmes MC, Burchell A, Houston PM, Schmoll D, Jamieson P, et al. 11 β -hydroxysteroid dehydrogenase type 1 knockout mice show attenuated glucocorticoid-inducible responses and resist hyperglycemia on obesity or stress. *PNAS.* 1997;94:14924–9.
- Mziaut H, Korza G, Hand AR, Gerard C, Ozols J. Targeting proteins to the lumen of endoplasmic reticulum using N-terminal domains of 11 β -hydroxysteroid dehydrogenase and the 50-kDa esterase. *J Biol Chem.* 1999;274:14122–9.
- Odermatt A, Arnold P, Stauffer A, Frey BM, Frey FJ. The N-terminal anchor sequences of 11 β -hydroxysteroid dehydrogenases determine their orientation in the endoplasmic reticulum membrane. *J Biol Chem.* 1999;274:28762–70.
- Hewitt KN, Walker EA, Stewart PM. Minireview: hexose-6-phosphate dehydrogenase and redox control of 11 β -hydroxysteroid dehydrogenase type 1 activity. *Endocrinology.* 2005;146:2539–43.
- Chou JY, Matern D, Mansfield BC, Chen YT. Type I glycogen storage diseases: disorders of the glucose-6-phosphatase complex. *Curr Mol Med.* 2002;2:121–43.
- Draper N, Walker EA, Bujalska IJ, Tomlinson JW, Chalder SM, Arlt W, et al. Mutations in the genes encoding 11 β -hydroxysteroid dehydrogenase type 1 and hexose-6-phosphate dehydrogenase interact to cause cortisone reductase deficiency. *Nat Genet.* 2003;34:434–9.
- Bujalska IJ, Hewitt KN, Hauton D, Lavery GG, Tomlinson JW, Walker EA, et al. Lack of hexose-6-phosphate dehydrogenase impairs lipid mobilization from mouse adipose tissue. *Endocrinology.* 2008;149:2584–91.
- Alberti L, Girola A, Gilardini L, Conti A, Cattaldo S, Micheletto G, et al. Type 2 diabetes and metabolic syndrome are associated with increased expression of 11 β -hydroxysteroid dehydrogenase 1 in obese subjects. *Int J Obes.* 2007;31:1826–31.
- Uçkaya G, Karadurmus N, Kutlu O, Corakçı A, Kizildağ S, Ural AU, et al. Adipose tissue 11 β -hydroxysteroid dehydrogenase type 1 and hexose-6-phosphate dehydrogenase gene expressions are increased in patients with type 2 diabetes mellitus. *Diabetes Res Clin Pract.* 2008;82:S135–40.
- Wang Y, Nakagawa Y, Liu L, Wang W, Ren X, Anghel A, et al. Tissue-specific dysregulation of hexose-6-phosphate dehydrogenase and glucose-6-phosphate transporter production in db/db mice as a model of type 2 diabetes. *Diabetologia.* 2011;54:440–50.
- Yan C, Yang H, Wang Y, Dong Y, Yu F, Wu Y, et al. Increased glycogen synthase kinase-3 β and hexose-6-phosphate dehydrogenase expression in adipose tissue may contribute to glucocorticoid-induced mouse visceral adiposity. *Int J Obes.* 2016;40:1233–41.
- Wang Y, Liu L, Du H, Nagaoka Y, Fan W, Lutfy K, et al. Transgenic overexpression of hexose-6-phosphate dehydrogenase in adipose tissue causes local glucocorticoid amplification and lipolysis in male mice. *Am J Physiol Endocrinol Metab.* 2014;306:E543–51.
- Stolic M, Russell A, Hutley L, Fielding G, Hay J, MacDonald G, et al. Glucose uptake and insulin action in human adipose tissue— influence of BMI, anatomical depot and body fat distribution. *Int J Obes Relat Metab Disord.* 2002;26:17–23.
- Rönquist-Nii Y, Edlund PO. Determination of corticosteroids in tissue samples by liquid chromatography-tandem mass spectrometry. *J Pharm Biomed Anal.* 2005;37:341–50.
- Szymczak J, Milewicz A, Thijssen JH, Blankenstein MA, Darsoszewski J. Concentration of sex steroids in adipose tissue after menopause. *Steroids.* 1998;63:319–21.
- Wang Y, Yan C, Liu L, Wang W, Du H, Fan W, et al. 11 β -hydroxysteroid dehydrogenase type 1 shRNA ameliorates glucocorticoid-induced insulin resistance and lipolysis in mouse abdominal adipose tissue. *Am J Physiol Endocrinol Metab.* 2015;308:E84–95.
- Sha H, He Y, Chen H, Wang C, Zenno A, Shi H, et al. The IRE1 α -XBP1 pathway of the unfolded protein response is required for adipogenesis. *Cell Metab.* 2009;9:556–64.
- Ozcan U, Yilmaz E, Ozcan L, Furuhashi M, Vaillancourt E, Smith RO, et al. Chemical chaperones reduce ER stress and restore glucose homeostasis in a mouse model of type 2 diabetes. *Science.* 2006;313:1137–40.
- Ozcan U, Cao Q, Yilmaz E, Lee AH, Ozdelen E, et al. Endoplasmic reticulum stress links obesity, insulin action, and type 2 diabetes. *Science.* 2004;306:457–61.
- Izumiya Y, Hopkins T, Morris C, Sato K, Zeng L, Viereck J, et al. Fast/glycolytic muscle fiber growth reduces fat mass and improves metabolic parameters in obese mice. *Cell Metab.* 2008;7:159–72.
- Ross SE, Hemati N, Longo KA, Bennett CN, Lucas PC, Erickson RL, et al. Inhibition of adipogenesis by Wnt signaling. *Science.* 2000;289:950–3.
- Morgan SA, McCabe EL, Gathercole LL, Hassan-Smith ZK, Larner DP, Bujalska IJ, et al. 11 β -HSD1 is the major regulator of the tissue-specific effects of circulating glucocorticoid excess. *Proc Natl Acad Sci USA.* 2014;111:E2482–91.
- Kalderon B, Mayorek N, Berry E, Zevit N, Bar-Tana J. Fatty acid cycling in the fasting rat. *Am J Physiol.* 2000;279:E221–7.
- Liu Y, Nakagawa Y, Wang Y, Sakurai R, Tripathi PV, Lutfy K, et al. Increased glucocorticoid receptor and 11 β -hydroxysteroid dehydrogenase type 1 expression in hepatocytes may contribute to the phenotype of type 2 diabetes in db/db mice. *Diabetes.* 2005;54:32–40.
- Yun SI, Yoon HY, Jeong SY, Chung YS. Glucocorticoid induces apoptosis of osteoblast cells through the activation of glycogen synthase kinase 3 β . *J Bone Miner Metab.* 2009;27:140–8.
- Rubio-Patiño C, Palmeri CM, Pérez-Perarnau A, et al. Glycogen synthase kinase-3 β is involved in ligand-dependent activation of transcription and cellular localization of the glucocorticoid receptor. *Mol Endocrinol.* 2012;26:1508–20.

34. Galliher-Beckley AJ, Williams JG, Collins JB, Cidlowski JA. Glycogen synthase kinase 3 β -mediated serine phosphorylation of the human glucocorticoid receptor redirects gene expression profiles. *Mol Cell Biol*. 2008;28:7309–22.
35. Chakraborty A, Koldobskiy MA, Bello NT, Maxwell M, Potter JJ, Juluri KR, et al. Inositol pyrophosphates inhibit Akt signaling, thereby regulating insulin sensitivity and weight gain. *Cell*. 2010;143:897–910.
36. Moon MH, Jeong JK, Lee JH, Park YG, Lee YJ, Seol JW, et al. Antiobesity activity of a sphingosine 1-phosphate analogue FTY720 observed in adipocytes and obese mouse model. *Exp Mol Med*. 2012;44:603–14.
37. Lin J, Yang R, Tarr PT, Wu PH, Handschin C, Li S, et al. Hyperlipidemic effects of dietary saturated fats mediated through PGC-1 β coactivation of SREBP. *Cell*. 2005;120:261–73.
38. Williams LJ, Lyons V, MacLeod I, Rajan V, Darlington GJ, Poli V, et al. C/EBP regulates hepatic transcription of 11 β -hydroxysteroid dehydrogenase type 1. A novel mechanism for cross-talk between the C/EBP and glucocorticoid signaling pathways. *J Biol Chem*. 2000;275:30232–9.
39. Galitzky J, Bouloumié A. Human visceral-fat-specific glucocorticoid tuning of adipogenesis. *Cell Metab*. 2013;18:3–5.
40. Yeh WC, Cao Z, Classon M, McKnight SL. Cascade regulation of terminal adipocyte differentiation by three members of the C/EBP family of leucine zipper proteins. *Genes Dev*. 1995;9:168–81.
41. Vidal-Puig A, Jimenez-Linan M, Lowell BB, Hamann A, Hu E, Spiegelman B, et al. Regulation of PPAR γ gene expression by nutrition and obesity in rodents. *J Clin Invest*. 1996;97:2553–61.
42. Carey AL, Vorlander C, Reddy-Luthmoodoo M, Natoli AK, Formosa MF, Bertovic DA, et al. Reduced UCP-1 content in in vitro differentiated beige/brite adipocyte derived from pre-adipocytes of human subcutaneous white adipose tissues in obesity. *PLoS ONE*. 2014;9:e91997.
43. Fu T, Seok S, Choi S, Huang Z, Suino-Powell K, Xu HE, et al. MicroRNA 34a inhibits beige and brown fat formation in obesity in part by suppressing adipocyte fibroblast growth factor 21 signaling and SIRT1 function. *Mol Cell Biol*. 2014;34:4130–42.
44. Singh R, Braga M, Reddy ST, Lee SJ, Parveen M, Grilalva L, et al. Follistatin targets distinct pathways to promote brown adipocyte characteristics in brown and white adipose tissues. *Endocrinology*. 2017;158:1217–30.
45. Zhou JY, Zhong HJ, Yang C, Yan J, Wang HY, Jiang JX. Corticosterone exerts immunostimulatory effects on macrophages via endoplasmic reticulum stress. *Br J Surg*. 2010;97:281–93.
46. Ron D, Walter P. Signal integration in the endoplasmic reticulum unfolded protein response. *Nat Rev Mol Cell Biol*. 2007;8:519–29.
47. Nekrutenko A, He J. Functionality of unspliced XBP1 is required to explain evolution of overlapping reading frames. *Trends Genet*. 2006;22:645–8.
48. Park SW, Zhou Y, Lee J, Lu A, Sun C, Chung J, et al. The regulatory subunits of PI3K, p85 α and p85 β , interact with XBP-1 and increase its nuclear translocation. *Nat Med*. 2010;16:429–37.