

Suppression of oxidative metabolism and mitochondrial biogenesis by the transcriptional corepressor RIP140 in mouse adipocytes

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Using an siRNA-based screen, we identified the transcriptional corepressor RIP140 as a negative regulator of insulin-responsive hexose uptake and oxidative metabolism in 3T3-L1 adipocytes. Affymetrix GeneChip profiling revealed that RIP140 depletion upregulates the expression of clusters of genes in the pathways of glucose uptake, glycolysis, TCA cycle, fatty acid oxidation, mitochondrial biogenesis, and oxidative phosphorylation in these cells. Conversely, we show that reexpression of RIP140 in mouse embryonic fibroblasts derived from RIP140-null mice downregulates expression of many of these same genes. Consistent with these microarray data, RIP140 gene silencing in cultured adipocytes increased both conversion of [¹⁴C]glucose to CO₂ and mitochondrial oxygen consumption. RIP140-null mice, previously reported to resist weight gain on a high-fat diet, are shown here to display enhanced glucose tolerance and enhanced responsiveness to insulin compared with matched wild-type mice upon high-fat feeding. Mechanistically, RIP140 was found to require the nuclear receptor ERR α to regulate hexose uptake and mitochondrial proteins SDHB and CoxVb, although it likely acts through other nuclear receptors as well. We conclude that RIP140 is a major suppressor of adipocyte oxidative metabolism and mitochondrial biogenesis, as well as a negative regulator of whole-body glucose tolerance and energy expenditure in mice.

Introduction

Type 2 diabetes mellitus is an increasingly prevalent metabolic disorder in which the combination of systemic insulin resistance and impaired insulin production by pancreatic β cells results in hyperglycemia. Insulin regulates plasma glucose levels by selectively increasing glucose uptake into fat and muscle and by inhibiting hepatic glucose production (1). Insulin-responsive glucose uptake is initiated by a PI3K-dependent signaling cascade that stimulates trafficking of GLUT4 glucose transporters to the plasma membrane, allowing removal of excess glucose from the bloodstream (2, 3). Up to 85% of glucose disposal in humans under conditions of glucose infusion can be accounted for by insulin-stimulated glucose uptake into muscle (4), which is impaired in type 2 diabetes mellitus (5).

Nonstandard abbreviations used: ACADM, medium chain acetyl-Coenzyme A dehydrogenase; Cox Vb, cytochrome oxidase subunit Vb; cyt c, cytochrome c; ERR α , estrogen related receptor α ; GABP α , GA repeat-binding protein α ; KRH, Krebs-Ringer-HEPES; MEF, mouse embryonic fibroblast; PGC-1 α , PPAR γ coactivator 1 α ; RIP140, receptor-interacting protein 140; SDHB, succinate dehydrogenase complex b; siRNA, small interfering RNA; UCP1, uncoupling protein 1; UQCRC2, ubiquinol-cyt c reductase core protein 2.

Conflict of interest: M.G. Parker is entitled to a share of revenue under Imperial College's discretionary rewards to inventors scheme. Imperial College London has received equity and may receive future consideration from CytRx Corp. under an existing license agreement. M.P. Czech, A.M. Powelka, X. Tang, A.D. Cherniack, and A. Guilherme own, or may be provided, equity in CytRx Inc., based on intellectual property licensed from the University of Massachusetts Medical School, which shares royalties with inventors. M.P. Czech is a consultant for CytRx Inc.

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Enhancing fatty acid oxidation or fatty acid removal from circulation through esterification to triglyceride in adipocytes appears to correlate with increased insulin sensitivity in various mouse models (6–9). Mitochondria may be key regulators of fatty acid levels since they catalyze fatty acid oxidation as well as produce ATP, which is required for fatty acid esterification to triglyceride. Indeed, mitochondrial abundance and protein expression are increased during adipogenesis in cell culture (10). Furthermore, the insulin sensitizer rosiglitazone, a ligand for the transcription factor PPAR γ , enhances mitochondrial mass and alters mitochondrial morphology in cultured adipocytes (10) and in the white adipose tissue of dogs and rats (11). Conversely, the onset of obesity and insulin resistance in *ob/ob* mice is accompanied by a decrease in the expression of mitochondrial genes in fat cells that is reversed by rosiglitazone treatment (12). Interestingly, microarray analysis of gene expression in human skeletal muscle revealed small decreases throughout a subset of mitochondrial genes in diabetic subjects (13, 14). Thus, mitochondrial function may be a critical regulator of glucose homeostasis in both mice and humans.

Multiple transcriptional regulators have been shown to both affect mitochondrial gene expression and to play a major role in whole-body energy balance. The PPAR family of nuclear receptors, which function as sensors for dietary lipids, play important roles in regulating lipid and carbohydrate metabolism in adipose tissue, liver, and muscle (15). The PPAR γ coactivator 1 α (PGC-1 α) stimulates adaptive thermogenesis and mitochondrial biogenesis (16, 17), while the related coactivator, PGC-1 β , regulates mitochondrial

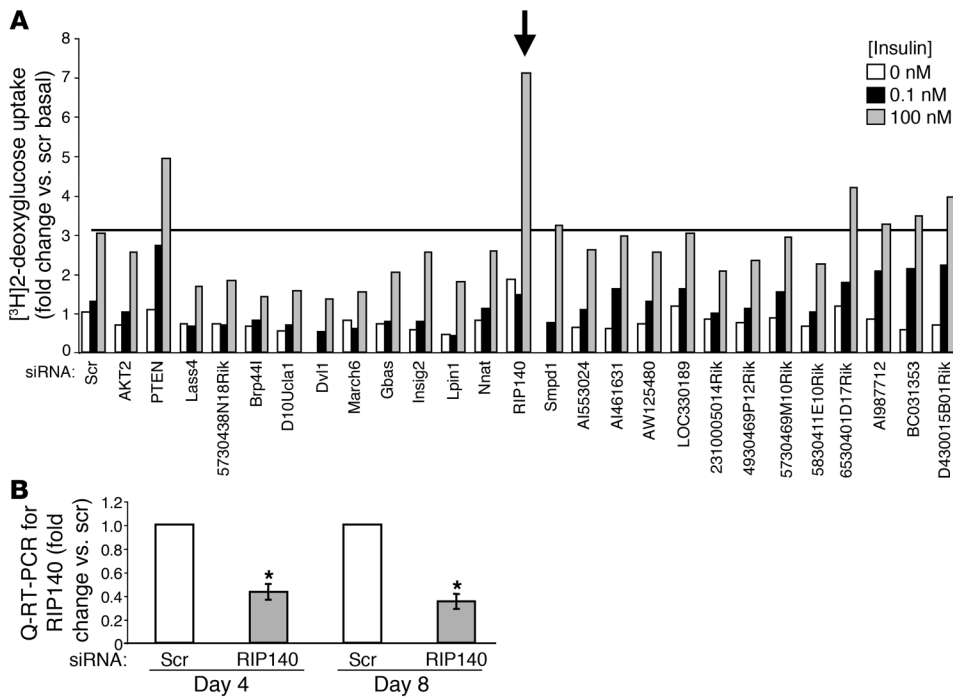


Figure 1
An siRNA-mediated screen identifies Nrip1 (RIP140) as a negative regulator of deoxyglucose uptake in 3T3-L1 adipocytes. **(A)** Four days after the induction of differentiation, 3T3-L1 adipocytes were transfected with smart pools of siRNA against a panel of genes (GenBank accession numbers are provided in Supplemental Table 1) selected as highly expressed in muscle and 3T3-L1 adipocytes compared with 3T3-L1 fibroblasts. The effect of each knockdown on glucose uptake was determined using a 2-deoxyglucose uptake assay. Arrow indicates insulin-stimulated 2-deoxyglucose uptake in RIP140-knockdown cells. Shown is the average of 2 independent experiments. **(B)** siRNA-mediated depletion of RIP140 mRNA was confirmed by real-time RT-PCR. The graph represents the mean \pm SEM of 3 independent experiments. Scr, scrambled; Q-RT-PCR, quantitative RT-PCR. * $P < 0.05$ compared with scrambled.

function by promoting the activity of the estrogen-related receptor α (ERR α) (18). In addition, the p160 coactivators, SRC1 and TIF2, modulate development and function of white and brown adipose tissue (19). Another important regulator is the cofactor receptor-interacting protein 140 (RIP140), a ligand-dependent corepressor for most, if not all, nuclear receptors (20, 21). It is essential for female fertility (22) and plays a crucial role in lipid metabolism (23, 24). RIP140-null mice are extremely lean and exhibit resistance to obesity and hepatic steatosis, which are normally induced in mice fed a high-fat diet (23). Adipogenesis is unaffected in these mice, but the expression of genes involved in energy dissipation and mitochondrial uncoupling, including uncoupling protein 1 (UCP1) and carnitine palmitoyltransferase 1b (CPT1b), are markedly increased in white adipose tissue. Consistent with these changes, the mice utilize more oxygen (23).

The aim of the present study was to identify novel proteins that influence insulin signaling and metabolism in adipocytes to better understand these processes. In a small interfering RNA-based (siRNA-based) screen of genes highly expressed in fat and muscle, we surprisingly identified RIP140 as a potent negative regulator of insulin-stimulated deoxyglucose uptake in 3T3-L1 adipocytes. Furthermore, RIP140 was found to regulate a broad program of metabolic gene expression by suppressing clusters of genes in the pathways of glycolysis, the TCA cycle, and mitochondrial biogenesis, in addition to its known role in regulating genes control-

ling fatty acid oxidation. Indeed, RIP140 depletion in cultured adipocytes increased glucose oxidation, mitochondrial protein expression, and oxygen consumption, while RIP140-null mice on a high-fat diet displayed enhanced glucose tolerance and increased responsiveness to insulin to lower blood glucose compared with wild-type mice. Thus, RIP140 functions in isolated adipocytes and in intact mice as both a major suppressor of oxidative metabolism and as a regulator of glucose homeostasis.

Results

To identify novel regulators of insulin signaling and glucose uptake, we developed a miniaturized siRNA-based gene silencing screen using 3T3-L1 adipocytes. Figure 1 depicts a screen of 24 genes identified in a subtractive cDNA library that enriched transcripts highly expressed in both skeletal muscle and 3T3-L1 adipocytes compared with 3T3-L1 fibroblasts (Supplemental Table 1; supplemental material available online with this article; doi:10.1172/JCI26040DS1). Insulin-stimulated 2-deoxyglucose uptake was assayed 72 hours after transfection of siRNA. Figure 1A

shows that depletion of the positive regulator Akt inhibited insulin stimulation of deoxyglucose uptake, as expected (25), while depletion of the known negative regulator PTEN (26) enhanced this process. While depletion of several of the candidate genes appeared to inhibit insulin action, the loss of 1 greatly enhanced insulin-responsive deoxyglucose uptake. This unique latter gene encodes the transcriptional corepressor RIP140.

RIP140 mRNA expression is increased 4-fold during adipocyte differentiation (Supplemental Table 2 and ref. 27), and it is effectively depleted by siRNA using real-time PCR of in vitro-transcribed message (Figure 1B). Commercial antibody did not detect RIP140 in 3T3-L1 cells. RIP140 mRNA was decreased by 57% and 65%, respectively, when 3T3-L1 cells were transfected with siRNA 4 days (day 4) or 8 days (day 8) after initiation of differentiation. Differentiation of 3T3-L1 fibroblasts into adipocytes is essentially complete 8 days after its initiation (27). We found that RIP140 depletion at either point during the differentiation process enhanced insulin-stimulated deoxyglucose uptake (Figure 2), indicating that RIP140 regulates hexose uptake even in mature adipocytes. Analysis of glucose transporter proteins by Western blotting showed that RIP140 depletion increased GLUT4, but not GLUT1, protein levels approximately 1.3-fold, regardless of the day of transfection. Other indicators of differentiation, such as lipid accumulation and expression of marker genes such as aP2, were not affected by RIP140 depletion (data not shown). These

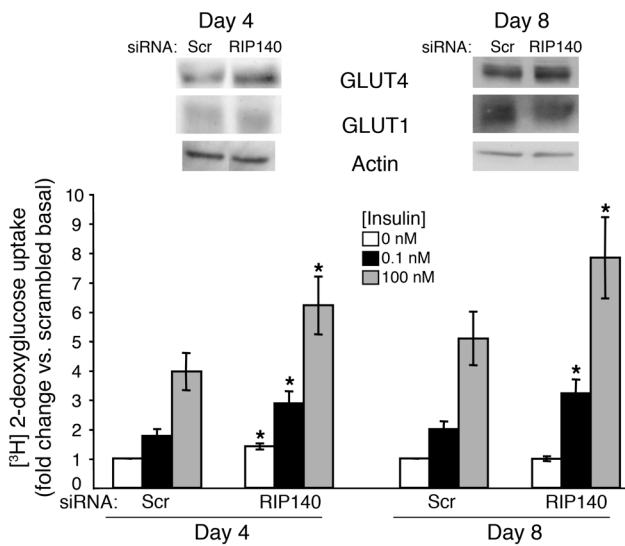


Figure 2

RIP140 depletion enhances GLUT4 expression and hexose uptake in both day-4 and day-8 3T3-L1 adipocytes. Four (day 4) or eight (day 8) days after the induction of differentiation, 3T3-L1 adipocytes were transfected with scrambled or RIP140 siRNA. Three days later, cell lysates were used for SDS-PAGE to assess GLUT4, GLUT1, and actin expression levels. Intact cells were used to examine the effect of RIP140 depletion on 2-deoxyglucose uptake. The graph shows the mean \pm SEM of 5 independent experiments. * $P < 0.05$ compared with scrambled, by Student's *t* test.

results are consistent with the observed normal development of white adipose tissue in RIP140-knockout mice and the ability of RIP140-null mouse embryonic fibroblasts (MEFs) to undergo adipogenesis in culture (23).

It is thought that insulin signaling to GLUT4 is mediated through activation of Akt2 (25), downstream of the PI3K signaling pathway (2). In order to test whether RIP140 might be a regulator of insulin signaling through this pathway, we tested the effect of its depletion on the rise in levels of phospho-Akt in response to insulin. No significant effect on stimulation of either Akt or MAPK due to the loss of RIP140 was observed when adipocytes were stimulated with submaximal or maximal concentrations of insulin (Figure 3). We did observe a small increase in the level of phospho-Akt in unstimulated cells in some experiments, although this effect was variable and not statistically significant.

We next applied Affymetrix GeneChip expression profiling to mature 3T3-L1 adipocytes transfected with scrambled or RIP140 siRNA and analyzed the 2,747 Affymetrix probes that changed significantly ($P < 0.05$) upon RIP140 depletion. These genes displayed a nonrandom distribution, with a preponderance of genes encoding proteins involved in carbohydrate and fatty acid catabolic pathways (see Figure 4 and Supplemental Table 3). The majority of the genes encoding enzymes within the glycolytic, TCA cycle, fatty acid oxidative, oxidative phosphorylation, and mitochondrial biogenesis pathways were upregulated in response to RIP140 depletion (Figure 4). Mitochondrial proteins that mediate respiratory chain expression and assembly, including mitochondrial ribosomal proteins required for translation of products of the mitochondrial genome and biosynthetic enzymes required for production of cytochrome heme, were also upregulated by RIP140 depletion (data not shown). To confirm that these changes in gene expression were due directly to the loss of RIP140, we also examined changes in gene expression upon lentivirus-mediated expression of RIP140 in adipocytes derived from RIP140-null MEFs (RIPKO-1 cells described in ref. 24). Strikingly, we found a pattern of gene downregulation due to RIP140 expression that corresponded closely to the pattern of upregulation due to RIP140 depletion in 3T3-L1 adipocytes, including GLUT4 modulation (compare Figures 4 and 5, and see Supplemental Table 3). Of the top 66 genes that were significantly upregulated in RIP140-depleted 3T3-L1

adipocytes, 54 were significantly downregulated by RIP140 expression in RIPKO-1 adipocytes (Table 1). These data support a role for RIP140 as a global regulator of adipocyte genes that control metabolic and mitochondrial pathways.

We confirmed by immunoblotting that several of the genes upregulated by RIP140 depletion showed increased expression at the protein level. The mitochondrial proteins cytochrome *c* (cyt *c*), cyt *c* oxidase subunit Vb (CoxVb), ubiquinol-cyt *c* reductase core protein 2 (UQCRC2), succinate dehydrogenase complex b (SDHB), and the fatty acid β oxidation protein medium chain acetyl-coenzyme A dehydrogenase (ACADM) were all modestly, but significantly ($P < 0.05$), upregulated in 3T3-L1 adipocytes upon loss of RIP140 (Figure 6).

Based on the genomics analyses described above, not only GLUT4 but also hexokinase 2 and other genes within glycolytic and downstream oxidative pathways are regulated by RIP140. We therefore examined the rates of glucose oxidation, triglyceride synthesis, and de novo fatty acid synthesis in adipocytes depleted of RIP140 using D-[6-¹⁴C]-glucose. The amount of glucose converted to carbon dioxide, which is a combined measurement of glycolysis and TCA cycling, was significantly enhanced in RIP140-depleted cells in the presence or absence of insulin (Figure 7). The amount of glucose converted to triglycerides and free fatty acids, however, was not significantly affected by the loss of RIP140. These results are consistent with the gene expression profiles observed upon RIP140 depletion (Figure 4), which showed greater changes in the glycolytic (69%) and TCA (81%) pathways compared with the triglyceride synthesis (50%) or fatty acid synthesis (40%) pathways.

To confirm that RIP140 depletion augments mitochondrial function in cultured adipocytes, fluorescence microscopy of these

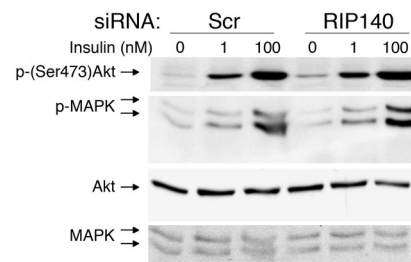


Figure 3

Insulin signaling is not significantly affected by RIP140 silencing. Four days after the induction of differentiation, 3T3-L1 adipocytes were transfected with scrambled or RIP140 siRNA. Three days later, the cells were starved for 2 hours and then stimulated with insulin for 30 minutes. Cell lysates were used for total and phospho-Akt and MAPK blots. The representative blot shows a small but insignificant increase in phospho-serine Akt levels.

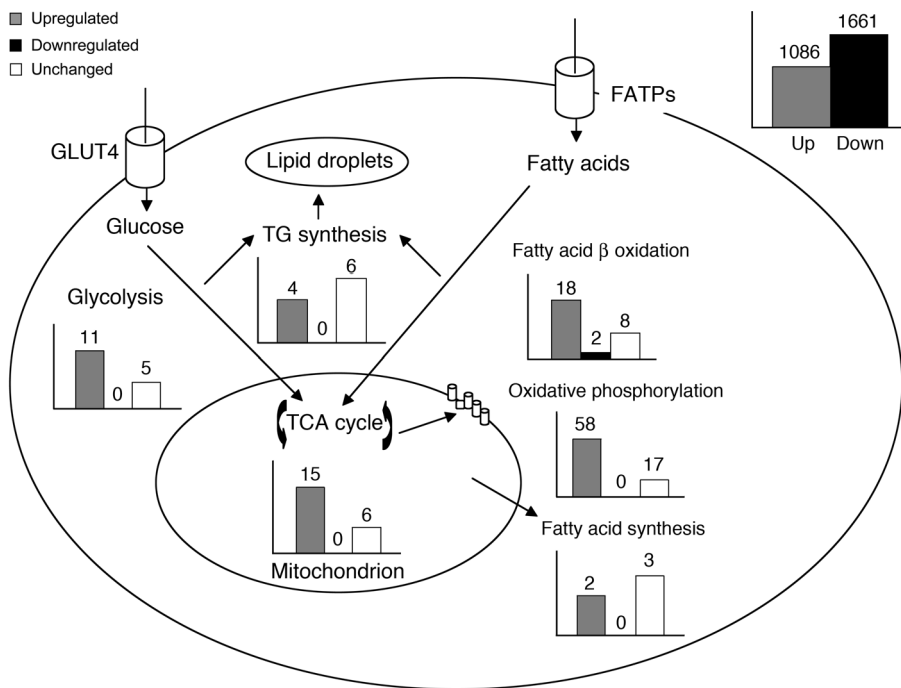


Figure 4

RIP140 silencing enhances multiple metabolic pathways in 3T3-L1 adipocytes. Eight days after the induction of differentiation, 3T3-L1 adipocytes were transfected with scrambled or RIP140 siRNA. Three days later, mRNA was harvested and used for Affymetrix GeneChip analysis. The numbers of genes that were significantly ($P < 0.05$) increased, decreased, or unchanged in expression within each of the pathways shown are depicted in bar graphs adjacent to the respective pathways. The total number of significantly ($P < 0.05$) changed probe sets from 3 independent experiments is shown in the upper-right panel. FATP, fatty acid transport protein; TG, triglycerides.

cells incubated with MitoTracker Red dye was performed. Significantly increased signal was observed in RIP140-depleted cells compared with controls, indicating an increase in actively respiring mitochondria (Figure 8A). Furthermore, RIP140-depleted adipocytes consumed more oxygen in the presence or absence of the mitochondrial uncoupler carbonyl cyanide 4-(trifluoromethoxy)phenylhydrazine (FCCP) compared with cells transfected with scrambled siRNA (Figure 8B). These data are consistent with the increased expression of genes encoding subunits within each of the respiratory complexes.

Because increased mitochondrial function is correlated with improved glucose metabolism, RIP140-null mice (23) were subjected to glucose tolerance tests. As shown in Figure 9A, young wild-type mice and RIP140-knockout mice fed a normal chow diet displayed similar increases in blood glucose concentrations after injection of a glucose load. In contrast, RIP140-null mice exhibited significantly improved glu-

ucose tolerance at 8 months of age (Figure 9B) or after a high-fat diet (Figure 9C) compared with wild-type mice matched for age and diet. We also tested whether insulin sensitivity is affected by RIP140 expression levels in mice. An insulin tolerance test demonstrated that RIP140-null mice have increased insulin responsiveness with respect to glucose lowering when on a high-fat diet (Figure 9D). These data indicate that RIP140-null mice show both higher insulin sensitivity and glucose tolerance under circumstances that normally inhibit these processes in wild-type mice. This phenotype is consistent with results described above in cul-

Figure 5

RIP140 reexpression in RIPKO-1 adipocytes results in downregulation of multiple metabolic pathways. Ten days after the induction of differentiation, RNA was isolated from RIPKO-1 adipocytes and RIPKO-1 adipocytes stably expressing RIP140 using a lentiviral vector and used for Affymetrix GeneChip analysis. The number of genes in the indicated metabolic pathways that were significantly ($P < 0.05$) upregulated, downregulated, or not changed is shown from left to right in the graphs.

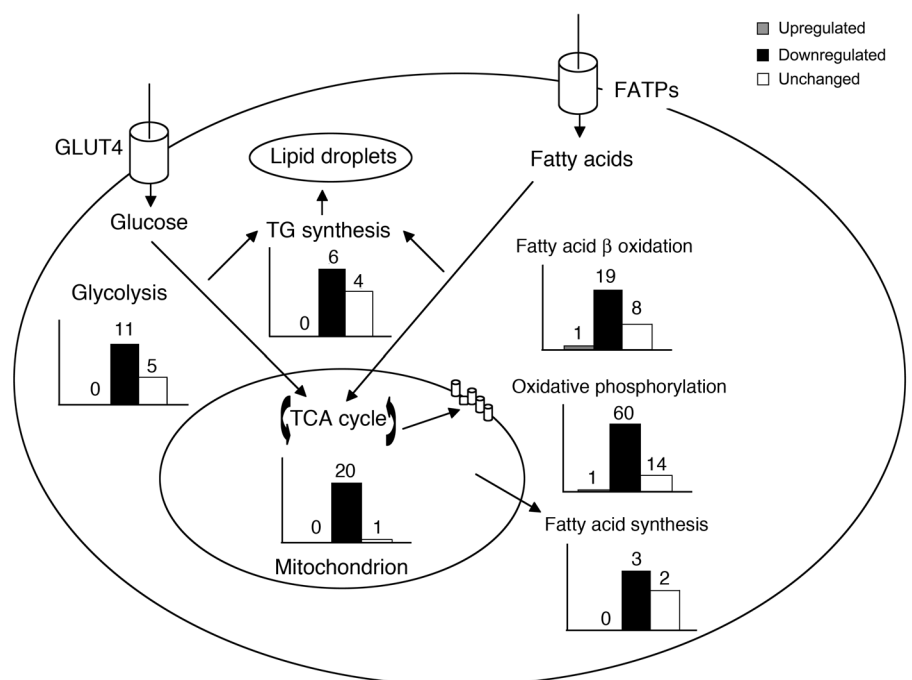




Table 1
RIP140 regulates the expression of metabolic and mitochondrial genes

Probe ID	Gene name	RIP140 siRNA (fold change)	P value	Lenti-RIP140 (fold change)	P value	Pathway
1418709_at	Cytochrome c oxidase, subunit VIIa 1	16.80	0.000	-3.43	0.000	Oxidative phosphorylation
1449218_at	Cytochrome c oxidase, subunit VIIIb	7.62	0.011	-4.06	0.001	Oxidative phosphorylation
1417956_at	Cell death-inducing DNA fragmentation factor, α subunit-like effector A	6.58	0.031	-11.54	0.000	Mitochondrial
1417482_at	Testis expressed gene 19	6.09	0.000	-1.46	0.155	
1416194_at	Cytochrome P450, family 4, subfamily b, polypeptide 1	4.72	0.000	-2.69	0.003	Oxidative phosphorylation
1429549_at	Procollagen, type XXVII, α 1	4.70	0.006	-2.13	0.001	
1427052_at	Acetyl-Coenzyme A carboxylase β	4.69	0.000	-1.43	0.223	Fatty acid oxidation
1448134_at	Paladin	4.35	0.000	-2.88	0.000	
1455061_a_at	Acetyl-Coenzyme A acyltransferase 2 (mitochondrial 3-oxoacyl-Coenzyme A thiolase)	4.23	0.001	-2.51	0.000	Fatty acid oxidation
1423944_at	Hemopexin	4.14	0.033	-3.48	0.173	
1417741_at	Liver glycogen phosphorylase	4.02	0.007	1.46	0.071	Glycolysis
1428146_s_at	Acetyl-Coenzyme A acyltransferase 2 (mitochondrial 3-oxoacyl-Coenzyme A thiolase)	3.98	0.000	-2.26	0.000	Fatty acid oxidation
1418708_at	Apollipoprotein C-IV	3.97	0.015	-3.42	0.008	
1453191_at	Procollagen, type XXVII, α 1	3.86	0.008	-2.04	0.024	
1417273_at	Pyruvate dehydrogenase kinase, isoenzyme 4	3.57	0.008	-2.26	0.000	TCA cycle
1417066_at	Chaperone, ABC1 activity of bc1 complex like (S. pombe)	3.51	0.000	-5.79	0.002	
1428145_at	Acetyl-Coenzyme A acyltransferase 2 (mitochondrial 3-oxoacyl-Coenzyme A thiolase)	3.46	0.000	-2.06	0.003	Fatty acid oxidation
1436590_at	Protein phosphatase 1, regulatory (inhibitor) subunit 3B	3.25	0.004	-3.05	0.000	Glycolysis regulator
1421681_at	Neuregulin 4	3.15	0.000	-1.89	0.001	
1419182_at	Polydomain protein	2.99	0.018	-1.60	0.004	
1451339_at	Sulfite oxidase	2.93	0.010	-1.74	0.000	Mitochondrial
1422501_s_at	Isocitrate dehydrogenase 3 (NAD+) α	2.79	0.000	-1.12	0.448	TCA cycle
1457123_at	Neuregulin 4	2.75	0.001	-1.45	0.031	
1436368_at	Solute carrier family 16 (monocarboxylic acid transporters), member 10	2.74	0.023	-1.29	0.021	Amino acid transporter
1448842_at	Cysteine dioxygenase 1, cytosolic	2.71	0.021	-3.56	0.000	
1422500_at	Isocitrate dehydrogenase 3 (NAD+) α	2.69	0.000	-1.99	0.008	TCA cycle
1424815_at	Glycogen synthase 2	2.62	0.003	-1.42	0.242	Glycogenesis
1452485_at	Phosphatase, orphan 1	2.61	0.003	-1.52	0.045	
1434082_at	PCTAIRE-motif protein kinase 2	2.58	0.001	-1.42	0.008	
1441944_s_at	G protein-coupled receptor 135	2.58	0.001	1.16	0.377	
1455112_at	Apoptosis-inducing factor (AIF)-like mitochondrion- associated inducer of death	2.55	0.001	-2.02	0.012	Mitochondrial
1423613_at	Sperm specific antigen 2	2.54	0.001	1.24	0.027	
1431142_s_at	Apoptosis-inducing factor (AIF)-like mitochondrion- associated inducer of death	2.54	0.001	-2.07	0.002	Mitochondrial
1432269_a_at	SH3-domain kinase binding protein 1	2.54	0.000	-1.79	0.001	
1422973_a_at	Thyroid hormone responsive SPOT14 homolog (Rattus)	2.50	0.009	-2.66	0.001	Fatty acid synthesis
1423109_s_at	Solute carrier family 25 (mitochondrial carnitine/acylcarnitine translocase), member 20	2.50	0.001	-2.36	0.001	Fatty acid oxidation
1448825_at	Pyruvate dehydrogenase kinase, isoenzyme 2	2.50	0.001	-3.06	0.001	Glycolysis
1424451_at	3-Ketoacyl-CoA thiolase B	2.48	0.001	-1.41	0.000	Fatty acid oxidation
1424316_at	Solute carrier family 25 (mitochondrial deoxynucleotide carrier), member 19	2.44	0.000	-2.21	0.001	Mitochondrial
1425753_a_at	Uracil-DNA glycosylase	2.38	0.044	-5.93	0.001	
1431592_a_at	SH3-domain kinase binding protein 1	2.38	0.000	-1.69	0.001	
1431143_x_at	Apoptosis-inducing factor (AIF)-like mitochondrion- associated inducer of death	2.36	0.004	-1.95	0.017	
1459274_at	G protein-coupled receptor 135	2.36	0.002	-1.14	0.572	
1452298_a_at	Myosin Vb	2.34	0.004	-3.67	0.015	
1448615_at	Copper chaperone for superoxide dismutase	2.31	0.000	-1.43	0.041	
1435872_at	Proviral integration site 1	2.29	0.009	-1.89	0.004	
1436934_s_at	Aconitase 2, mitochondrial	2.27	0.001	-1.42	0.001	TCA cycle
1460337_at	SH3-domain kinase binding protein 1	2.26	0.000	-1.77	0.002	
1418328_at	Carnitine palmitoyltransferase 1b, muscle	2.24	0.019	-3.98	0.000	Fatty acid oxidation
1430128_a_at	Deleted in polyposis 1-like 1	2.22	0.000	-1.99	0.003	
1424359_at	5-Oxoprolinase (ATP-hydrolysing)	2.21	0.001	-4.87	0.000	Glutathionine metabolism
1436521_at	Solute carrier family 36 (proton/amino acid symporter), member 2	2.20	0.003	-3.04	0.000	Amino acid transporter
1419714_at	Programmed cell death 1 ligand 1	2.18	0.002	-3.88	0.007	
1448237_x_at	Lactate dehydrogenase 2, B chain	2.17	0.000	-1.52	0.001	
1448830_at	Dual specificity phosphatase 1	2.17	0.005	2.71	0.113	
1424737_at	Thyroid hormone responsive SPOT14 homolog (Rattus)	2.16	0.006	-2.69	0.001	Fatty acid synthesis
1426989_at	Calsyntenin 3	2.13	0.001	1.45	0.375	
1416183_a_at	Lactate dehydrogenase 2, B chain	2.11	0.000	-1.65	0.000	Glycolysis
1430896_s_at	Nudix (nucleoside diphosphate linked moiety X)-type motif 7	2.11	0.000	-2.55	0.000	
1434499_a_at	Lactate dehydrogenase 2, B chain	2.10	0.000	-1.57	0.020	Glycolysis
1451899_a_at	General transcription factor II repeat domain-containing 1	2.09	0.011	-1.43	0.002	
1423108_at	Solute carrier family 25 (mitochondrial carnitine/ acylcarnitine translocase), member 20	2.07	0.001	-2.18	0.000	Fatty acid oxidation
1455235_x_at	Lactate dehydrogenase 2, B chain	2.05	0.000	-1.59	0.001	Glycolysis
1418472_at	Aspartoacylase (aminoacylase) 2	2.03	0.000	-2.91	0.002	
1448681_at	Interleukin 15 receptor, α chain	2.03	0.001	-1.35	0.078	
1435339_at	Potassium channel tetramerisation domain containing 15	2.01	0.002	-2.83	0.003	

3T3-L1 adipocytes transfected with scrambled or RIP140 siRNA were used for Affymetrix GeneChip analysis. Shown are all named genes significantly upregulated 2-fold or more upon RIP140 depletion. Corresponding changes in gene expression in RIPKO cells, in which RIP140 was reexpressed by lentivirus-RIP140 (lenti-RIP140) infection, are also shown. Boldface indicates statistical significance ($P < 0.05$).

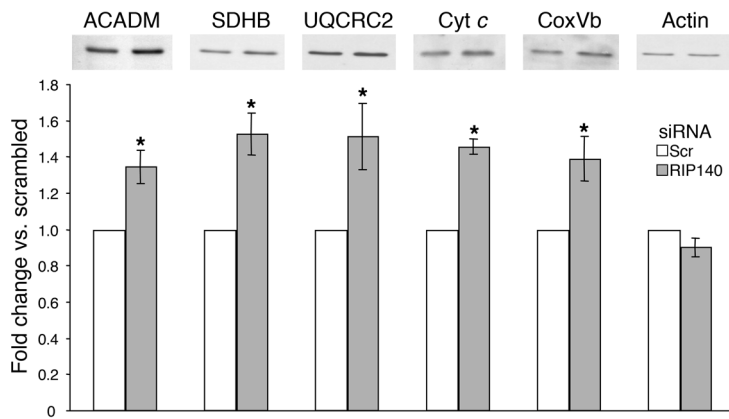


Figure 6

Metabolic and mitochondrial protein expression is enhanced in 3T3-L1 adipocytes depleted of RIP140. Eight days after the induction of differentiation, 3T3-L1 adipocytes were transfected with scrambled or RIP140 siRNA. After 72 hours, post-nuclear supernatants were used for immunoblotting selected proteins in oxidative phosphorylation and fatty acid oxidation pathways. Shown is a representative blot and densitometric quantification of 3 independent experiments (mean \pm SEM). All of the changes shown (except those for actin) were significant at $P < 0.05$ (indicated by asterisk).

tured adipocytes showing that RIP140 is a negative regulator of oxidative metabolism, although more work is needed to determine whether the role of RIP140 in adipocytes is sufficient to explain the whole-body effects.

RIP140 could regulate the expression of metabolic genes by repressing nuclear hormone receptors that act directly at the promoters of these genes. Alternatively, RIP140 could repress nuclear receptors that act indirectly by regulating expression levels of transcription factors that in turn regulate metabolic gene promoters. To evaluate the latter possibility, we examined the microarray-based gene expression database described above for expression profiles of transcription factors and cofactors known to regulate genes encoding mitochondrial proteins. The PPAR γ coactivator PGC-1 α , which has been shown to regulate mitochondrial biogenesis in brown adipose tissue, muscle, and liver (16, 28), is expressed at very low levels in white adipose tissue and 3T3-L1 adipocytes, and its expression did not change significantly upon RIP140 depletion. Expression of nuclear respiratory factor 1 (28) and thyroid hormone receptor (29), both reported to be important in mitochondrial biogenesis, was unchanged as well. However, a modest but significant increase in the expression of the PPAR γ coactivator PGC-1 β in RIP140-depleted cells was evident (Table 2). PGC-1 β is known to enhance expression of genes related to energy metabolism (18, 30). Furthermore, increases in the expression of ERR α and GA repeat-binding protein α (GABP α), which have been reported to act downstream of PGC-1 α to regulate oxidative phosphorylation in skeletal muscle (31, 32), were observed (Table 2). Finally, the absence of changes in expression of Tfam, Tfb1, and Tfb2 might reflect the absence of increased transcription or replication of the mitochondrial genome. However, the elevated levels of mitochondrial ribosomal proteins observed in RIP140-depleted cells might increase translation of the 13 subunits of the respiratory chain encoded in mitochondrial DNA.

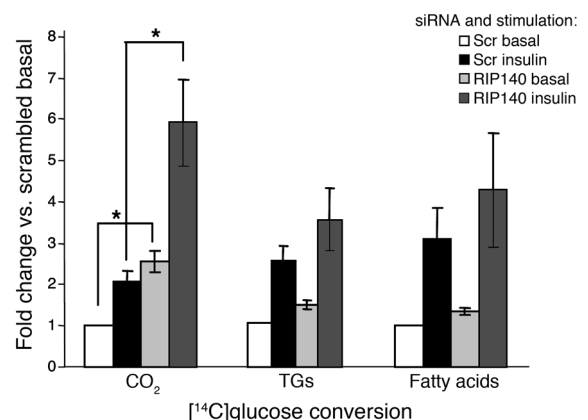
Figure 7

RIP140 depletion enhances glycolysis and TCA cycling, but not triglyceride synthesis or de novo free fatty acid synthesis. Eight days after the induction of differentiation, 3T3-L1 adipocytes were transfected with scrambled or RIP140 siRNA. After 72 hours, cells were starved and glucose metabolism (with continued starvation [basal] or 1 μ M insulin stimulation) was measured by [6- 14 C]-glucose conversion into carbon dioxide (CO $_2$), triglycerides, and fatty acids as described in Methods. Graphs show the mean \pm SEM of 3–4 independent experiments. * $P < 0.05$ compared with scrambled, by Student's t test.

Taken together, these results indicate that RIP140 suppresses the expression of several positive regulators of mitochondrial protein expression, providing 1 potential mechanism for its action to increase mitochondrial protein levels.

The enhancement in insulin-responsive deoxyglucose uptake and mitochondrial protein expression caused by RIP140 depletion resembles the increased insulin sensitivity (33), glucose uptake (34–36), and mitochondrial biogenesis (10) elicited by treatment of adipocytes with rosiglitazone, a PPAR γ agonist. RIP140 was shown to interact with PPAR γ in a yeast 2-hybrid assay and in vitro (37). To investigate whether RIP140 regulates GLUT4 expression and glucose uptake by modulating PPAR γ activity, we used siRNA to deplete RIP140, PPAR γ , or both proteins in fully differentiated 3T3-L1 adipocytes. As expected, depletion of RIP140 enhanced GLUT4 expression and deoxyglucose uptake (Figure 10). PPAR γ depletion resulted in reduced GLUT4 levels, which is consistent with the requirement of PPAR γ for the maintenance of the adipocyte phenotype (38). Interestingly, depletion of RIP140 in combination with PPAR γ depletion caused an enhancement of GLUT4 expression and deoxyglucose uptake compared with PPAR γ depletion alone. This suggests that RIP140 may be able to regulate GLUT4 expression independently of PPAR γ and that other nuclear receptors are involved.

We thus investigated whether RIP140 uses an ERR α -dependent mechanism to regulate GLUT4 and mitochondrial gene expression, as ERR α is thought to be an important regulator of energy balance (39) and oxidative phosphorylation-related gene expression (32) and its expression is increased in RIP140-depleted adi-



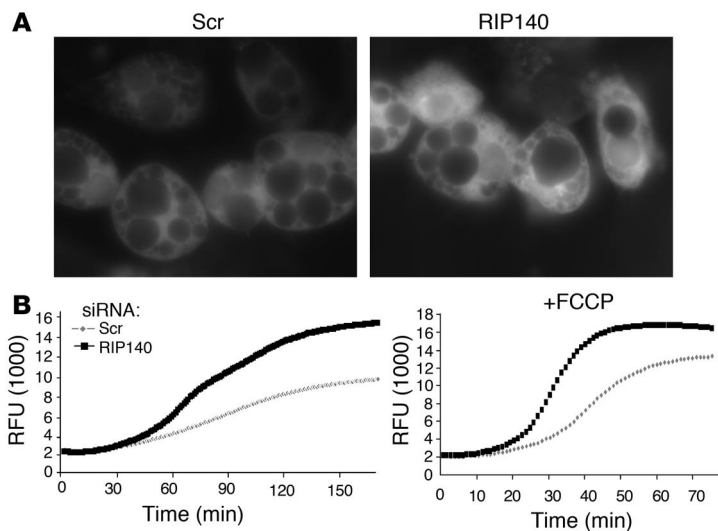


Figure 8 RIP140 is a negative regulator of cellular respiration in 3T3-L1 adipocytes. Eight days after the induction of differentiation, 3T3-L1 adipocytes were transfected with scrambled or RIP140 siRNA. (A) After 72 hours, cells were incubated with MitoTracker Red to stain actively respiring mitochondria. Fluorescence microscopy was performed as described in Methods. Shown are representative photographs with identical exposure times. (B) Cells were assayed for respiration using the oxygen biosensor fluorescence assay described in Methods. Basal respiration is indicated on the left, and uncoupled respiration, induced by addition of the mitochondrial uncoupler carbonyl cyanide 4-(trifluoromethoxy)phenylhydrazone (FCCP; 33 μ M), is shown on the right. Illustrated are the averages of 3–4 independent experiments. The respiration with or without FCCP of cells transfected with scrambled or RIP140 siRNA are considered statistically different based on the fluorescence intensity at the time when fluorescence of the leading curve was equal to half-maximum for each experiment. RFU, raw fluorescence units.

pocytes (Table 2). ERR α depletion alone in 3T3-L1 adipocytes modestly decreased GLUT4 and SDHB levels and inhibited deoxyglucose uptake compared with cells treated with scrambled siRNA (Figure 11). Strikingly, the loss of ERR α prevented the increased GLUT4 and SDHB expression and deoxyglucose uptake due to RIP140 depletion. Similar results were seen with the mitochondrial protein CoxVb (data not shown). These data suggest that RIP140 regulates GLUT4 and mitochondrial gene expression by a mechanism that is dependent upon ERR α .

Discussion

The findings presented here demonstrate that the transcriptional corepressor RIP140 negatively regulates major gene clusters in the pathways of carbohydrate and fatty acid catabolism and mitochondrial biogenesis in cultured adipocytes. Depletion of RIP140 in 3T3-L1 adipocytes caused increased expression of clusters of genes in these metabolic pathways, whereas its reintroduction into RIPKO-1 adipocytes caused decreased expression of these same genes (Figures 4 and 5 and Supplemental Table 3). Expression of large numbers of genes encoding proteins that function in all of the mitochondrial electron transport complexes was increased by RIP140 loss in adipocytes and resulted in an increase in overall mitochondrial functioning (Figures 7 and 8). Specifically, RIP140 suppression enhanced hexose uptake (Figures 1 and 2) and conversion of labeled glucose to CO₂ (Figure 7) and markedly elevated mitochondrial oxygen consumption in cultured adipocytes (Figure 8). Our results also agree with those of Leonardsson et al. (23) and Soukas et al. (27), demonstrating that RIP140 expression is markedly increased during adipocyte differentiation from preadipocytes (Supplemental Table 2). Taken together, these findings

indicate that in adipocytes, the highly expressed protein RIP140 exerts a major suppressive effect on oxidative metabolism in the glycolytic and TCA cycle pathways as well as mitochondrial biogenesis, in addition to its known role in fatty acid oxidation.

The dramatic effect of RIP140 depletion in upregulating genes encoding metabolic enzymes and proteins that catalyze mitochondrial respiration in adipocytes raised interesting questions about the potential consequences of RIP140 deletion on whole-body glucose homeostasis. Adipogenesis itself enhances insulin sensitivity of glucose utilization as well as mitochondrial gene expression (10), both of which are expected to enhance triglyceride synthesis. Lipogenesis in adipocytes in turn is thought to be 1 pathway whereby fatty acids are sequestered away from other tissues such as muscle, reducing insulin resistance (33). Interestingly, obesity and insulin resistance in *ob/ob* mice are accompanied by a decrease in mitochondrial gene expression in white adipose tissue (12).

Figure 9

Absence of RIP140 protects against age- and diet-induced glucose intolerance and enhances insulin responsiveness on a high-fat diet. Glucose tolerance tests were performed on 3-month-old female mice fed a normal chow diet (A), 8-month-old female mice fed a normal chow diet (B), or 3-month-old female mice fed a high-fat diet (C) for 10 weeks. Values are mean \pm SEM; *n* = 6–12. (D) Insulin tolerance tests were performed on 3-month-old female mice fed a high-fat diet for 9 weeks. Values are mean \pm SEM; *n* = 9.

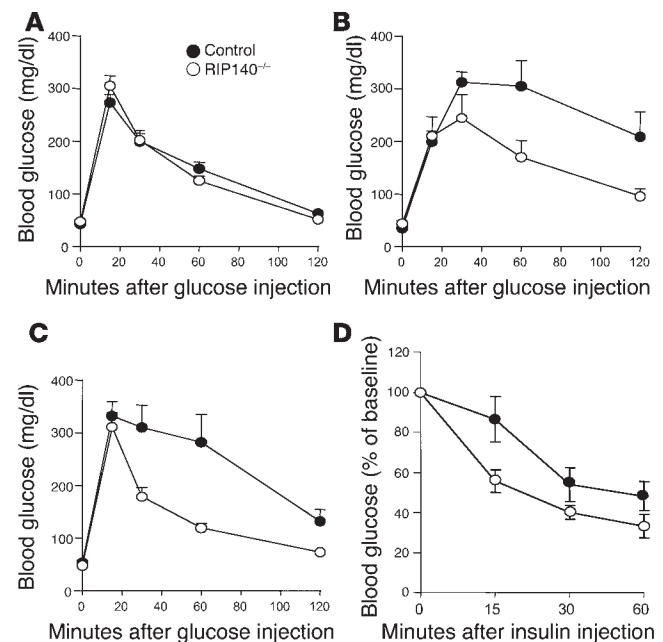




Table 2

Increased expression of transcription factors that regulate metabolism, including PGC-1 β , ERR α , and GABP α , is caused by RIP140 depletion

Probe ID	Symbol	Fold change	P value	Gene name
1418982_at	<i>Cebpa</i>	-1.03	0.746	<i>CCAAT/enhancer binding protein (C/EBP), α</i>
1418901_at	<i>Cebpb</i>	1.49	0.002	<i>CCAAT/enhancer binding protein (C/EBP), β</i>
1423233_at	<i>Cebpd</i>	-1.29	0.609	<i>CCAAT/enhancer binding protein (C/EBP), δ</i>
1425261_at	<i>Cebpg</i>	-1.26	0.002	<i>CCAAT/enhancer binding protein (C/EBP), γ</i>
1420497_a_at	<i>Cebpz</i>	-1.37	0.009	<i>CCAAT/enhancer binding protein (C/EBP), ζ</i>
1460652_at	<i>Esrra</i>	1.61	0.002	<i>Estrogen-related receptor, α</i>
1441921_x_at	<i>Esrrb</i>	1.01	0.937	<i>Estrogen-related receptor, β</i>
1418929_at	<i>Esrrb1</i>	-1.16	0.421	<i>Estrogen-related receptor β like 1</i>
1450665_at	<i>Gabpa</i>	1.38	0.018	<i>GA repeat binding protein, α (NRF-2)</i>
1436232_a_at	<i>Gabpb1</i>	-1.15	0.238	<i>GA repeat binding protein, β 1</i>
1453682_at	<i>Gabpb2</i>	-1.10	0.252	<i>GA repeat binding protein, β 2</i>
1443952_at	<i>Nr1d1</i>	-1.15	0.023	<i>Rev-Erba α</i>
1416958_at	<i>Nr1d2</i>	1.06	0.483	<i>Rev-Erba β</i>
1416353_at	<i>Nr1h2</i>	1.19	0.001	<i>LXR β</i>
1450444_a_at	<i>Nr1h3</i>	-1.18	0.005	<i>LXR α</i>
1416159_at	<i>Nr2f2</i>	-1.23	0.039	<i>COUP/TFII β</i>
1460648_at	<i>Nr2f6</i>	1.17	0.047	<i>EAR2</i>
1416505_at	<i>Nr4a1</i>	1.82	0.001	<i>Nur77</i>
1455034_at	<i>Nr4a2</i>	-1.32	0.011	<i>Nurr1</i>
1417098_s_at	<i>Nrbf1</i>	1.58	0.018	<i>Nuclear receptor binding factor 1</i>
1448758_at	<i>Nrbf2</i>	-1.06	0.231	<i>Nuclear receptor binding factor 2</i>
1423864_at	<i>Nrbp</i>	1.05	0.575	<i>Nuclear receptor binding protein</i>
1424787_a_at	<i>Nrf1</i>	-1.01	0.685	<i>Nuclear respiratory factor 1</i>
1418469_at	<i>Nrip1</i>	-3.53	0	<i>Nuclear receptor interacting protein 1</i>
1423451_at	<i>Pgrmc1</i>	-1.13	0.031	<i>Progesterone receptor membrane component 1</i>
1452882_at	<i>Pgrmc2</i>	-1.22	0.027	<i>Progesterone receptor membrane component 2</i>
1449051_at	<i>Ppara</i>	1.61	0.012	<i>Peroxisome proliferator activated receptor α</i>
1448708_at	<i>Pparbp</i>	1.09	0.044	<i>PPAR binding protein</i>
1439797_at	<i>Ppard</i>	-1.04	0.655	<i>Peroxisome proliferator activated receptor δ</i>
1420715_a_at	<i>Pparg</i>	1.01	0.94	<i>Peroxisome proliferator activated receptor γ</i>
1460336_at	<i>Ppargc1a</i>	1.05	0.838	<i>PPAR gamma, coactivator 1 α</i>
1449945_at	<i>Ppargc1b</i>	1.33	0.006	<i>PPAR gamma, coactivator 1 β</i>
1425762_a_at	<i>Rxra</i>	-1.26	0.018	<i>Retinoid X receptor α</i>
1416990_at	<i>Rxrb</i>	1.13	0.062	<i>Retinoid X receptor β</i>
1418782_at	<i>Rxrg</i>	1.15	0.313	<i>Retinoid X receptor γ</i>
1426690_a_at	<i>Srebf1</i>	-1.13	0.144	<i>Sterol regulatory element binding factor 1</i>
1452174_at	<i>Srebf2</i>	-1.03	0.532	<i>Sterol regulatory element binding factor 2</i>
1456215_at	<i>Tfam</i>	1.01	0.858	<i>Transcription factor A, mitochondrial</i>
1435924_at	<i>Tfb1m</i>	1.05	0.627	<i>Transcription factor B1, mitochondrial</i>
1423441_at	<i>Tfb2m</i>	1.16	0.133	<i>Transcription factor B2, mitochondrial</i>
1454675_at	<i>Thra</i>	-1.07	0.486	<i>Thyroid hormone receptor α</i>

3T3-L1 adipocytes transfected with scrambled or RIP140 siRNA were used for Affymetrix GeneChip analysis. Fold change indicates the difference in expression between scrambled and RIP140 siRNA-transfected cells, with boldface indicating significance ($P < 0.05$).

Long-term calorie restriction, which results in enhanced insulin sensitivity, causes an increase in the expression of mitochondrial genes in white adipose tissue (40). Also, subcutaneous fat, which is more insulin sensitive than visceral fat, has a higher mitochondrial density, mitochondrial DNA content, and mitochondrial activity than visceral fat (41). Furthermore, rosiglitazone treatment of *ob/ob* mice enhances expression of genes encoding mitochondrial proteins in adipose tissue while at the same time enhancing muscle insulin sensitivity and glucose tolerance (10). These data, coupled with results from experiments showing inverse relationships between the amount of GLUT4 glucose transporters in adipocytes and whole-body insulin resistance (4, 42, 43), reinforce the

hypothesis that oxidative metabolism in adipocytes may indirectly control muscle insulin sensitivity and whole-body glucose homeostasis.

The above considerations prompted us to test whether RIP140-null mice might exhibit increased glucose tolerance and insulin sensitivity. RIP140-null mice were previously shown to be lean and resistant to high-fat diet-induced obesity (23). However, these mice did not initially show a change in glucose tolerance at early ages on a normal diet. Here we show that RIP140-null mice are resistant to the impaired glucose tolerance and poor insulin responsiveness observed in wild-type mice fed a high-fat diet (Figure 9, C and D). Enhanced glucose tolerance in aged RIP140-null mice was also observed compared with matched wild-type mice (Figure 9B). Taken together, these data indicate that the increased glucose tolerance observed in RIP140-null mice on a high-fat diet is due to increased insulin responsiveness. Thus, RIP140 may be an important modulator of potential linkages between adipocyte metabolism and whole-body glucose tolerance. Our data show only a correlation between these processes at this stage, which should provoke further work to rigorously test this hypothesis.

The expression of genes encoding mitochondrial proteins in skeletal muscle has recently been correlated with insulin sensitivity in humans (44). Many studies focused on skeletal muscle have indicated the importance of mitochondrial size and number (45, 46) and expression of genes encoding proteins that catalyze oxidative phosphorylation and mitochondrial capacity (13, 14, 47–50). A recent study also found that depletion of mitochondrial DNA leads to a dramatic decrease in glucose uptake

in skeletal muscle (51). It will thus be interesting to determine whether RIP140 modulates energy metabolism in skeletal muscle, as it does in fat. The degree to which RIP140 depletion in muscle or other tissues contributes to the enhanced glucose tolerance and insulin sensitivity of RIP140-null mice is an important question and remains to be determined.

While our data show that many proteins of the mitochondrial electron transport chain and oxidative phosphorylation are negatively controlled by RIP140, the expression of mitochondrial uncoupling proteins was not apparently modulated by RIP140 depletion in 3T3-L1 adipocytes (data not shown). In contrast, Leonardsson et al. did observe expression of the uncoupling

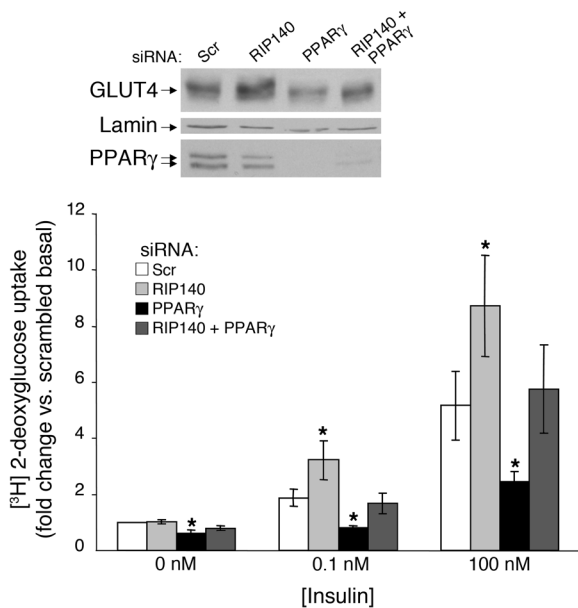


Figure 10

RIP140 depletion does not require normal PPAR γ levels to enhance GLUT4 expression. Eight days after the induction of differentiation, cells were transfected with scrambled, RIP140, PPAR γ , or RIP140 plus PPAR γ siRNA. Three days later, cell lysates were used for immunoblotting GLUT4 and actin, while nuclear extracts were used to immunoblot PPAR γ and lamin. A 2-deoxyglucose uptake assay was performed with the indicated knockdowns. The graph shows the average and SE of 5 independent experiments. * $P < 0.05$ compared with scrambled siRNA, by Student's t test.

protein UCP1 in white adipose tissue of RIP140-null mice (23), a protein normally expressed only in brown fat, as did Christian et al. in differentiated RIP140-null MEFs (24). This discrepancy might be due to the presence of RIP140 in 3T3-L1 adipocytes prior to its depletion by siRNA and during a critical time period when UCP1 expression is determined, whereas in RIP140-null mice and MEFs, RIP140 was never present. UCP1 is thought to divert the energy derived from mitochondrial electron transport that would normally generate ATP to produce heat. This uncoupling effect greatly accelerates oxygen consumption. In intact mice, RIP140 might act to divert white adipocytes from a brown fat phenotype by attenuating the expression of UCP1 and genes involved in fatty acid oxidation and mitochondrial respiration, all of which are abundantly expressed in brown fat. Interestingly, conversion of white adipocytes to a brown fat phenotype has been achieved by ectopic expression of PGC-1 α in human adipocytes (52), suggesting that RIP140 and the PGC family coactivators may have antagonizing roles in adipocyte development.

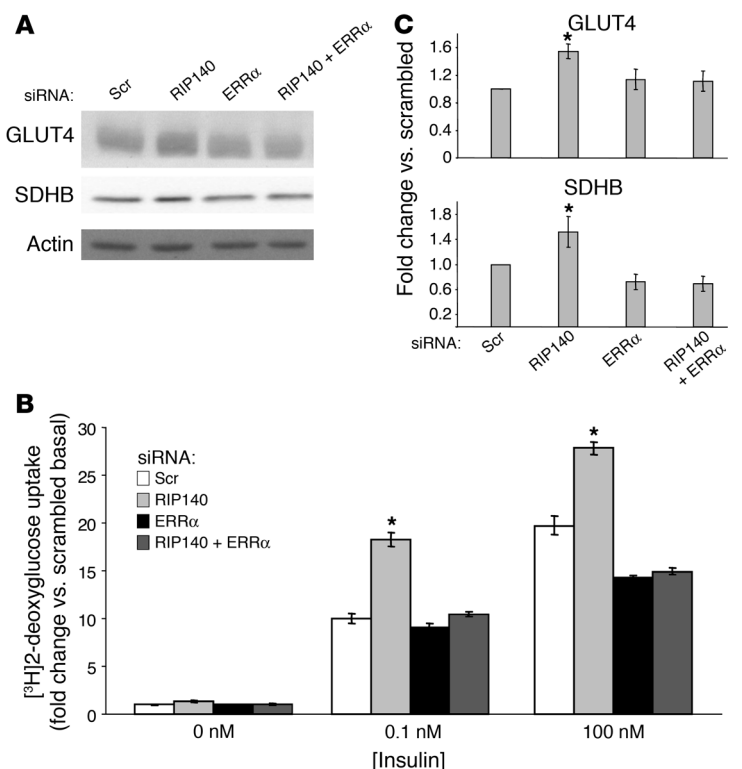
RIP140 appears to interact with multiple nuclear receptors through its LXXLL motifs, including PPAR γ (37), which promotes and maintains the adipocyte phenotype (53). Both RIP140 depletion and the treatment of 3T3-L1 adipocytes or obese *ob/ob* mice with the PPAR γ agonist rosiglitazone cause an increase in the expression of genes

encoding mitochondrial proteins (12). We therefore tested whether RIP140 might act by a mechanism involving direct repression of PPAR γ to regulate glucose uptake and GLUT4 expression. PPAR γ depletion in mature adipocytes by siRNA transfection did decrease deoxyglucose transport and GLUT4 expression, as expected. However, the loss of RIP140 in these cells was still able to enhance these processes, even with greatly reduced amounts of PPAR γ (Figure 10). Taken together, these data indicate that PPAR γ is either not a major target of RIP140-mediated repression responsible for regulating GLUT4 expression or that other pathways can also mediate this response in parallel. Further experiments will be required to determine whether other genes responsive to RIP140 involve PPAR γ regulation by the corepressor.

RIP140 depletion may increase gene expression in cultured adipocytes by upregulating the expression of ERR α (Table 2). We found that RIP140 knockdown requires the presence of ERR α to enhance GLUT4 expression and deoxyglucose uptake (Figure 11), suggesting a potential mechanism in which RIP140 represses ERR α -mediated transcription of GLUT4 directly or indirectly through other regulatory genes. ERR α is also required for RIP140 regulation of other genes, including oxidative phosphorylation

Figure 11

RIP140 depletion requires ERR α to enhance GLUT4 expression, deoxyglucose uptake, and mitochondrial protein expression. Eight days after the induction of differentiation, cells were transfected with scrambled, RIP140, ERR α , or RIP140 plus ERR α siRNA. (A) Three days later, cell lysates were used for immunoblotting GLUT4, SDHB, and actin. A representative blot and the mean \pm SEM of 4–5 independent experiments are shown. (B) A 2-deoxyglucose uptake assay was performed with the indicated knockdowns. A representative blot and the mean \pm SEM of 3 independent experiments are shown. * $P < 0.05$ compared with scrambled siRNA, by Student's t test.





proteins such as CoxVb and SDHB. Interestingly, expression of a transcriptionally active $ERR\alpha$ was sufficient to induce mitochondrial biogenesis in an osteoblast cell line with adipocyte differentiation capacity (31). Thus, RIP140 depletion may activate $ERR\alpha$, leading to an increase in mitochondrial content and GLUT4 expression. The activation of $ERR\alpha$ might also play a role in the increases in expression of genes that mediate fatty acid oxidation (18), although further work is needed to test this hypothesis.

The repression of genes related to oxidative phosphorylation and fatty acid oxidation by RIP140 is likely to involve additional factors. In skeletal muscle, for example, the coactivator PGC-1 α induces $ERR\alpha$ and GABP α , which then turn on the transcription of genes that encode a subset of the proteins that mediate oxidative phosphorylation (32). Thus, RIP140 depletion may require GABP α and the related coactivator PGC-1 β , a coactivator for $ERR\alpha$ (18), to enhance oxidative phosphorylation gene expression. Similarly, PPAR α may be required for $ERR\alpha$ -induced increases in fatty acid oxidation (54), although both PGC-1 β (55, 56) and $ERR\alpha$ (54) have been shown to activate PPAR α . Interestingly, each of these additional regulators (GABP α , PGC-1 β , and PPAR α) exhibit increased expression upon RIP140 depletion (Table 2). Finally, several other transcription factors are increased in RIP140-depleted adipocytes and may contribute to the enhanced insulin sensitivity and metabolism of these cells.

In summary, we have shown here that the metabolic pathways suppressed by RIP140 in adipocytes include glycolysis, triglyceride synthesis, the TCA cycle, fatty acid β oxidation, mitochondrial electron transport, and oxidative phosphorylation. Indeed, glucose oxidation and oxygen consumption are significantly increased in adipocytes when RIP140 function is attenuated by siRNA silencing. RIP140 regulates GLUT4 and mitochondrial protein expression by a mechanism that is dependent upon $ERR\alpha$ in cultured adipocytes, although GLUT4 is not regulated in the adipose tissue of RIP140-null mice. Finally, RIP140-null mice display enhanced glucose and insulin tolerance after high-fat feeding compared with matched wild-type mice. Since RIP140 exerts broad negative regulation of multiple metabolic pathways in adipocytes, it may in part mediate changes in adipocyte metabolism in response to a variety of physiological conditions in order to effectively alter whole-body glucose metabolism. Furthermore, our results suggest that RIP140 may be a candidate therapeutic target for the syndromes of obesity and type 2 diabetes.

Methods

Materials and chemicals. Human insulin was obtained from Eli Lilly and Co. Rabbit polyclonal antibodies against phospho-Akt (Ser473), phospho-Erk1/2, anti-Akt, and anti-Erk1/2 were from Cell Signaling Technology. Other antibodies included goat anti-GLUT4 (Santa Cruz Biotechnology Inc.), mouse anti-actin (Sigma-Aldrich), rabbit anti-GLUT1 (gift from Paul Pilch, Boston University, Boston, Massachusetts, USA), mouse anti-cyt *c* (BD Biosciences – Pharmingen), and mouse anti-CoxVb (MitoSciences). Goat anti-ACADM, mouse anti-UQCRC2, and mouse anti-SDHB antibodies were from Abcam. The HRP-conjugated goat anti-rabbit IgG and rabbit anti-goat IgG were from Chemicon International.

Subtractive cDNA library. The Muscle/Adipocyte Union cDNA library was made using a modification of the subtractive suppression hybridization technique of the Clontech PCR-Select cDNA Subtraction Kit (57).

siRNA. siRNA was purchased from Dharmacon. Smart pools of siRNA consisting of a mix of 4 individual oligonucleotides were used for the screen and to deplete $ERR\alpha$. Individual siRNA sequences included: scrambled,

5'-CAGTCGCGTTTGGCGACTGG-3'; RIP140, 5'-GGAATGAGCTCGAT-TATAA-3'; and PPAR γ , 5'-GACATGAATTCCTTAATGA-3'. Smart pool and individual siRNA-mediated depletion of RIP140 and PPAR γ produced similar experimental results.

Cell culture and electroporation of 3T3-L1 adipocytes. 3T3-L1 fibroblasts and RIP140-null MEFs, referred to as RIPKO cells (24), were grown in DMEM supplemented with 10% FBS, 50 μ g/ml streptomycin, and 50 U/ml penicillin and differentiated into adipocytes as described previously (25). 3T3-L1 adipocytes were transfected with siRNA duplexes by electroporation on the fourth or eighth day following the induction of differentiation (day 4 or day 8) as described previously (25). The efficiency of siRNA electroporation using this method is greater than 95%, based on fluorescence microscopy of cells electroporated with Cy3-siRNA.

2-Deoxyglucose uptake assay. Insulin-stimulated glucose transport in 3T3-L1 adipocytes was estimated by measuring 2-deoxyglucose uptake using a miniaturized version of the typical assay with cells seeded into 96-well plates (25). Cells were solubilized with 0.1 ml of 1% SDS, and 3 H uptake was quantitated by scintillation counting. Nonspecific deoxyglucose uptake was measured in the presence of 20 μ M cytochalasin B and subtracted from each determination to obtain specific uptake.

Quantitative RT-PCR. Adipocytes were transfected as described above. Total RNA was prepared 24 hours after transfection using TRIzol reagent (Invitrogen Corp.). cDNA was synthesized using oligo-dT primers and AMV reverse transcriptase (Roche Diagnostics Corp.) according to manufacturer's instructions. Quantitative real-time PCR was performed in a LightCycler (Roche Diagnostics Corp.) using a SYBR Green I PCR kit (Roche Diagnostics Corp.) and specific primers to amplify a 553-bp fragment from RIP140 or a 500-bp fragment from hypoxanthine phosphoribosyl transferase.

Immunoblotting. Cell lysates were harvested by the addition of SDS lysis buffer (2% SDS, 30 mM NaCl, 10 mM HEPES, pH 7.4, 20 mM NaF, 1 mM NaPPI). For detecting phospho-proteins, cells were starved overnight in serum-free DMEM medium with 0.5% BSA. Cells were then incubated with 0, 1, or 100 μ M insulin for 30 minutes and harvested with 1% SDS lysis buffer as described above. For enrichment of nuclear proteins (PPAR γ , lamin A/C blots), cells were lysed in hypotonic lysis buffer (20 mM Tris-HCl, pH 7.5, 10 mM NaCl, 3 mM MgCl $_2$, 1 mM DTT, 1 mM sodium orthovanadate, 30 mM β -glycerophosphate, 1.5% NP-40) using a Dounce homogenizer. Nuclei were pelleted (3,500 g for 5 minutes at 4°C), washed in hypotonic lysis buffer, and pelleted again. They then were resuspended in NUN buffer (1.1 M urea, 330 mM NaCl, 1.1% NP-40, 17.5 mM HEPES, pH 7.6, 100 mM DTT, 100 mM sodium orthovanadate, 3 M β -glycerophosphate) and incubated on ice for 30 minutes. A final spin at 16,000 g for 10 minutes at 4°C was performed, and the extracted nuclear proteins in the supernatant were mixed with glycerol and stored at -80°C. Postnuclear supernatants were used for immunoblotting ACADM, cyt *c*, CoxVb, UQCRC2, and SDHB. Equal amounts of protein from lysates were resolved by SDS-PAGE, immunoblotted, and detected with an enhanced chemiluminescence kit (PerkinElmer).

Affymetrix GeneChip analysis. Isolation of RNA and Affymetrix GeneChip analysis were performed as described for fibroblasts, 3T3-L1 adipocytes, and primary fat cells (10). For RIP140 3T3-L1 Affymetrix GeneChip analysis, cells were transfected as described above. RNA was prepared from TRIzol following manufacturer's instructions, and cDNA and cRNA were made using Affymetrix's One-Cycle cDNA Synthesis and IVT Labeling Kits. cRNA was hybridized to Affymetrix GeneChip Mouse Genome 430 2.0 array. All microarray experiments were performed in triplicate using separate plates of cells.

3T3-L1 GeneChip data was analyzed using the robust multiarray average algorithm as a function of the BioConductor Affy R package. The pre-processed data is stored as log $_2$ -transformed real signal numbers. The fold



change of expression of a gene in 2 experiments is the ratio of mean signal values from these experiments. To determine differential expression of genes in 2 hybridization experiments, we conducted a 2-tailed Student's *t* test with the expression signal values of the 2 hybridizations for all genes in the set. The *P* values were not adjusted for multiple testing. *P* < 0.05 was considered statistically significant.

For Affymetrix analysis of gene expression in MEFs derived from RIP140-null mice, with or without RIP140 lentiviral reexpression, total RNA was isolated from 3 samples each after adipocyte differentiation (10 days) in the presence of the PPAR γ ligand GW1929 (2.5 μ M; Sigma-Aldrich). Two RIP140-lentivirus transduced cell lines were used for profiling in order to avoid variation due to different sites of incorporation of the virus. Affymetrix array hybridization and scanning were performed by the CSC/IC Microarray Centre, Imperial College London, Hammersmith Campus, using Murine 430 2.0 chips. Array data were analyzed with d-CHIP software and are available at <http://www.ebi.ac.uk/arrayexpress/>, accession number E-MIMR-42.

Tables list only genes indicated as present by Affymetrix, except for Supplemental Table 2. Only 1 representative probe set per gene is shown if more than 1 probe set was present. Boldface in Tables 1 and 2 and shading in supplemental tables indicate that gene expression was significantly changed, with *P* < 0.05.

Glucose conversion to CO₂, triglycerides, and free fatty acids. Adipocytes were transfected as described above and reseeded into 15-cm dishes. Cells were washed twice and starved for 2 hours with Krebs-Ringer-HEPES buffer (KRH buffer; 130 mM NaCl, 5 mM KCl, 1.3 mM CaCl₂, 1.3 mM MgSO₄, 25 mM HEPES, pH 7.4) supplemented with 2.5% BSA and 2 mM sodium pyruvate. Detached cells were aliquoted in triplicate to tubes containing supplemented KRH buffer (basal), 1 μ M insulin, or 20 μ M cytochalasin B, and D-[6-¹⁴C]-glucose was added. After 2 hours at 37°C, [³H]CO₂ was released by the addition of 2 M HCl, and collection filter papers were scintillation counted to measure glucose conversion to CO₂. Five milliliters of Dole's solution (80 ml isopropanol, 20 ml hexane, 2 ml 0.5 M H₂SO₄) was added and incubated for 15 minutes. Hexane was used to extract the lipids, and 1 ml was transferred to a scintillation vial, evaporated, and used to measure glucose conversion to triglycerides. Another 1 ml of the hexane phase was evaporated, and 1 ml of ethanolic KOH was added. Vials were heated at 80°C for 1 hour to release fatty acids from triglycerides and washed with water, and the pH was adjusted with 3 M sulfuric acid. Hexane was added, evaporated, and used to measure glucose conversion to fatty acids. DNA content was assessed using the QIAGEN DNeasy kit, and counts per minute were converted to picomoles glucose and normalized to the number of cells per tube.

Oxygen consumption assays. Three days following transfection with siRNA, adipocytes were resuspended in KRH buffer supplemented with 2.5% BSA and 2 mM sodium pyruvate. Equal numbers of cells were seeded in triplicate into BD Oxygen Biosensor System plates (BD Biosciences). Plates were pre-read to determine baseline fluorescence and were read on a Safire multimode microplate spectrophotometer (Tecan) at 1-minute intervals for 150 minutes. An excitation wavelength of 485 nm and emission wavelength of 630 nm were used. The mitochondrial uncoupler FCCP (Sigma-

Aldrich) was used at a final concentration of 33 μ M. Data were analyzed by averaging the triplicates in each experiment and determining the maximum fluorescence value for the leading curve. Fluorescence intensities were collected at the time point corresponding to half-maximal fluorescence of the leading curve.

Fluorescence microscopy. Adipocytes transfected with siRNA were seeded directly onto coverslips. After 3 days, mitochondrial respiration was measured by incorporation of 1 μ M MitoTracker Red (Molecular Probes; Invitrogen Corp.) for 30 minutes at 37°C. Cells were washed and fixed for 10 minutes using 4% formaldehyde. Coverslips were mounted onto DABCO on slides and sealed. Imaging was performed using a wide-field microscope with a \times 100 Nikon PlanApo objective. Image collection and quantitation were performed blindly. The total fluorescence intensity of a minimum of 50 cells was determined for each experiment by outlining each cell in Adobe Photoshop CS, version 8.0 (Adobe Systems Inc.) and multiplying the number of pixels by the average pixel intensity from the histogram function.

RIP140-null mice and glucose and insulin tolerance tests. RIP140-null mice were generated as previously described (22). Mice used in this study were backcrossed 8 generations to a C57BL/6J background. Mice were maintained under standard conditions, with controlled light and temperature, and fed a chow diet ad libitum, except in high-fat diet experiments, wherein mice were fed a 45% kcal fat diet (Research Diets). All animal work was performed with approval of the Home Office (London, United Kingdom). Glucose and insulin tolerance tests were performed on mice after a 16-hour overnight fast. Animals were injected intraperitoneally with D-glucose (2 g/kg) or insulin and blood glucose levels measured at the time points indicated. Blood glucose was measured using a Glucometer Elite (Bayer Corp).

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