Enhanced Metabolic Flexibility Associated with Elevated Adiponectin Levels

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Metabolically healthy individuals effectively adapt to changes in nutritional state. Here, we focus on the effects of the adipocyte-derived secretory molecule adiponectin on adipose tissue in mouse models with genetically altered adiponectin levels. We found that higher adiponectin levels increased sensitivity to the lipolytic effects of adrenergic receptor agonists. In parallel, adiponectin-overexpressing mice also display enhanced clearance of circulating fatty acids and increased expansion of subcutaneous adipose tissue with chronic high fat diet (HFD) feeding. These adaptive changes to the HFD were associated with increased mitochondrial density in adipocytes, smaller adipocyte size, and a general transcriptional up-regulation of factors involved in lipid storage through efficient esterification of free fatty acids. The physiological response to adiponectin overexpression resembles in many ways the effects of chronic exposure to β3-adrenergic agonist treatment, which also results in improvements in insulin sensitivity. In addition, using a novel computed tomography-based method for measurements of hepatic lipids, we resolved the temporal events taking place in the liver in response to acute HFD exposure in both wild-type and adiponectin-overexpressing mice. Increased levels of adiponectin potently protect against HFD-induced hepatic lipid accumulation and preserve insulin sensitivity. Given these profound effects of adiponectin, we propose that adiponectin is a factor that increases the metabolic flexibility of adipose tissue, enhancing its ability to maintain proper function under metabolically challenging conditions. (Am J Pathol 2010, 176:1364–1376; DOI: 10.2353/ajpath.2010.090647)

Ectopic storage of lipids in nonadipose tissue is widely accepted as a cause of metabolic dysregulation and occurs at variable degrees in obese or lipodystrophic individuals. This phenomenon is commonly referred to as lipotoxicity, and a large body of literature has revealed a variety of mechanisms underlying fatty acid-induced metabolic dysregulation. Furthermore, reduction of ectopic lipid storage has been proven successful in the prevention or reversal of metabolic disease. For instance, treatment with peroxisome proliferator-activated receptor (PPAR)-γ agonists (eg, thiazolidinediones) has been shown to decrease lipotoxicity through facilitating deposition of fatty acids in the adipose tissue and thereby sequestering the fatty acids away from nonadipose tissues.

Improved insulin sensitivity in rodents can also be achieved through chronic treatment with β3-adrenergic receptor (AR) agonists. The underlying mechanism behind the beneficial effects of β3-AR agonists probably involves reduction of circulating free fatty acids (FFAs) through increased capacity for fatty acid catabolism within the adipose tissue. Although human adipocytes have a negligible β3-adrenergic response, intact adrenergic signaling in adipose tissue may still be important for the maintenance of metabolic health. On the one hand, insulin resistance increases lipolysis, but on the other hand, it has also been reported that adipocytes from insulin-resistant subjects are less sensitive to catecholamine-induced lipolysis. Adrenergic signaling may also regulate adipose tissue biology through generation of cAMP, which is crucial for adipocyte differentiation. Thus, reduced sensitivity to both insulin and ad-
Adiponectin is a hormone that is secreted almost exclusively by adipose tissue. The circulating levels of adiponectin are suppressed in insulin-resistant states, whereas administration of insulin-sensitizing thiazolidinediones increases circulating adiponectin levels. Adiponectin is not only a marker for insulin sensitivity, but plays an active role in enhancing insulin sensitivity. Numerous studies suggest that adiponectin acts on the liver and suppresses glucose output. Moreover, adiponectin knockout (adipo−/−) mice have decreased hepatic insulin sensitivity and an impaired response to thiazolidinediones. In contrast, a mouse model with two- to threelfold elevated adiponectin levels (adiponectin transgenic [adipo tg] mice) is protected against HFD-induced insulin resistance.

Most studies on the effects of adiponectin have mainly explored the endocrine properties of adiponectin, ie, how circulating adiponectin improves insulin signaling in target tissues such as the liver and muscle. Little effort has been directed to the possibility that adiponectin is of importance for adipose tissue physiology directly. However, we have recently shown that overexpression of adiponectin enhances adiposity and protects against lipotoxicity in the metabolically compromised ob/ob mouse model. Given that complete leptin deficiency represents an extreme case, which may be less relevant for most cases of human obesity, we wanted to extend our studies of adiponectin in this regard. We first established an experimental setting in which vulnerability to lipotoxicity can easily be assessed in vivo. In a second step, taking advantage of transgenic and knockout mice with altered adiponectin levels, we demonstrate that adiponectin protects against lipid deposition in the liver. Finally, we show that the underlying mechanism is likely to involve enhanced adrenergic sensitivity in the adipose tissue, resembling the effects of chronic β3-AR agonist treatment. Therefore, we suggest that the actions of adiponectin on metabolism are well beyond the previously demonstrated endocrine effects and involve remodeling of the adipose tissue into a more efficient "metabolic sink." Given the profound effects on both FFA re-esterification as well as lipolysis, we propose that adiponectin confers metabolic flexibility to the system by balancing the ability to enhance expansion of adipose tissue while also enhancing the ability to respond to lipolytic stimuli.

**Materials and Methods**

**Animals**

Wild-type FVB and C57B6 mice were used to establish a computed tomography (CT)-based method for assessing liver lipids as well as for characterizing the acute metabolic effects of a HFD. Adiponectin-deficient mice (adipo−/−) on the FVB background and adipo tg on both FVB and C57B6 backgrounds were used to explore the actions of adiponectin. All mice were backcrossed more than 10 times to their respective background. Mice were maintained on a 12-hour dark/light cycle and housed in groups of two to four with unlimited access to water, chow (no. 5058, LabDiet, St. Louis, MO), or the HFD (D12492, Research Diets Inc., New Brunswick, NJ) as indicated for the individual experiments. The Institutional Animal Care and Use Committee of the University of Texas Southwestern Medical Center, Dallas approved all animal experiments.

**Computed Tomography**

Mice were anesthetized with isoflurane and a CT scan was performed at a resolution of 93 μm using the short scan mode (180°) on a eXplore Locus in vivo MicroCT scanner from GE Healthcare (Little Chalfont, Buckinghamshire, UK). The CT scanner was calibrated according to protocols provided by the manufacturer. Each scan took about 6 minutes, and the mice showed no sign of discomfort during the procedure. The images obtained were analyzed using Microview software (GE Healthcare). Adipose tissue distribution was assessed from the base of the lungs to the distal side of the hip joint. Histogram analysis of CT images obtained showed that voxels with CT values between −300 and −100 Hounsfield units (H.U.) may be defined as fat tissue. Visceral adipose tissue was analyzed by assessing the fat content within the peritoneal cavity using the clearly visible peritoneum as guidance. Liver lipid content was analyzed by obtaining the average CT value in multiple regions well within the liver. The average variability between analyses of the same liver was no higher than 1%. Because increased lipid content reduces the density, we assumed that an increased lipid content would result in a decreased average liver CT value. This assumption was tested and proven to be valid because a clearly significant correlation between average liver CT values and extracted liver triglycerides was obtained (Figure 1A). Thus, our CT-based method for estimation of hepatic lipid content is reproducible and accurate and greatly facilitates longitudinal measurements in response to metabolic challenges.

**Triglyceride Extraction**

Mice were anesthetized with ketamine (100 mg/kg)/xylazine (10 mg/kg). Livers were harvested, and 100- to 200-mg pieces were snap-frozen in liquid nitrogen until analysis. Lipids were extracted according to the method of Folch et al. The chloroform phase was brought up to 5 ml, and triplicates of 50 ml together with standards were dried down by the addition of 10 ml of 2:1 chloroform-methanol. Triglyceride levels were measured with Infinity reagent (Thermo Fisher Scientific, Waltham, MA).

**FFA Clearance**

Tail vein serum samples were obtained before and 2 hours after an oral administration of olive oil (10 μl/g).
β3AR-Agonist Sensitivity Test
Tail vein serum samples were obtained before and 5, 15, and 60 minutes after an intraperitoneal injection of 1 mg/kg CL 316,243 (Sigma-Aldrich, St. Louis, MO).

Blood Chemistry
Insulin, leptin, and adiponectin levels were measured by commercial enzyme-linked immunosorbent assay kits (Linco Research, St. Charles, MO). Glucose and glycerol levels were determined by, respectively, Sigma Diagnostics glucose reagents and free glycerol reagent (Sigma-Aldrich). Triglycerides and cholesterol levels were measured with Infinity reagents (Thermo Fisher Scientific). Free fatty acid levels were measured with NEFA-HR(2) (Wako Pure Chemical Industries, Tokyo, Japan).

DNA Array Analysis and Quantitative Real-Time RT-PCR
Tissues were collected in RNA later (Ambion, Austin, TX) and stored at −80°C until TRIzol reagent (Invitrogen, Carlsbad, CA) extraction followed by RNA purification using an RNeasy Mini Kit and RNase-Free DNase (Qiagen, Valencia, CA). The integrity of the RNA was estimated with an Agilent 2100 bioanalyzer instrument. All samples had an RNA integrity number score > 8. The Mouse WG-6 v2.0 Illumina bead chip platform (Illumina, Carlsbad, CA) was used for DNA array analysis. RNA was reverse-transcribed to cDNA by an iScript cDNA synthesis kit (Bio-Rad Laboratories, Hercules, CA). Gene expression was subsequently measured for individual genes (β-ARs, ATP6, mDIC, GPD2, PPARα, and PPARγ) to confirm results obtained by DNA array. IQ SYBR Green Supermix (Bio-Rad) was used for the quantitative PCR reactions and HPRT was used as an endogenous control. The relative expression level was calculated by the comparative Ct method. Primer sequences are indicated in Table 1.

Histology and Electron Transport Activity
Adipose tissue was collected in Histochoice and stored at −80°C until TRIzol reagent (Invitrogen, Carlsbad, CA) extraction followed by RNA purification using an RNeasy Mini Kit and RNase-Free DNase (Qiagen, Valencia, CA). Tissues were collected in RNA later (Ambion, Austin, TX) and stored at −80°C until TRIzol reagent (Invitrogen, Carlsbad, CA) extraction followed by RNA purification using an RNeasy Mini Kit and RNase-Free DNase (Qiagen, Valencia, CA). The integrity of the RNA was estimated with an Agilent 2100 bioanalyzer instrument. All samples had an RNA integrity number score > 8. The Mouse WG-6 v2.0 Illumina bead chip platform (Illumina, Carlsbad, CA) was used for DNA array analysis. RNA was reverse-transcribed to cDNA by an iScript cDNA synthesis kit (Bio-Rad Laboratories, Hercules, CA). Gene expression was subsequently measured for individual genes (β-ARs, ATP6, mDIC, GPD2, PPARα, and PPARγ) to confirm results obtained by DNA array. IQ SYBR Green Supermix (Bio-Rad) was used for the quantitative PCR reactions and HPRT was used as an endogenous control. The relative expression level was calculated by the comparative Ct method. Primer sequences are indicated in Table 1.

Table 1. Primer Sequences for Quantitative Real-Time RT-PCR

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<thead>
<tr>
<th>Primer</th>
<th>Forward primer</th>
<th>Reverse primer</th>
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<td>HPRT</td>
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<td>5’-GCCGCTGAGAAATCTACGTTTC-3’</td>
<td>5’-GAGTACTGTTGAGAGACCC-3’</td>
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<tr>
<td>PPARα2</td>
<td>5’-AAGCAGTAAGATCTCCTCAG-3’</td>
<td>5’-GAGTGTGTCATAGGATCTGAC-3’</td>
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<td>mDIC</td>
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<td>5’-TTTGCGCTTTCTCCAGTATTCA-3’</td>
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<td>5’-GGCAGGTAGGCTCCCATGGC-3’</td>
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<td>β1-AR</td>
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<td>5’-GCTCGACAGCTCTCAGCA-3’</td>
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<tr>
<td>GPD2</td>
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<td>5’-GAGTACTGTTGAGAGACCC-3’</td>
</tr>
<tr>
<td>ATP6</td>
<td>5’-ACTTCAGGACCTCCCTCCCCAAA-3’</td>
<td>5’-TTTGCGCTTTCTCCAGTATTCA-3’</td>
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</table>
Metabolic Cage Studies

Body composition was assessed by nuclear magnetic resonance (Minispec mq10, Bruker, Newark, DE). Indirect calorimetry, food intake, and activity measurements were performed using the CLAMS system (Columbus Instruments, Columbus, OH). The mice were allowed to acclimatize to the metabolic cages for 2 days. Variability of the respiratory exchange ratio (RER) for each condition was estimated by calculating the difference between the average of the six highest values and the average of the six lowest values.

Statistical Methods

Data are generally expressed as means ± SEM. The nonparametric Wilcoxon test was used for comparisons between groups. P < 0.05 was considered as significant.

Results

HFD-Induced Hepatic Steatosis in C57B6 and FVB Mice

Hepatic steatosis is a key event in the pathogenesis of the metabolic syndrome and can exert lipotoxic effects under some conditions. We therefore explored the detailed time course of HFD-induced hepatic steatosis in C57B6 and FVB mice as these strains are commonly used in metabolic studies. Although rodents are widely used clinically, there is no established protocol available for longitudinal measurements of hepatic lipids in them using micro-CT scanners. We therefore established this methodology as described in Materials and Methods and validated all measurements on a CT scanner with conventional biochemical measurements of triglycerides. The CT measurements were in excellent agreement with the biochemical assessment (Figure 1A), providing strong support that this methodology is suitable for longitudinal measurements in vivo.

We found that within as short a time frame as 48 hours, hepatic lipid content was dramatically increased, especially in the C57B6 strain. This finding was confirmed by conventional measurements of liver triglycerides in another set of mice (12 ± 20.0 versus 36 ± 9.5 mg/g in, respectively, normal chow-fed versus 48 hours HFD-fed mice; P < 0.05). The FVB mice were less sensitive to this acute effect of a HFD but developed more severe hepatic steatosis over the long-term exposure as the high fat diet regimen was continued for several weeks and followed with repeated CT measurements (Figure 1B). After the acute phase of the HFD, an “adaptation phase” followed during which the hepatic steatosis improved in the C57B6 mice and remained unchanged in the FVB mice until approximately 2 weeks (Figure 1B). Thereafter, the gain in hepatic lipids was highly correlated with the body weight gain (data not shown).

Because the 48-hour HFD regimen caused a robust effect on hepatic steatosis and may serve as an excellent model system for a lipotoxic environment in the liver, we studied the mice under these conditions more extensively. Blood chemistry analysis showed that C57B6 and FVB mice responded slightly differently to this acute metabolic challenge, although both strains showed signs of compromised insulin sensitivity (Table 2). Hyperinsulinemia was very pronounced in the C57B6 mice, whereas FVB mice displayed a more variable response. Nevertheless, both strains and genders developed either hyperglycemia or hyperinsulinemia. Total cholesterol levels did not change in the C57B6 mice, whereas female but not male FVB mice exhibited increased cholesterol levels. Surprisingly, total triglyceride levels decreased in both FVB and C57B6 mice in both genders (Table 2 and female data not shown). These may reflect either decreased VLDL secretion and/or increased triglyceride clearance because of the hyperinsulinemic conditions established under these circumstances.

Adipo tg Mice Are Protected Against the Deleterious Effects of a HFD

As demonstrated above, acute exposure to the HFD caused a major metabolic challenge in the mice. In particular, C57B6 mice exhibited dramatic increases in liver lipids as well as hyperinsulinemia. Adiponectin has previously been associated in clinical studies with de-
creased levels of hepatic lipids. Therefore, we wanted to investigate whether mice overexpressing higher levels of adiponectin (adipo tg) mice on the C57B6 background would be protected against the deleterious effects of acute exposure to a HFD. Indeed, increases in hepatic lipids after 48 hours of the HFD were significantly reduced in adipo tg mice despite similar overall weight gain (2.1 ± 0.2 versus 2.2 ± 0.3 in, respectively, wild-type [WT] and adipo tg mice) (Figure 2A). The adipo tg mice were also able to maintain their baseline insulin levels, whereas the levels were approximately twofold increased in both female and male wild-type mice (1.8- and 2.3-fold in wild-type males and females, respectively; P < 0.05). Cholesterol levels were reduced both at baseline and after 48 hours of the HFD in the adipo tg mice (data not shown). These observations strongly suggest that adiponectin reduces the lipotoxic effects of a HFD. This reduction was further supported by the lower levels of circulating FFAs in the adipo tg mice relative to controls when challenged with an oral load of olive oil (Figure 2B).

With chronic HFD feeding, the adipo tg mice are partially protected against hepatic steatosis and have consistently lower insulin levels (Figure 2C). Neither body weight nor leptin levels were significantly different between genotypes, whereas adiponectin levels were two to threefold higher in the adipo tg mice (data not shown). Body weight gain and liver lipid gain are positively correlated after 10, 20, and 40 days, but no longer after 80 days with HFD in wild-type mice (Figure 3, A–D). In adipo tg mice, this correlation is significant after 40 days with the HFD, but the slope is not as steep as in the wild-type mice. Thus, the adipo tg mice have to gain more weight to develop the same degree of HFD-induced hepatic steatosis. After 80 days of the HFD, these mice were again fed normal chow. Already within the 10 subsequent days, the normal chow feeding dramatically improved hyperinsulinemia and hepatic steatosis, but weight loss was achieved at a slower pace (Figure 2, C–E). This result indicates that HFD-induced lipotoxicity may depend both on obesity and on a constant supply of exogenous lipids.

Fat distribution, CT analysis, and dissection of individual fat pads demonstrates that the adipo tg mice have significantly less visceral fat than wild-type littermate controls (Figure 2F, Table 3). The subcutaneous fat mass was increased in adipo tg males after 80 days of an HFD, and a similar tendency was also seen in adipo tg females (Figure 2F and data not shown). Moreover, liver size was significantly decreased in the adipo tg mice (Table 3). Thus, the adipo tg mice are less prone to visceral obesity and effectively expand the subcutaneous fat mass, which is likely to be the underlying reason for the potent protective effects against metabolic dysfunction.

**Adiponectin Increases Adrenergic Sensitivity and Causes Remodeling of Adipose Tissue**

In light of these findings, we wanted to further investigate the baseline phenotype of the adipo tg mice on the C57B6 background in the hope of identifying the underlying mechanism by which adiponectin exerts these potent protective effects against lipotoxicity. Analysis of tail vein serum samples obtained during the light phase showed that the glycerol levels are increased in the transgenic mice, whereas FFA levels were comparable in the adipo tg and wild-type littermates. Surprisingly, this difference in glycerol levels was no longer apparent in additional samples obtained under the same conditions (but without the added stress of re-cutting the tail) (Supplemental Figure S1, see http://ajp.amipathol.org). When sampling was performed under anesthetized conditions, we saw no inducible lipolysis as expected in both genotypes. Instead, anesthetized female adipo tg mice displayed decreased FFA levels relative to wild-type mice (Figure 4, A and B). These data strongly suggest that the increased glycerol levels in the adipo tg mice are induced by an enhanced lipolytic response to handling stress. Consistent with our previous reports, the adipo tg mice rapidly became hypoglycemic on fasting and had reduced insulin levels (Figure 5, A and B).

Glycerol levels, obtained in tail vein samples from FVB mice during the light phase, were in general higher than those in the C57B6 mice (data not shown) but were comparable between wild-type and adipo tg mice. FVB adipo tg mice had significantly decreased levels of FFAs relative to those of littermate controls. Consistent with that finding, an opposite phenotype was seen in male FVB adipo–/– mice. These mice displayed similar glycerol but increased FFA levels compared with those of wild-type controls (Figure 5D). These data are consistent with a

<p>| Table 2. Body Weight and Blood Chemistry of Male FVB and C57B6 Mice Fed Normal Chow and after 48 Hours of the HFD |
|---------------------------------|------------------|------------------|</p>
<table>
<thead>
<tr>
<th></th>
<th>FVB</th>
<th>C57B6</th>
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<tbody>
<tr>
<td>Body weight (g)</td>
<td>NC</td>
<td>HFD 48 hours</td>
</tr>
<tr>
<td>Glucose (mg/dl)</td>
<td>262 ± 28</td>
<td>382 ± 31†</td>
</tr>
<tr>
<td>Insulin (ng/ml)</td>
<td>0.48 ± 0.04</td>
<td>0.52 ± 0.1</td>
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<tr>
<td>Triglycerides (mg/dl)</td>
<td>143 ± 10</td>
<td>77 ± 9§</td>
</tr>
<tr>
<td>Cholesterol (mg/dl)</td>
<td>196 ± 11</td>
<td>217 ± 15</td>
</tr>
<tr>
<td></td>
<td>NC</td>
<td>HFD 48 hours</td>
</tr>
<tr>
<td>Glucose (mg/dl)</td>
<td>25.8 ± 0.5†</td>
<td>27.1 ± 0.3††</td>
</tr>
<tr>
<td>Insulin (ng/ml)</td>
<td>260 ± 20</td>
<td>269 ± 21†</td>
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<tr>
<td>Triglycerides (mg/dl)</td>
<td>0.34 ± 0.04</td>
<td>0.87 ± 0.23††</td>
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<tr>
<td>Cholesterol (mg/dl)</td>
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<td>54 ± 2††</td>
</tr>
<tr>
<td></td>
<td>98 ± 12*</td>
<td>112 ± 9*</td>
</tr>
</tbody>
</table>

n = 5 to 7/group. NC, normal chow.  
*P < 0.01 for C57B6 vs. FVB mice.  
†P < 0.05 for C57B6 vs. FVB mice.  
§P < 0.05 for NC versus 48 hours of a HFD.  
¶P < 0.01 for NC versus 48 hours of a HFD.  
*P < 0.01 for C57B6 versus FVB mice.
model in which adiponectin promotes dose-dependent efficient fatty acid re-esterification and/or fatty acid oxidation in light of the fact that the FFA/glycerol ratio was lower in adipo tg and higher in adipo−/− compared with wild-type mice.

Adipo tg and littermate controls on the FVB background had similar glucose and insulin levels during the light phase as well as after a 6-hours fast (Figure 5C). Therefore, we challenged them further with a more extended fast to investigate whether adipo tg on the FVB background eventually also become hypoglycemic. To our big surprise, fasting for 24 hours caused an increase in insulin levels in the adipo tg mice in both genders and strains (FVB females in Figure 5C shown as a representative example). This effect is reminiscent of the effects that can be induced pharmacologically by β3-AR agonists.24,25 Therefore, we were prompted to test whether adipo tg mice have an increased sensitivity to CL 316,243, a highly specific β3-AR agonist.24,25 Indeed, the adipo tg mice were more sensitive to CL 316,243 as judged by the increased glycerol and insulin levels relative to those of controls (Figure 6, A–E). In fact, the adipo tg mice on the FVB background had to be given glucose to survive the severe hypoglycemia that they developed. In contrast, the adipo−/− mice were, as expected, less sensitive than wild-type controls (Figure 6C), consistent with an adiponectin dose-dependent effect on this process.

The increased sensitivity to adrenergic agonist-induced lipolysis in adipo tg mice may not be selective to just β3-AR agonist, because the adipo tg mice were also more sensitive to isoproterenol (data not shown) and norepinephrine-induced lipolysis. The maximum increase in glycerol levels 15 minutes after injection was seen at a dose of 1 mg/kg norepinephrine for the adipo tg mice, whereas 10 mg/kg was required for the littermate controls (Figure 6E). Consistent with the increased sensitivity to adrenergic stimuli, the adipo tg mice displayed increased expression of β1-, β2- and β3-AR mRNAs in the gonadal adipose tissue (1.7-, 1.6-, and 3-fold increase for β1-, β2-, and β3-AR mRNA in adipo tg mice; \( P < 0.05 \)). Similar trends were observed in the inguinal depots.

To further test whether the adrenergic sensitivity was specific to adipose tissue, tissue levels of CREB and phospho-CREB were measured in adipose tissue and liver after a 30-hour fast. The levels of phospho-CREB in liver were similar in wild-type and adipo tg mice. However, the phospho-CREB levels in both gonadal and inguinal...
adipose depots were increased in the adipo tg mice (Figure 7A). These differences in the degree of phosphorylation of CREB were not observed under fed conditions (not shown). This finding suggests the presence of elevated levels of cAMP in adipose tissue of the adipo tg mice in the fasted state.

Chronic treatment with β3-AR agonists has been shown to improve metabolism in rodents at multiple levels. Thus, increased β3-AR adrenergic signaling in adipose tissue may be one of the underlying mechanisms that enable adiponectin to exert its positive effects on metabolic health.

Table 3. Body, Gonadal Adipose Tissue, Inguinal Adipose Tissue, and Liver Weights

<table>
<thead>
<tr>
<th>Weight</th>
<th>Females</th>
<th>Males</th>
</tr>
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<tbody>
<tr>
<td>Body (g)</td>
<td>WT Adipo tg</td>
<td>WT Adipo tg</td>
</tr>
<tr>
<td>Liver (mg/g)</td>
<td>21.5 ± 0.5</td>
<td>26.7 ± 0.7</td>
</tr>
<tr>
<td>GWAT (mg/g)</td>
<td>9.02 ± 0.8</td>
<td>16.1 ± 1.9</td>
</tr>
<tr>
<td>IWAT (mg/g)</td>
<td>9.98 ± 0.6</td>
<td>7.5 ± 0.8</td>
</tr>
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n = 5 to 7/group. GWAT, gonadal white adipose tissue; IWAT, inguinal white adipose tissue.
*P < 0.05 for wild-type versus adipo tg mice.
†P < 0.001 for wild-type versus adipo tg mice.
‡P < 0.01 for wild-type versus adipo tg mice.

Figure 3. Body weight (BW) gain is plotted against liver lipid gain after 10 (A), 20 (B), 40 (C), and 80 (D) days on HFD using both male and female C57B6 adipo tg (white squares) and littermate control mice (black squares). — — —, nonsignificant trend; — — —, significant correlation.

Figure 4. Serum glycerol (A) and free fatty acid (B) levels were measured in awake and chloral hydrate-anesthetized male (M) and female (F) C57B6 adipo tg and littermate control mice. n = 5 to 7/group. *P < 0.05.
Consequently, we examined the adipose tissue of the mitochondrial density and function in adipose tissue. Mitochondrial electron transport activity as judged by exposure to 2,3,5-triphenyltetrazoliumchloride was increased in both inguinal and gonadal fat pads (Figure 7B). Histological examination also revealed increased adipocyte size in the adipose tissue (Figure 7C). Consistent with these morphological changes, the levels of the mitochondrial protein HSP60 (Supplemental Figure S2, see http://ajp.amipathol.org), and the mRNA expression of mitochondrial markers was up-regulated in the adipose tissue of the adipotg mice (Table 4). However, mRNA levels of UCP-1 were similar in the adipotg and littermate controls.

DNA microarray analysis of gonadal and inguinal adipose depots showed that genes were involved in oxidative phosphorylation, accumulation of triglyceride, and fatty acid catabolism and were consistently up-regulated in the adipotg mice. In contrast, a module of genes that is negatively affected comprises the general category of extracellular matrix proteins that are usually associated with a fibrotic phenotype. These genes were reduced in the adipotg mice (Table 4). The most up-regulated gene in both gonadal and inguinal fat pads is the lipid-droplet protein cidea (7.0- and 7.2-fold in gonadal and inguinal fat pads). In general, gene expression in the two fat depots showed similar patterns of regulation with respect to metabolic pathways, but the difference between wild-type and adipotg mice was in many cases more pronounced in the gonadal depot. On the other hand, mRNA expression in the inguinal depots of adipotg mice indicated further enhanced adipogenesis (Figure 7D), which is consistent with the large clusters of smaller adipocytes together with fibroblast-like cells in this depot (Figure 7, E and F). The mRNA expression of \( \beta \)-AR in the liver or downstream targets of adrenergic signaling were not regulated differently in adipotg than in littermate control mice.

Collectively our data indicate that the adipotg adipose tissues display an enhanced lipolytic response on adrenergic stimulation. At the same time, they display an enhanced capacity to store excess lipids with fed conditions. To further examine these aspects, serum samples were obtained from female FVB mice under fed conditions during the dark phase and after a 24-hour fast. Great care was taken to avoid the anticipated stress influence from the sampling procedure by cutting the tail 1 hour before the blood collection. These conditions revealed a clear reduction in fed insulin and FFAs levels, whereas glucose and glycerol levels were unchanged in the adipotg mice (Figure 8, A–D). The 24-hour fast induced an increase in glycerol levels and more severe hypoglycemia in the adipotg mice than in littermate controls. Fasting FFA and insulin levels were similar in wild-type and adipotg mice (Figure 8). Thus, prolonged fasting in the absence of the stressful tail cutting procedure does not lead to the paradoxical increase in insulin levels in the adipotg mice. This observation highlights the importance of appropriate sampling procedures that becomes particularly relevant in the context of models that are differentially susceptible to \( \beta \)-AR stimulation.

Subsequently, we aimed to estimate how quickly the adipotg mice are able to switch from fasted to fed state by giving the mice an oral load of glucose after the 24-hour fast. This induced a similar increase in circulating glucose and insulin levels, but the reduction of the FFA level was enhanced in the adipotg mice (Figure 8). Thus, prolonged fasting in the absence of the stressful tail cutting procedure does not lead to the paradoxical increase in insulin levels in the adipotg mice. This observation highlights the importance of appropriate sampling procedures that becomes particularly relevant in the context of models that are differentially susceptible to \( \beta \)-AR stimulation.

Chronic \( \beta_3 \)-AR agonist improves the capacity for catabolism and re-esterification of fatty acids by increasing the mitochondrial density and function in adipose tissue. Consequently, we examined the adipose tissue of adipotg mice in more detail with respect to mitochondrial content and function. Mitochondrial electron transport activity as judged by exposure to 2,3,5-triphenyltetrazoliumchloride was increased in both inguinal and gonadal fat pads (Figure 7B). Histological examination also revealed increased mitochondrial density as well as smaller average adipocyte size in the adipotg mice (Figure 7C). Consistent with these morphological changes, the levels of the mitochondrial protein HSP60 (Supplemental Figure S2, see http://ajp.amipathol.org), and the mRNA expression of mitochondrial markers was up-regulated in the adipose tissue of the adipotg mice (Table 4). However,
0.4 g for wild-type versus adipo tg mice; $P = 0.87$) and body composition (9.1 ± 1.6 versus 8.1 ± 0.8% fat, wild-type versus adipo tg mice; $P = 0.59$) to avoid possible interference from these parameters. Neither oxygen consumption nor food intake was altered in the adipo tg mice, although the level of activity was reduced under chow-fed conditions and in the fasted state (Supplemental Figure S3, see http://ajp.amjpathol.org). However, the

Figure 6. Female FVB adipo tg, adipo$^{-/-}$, and control (WT) mice were injected with $\beta_3$AR agonist (1 mg/kg CL 316,243 i.p.), and serum levels of glycerol (A), FFA (B), insulin (C), and glucose (D) were measured in tail vein samples after 0, 5, 10, and 60 minutes. $n = 4$ to 6/group. *$P < 0.05$; **$P < 0.01$, ***$P < 0.001$ for WT versus adipo tg mice. $^5P < 0.05$ for WT versus adipo$^{-/-}$ mice. E: Increase in serum glycerol levels 15 minutes after norepinephrine (NE) i.p. injection at the indicated dose. $n = 2$ to 5/dose and group. *$P < 0.05$ for WT versus adipo tg.

Figure 7. A: Western blot analysis for total and phosphorylated CREB protein in liver, inguinal (IWAT) and gonadal adipose tissue (GWAT). B: Whole fat pads stained with 1% 2,3,5-triphenyltetrazoliumchloride to assess electron transport activity. C: Fat pad minces stained with MitoTracker Orange. D: Regulation of genes involved in adipogenesis in inguinal and gonadal adipose tissue of adipo tg mice. Gene expression data were obtained with DNA array analysis and are expressed as a percentage of wild-type littermate controls (ie, 100% indicates no change in expression compared with controls). *$P < 0.05$, **$P < 0.01$, ***$P < 0.001$ for WT versus adipo tg mice. E and F: Representative examples of H&E-stained IWAT. All data presented in this figure were obtained in male C57B6 adipo tg with littermate wild-type control mice. Pref1, preadipocyte factor 1; Srebp1, sterol regulatory element binding transcription factor 1.
analysis of RERs revealed significant changes between the genotypes. *Adipo tg* mice had a larger variability in RER under both chow-fed and HFD-fed conditions, reached a higher average maximal RER with refeeding, and had a lower average RER when fed the HFD compared with controls (Figure 10, A–C). These data indicate a higher metabolic flexibility in the *adipo tg* mice as judged by an increased lipogenic response on refeeding as well as enhanced fatty acid oxidation with the HFD.

**Discussion**

We have established a novel experimental paradigm for the *in vivo* analysis of lipotoxicity using a short-term HFD feeding protocol and CT-based measurements of hepatic lipids in rodents. The HFD rapidly induced hepatic steatosis and insulin resistance in mice of both C57B6 and FVB backgrounds. A similar phenomenon has also been reported in human subjects. The increase in hepatic lipids slows down during the adaptive phase to exposure to the HFD, but after this initial counterregulatory response, the degree of hepatic lipid accumulation resumes during chronic exposure to the HFD.

**Table 4.** Significantly Regulated Genes (*P < 0.05*) in Both Inguinal and Gonadal Adipose Tissue of C57B6 *Adipo tg* versus Littermate Control Male Mice

<table>
<thead>
<tr>
<th>Function</th>
<th>Examples of regulated genes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mitochondrial biogenesis +</td>
<td>PGC-1α, PGC-1β, PPARα</td>
</tr>
<tr>
<td>Electron transport chain +</td>
<td>Cox8b, GPD2, CYC1, NDUFB5, ATP5G2</td>
</tr>
<tr>
<td>Fatty acid catabolism +</td>
<td>CPT2, Crat, Acadl</td>
</tr>
<tr>
<td>Accumulation of triglycerides +</td>
<td>LPL, DGAT, GLUT4</td>
</tr>
<tr>
<td>Fibrosis –</td>
<td>LOX, Col6A3, SPARC, FN1</td>
</tr>
</tbody>
</table>

**Figure 8.** Serum insulin (A), glucose (B), glycerol (C), and FFA (D) levels in the fed state at the dark phase and after a 24-hour fast in female *adipo tg* FVB and littermate *wild-type* control mice. *n = 4 + 5. *P < 0.05; **P < 0.01 for WT versus *adipo tg*.

**Figure 9.** Change in serum glucose (A), insulin (B), and FFA (C) after an oral load of glucose (2.5 g/kg) to 24-hour fasted mice. *n = 4 + 5. *P < 0.05 for WT versus *adipo tg*.

In line with our previous studies, we found that overexpression of adiponectin protects against both the acute and the chronic effects of HFD-induced lipotoxic effects of lipid accumulation. We noticed that a relatively short fast causes hypoglycemia and hypoinsulinemia in *adipo tg* mice. In contrast, when animals were fasted for 24 hours, we initially found that insulin levels were paradoxically increased compared with baseline levels in the *adipo tg* mice. However, this increase was not present when care was taken to avoid the stress influence from the tail cut procedure. However, this increase was not present when care was taken to avoid the stress influence from the tail cut procedure. These observations were highly reproducible and highlight the importance of well controlled sampling procedures. We hypothesized that this unexpected increase in insulin levels may be due to increased adrenergic sensitivity in adipose tissue of the *adipo tg* mice. Several of our observations support this model. First, fasting-induced hypoglycemia and hy-
poininsulinemia increase both sympathetic outflow and signaling, and second, adrenergic signaling (via β2-AR) in the adipose tissue causes release of insulin. Third, injection of a β2-AR agonist causes higher insulin release in the adipose tissue than lower insulin release in the adipose tissue of wild-type controls.

The increased sensitivity to adrenergic agonist-induced lipolysis is further enhanced by hypoinsulinemia in the adipose tissue as chronic thiazolidinedione or β-3AR agonist treatment. Although, mitochondria are classically associated with oxidative phosphorylation, they are also critically involved in providing substrates for gluconeogenesis and fatty acid synthesis. Thus, mitochondrial function in adipose tissue is essential for the reduction of local and systemic lipotoxicity, thereby maintaining metabolic health and flexibility. Gene expression analyses and measurements of mitochondrial density and electron transport function show that adiponectin induces a similar remodeling of adipose tissue as chronic β2-AR agonist exposure. In addition, the reduced FFA/glycerol ratio in young unchallenged adipose tissue as well as the improved FFA clearance in the adipose tissue of young unchallenged adipose tissue support a direct role for adiponectin in reduction of circulating fatty acids through more efficient re-esterification steps and fatty acid oxidation.

Figure 10. RER analysis of data obtained from female adipose type C57B6 and littermate wild-type controls fed normal chow, during a fast, on the second day after refeeding and on the second day after the switch form normal chow to the HFD. The average (mean RER, A), the average of the six highest (max RER, B), and the average of the six lowest (min RER) RER values were calculated for each mouse and condition using data collected for 24 hours (12 hours light and 12 hours dark phase). C: Difference (ΔRER) between the averages of the six highest and the six lowest RER values. *P < 0.05 for WT versus adipose type mice.
The transient increase in fatty acids may be central in β₃-AR agonist-induced remodeling of the adipose tissue as this can activate PPARγ and PPARα, key regulators of adipogenesis and fatty acid catabolism. Once remodeling has occurred, adipose tissue is more efficient in clearing all sources of fatty acids including dietary lipids. The beneficial effect of adrenergic signaling may also critically depend on cAMP-mediated signaling. Notably, proliferator-activated receptor-γ coactivator-1β, the master regulator of mitochondrial biogenesis, is regulated by cAMP. Moreover, adipocyte differentiation from preadipocytes/fibroblasts is also driven by cAMP. The over-expression of adiponectin in our adipotg mice is driven by the aP2 promoter, which is induced at early stages of adipogenesis. Thus, increased expression of adiponectin at this stage may enhance adipogenesis through an increase in cAMP levels. The adiponectin-induced phenotype of specific fat pads may therefore be a consequence of the degree of sympathetic innervation as well as the local availability of adipogenic precursor cells. The gonadal fat pad remains very small in the adipotg mice, indicating that the rate of adipogenesis is lower than the rate of fatty acid catabolism in this location. In contrast, overexpression of adiponectin in the inguinal fat pad seems to have a profound stimulatory effect on adipogenesis as judged from histological examination, gene expression profiling, and the larger size of this fat pad. This could be a reflection of enhanced adrenergic signaling and cAMP levels in preadipocytes, which overrules the effects of adrenergic signaling in mature adipocytes. Of particular interest is the observation that pref-1 (a preadipocyte marker) is strongly up-regulated in the inguinal but not in the gonadal depots of the adipotg mice. The size and the appearance of the inguinal fat pad together with the increased expression of pro-adipogenic genes indicate increased inguinal adipogenesis. Therefore, the increased presence of pref-1-positive cells is a reflection of a larger number of preadipocytes in this depot. Of note is the fact that in this context patients with lipodystrophic HIV exhibit a nearly complete absence of pref-1 expression in adipose tissue. Thus, a deficiency of preadipocytes is a reflection of limited expansion potential.

The relative expression pattern of genes involved in metabolism is highly dependent on whether the tissues are obtained during an anabolic or catabolic state. Our in vivo studies are mostly performed during the daytime, ie, when the mice are usually inactive and in a catabolic state. This is a state in which the effects of adrenergic signaling pathways are of particular importance. However, analysis of RER throughout the day and night and in response to refeeding as well as the measurement of serum parameters indicate that adiponectin also promotes lipogenesis in the presence of insulin under fed conditions. We therefore propose that adiponectin plays an important role in promoting metabolic flexibility by making adipose tissue more sensitive to changes in metabolic state. Thus, lipid catabolism is enhanced in the fasted state, whereas lipid synthesis and storage are enhanced in the fed state. A similar phenomenon is observed during caloric restriction and intermittent fasting, both of which are associated with improved systemic insulin sensitivity. For instance, Varady et al show that intermittent fasting induces increased lipolysis, increased lipogenesis, decreased adipocyte size, but no change in total fat mass. Furthermore, the concept of metabolic flexibility (or loss thereof) in muscle in the context of type 2 diabetes has found widespread acceptance through the elegant work of Kelley.

How well can these observations from rodents be translated to human physiology? The cause-effect relationships between altered levels of adiponectin (brought about either through genetic or pharmacological interventions) established in murine studies have held up remarkably well at the correlational level in humans. Adrenergic receptors are, however, not expressed to the same high levels in human adipose tissue compared with mice. Nevertheless, adiponectin may increase sensitivity to adrenergic stimulation originating from other β-ARs as well. In fact, adrenergic sensitivity in adipose tissue is reduced in obese as well as in nonobese insulin-resistant subjects, both states that are associated with reduced adiponectin levels. It may be counterintuitive that increased sensitivity to adrenergic stimuli improves metabolism, as increased lipolysis in insulin-resistant states is viewed as a reason for disproportionate elevations of FFAs and a major cause of lipotoxicity. However, under metabolically healthy conditions, activation of the sympathetic nervous system is only induced during fasting or during “fight and flight responses,” ie, when the fatty acids are needed. Thus, increased sensitivity to adrenergic stimuli in adipose tissue will not cause metabolic dysregulation per se but rather will be beneficial for the maintenance of metabolic flexibility in the long run.

Combining our results, we propose a model in which adiponectin exerts a crucial role in the maintenance of adaptation to rapidly changing metabolic conditions as we go from the fed postabsorptive state to fasting conditions (Figure 11, A and B). Although the effects of adiponectin on target tissues are well established, our
observations here also point to an important role of adiponectin in adipose tissue proper.

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References
15. Ek I, Aper M, Bergqvist A, Carlstrom K, Wahrenberg H: Impaired adipocyte lipolysis in nonobese women with the polycystic ovary syndrome: a possible link to insulin resistance?. J Clin Endocrinol Metab 1997, 82:1147–1153