Altered Interleukin-10 Signaling in Skeletal Muscle Regulates Obesity-Mediated Inflammation and Insulin Resistance

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Altered Interleukin-10 Signaling in Skeletal Muscle Regulates Obesity-Mediated Inflammation and Insulin Resistance

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Skeletal muscle insulin resistance is a major characteristic of obesity and type 2 diabetes. Although obesity-mediated inflammation is causally associated with insulin resistance, the underlying mechanism is unclear. Here, we examined the effects of chronic obesity in mice with muscle-specific overexpression of interleukin-10 (M110). After 16 weeks of a high-fat diet (HFD), M110 mice became markedly obese but showed improved insulin action compared to that of wild-type mice, which was largely due to increased glucose metabolism and reduced inflammation in skeletal muscle. Since leptin regulates inflammation, the beneficial effects of interleukin-10 (IL-10) were further examined in leptin-deficient ob/ob mice. Muscle-specific overexpression of IL-10 in ob/ob mice (MCK-IL10ob/ob) did not affect spontaneous obesity, but MCK-IL10ob/ob mice showed increased glucose turnover compared to that in ob/ob mice. Last, mice with muscle-specific abduction of IL-10 receptor (M-IL10R–/–) were generated to determine whether IL-10 signaling in skeletal muscle is involved in IL-10 effects on glucose metabolism. After an HFD, M-IL10R–/– mice developed insulin resistance with reduced glucose metabolism compared to that in wild-type mice. Overall, these results demonstrate IL-10 effects to attenuate obesity-mediated inflammation and improve insulin sensitivity in skeletal muscle, and our findings implicate a potential therapeutic role of anti-inflammatory cytokines in treating insulin resistance and type 2 diabetes.

Obesity has emerged as a global issue in recent decades and is associated with numerous human diseases, including insulin resistance, type 2 diabetes, and cardiovascular diseases (1,2). The underlying mechanism by which obesity induces numerous health problems remains poorly understood. In that regard, increasing evidence suggests an important role of a dysregulated immune system in obesity-mediated insulin resistance (3,4). Obesity is characterized by altered levels of circulating cytokines, and adipose tissue macrophage accumulation and inflammation have been causally associated with insulin resistance (5,6). However, recent studies have challenged this earlier notion on the causal role of adipose tissue macrophages and inflammation in the development of insulin resistance (7,8).

While adipose tissue is widely viewed as the epicenter of obesity-mediated inflammation, it is not the only organ shown to develop macrophage infiltration and inflammation in obesity. In fact, recent studies indicate that obesity-mediated inflammation and macrophage accumulation develop in multiple organs, including skeletal muscle, liver, pancreas, heart, and brain (9–13). In that regard, our recent study found that exercise-mediated weight loss improved insulin action without affecting adipose tissue inflammation in mice with diet-induced obesity (7). Additionally, improved insulin action following weight loss was associated with reduced local inflammation in skeletal muscle, suggesting an important role of muscle inflammation in obesity-mediated insulin resistance (7). These findings clearly contest the adipocentric view of insulin resistance.

Interleukin-10 (IL-10) is a Th2-type cytokine that inhibits the synthesis and activity of proinflammatory cytokines and counteracts Toll-like receptor-mediated inflammation (14–16). We have previously shown that transgenic overexpression of IL-10 selectively in skeletal muscle improved glucose metabolism in mice after 3 weeks of a high-fat diet (HFD) (9). While our previous data suggest a potential therapeutic role of IL-10 in type 2 diabetes, our interpretation is limited due to the short-term feeding of an HFD not resulting in mice having developed type 2 diabetes phenotypes (i.e., hyperglycemia). Therefore, the current study was designed to specifically examine the role of IL-10 in markedly obese and diabetic mice after 16 weeks of an HFD (chronic HFD), a better representation of obese type 2 diabetic human subjects. Additionally, leptin is an important adipocyte-derived hormone that is elevated in obesity and regulates numerous physiological functions, including energy balance and inflammation (17). Thus, the present study also examined the effects of muscle-specific transgenic expression of IL-10 on glucose metabolism in leptin-deficient ob/ob mice. Last, we examined whether IL-10 signaling in...
skeletal muscle is directly responsible for IL-10 effects on muscle glucose metabolism using a newly generated mouse model lacking the IL-10 receptor 1 type chain selectively in skeletal muscle. Our findings indicate that selective targeting of IL-10 signaling in skeletal muscle improves glucose metabolism in obese and diabetic mice following a chronic HFD or with a deficiency in leptin and further demonstrate that these effects are mediated by direct activation of IL-10 signaling in skeletal muscle.

MATERIALS AND METHODS

Chronic HFD in M\textsuperscript{IL10}\textsubscript{obob} mice. Male transgenic mice with muscle-specific overexpression of IL-10 (M\textsuperscript{IL10}\textsubscript{obob}) and wild-type (WT) littermates were fed an HFD (55% fat by calories [Teklad TD93075; Harlan, Madison, WI]) or standard chow diet (Prolab Isopro RMH 3000 SP75; LabDiet, St. Louis, MO) ad libitum for 16 weeks (n = 6/group). During chronic high-fat feeding, we performed a weekly measurement of body composition to determine the changes in whole-body fat and lean masses.

Generation of MCK-IL10\textsubscript{obob} mice. We generated leptin-deficient mice with muscle-specific overexpression of IL-10 (MCK-IL10\textsuperscript{obob}) by cross-breeding M\textsuperscript{IL10}\textsubscript{obob} mice with ob/ob heterozygous mice (purchased from the Jackson Laboratory, Bar Harbor, ME). The F1 female MCK-IL10\textsuperscript{obob} mice were then intercrossed with male ob/+ mice to generate MCK-IL10\textsuperscript{obob} mice. The metabolic studies were conducted in MCK-IL10\textsuperscript{obob} mice that had been backcrossed for more than five generations (ob/ob, n = 7; MCK-IL10\textsuperscript{obob}, n = 12).

Generation of M-IL10R\textsuperscript{-/-} mice. Mice lacking IL-10 signaling in skeletal muscle (M-IL10R\textsuperscript{-/-}) were generated by cross-breeding MCK-Cre-expressing mice (kindly donated by Roger J. Davis) and floxed IL10R\textsuperscript{+/–} mice (kindly donated by Werner Muller). The metabolic studies were conducted in M-IL10R\textsuperscript{-/-} mice that had been backcrossed for several generations. M-IL10R\textsuperscript{-/-} mice and MCK-Cre mice (referred to as WT) serving as controls were fed with an HFD or standard chow diet for 6 weeks (n = 6 to 11/group). All mice were housed under a controlled temperature (23°C) and light/dark cycle with free access to food and water. The animal studies were approved by the Institutional Animal Care and Use Committee of the University of Massachusetts Medical School.

Body composition and energy balance. Whole-body fat and lean masses were noninvasively measured using proton magnetic resonance spectroscopy (\textsuperscript{1}H-MRS) (Echo Medical Systems, Houston, TX). Indirect calorimetry and energy balance parameters, including food/water intake, energy expenditure, respiratory exchange ratio, and physical activity were noninvasively assessed for 3 days using metabolic cages (TSE-Systems, Bad Homburg, Germany). We used the TSE-Systems LabMaster platform with easy-to-use calorimetry featuring fully automated monitoring for food and water and activity in the x, y, and z planes. LabMaster cages that are most similar to facility home cages were used, thereby allowing the use of bedding in the cage and minimizing any animal anxiety during the experimental period. The system provides intuitive software with flexibility for experimental setup and data utilization.

Hyperinsulinemic-euglycemic clamp experiments. Following a standard chow diet or an HFD, survival surgery was performed at 5 to 6 days before clamp experiments to establish an indwelling catheter in the jugular vein. On the day of the clamp experiment, mice were fasted overnight (~17 h), and a 2-h hyperinsulinemic-euglycemic clamp was conducted in conscious mice with a primed and continuous infusion of human insulin (priming at 150 mU/kg body weight, followed by 2.5 mU/kg/min [Humulin; Eli Lilly, IN]) (18). To maintain euglycemia, 20% glucose was infused at variable rates during the clamps. Whole-body glucose turnover was assessed with a continuous infusion of [\textsuperscript{3}H]glucose (PerkinElmer, Waltham, MA), and 2-deoxy-\textsuperscript{[1-14]C}glucose (2-\textsuperscript{[1-14]C}DG) (PerkinElmer, Waltham, MA) was administered as a bolus (10 μCi) at 75 min after the start of the clamps to measure insulin-stimulated glucose uptake in individual organs. At the end of the clamps, mice were anesthetized, and tissues were taken for biochemical analysis.

Biochemical analysis and calculation. Glucose concentrations during the clamps were analyzed using 10 μl of plasma by a glucose oxidase method on an Analox GM9 Analyser (Analox Instruments, Ltd., London, United Kingdom). Plasma concentrations of [\textsuperscript{3}H]glucose, 2-\textsuperscript{[14]C}DG, and \textsuperscript{3}H\textsubscript{2}O were determined following deproteinization of plasma samples as previously described (18). For the determination of tissue 2-\textsuperscript{[14]C}DG-6-phosphate (2-\textsuperscript{[14]C}DG-6-P) content, tissue samples were homogenized, and the supernatants were subjected to an ion exchange column to separate 2-\textsuperscript{[14]C}DG-6-P from 2-\textsuperscript{[14]C}DG. Plasma insulin levels were measured using an enzyme-linked immunosorbent assay (ELISA) kit (Alpco Diagnostics, Salem, NH). Intramuscular triglyceride concentrations were determined by homogenizing muscle samples (quadriiceps) in chloroform-methanol and using a triglyceride assay kit (Sigma, St. Louis, MO). Plasma nonesterified fatty acid (NEFA) levels were measured by an NEFA kit (Zenbio, Durham, NC), according to the manufacturer’s protocol.

Rates of basal hepatic glucose production (HGP) and insulin-stimulated whole-body glucose turnover were determined as previously described (18). The insulin-stimulated rate of HGP was determined by subtracting the glucose infusion rate from whole-body glucose turnover. Whole-body glycolysis and glycogen synthesis from glucose were calculated as previously described (18). Insulin-stimulated glucose uptake in individual tissues was assessed by determining the tissue (e.g., skeletal muscle) content of 2-\textsuperscript{[14]C}DG-6-phosphate and the plasma 2-\textsuperscript{[14]C}DG profile.

Molecular analysis for insulin signaling and inflammation. Skeletal muscle (quadriiceps) and liver samples were collected at the end of the clamp experiments to assess insulin signaling by immunoblotting with rabbit monoclonal antibodies against Akt and phospho-Akt-Ser\textsuperscript{473} (p-Akt-Ser\textsuperscript{473}) (Cell Signaling, Danvers, MA). Muscle samples were homogenized, and plasma and homogenized muscle samples were used to measure the levels of interleukin-6 (IL-6), IL-10, gamma interferon (IFN-γ), and IL-1α using an enzyme-linked immunosorbent assay (ELISA) with a Luminex 200 Multiplex system (Millipore, Darmstadt, Germany). Plasma IL-10 levels were measured using an ELISA kit (Abcam, Cambridge, United Kingdom).

For quantitative real-time PCR (qRT-PCR), RNA isolation was performed with homogenized muscle (gastrocnemius and quadriiceps), liver, and white adipose tissue samples using TRIzol (Life Sciences, Carlsbad, CA) according to the manufacturer’s protocols. cDNA was synthesized from 2 μg of total RNA by use of an Omniscript cDNA synthesis kit (Qiagen, Venlo, Netherlands), cDNA and Sybr Green supermix (Bio-Rad, Hercules, CA) were run in a Bio-Rad-CFX96 real-time system with the primers listed in Table S1 in the supplemental material. Relative gene expression was calculated in comparison to the level of the housekeeping gene.

Histologic analysis. Skeletal muscle samples (gastrocnemius) were collected, fixed in 10% neutral formalin for 48 h, and embedded in paraffin blocks. Sections (5 μm) were taken and stained with hematoxylin-eosin. Images were taken under a magnification of ×20.

Statistical analysis. Data are expressed as means ± standard errors (SE). The significance of difference in mean values was determined using two-way analysis of variance (ANOVA) with Newman-Keuls and Games-Howell tests for post hoc analysis and Student’s t test where applicable. The statistical significance was set at a P value of <0.05.

RESULTS

Chronic feeding of an HFD in M\textsuperscript{IL10} mice. Starting at 6 to 7 weeks of age, male M\textsuperscript{IL10} and WT mice were fed with a standard chow diet or an HFD ad libitum for 16 weeks. At the end of the feeding period, all mice were age matched for metabolic studies. M\textsuperscript{IL10} and WT mice showed similar body weights on the standard chow diet, and after 16 weeks of an HFD, both groups of mice became obese with comparable increases in body weights (Fig. 1A). Consistent with this, whole-body fat mass, measured using \textsuperscript{1}H-MRS, was not
different between standard-chow-fed M<sup>IL10</sup> and WT mice, and fat mass increased by 3- to 4-fold after an HFD in both groups of mice (Fig. 1B). Although whole-body lean mass was statistically different between the groups on the standard chow diet (25.1 ± 0.3 g in WT mice versus 23.5 ± 0.6 g in M<sup>IL10</sup> mice; P = 0.03), this difference of 1.5 g of lean mass is within the range of variability for the lean mass of C57BL/6 mice. Whole-body lean mass measurements after an HFD did not differ between groups (Fig. 1C). Metabolic cage analysis showed no significant difference in values for daily food intake, VO2 consumption, and physical activity between M<sup>IL10</sup> and WT mice after 16 weeks of an HFD (see Fig. S1 in the supplemental material).

**M<sup>IL10</sup> mice are protected from HFD-induced insulin resistance.** Basal glucose levels increased after 16 weeks of an HFD in both groups of mice (i.e., hyperglycemia), and plasma insulin levels were also elevated by more than 6-fold in both groups of mice after an HFD (Fig. 1D and E). To determine the effects on whole-body glucose metabolism, a 2-h hyperinsulinemic-euglycemic clamp was conducted in awake mice. During the clamp, plasma glucose levels were maintained at euglycemia (~7 mM), and plasma insulin levels were raised to ~130 pM and ~230 pM in both groups of standard-chow- and HFD-fed mice, respectively (Fig. 2A and B).

After 16 weeks of an HFD, the WT mice developed insulin resistance, as shown by a ~60% decrease in glucose infusion rates during the clamp compared to the level in standard-chow-fed WT mice (Fig. 2C). Although glucose infusion rates also decreased in the M<sup>IL10</sup> mice after an HFD, the glucose infusion rates in the HFD-fed M<sup>IL10</sup> mice were significantly higher than in HFD-fed WT mice (Fig. 2C). Radioactive isotope labeling data during the clamp showed a markedly increased whole-body glucose turnover in the HFD-fed M<sup>IL10</sup> mice compared to that of the HFD-fed WT mice (Fig. 2D), indicating that M<sup>IL10</sup> mice were more insulin sensitive than WT mice after 16 weeks of an HFD.

Increased insulin sensitivity in HFD-fed M<sup>IL10</sup> mice was largely due to a 30% increase in glucose uptake in skeletal muscle (quadriceps) (Fig. 2E). Insulin-stimulated glucose uptake in gastrocnemius muscle also increased in HFD-fed M<sup>IL10</sup> mice compared to that in HFD-fed WT mice, but this difference did not reach statistical significance (Fig. 2F). To further examine muscle insulin action, we performed Western blotting using skeletal muscle to assess insulin signaling and found that Ser-473 phosphorylation of Akt did not differ between WT and M<sup>IL10</sup> mice on a standard chow diet, consistent with comparable muscle glucose uptake in these mice (see Fig. S2A in the supplemental material). After 16 weeks of an HFD, muscle Akt phosphorylation was decreased by ~70% in WT mice (P = 0.1), and muscle Akt phosphorylation tended to be higher in HFD-fed M<sup>IL10</sup> mice than in HFD-fed WT mice (Fig. 3A). Hematoxylin-eosin-stained sections of skeletal muscle from WT and M<sup>IL10</sup> mice showed no obvious anomaly in the overall structure (Fig. 3B). Additionally, intramuscular triglyc-
eride levels tended to be elevated in standard-chow-fed M<sub>IL10</sub> mice compared to the levels in standard-chow-fed WT mice (Fig. 3C). After the HFD, intramuscular triglyceride levels were comparable between M<sub>IL10</sub> and WT mice. The IL-10 mRNA levels in skeletal muscle were approximately 2-fold higher in HFD-fed M<sub>IL10</sub> mice than in HFD-fed WT mice (Fig. 3D). Obesity-mediated inflammation in skeletal muscle is attenuated in M<sub>IL10</sub> mice. We have previously shown that inflammation develops in skeletal muscle after 3 weeks of an HFD and that this local inflammation may be causally associated with insulin resistance (9). To that end, we assessed the inflammation profile in skeletal muscle by performing quantitative RT-PCR using samples obtained from HFD-fed M<sub>IL10</sub> and WT mice. Skeletal muscle mRNA levels of F4/80 and CD68, as markers of macrophage infiltration, were decreased significantly in HFD-fed M<sub>IL10</sub> mice compared to levels in HFD-fed WT mice (Fig. 4A and B). Muscle monocyte chemotactic protein 1 (MCP-1) mRNA levels also decreased in HFD-fed M<sub>IL10</sub> mice (Fig. 4C). Using Luminex analysis, we found that local levels of IFN-γ, IL-1α, and IL-6 were increased by 2- to 4-fold in muscle samples from WT mice after 16 weeks of an HFD (Fig. 4D to F). In contrast, muscle samples from HFD-fed M<sub>IL10</sub> mice showed completely normal levels of IFN-γ, IL-1α, and IL-6, indicating that muscle IL-10 overexpression protected against diet-induced inflammation in skeletal muscle (Fig. 4D to F). Plasma levels of IL-10 and IL-6 did not significantly differ between WT and M<sub>IL10</sub> mice on standard chow or an HFD (see Fig. S2B and C in the supplemental material). Plasma IL-1α levels tended to increase by more than 2-fold in WT mice after 16 weeks of an HFD, but levels in M<sub>IL10</sub> mice were not affected by an HFD (see Fig. S2D).

Obesity-mediated inflammation and insulin resistance in adipose tissue and liver. Insulin-stimulated glucose uptake in white and brown adipose tissues did not differ between standard-chow-fed WT and M<sub>IL10</sub> mice (Fig. 5A and B). After 16 weeks of an HFD, white and brown adipose tissue developed insulin resistance in both groups of mice (Fig. 5A and B). mRNA levels of the macrophage markers F4/80 and CD68 were also increased in white adipose tissue by a chronic HFD but were not different between HFD-fed WT and M<sub>IL10</sub> mice (Fig. 5C and D). Basal HGP levels did not differ between WT and M<sub>IL10</sub> mice on a standard chow diet or an HFD (Fig. 6A). During the clamp, insulin decreased HGP in both standard-chow-fed WT and M<sub>IL10</sub> mice (Fig. 6B). HFD-fed WT mice developed insulin resistance in liver, as indicated by increased clamp HGP, but HFD-fed M<sub>IL10</sub> mice showed lower clamp HGP than HFD-fed WT mice (Fig. 6B). Western blot analysis showed that neither phospho-Akt nor total Akt protein levels were altered between WT and M<sub>IL10</sub> mice on standard chow or an HFD (Fig. 6C). However, liver mRNA levels of glucose-6-phosphatase (G6Pase) and phosphoenolpyruvate carboxykinase (PEPCK) tended to be lower in HFD-fed M<sub>IL10</sub> mice than in HFD-fed WT mice (Fig. 6D and E). mRNA levels in liver of the macrophage markers CD68 and F4/80 were not significantly altered in HFD-fed M<sub>IL10</sub> mice, indicating that local inflammation was selectively suppressed in skeletal muscle (see Fig. S2E and F in the supplemental material).

IL-10-expressing leptin-deficient mice are protected from insulin resistance. Recent studies have shown deleterious effects of an HFD on the intestinal epithelium and the role of gut microbes on obesity-mediated inflammation (19). Thus, we have generated muscle-selective IL-10-expressing, spontane-
ously obese mice by cross-breeding $\text{M}^{\text{IL10}}$ mice with heterozygous leptin-deficient $\text{ob}^{+}$ mice. The offspring from this cross were further mated to obtain $\text{MCK-IL10}^{\text{ob}^{+}}$ mice. As expected, $\text{ob}^{+}$ mice became obese spontaneously on a standard chow diet, and by 16 weeks of age, whole-body fat mass accounted for more than 50% of body weight in $\text{ob}^{+}$ mice (Fig. 7A and B). $\text{MCK-IL10}^{\text{ob}^{+}}$ mice also became spontaneously obese on a standard chow diet and reached a degree of obesity comparable to that of $\text{ob}^{+}$ mice at 16 weeks of age (Fig. 7A and B). Basal plasma glucose and fatty acid levels were similar in both groups of mice (Fig. 7C; see also Fig. S3A in the supplemental material). A 3-day metabolic cage analysis showed that daily food intake and physical activity did not differ between groups (Fig. 7D and E). In contrast, $\text{MCK-IL10}^{\text{ob}^{+}}$ mice showed a modest but significant increase in energy expenditure that was largely due to elevated VO$_2$ consumption selectively in the night cycle compared to levels in $\text{ob}^{+}$ mice at 16 weeks of age (Fig. 7F and G).

The hyperinsulinemic-euglycemic clamp study showed a significantly increased whole-body glucose turnover in MCK-IL10$^{\text{ob}^{+}}$ mice compared to that in $\text{ob}^{+}$ mice, suggesting increased insulin sensitivity in these mice (Fig. 8A). Consistent with this notion, insulin-stimulated glucose uptake in white adipose tissue and hepatic insulin action did not differ between groups (Fig. 8C and D). Glucose uptake in skeletal muscle tended to be lower in MCK-IL10$^{\text{ob}^{+}}$ mice than in WT mice after an HFD (see Fig. S3D in the supplemental material).

FIG 3 Insulin signaling, muscle morphology, and triglyceride and IL-10 levels in skeletal muscle. (A) Insulin-stimulated Akt phosphorylation in quadriceps muscles. (B) Hematoxylin-eosin staining of gastrocnemius muscles of $\text{M}^{\text{IL10}}$ and WT mice fed standard chow or an HFD ($n = 3$ to 5/group). (C) Intramuscular triglyceride content in quadriceps muscles of $\text{M}^{\text{IL10}}$ and WT mice fed standard chow or an HFD ($n = 3$ to 5/group). (D) IL-10 mRNA levels in gastrocnemius muscles of HFD-fed $\text{M}^{\text{IL10}}$ and WT mice as measured by qRT-PCR ($n = 4$ or 5/group). Values are means ± SE for each group. *, $P < 0.05$. GAPDH, glyceraldehyde-3-phosphate dehydrogenase.
Furthermore, local IL-10 levels in skeletal muscle tended to decrease in M-IL10R−/− mice as a possible feedback loop of IL-10 signaling (see Fig. S3E). Muscle mRNA levels of F4/80 (macrophage marker) and tumor necrosis factor alpha (TNF-α) were markedly increased in WT mice after an HFD (Fig. 9F and G). Importantly, HFD-fed M-IL10R−/− mice showed a significant further increase in muscle mRNA levels of F4/80, TNF-α, IL-6, and IL-1β compared to levels in HFD-fed WT mice, indicating that muscle deletion of IL-10 receptor exacerbated HFD-induced local inflammation in skeletal muscle (Fig. 9F to I).

![Figure 4](http://mcb.asm.org/)

**FIG 4** Local inflammation in skeletal muscles of MIL10 and WT mice on standard chow or after 16 weeks of an HFD. (A to C) F4/80, CD68, and MCP-1 mRNA levels in quadriceps muscles of HFD-fed MIL10 and WT mice as measured by qRT-PCR (n = 3/group). (D to F) IFN-γ, IL-6, and IL-1α levels in gastrocnemius muscles of HFD- and standard-chow-fed MIL10 and WT mice as measured by multiplex ELISA (n = 4 to 6/group). Values are means ± SE for each group. *, P < 0.05. AU, arbitrary units.

![Figure 5](http://mcb.asm.org/)

**FIG 5** Insulin-stimulated glucose uptake and inflammation in adipose tissues. (A and B) Insulin-stimulated glucose uptake in white (epidydimal) and brown (intrascapular) adipose tissue in MIL10 and WT mice fed standard chow or an HFD (n = 4 to 7/group). (C and D) F4/80 and CD68 mRNA levels in white adipose tissues of MIL10 and WT mice fed standard chow or an HFD (n = 5 or 6/group). Values are means ± SE for each group. *, P < 0.05.
DISCUSSION

Although adipose tissue macrophage accumulation and inflammation are well-described characteristics of obesity-mediated insulin resistance, recent studies refute the cause-and-effect relationship between adipose tissue inflammation and insulin resistance (5–8). Lee et al. have shown that insulin resistance develops in the absence of adipose tissue inflammation after 3 days of high-fat feeding (8). Our recent study also found that a short-term weight loss induced by a low-calorie diet or by exercise improves insulin sensitivity without altering adipose tissue inflammation (7). These findings indicate that adipose tissue inflammation develops in obesity, but it may not be causally associated with insulin resistance in skeletal muscle, a major organ responsible for glucose disposal.

Obesity is a state of systemic inflammation, and local inflammation develops in multiple organs, including skeletal muscle and liver (9, 10). Importantly, recent studies have suggested a direct and causal role of skeletal muscle inflammation and muscle-derived cytokines in the development of insulin resistance (20–25). Our previous study has found that after 3 weeks of high-fat feeding, skeletal muscle was characterized by inflammation, insulin resistance, and reduced glucose metabolism (9). Thus, these findings are consistent with a notion that an HFD-mediated local inflammation in skeletal muscle is causally associated with insulin resistance. However, the metabolic and inflammatory process developing in metabolic organs after a short-term high-fat feeding may differ from the events following chronic obesity. Additionally, chronic obesity models (e.g., a long-term HFD) that are better reflecting type 2 diabetes conditions potentially involve increased insulin from immune cells and inflammatory signaling in skeletal muscle, adipose tissue, and kidney as well as more compromised lipid metabolism and inflammation in the liver (8, 26, 27). Therefore, our current study addresses the effects of anti-inflammatory cytokine IL-10 under chronically obese conditions induced by a 16-week feeding of an HFD. Moreover, leptin is known to regulate inflammation and immunity with its effects on T cell and macrophage-secreted cytokines (17, 28, 29). Thus, the leptin-deficient model used in the current study rules out the potential leptin effects on inflammation. Our newly generated MCK-IL10ob mice also circumvent direct effects of excess dietary lipid uptake and lipid-induced inflammation/insulin resistance because these mice develop spontaneous obesity on a standard chow diet.

As a model of diet-induced obesity and type 2 diabetes, we examined the effects of a long-term HFD on glucose metabolism and local inflammation in MCK-IL10ob mice. After 16 weeks of an HFD, WT mice became markedly obese, with a 4-fold increase in whole-body fat mass. Intramuscular levels of IL-6, IFN-γ, and IL-1α were low in chow-fed mice as expected, but after 16 weeks of an HFD, local cytokine levels in skeletal muscle were elevated by 3- to 4-fold in WT mice. Some of these inflammatory cytokines may be released by locally infiltrating macrophages and/or surrounding adipocytes in skeletal muscle fiber. IL-6 was previously shown to induce insulin resistance by activating STAT3 and increasing intracellular levels of SOCS3, which may target insulin-signaling proteins for ubiquitin-mediated degradation, causing insulin re-

FIG 6 Hepatic glucose metabolism and insulin signaling in MCK-IL10ob and WT mice fed standard chow or an HFD for 16 weeks. (A and B) Hepatic glucose production (HGP) at the basal state and during a hyperinsulinemic-euglycemic clamp in mice (n = 5 or 6/group). (C) Insulin-stimulated Akt phosphorylation in liver. (D and E) Liver G6Pase and PEPCK mRNA levels (n = 4 to 6/group). Values are means ± SE for each group. *, P < 0.05.
IL-10 Regulates Skeletal Muscle Glucose Metabolism

FIG 7 Metabolic effects of muscle-specific IL-10 overexpression in leptin-deficient ob/ob mice (MCK-IL10ob/ob). (A to C) Body weight, whole-body fat mass using 1H-MRS, and fasting glucose levels were measured from 7 MCK-IL10ob/ob and 11 ob/ob mice. (D to G) Daily food intake, physical activity, energy expenditure, and average of hourly VO2 consumption were measured during a 3-day analysis of metabolic cages in 3 MCK-IL10ob/ob and 4 ob/ob mice. Values are means ± SE for each group. *, P < 0.05.

FIG 8 A 2-h hyperinsulinemic-euglycemic clamp in awake MCK-IL10ob/ob and ob/ob mice. (A) Whole-body glucose turnover in MCK-IL10ob/ob (n = 7) and ob/ob (n = 11) mice. (B) Insulin-stimulated glucose uptake in gastrocnemius and quadriceps muscles. (C) White adipose tissue glucose uptake in MCK-IL10ob/ob (n = 7) and ob/ob (n = 12) mice. (D) Hepatic insulin action reflected as insulin-mediated percent suppression of HGP in MCK-IL10ob/ob (n = 7) and ob/ob mice (n = 11). Values are means ± SE for each group. *, P < 0.05.

In contrast, IL-6 has also been shown to be released by postexercise skeletal muscle and to promote glucose metabolism, but this cellular effect involves a much higher level of IL-6 than is typically observed in obesity (32). Furthermore, IL-1α was shown to inhibit insulin signaling by inducing serine phosphorylation of insulin receptor substrate 1 (IRS-1) in adipocytes (33). Consistent with this notion, insulin-stimulated Akt phosphorylation was reduced in the skeletal muscle of HFD-fed
WT mice, supporting the cause-and-effect relationship between obesity-mediated increases in local macrophages and inflammatory cytokines and skeletal muscle insulin resistance.

Despite marked obesity after 16 weeks of an HFD, M<sub>IL10</sub> mice were significantly more insulin sensitive than WT mice, which was largely due to increased insulin signaling and glucose metabolism in skeletal muscle. Intramuscular lipid levels were also similar between WT and M<sub>IL10</sub> mice after an HFD, indicating that IL-10-mediated suppression of muscle inflammation can improve insulin sensitivity without altering intramuscular lipid content in obese mice. IL-10 is an anti-inflammatory cytokine previously known as cytokine synthesis inhibiting factor (CSIF) and produced by many immune cell types, including CD4 T helper cells and macrophages (34, 35). Similar to other cytokines, such as IL-6 and tumor necrosis factor alpha (TNF-α), myocytes have also been shown to express IL-10 and IL-10 receptors (9, 36). IL-10 suppresses local inflammation by inhibiting the synthesis and action of proinflammatory cytokines, including TNF-α, IL-1β, and IL-6, as well as by inhibiting macrophage activation (37, 38). To that end, intramuscular injection of IL-10 DNA was shown to be effective in suppressing inflammation and preventing autoimmune diabetes in mice (39).

Our findings that an obesity-mediated increase in IL-6, IFN-γ, and IL-1α in skeletal muscle was normalized in M<sub>IL10</sub> mice support the anti-inflammatory action of IL-10 in these mice. Also, HFD-fed M<sub>IL10</sub> mice had decreased levels of the macrophage markers F4/80 and CD68, as well as reduced MCP-1 mRNA levels. Importantly, these data indicate that improved muscle glucose metabolism may be due to IL-10 effects to counteract local inflammation in response to chronic high-fat feeding in mice. Additionally, the inflammation was suppressed only in skeletal muscle of HFD-fed M<sub>IL10</sub> mice; there was no difference in liver and adipose tissue inflammation levels in these mice. Plasma IL-10 levels were also not different in HFD-fed M<sub>IL10</sub> mice. Furthermore, in contrast to skeletal muscle, white and brown adipose tissues of M<sub>IL10</sub> mice remained insulin resistant after an HFD, supporting the muscle-specific expression of IL-10 and its effects on muscle glucose metabolism. However, we found that HGP during the clamp was significantly reduced in M<sub>IL10</sub> mice compared to the level in WT mice after 16 weeks of an HFD, suggesting improved hepatic insulin action in HFD-fed M<sub>IL10</sub> mice. Since IL-10 has been shown to be released by myocytes and since its level has been shown to modestly increase in circulation (9), IL-10 may also be responsible for suppressing gluconeogenesis in these mice. In that regard, we did not see alterations in liver Akt phosphorylation in HFD-fed M<sub>IL10</sub> mice, but gluconeogenic gene expression tended to decrease in M<sub>IL10</sub> mice, consistent with reduced HGP in these mice.

Recent studies have shown that an HFD with selective composition of fatty acids may be directly responsible for systemic inflammation, possibly by altering intestinal permeability or the gut microbe population (19). We along with others have also suggested that obesity-mediated inflammation is due to excess nutri-
ent availability, imbalanced nutrient flux and metabolism, and activation of intracellular endoplasmic reticulum and oxidative stress (3, 12, 30). In order to delineate the important question pertaining to the source of obesity-mediated inflammation, we determined the effects of IL-10 in spontaneously obese mice by cross-breeding M^IL10^/ob mice with leptin-deficient ob/ob mice. As expected, IL-10-expressing ob/ob mice (MCK-IL10^ob/ob^) became profoundly obese while on a standard chow diet, with a fat mass accounting for more than 50% of their body weight. Despite being markedly obese and having the same level of plasma free fatty acids (FFA), MCK-IL10^ob/ob^ mice were more insulin sensitive than ob/ob mice, further demonstrating IL-10’s potent insulin-sensitizing effects associated with its anti-inflammatory effect in skeletal muscle. Thus, these data indicate that obesity-mediated inflammation and insulin resistance in skeletal muscle are not dependent on dietary composition. Our findings also suggest that IL-10’s insulin-sensitizing effects are independent of leptin signaling.

The beneficial effects of IL-10 to protect against obesity-mediated insulin resistance may be due to IL-10’s ability to suppress local inflammation and the deleterious action of proinflammatory cytokines in skeletal muscle. Gao et al. have recently shown that hydrodynamic delivery of mouse IL-10 protects against the HFD-mediated glucose intolerance that was associated with reduced inflammation in adipose tissue (40). However, since IL-10 receptors are expressed in multiple cell types, including immune cells and myocytes (9, 41), IL-10’s direct action on myocytes cannot be ruled out. To further determine the mechanism of IL-10’s insulin-sensitizing effects, we generated mice with muscle-specific ablation of IL-10 receptor. Since deletion was chronically high-fat feeding or leptin deficiency causes local inflammation and the deleterious action of proinflammatory cytokines were profoundly elevated in HFD-fed M^IL10R^-/- mice, indicating local activation of inflammatory action of IL-10 protects against muscle insulin resistance. Also, mRNA levels of macrophage markers and inflammatory cytokines were profoundly elevated in HFD-fed M^IL10R^-/- mice. These results suggest that IL-10 effects may be mediated by intracellular IL-10 signaling in skeletal muscle. To that end, we have previously found that the obesity-induced signal to activate local inflammation may involve oxidative stress and altered Ca^2+ homeostasis in skeletal muscle (42). Muscle IL-10 levels also tended to be lower in M^IL10R^-/- mice, which might be the result of a feedback mechanism. Thus, IL-10 signaling in myocytes may relieve oxidative stress and suppress local inflammation, resulting in improved glucose metabolism in skeletal muscle. Further studies are needed to understand how IL-10 may affect oxidative stress in skeletal muscle.

In conclusion, our findings demonstrate that obesity due to chronic high-fat feeding or leptin deficiency causes local inflammation with marked increases in IL-6, IL-1α, and IFN-γ levels in skeletal muscle. Obesity-mediated muscle inflammation is completely attenuated by local expression of IL-10, and this anti-inflammatory action of IL-10 protects against muscle insulin resistance. Additionally, IL-10 signaling in myocytes may relieve obesity-induced oxidative stress and inflammation in skeletal muscle. Taken together, these results support our previous notion that IL-10 may be a potential therapeutic target to treat insulin resistance and type 2 diabetes. Although IL-10 has a short half-life in vivo, local delivery of IL-10 is safe and may have positive effects in skeletal muscle insulin sensitivity (35). While the eventual clinical application and method of delivery need further investigation, the role of IL-10 as an anti-inflammatory and insulin-sensitizing agent may open new doors in type 2 diabetes research.

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