Increased oxidative stress in obesity and its impact on metabolic syndrome.

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Abstract

Obesity is an important causative factor in the development of metabolic syndrome. Here we report that increased oxidative stress in accumulated fat is an important pathogenic mechanism of obesity-associated metabolic syndrome. Fat accumulation correlated with systemic oxidative stress in humans and mice. Production of ROS increased selectively in adipose tissue of obese mice, accompanied by augmented expression of NADPH oxidase and decreased expression of antioxidative enzymes. In cultured adipocytes, elevated levels of fatty acids increased oxidative stress via NADPH oxidase activation, and oxidative stress caused dysregulated production of adipocytokines (fat-derived hormones), including adiponectin, plasminogen activator inhibitor-1, IL-6, and monocyte chemotactic protein-1. Finally, in obese mice, treatment with NADPH oxidase inhibitor reduced ROS production in adipose tissue, attenuated the dysregulation of adipocytokines, and improved diabetes, hyperlipidemia, and hepatic steatosis. Collectively, our results suggest that increased oxidative stress in accumulated fat is an early instigator of metabolic syndrome and that the reduct state in adipose tissue is a potentially useful therapeutic target for obesity-associated metabolic syndrome.

Oxidative stress and plasma adiponectin levels in human obese subjects.

To investigate whether oxidative stress is increased in obese subjects, we measured lipid peroxidation, a marker of oxidative injury, in nondiabetic human subjects. Lipid peroxidation, represented by plasma thiobarbituric acid reactive substance (TBARS) and urinary 8-epi-prostaglandin-F2α (8-epi-PGF2α), significantly correlated with BMI and waist circumference (Figure 1A,1A). Plasma adiponectin levels correlated inversely with BMI and waist circumference (Figure 1A,1A), as we reported previously (21). We also found significant inverse correlations between plasma adiponectin and plasma TBARS and between plasma adiponectin and urinary 8-epi-PGF2α (Figure 1B,1B). These results of human studies allowed us to hypothesize that fat accumulation itself could increase systemic oxidative stress independent of hyperglycemia, and that increased oxidative stress in obesity might relate to the dysregulated production of adipocytokines.

Increased oxidative stress in plasma and white adipose tissue of KKAy mice.

To determine whether fat accumulation is primarily involved in increased oxidative stress, we analyzed 7-week-old KKAy mice as a model of nondiabetic obesity, and 13-week-old KKAy mice as a model of diabetic obesity. KKAy mice exhibit severe obesity, hyperglycemia, and insulin resistance. Compared with age-matched control C57BL/6 mice, KKAy mice at 7 and 13 weeks of age were significantly heavier (Figure 2A,2A) and had a heavier parametrical fat pad (Figure 2A,2A). The increase in plasma glucose was marginal in 7-week-old KKAy mice, but 13-week-old KKAy mice showed significant hyperglycemia relative to C57BL/6 mice (Figure 2A,2A). Surprisingly, plasma lipid peroxidation in nondiabetic 7-week-old KKAy mice was significantly higher than in control mice, and was similar to that in diabetic 13-week-old KKAy mice (Figure 2A,2A). Moreover, plasma levels of H2O2, a hazardous ROS against tissues and cells, were also elevated in nondiabetic and diabetic KKAy mice compared with C57BL/6 mice (Figure 2A,2A). These results demonstrated that oxidative stress in blood was augmented in obesity, that is, fat accumulation, independent of hyperglycemia.

Síndrome metabólica. A obesidade aumenta o estresse oxidativo e pode desencadear a síndrome metabólica


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Abstract

Obesity is a principal causative factor in the development of metabolic syndrome. Here we report that increased oxidative stress in accumulated fat is an important pathogenic mechanism of obesity-associated metabolic syndrome. Fat accumulation correlated with systemic oxidative stress in humans and mice. Production of ROS increased selectively in adipose tissue of obese mice, accompanied by augmented expression of NADPH oxidase and decreased expression of antioxidative enzymes. In cultured adipocytes, elevated levels of fatty acids increased oxidative stress via NADPH oxidase activation, and oxidative stress caused dysregulated production of adipocytokines (fat-derived hormones), including adiponectin, plasminogen activator inhibitor-1, IL-6, and monocyte chemotactic protein-1. Finally, in obese mice, treatment with NADPH oxidase inhibitor reduced ROS production in adipose tissue, attenuated the dysregulation of adipocytokines, and improved diabetes, hyperlipidemia, and hepatic steatosis. Collectively, our results suggest that increased oxidative stress in accumulated fat is an early instigator of metabolic syndrome and that the reduct state in adipose tissue is a potentially useful therapeutic target for obesity-associated metabolic syndrome.

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Increased oxidative stress in plasma and WAT of obese KKAy mice. (A) Body weight, parametrial WAT weight, plasma levels of glucose, lipid peroxidation (TBARS), and H2O2 in C57BL/6 and KKAY mice at 7 and 13 weeks of age. Values are expressed as mean ± (more ...)

Next, we determined the tissue type that could be responsible for the increased oxidative stress in plasma of obese mice. Lipid peroxidation was markedly elevated in white adipose tissue (WAT) of 7- and 13-week-old KKAY mice compared with C57BL/6 mice (Figure 2B). In contrast, the levels of lipid peroxidation in the liver and skeletal muscle were similar between KKAY and C57BL/6 mice at both 7 and 13 weeks of age (Figure 2B). Furthermore, H2O2 production from WAT was significantly higher in 7-week-old KKAY mice than in the control mice (Figure 2C). In contrast, H2O2 production from skeletal muscle and aorta was not altered in KKAY mice (Figure 2C).

Similarly, lipid peroxidation were elevated in other mouse models of obesity, including the diet-induced obesity (DIO) model and another genetic model of obesity (db/db mice) (Figure 2C). These results suggest that plasma lipid peroxidation in mice is due to increased ROS production from accumulated fat.

Dysregulated mRNA expressions of adipocytokines and PPARy in WAT of KKAY mice. The mRNA expressions of adiponectin and PPARy were lower in WAT of 7- and 13-week-old KKAY mice compared with C57BL/6 mice (Figure 3A). In contrast, the mRNA expressions of PAI-1 and TNF-α were high relative to the corresponding levels in the control mice (Figure 3A). We also found that plasma adiponectin levels were lower in KKAY mice than in C57BL/6 mice at both 7 and 13 weeks of age (15.7 ± 0.8 µg/ml vs. 26.2 ± 1.0 µg/ml, P < 0.001 at 7 weeks of age, 16.0 ± 0.9 µg/ml vs. 25.5 ± 1.8 µg/ml, P < 0.001 at 13 weeks of age). Thus, dysregulated expressions of adipocytokines already existed in the nondiabetic obesity stage.

Figure 3 Dysregulated expressions of adiponecin and PAI-1 in WAT of obese KKAY mice. The mRNA expressions of adiponectin, TNF-α, PAI-1, and PPARy in WAT of C57BL/6 (white bars) and KKAY (black bars) (more ...)

Increased mRNA expression of NADPH oxidase in WAT of KKAY mice. NADPH oxidase complex is a major source of ROS in various cells (35, 36). Increased NADPH oxidase activity in vascular cells has been reported to be important in the pathogenesis of hypertension and atherosclerosis by increasing oxidative stress (36). In order to investigate the possible role of augmented NADPH oxidase in increased ROS production, we determined the mRNA expression of NADPH oxidase in WAT of KKAY mice.

The NADPH oxidase complex consists of membrane-associated flavocytochrome b588 protein, which is composed of gp91phox and p22phox, and cytosolic components p47phox, p67phox, and p40phox. In C57BL/6 mice, the mRNA expression of each oxidase subunit was detected in WAT, and the expression levels of most subunits were quite high in WAT compared with other tissues tested (Figure 3B). The mRNA expression levels of these NADPH oxidase subunits were significantly augmented in WAT of nondiabetic 7-week-old KKAY mice, and they were even higher in WAT of diabetic 13-week-old KKAY mice compared with the control mice (Figure 3C). In contrast, the mRNA expression levels of the NADPH oxidase subunits in the liver and skeletal muscle of 7- and 13-week-old KKAY mice were similar to those of control mice (Figure 3D). We also found that the mRNA expression level of transcription factor PU.1, which is known to upregulate the transcription of NADPH oxidase subunits in myeloid cells (37), was also elevated in WAT (Figure 3C), but not in the liver or in skeletal muscle (Figure 3D) of KKAY mice compared with C57BL/6 mice. Similar mRNA changes of NADPH oxidase subunits were observed in WAT, but not in liver or skeletal muscle, of both DIO and db/db mice (Supplemental Figure 1, F and G). These results indicate that the NADPH oxidase pathway is specifically induced in WAT of obese mice.

Decreased mRNA expressions and activities of antioxidant enzymes in WAT of KKAY mice. In the next step, we measured the expressions of antioxidant enzymes including superoxide dismutase (SOD), glutathione peroxidase (GPx), and catalase. At 7 weeks of age, the mRNA expression levels of cytoplasmic Cu/Zn-SOD, GPx, and catalase were significantly lower in WAT of KKAY mice, compared with C57BL/6 mice (Figure 4A). The amount of Cu/Zn-SOD protein was also lower in WAT, but not in the liver and skeletal muscle, of KKAY mice compared with C57BL/6 mice (Figure 4B). Furthermore, total SOD activities (Figure 4C) and GPx activities (Figure 4D) were also significantly and specifically lower in WAT of KKAY mice than in the control mice. Similar mRNA decreases of Cu/Zn-SOD and GPx were observed in WAT, but not in liver or skeletal muscle, of DIO and db/db mice (Supplemental Figure 1, F and G). Taken together, these results indicate that increased ROS production in accumulated fat is due to the activated NADPH oxidase pathway and impaired antioxidant defense system (Figure 4D).

Figure 4 Decreased mRNA expressions and activities of antioxidant enzymes in WAT of obese KKAY mice. (A) The mRNA expressions of Cu/Zn-SOD, Gpx, and catalase in WAT, liver, and skeletal muscle of C57BL/6 (white bars) and KKAY (black and gray bars) mice at 7 and (more ...)

ROS production in adipocytes. We next examined the significance of ROS in cultured adipocytes. ROS production was markedly increased during differentiation of 3T3-L1 cells into adipocytes (Figure 5A), suggesting that ROS production increases in parallel with fat accumulation in adipocytes. We then determined the cellular pathway involved in increased ROS production in mature adipocytes, including NADPH oxidase, xanthine oxidase (XO), and mitochondria-mediated pathways. ROS production in fully differentiated 3T3-L1 adipocytes was markedly suppressed by 2 structurally unrelated inhibitors of NADPH oxidase, diphenyleneiodonium (DPI) and apocynin, as well as the general antioxidant N-acetyl-cysteine (NAC) (Figure 5B). In contrast, ROS production in 3T3-L1 adipocytes was not suppressed by oxypurinol, an inhibitor of xanthine oxidase, rotenone, an inhibitor of mitochondrial electron transport chain complex I, or thenoyltrifluoroacetone, an inhibitor of complex II (Figure 5B). These results suggest that NADPH oxidase is the main source of ROS in adipocytes, and that augmented NADPH oxidase seems to contribute to increased ROS production in adipose tissue in obesity.

Figure 5 Production of ROS in 3T3-L1 adipocytes. (A) ROS production during differentiation of 3T3-L1 cells into adipocytes. ROS production was measured by MBT reduction. Oil Red O staining (top) and NBT treatment (middle) of the cells. Dark-blue formazan was dissolved (more ...)
Oxidative stress also underlies the pathophysiology of hepatic steatosis (51). Thus, oxidative stress locally produced in each of the liver, skeletal muscle, and aorta. These results suggest that adipose tissue is the major source of the elevated plasma ROS.

In the present study, H2O2 production was increased only in adipose tissue of obese mice, but not in other tissues examined, including liver, skeletal muscle, and heart. Expression and smaller amount of nuclear PPARγ under conditions of oxidative stress.

Why is oxidative stress increased only in accumulated fat? We found that in WAT but not other tissues of obese mice, the mRNA expression levels of NADPH oxidase subunits and PPARγ (41). Recent studies of adipocytes found that NOX4, a member of the NOX family, plays a key role in the generation of H2O2 (56). The expression of NOX4 was not detected in macrophages (57, 58). In contrast, ROS increased in adipose tissue of obese mice. Recently, Weisberg et al. (52) and Xu et al. (53) reported that macrophages produce ROS, it is possible that infiltrated macrophages are involved in augmented NADPH oxidase and elevated ROS production in the adipose tissue.

Other Sections

Discussion

Obesity is closely associated with metabolic syndrome (1–3). Recent studies have shown that the dysregulated production of "offensive" adipocytokines, such as PAI-1 (10), TNF-α (11, 12), IL-6 (45), MCP-1 (46), and angiotensinogen (47), and of "defensive" adipocytokines, such as adiponectin (18–20) and leptin (15–17), is critically involved in the pathogenesis of metabolic syndrome. However, the mechanisms by which fat accumulation leads to abnormal expression of adipocytokines and development of metabolic syndrome have not been fully elucidated.

In the present study, we have demonstrated that, in nondiabetic human subjects, fat accumulation closely correlated with the markers of systemic oxidative stress. These data are in good agreement with recent studies suggesting that systemic oxidative stress correlates with BMI (48, 49). In addition, we demonstrated that plasma adiponectin levels correlated inversely with systemic oxidative stress. We then reproduced the results of our human studies in several mouse models of obesity, including KKAy, db/db, and DIO mice. The main finding of the present study is that oxidative stress in accumulated fat mediates the obesity-associated development of metabolic syndrome by the following potential mechanisms: (a) increased oxidative stress in accumulated fat leads to dysregulated production of adipocytokines, and (b) the selective increase in ROS production in accumulated fat leads to elevation of systemic oxidative stress. Plasma adiponectin levels correlated inversely with the markers of systemic oxidative stress in nondiabetic human subjects. In cultured adipocytes, addition of oxidative stress suppressed mRNA expression and secretion of adiponectin, and increased PAI-1, IL-6, adipocytokines, and (b) the selective increase in ROS production in accumulated fat leads to elevation of systemic oxidative stress. These results indicate that local increase in oxidative stress in accumulated fat causes dysregulated production of adipocytokines. Recently, we reported that PPARY positively regulates the transcription of the adiponectin gene via PPARY-responsive element in the promoter (40). We also found that nuclear translocation of PPARY was inhibited by nitrination associated with oxidative stress (50). Therefore, downregulation of adiponectin expression may be partially attributed to the decreased gene expression and smaller amount of nuclear PPARY under conditions of oxidative stress.

In the present study, H2O2 production was increased only in adipose tissue of obese mice, but not in other tissues examined, including liver, skeletal muscle, and heart. Oxidative stress is known to impair both insulin secretion by pancreatic β cells (32) and glucose transport in muscle (30) and adipose tissue (31). Increased oxidative stress in vascular walls is involved in the pathogenesis of hypertension (33) and atherosclerosis (34). Oxidative stress also underlies the pathophysiology of hepatic steatosis (51). Thus, oxidative stress locally produced in each of the above tissues seems to be involved in the pathogenesis of these diseases. Our results suggest that increased ROS secretion into adipose tissue is the result of increased oxidative stress in adipocytes, which is associated with oxidative stress.

Why is oxidative stress increased only in accumulated fat? We found that in WAT but not other tissues of obese mice, the mRNA expression levels of NADPH oxidase subunits increased, and mRNA expression levels and activities of antioxidant enzymes decreased. We also found a high level of mRNA expression of the transcription factor PU.1, which upregulates the transcription of the NADPH oxidase subunits (32) in adipose tissue of obese mice. Recently, Weisberg et al. (52) and Xu et al. (53) reported that macrophages infiltrated the obese adipose tissues and were important source of inflammatory cytokines. Since macrophages are also known to produce ROS, it is possible that infiltrated macrophages are involved in augmented NADPH oxidase and elevated ROS production in the obese adipose tissue. In this regard, a family of gp91phox homologs, termed NOX (NADPH oxidase) proteins, has been reported to be expressed in nonphagocytic cells, not in macrophages (54, 55). Recent studies of adipocytes found that NOX4, a member of the NOX family, plays a role in the generation of H2O2 (56). The expression of NOX4 was not detected in macrophages (57, 58). In contrast,
we found high expression levels of NOX4 in WAT, as well as gp91phox, and mRNA expression of NOX4 was significantly increased in WAT of obese mice (Supplemental Figure 3, A and B). These results suggest that adipose NADPH oxidase is elevated and contributes to ROS production in accumulated fat.

We demonstrated that ROS production was increased in 3T3-L1 adipocytes, in parallel with fat accumulation and by incubation with linoleic acid, in a NADPH oxidase–dependent manner. These results suggest that in accumulated fat, elevated fatty acids activate NADPH oxidase and induce ROS production. We also demonstrated that ROS itself augmented mRNA expressions of NADPH oxidase subunits, including NOX4 (Supplemental Figure 3C) and PU.1 in adipocytes. Therefore, in accumulated fat of obesity, elevated ROS appear to upregulate mRNA expression of NADPH oxidase, establishing a vicious cycle that augments oxidative stress in WAT and blood. We found that ROS increased the expression of MCP-1, a chemoattractant for monocytes and macrophages, in adipocytes. Byproducts of lipid peroxidation by ROS, such as trans-4-hydroxy-2-nonenal and malondialdehyde, are themselves potent chemoattractants (59). Hence, it is possible that increased ROS production and MCP-1 secretion from accumulated fat could cause infiltration of macrophages and inflammation in adipose tissue of obesity.

Importantly, our in vivo study revealed that treatment with the NADPH oxidase inhibitor apocynin reduced ROS production in adipose tissue of KKAy mice. It also improved hyperinsulinemia, hyperglycemia, hypertriglyceridemia, and hepatic steatosis. Increased expression of adipocytokines and decreased expression of TNF-α were observed in WAT and in apocynin-treated KKAy mice. These results demonstrate that reduction of oxidative stress in accumulated fat could improve the dysregulation of adipocytokines in vivo. Our results demonstrate that treatment with NADPH oxidase inhibitor is effective in ameliorating the development of obesity-associated metabolic syndrome. It has been reported that prolonged exposure of 3T3-L1 adipocytes to ROS results in impairments of insulin-induced activation of PI3-kinase and Akt, insulin-stimulated lipogenesis, glucose uptake, and GLUT4 translocation to the plasma membrane (31, 60). Thus, NADPH oxidase inhibitors might improve insulin sensitivity via attenuation of these effects induced by chronic exposure to ROS. Our current results are consistent with several previous studies demonstrating that antioxidant treatment improves insulin function in diabetic subjects (61, 62).

Recent studies, on the other hand, have proposed that ROS such as H2O2 are produced transiently in response to insulin stimulation and also act as a second messenger for insulin signaling in adipocytes (63, 64). NADPH oxidase is thought to be involved in insulin-induced ROS generation (63, 64). We assume that a transient increase of intracellular ROS is important for the insulin signaling pathway, while excessive and long-term exposure to ROS reduces insulin sensitivity and impairs glucose and lipid metabolism.

In the present study, we observed stimulation of ROS production by fatty acids via NADPH oxidase activation. In addition, we also found that ROS suppressed the mRNA expressions of lipogenic genes, such as fatty acid synthase and sterol regulatory element binding protein-1c, in 3T3-L1 adipocytes (data not shown). These results suggest that increased ROS production caused by fat accumulation may prevent further lipid storage, but may simultaneously cause dysregulated expression of adipocytokines and insulin resistance. Conceivably, inhibition of NADPH oxidase should improve the dysregulation of adipocytokines and insulin sensitivity via restoration of normal ROS production in obese adipocytes.

Figure 8 illustrates our working hypothesis regarding the role of ROS in metabolic syndrome. Increased oxidative stress in accumulated fat, via increased NADPH oxidase and decreased antioxidant enzymes, causes dysregulated production of adipocytokines, leading to increased ROS production from accumulated fat, which also leads to increased oxidative stress in blood, hazardously affecting other organs including the liver, skeletal muscle, and aorta. We propose that increased oxidative stress in accumulated fat is an early instigator and one of the important underlying causes of obesity-associated metabolic syndrome, hence, the redox state in adipose tissue is a potentially useful target in new therapies against obesity-associated metabolic syndrome, as demonstrated in the mouse in vivo study.

Figure 8
A working model illustrating how increased ROS production in accumulated fat contributes to metabolic syndrome.

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Methods
Subjects.
Plasma and urinary samples were obtained after overnight fasting from 140 subjects (69 men and 71 women; mean age 56 ± 13 SD) who visited the University hospital for a healthy checkup. Patients with a history of diabetes, cardiovascular or cerebrovascular disease, hepatic or renal disease, tobacco abuse, or those on hormone replacement therapy were excluded. The study protocols complied with the Guidelines of the Ethical Committees of Osaka University and University of the Ryukyus. Informed consent was obtained from all subjects. Samples were stored at –80°C until use.

Animals.
All animals were purchased from Clea Japan and housed in a room under controlled temperature (23 ± 1°C) and humidity (45–65%) and had free access to water and chow (Oriental Yeast Co.). All animal experiments were conducted in accordance with the aforementioned institutional guidelines for the care and use of laboratory animals. Female C57BL/6J and KKAy mice at 7 or 13 weeks of age were anesthetized using pentobarbital sodium, blood was collected, and parametrial WAT, liver tissue, and gastrocnemius muscle were dissected out and frozen in liquid nitrogen. Samples were stored at ~80°C until use. For DIO studies, male C57BL/6J mice were divided at random into 2 groups at 10 weeks of age. The first group was fed a high-fat diet containing 30% fat by weight (AIN93G) while the second group was fed normal chow containing 5.9% fat by weight (CRF-1; Oriental Yeast Co.) for 9 weeks. After an overnight fast, mice were anesthetized, and blood and tissue samples were obtained as described above.

Biochemical measurements.
Urinary 8-epi-PGF2α was determined using an enzyme immunoassay kit (Assay Designs Inc.). Plasma levels of glucose and TG were measured using the Glucose-test and TG E-test, respectively, from Wako Pure Chemical Industries. Plasma adiponectin levels were determined using the Adiponectin ELISA kit (Otsuka). Plasma insulin levels were assessed using an insulin ELISA kit (Shibayagi).

Lipid peroxidation and hydrogen peroxide concentration.
Tissue samples were homogenized in a buffer solution containing 50 mM Tris-HCl (pH 7.4) and 1.15% KCl, and then centrifuged. The supernatant was used for the assay. The levels of lipid peroxidation in plasma and tissue homogenate were measured as TBARS using the LPO-test (Wako Pure Chemical Industries). Hydrogen peroxide concentration in plasma was measured using an Amplex Red hydrogen peroxide assay kit (Invitrogen Corp.).

H2O2 production.
Female C57BL/6J and KKAy mice at 7 weeks of age were anesthetized. Parametral WAT and gastrocnemius muscle were dissected out and placed in Krebs-Ringer phosphate buffer containing (in mM) 145 NaCl, 5.7 sodium phosphate, 4.86 KCl, 0.54 CaCl2, 1.22 MgSO4, and 5.5 glucose (pH 7.4). The aorta was carefully removed, cleaned of excess fat and adventitia, and placed in Krebs-Ringer phosphate buffer. WAT, muscle, and aorta were cut into 2-mm square pieces and incubated at 37°C for 90 minutes. H2O2 released from the tissue was detected using the Amplex Red hydrogen peroxide assay kit (Invitrogen Corp.).

Quantitative RT-PCR.
Total RNA from WAT, liver, and skeletal muscle were prepared with an RNA STAT-60 kit (Tel-Test "B" Inc.). The cDNA was synthesized using the ThermoScript RT-PCR System (Invitrogen Corp.). Real-time PCR was performed on a LightCycler using the FastStart DNA
Master SYBR Green 1 (Roche Diagnostics) according to the protocol provided by the manufacturer. Sequences of primers used for real-time PCR were as follows: Adiponectin, 5'-GATGGCAGAGATGGCACTCC-3' and 5'-CTTGCGAGCTGCTCCTAT-3'; TNF-α, 5'-GCCACCACCTTCTTG-3' and 5'-GTTGTTGGGTAGAAGGACA-3'; PAI-1, 5'-TCCAGCTTTGCGGCTCT-3' and 5'-GCATAGCCGACGCCGAG-3'; IL-6, 5'-ACAACACCGGCTCTCCCTACT-3' and 5'-CACAGTTTCCAGAAACATGG-3'; MCP-1, 5'-CCACTACCTGGCTCTACCT-3' and 5'-TTGGATCTCTTCATGCTTCC-3'; PPARy, 5'-CCAGAGTCTGGCAGTCTCG-3' and 5'-TGCGCGGTTGTCAGTGCTC-3'; p22phox, 5'-GTTCCACCTGAGGCGATGG-3' and 5'-CAATGGCAAGACGAGGCG-3'; p67phox, 5'-CCACTCTTCAGGAATCGCC-3'; cyclophilin, 5'-CAGACGCACAGTCGCTT-3' and 5'-TGTTCTTGGGAATTCGTCGCAA-3'; 18S ribosomal RNA, 5'-CGGCTACCACATCCAAGGAA-3' and 5'-GCTGGAATTACCGCGGCT-3'.

Antioxidant enzyme activity.

Tissue homogenates were prepared as above. SOD activity was measured using the BIDYTECH SOD-525 kit (Oxir). GPx activity was measured using the method described previously (65).

Western blot analysis.

Tissue homogenates (20 µg protein) were subjected to SDS-PAGE, and immunoblotting was performed with anti-Cu,Zn-SOD polyclonal antibody (Upstate Biotechnology).

ROS production in 3T3-L1 adipocytes.

At 5 weeks of age, female C57BL/6J and KKAy mice were treated for 6 weeks with 5 mM apocynin added to the drinking water. At 11 weeks of age, the mice, fed ad libitum, were anesthetized using pentobarbital sodium, blood was collected, and parametrial WAT, liver, and skeletal muscle were dissected out for physiologic analyses. The liver TG contents were measured as described previously (66).

In the human study, linear regression analysis was used to evaluate the relationship between 2 variables. All data are presented as mean ± SEM and were analyzed using unpaired Student's t tests. P values less than 0.05 were considered statistically significant.

In the human adipose tissue, ROS production was detected by nitroblue tetrazolium (NBT) assay (35). NBT is reduced by ROS to a dark-blue, insoluble form of NBT called formazan. At days 0, 2, 4, and 8 after induction, 3T3-L1 cells were incubated for 90 minutes in PBS containing 0.2% NBT. Formazan was dissolved in 50% acetic acid, and the absorbance was determined at 560 nm. Fully differentiated 3T3-L1 adipocytes were incubated with or without 200 µM linoleic acid for 24 hours. Various inhibitors were added in the last hour of incubation, and the cells were incubated for 90 minutes in PBS containing 0.2% NBT with or without inhibitors. NBT, DPI, NAC, oxyurinol, rotenone, and thienoylfluorocetone were purchased from Sigma-Aldrich. Apocynin was purchased from Calbiochem.

Effect of ROS on expression of genes in 3T3-L1 adipocytes.

At day 8 after induction, fully differentiated 3T3-L1 adipocytes were exposed to ROS by incubation with fresh medium containing xanthine oxidase plus hypoxanthine or H2O2 for 24 hours, with or without 10 mM NAC. Cells were harvested, total RNAs were extracted, and the mRNA amounts were quantified by real-time PCR, as described above. The levels of adiponectin in medium were determined by Western blotting (40). Transfection experiments were performed as described previously (40). At day 4 after induction, 3T3-L1 cells in 12-well plates were transfected with 1 µg of the reporter construct containing the fragment corresponding to base pairs –908 to +14 of the human adiponectin promoter linked to a luciferase reporter gene, along with 0.5 µg of pCMX-β-gal (internal control). Four hours later, cells were exposed to ROS for 20 hours, with or without 10 mM NAC. Luciferase activities were assayed using the luciferase assay system (Promega Corp.). Values were normalized by an internal β-galactosidase control and expressed as the relative luciferase activity.

NADPH oxidase inhibitor treatment.

At 5 weeks of age, female CS7BL/6J and KKAY mice were treated for 6 weeks with 5 mM apocynin added to the drinking water. At 11 weeks of age, the mice, fed ad libitum, were anesthetized using pentobarbital sodium, blood was collected, and parametrial WAT, liver, and skeletal muscle were dissected out for physiologic analyses. The liver TG contents were measured as described previously (66).

Statistical analysis.

In the human study, linear regression analysis was used to evaluate the relationship between 2 variables. All data are presented as mean ± SEM and were analyzed using unpaired Student’s t tests. P values less than 0.05 were considered statistically significant.

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