Catalase deficiency facilitates the shuttling of free fatty acid to brown adipose tissue through lipolysis mediated by ROS during sustained fasting

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Abstract

Background

Fatty acids (FA) derived from adipose tissue and liver serve as the main fuel in thermogenesis of brown adipose tissue (BAT). Catalase, a peroxisomal enzyme, plays an important role in maintaining intracellular redox homeostasis by decomposing hydrogen peroxide to either water or oxygen that oxidize and provide fuel for cellular metabolism. Although the antioxidant enzymatic activity of catalase is well known, its role in the metabolism and maintenance of energy homeostasis has not yet been revealed. The present study investigated the role of catalase in lipid metabolism and thermogenesis during nutrient deprivation in catalase-knockout (KO) mice.

Results

We found that hepatic triglyceride accumulation in KO mice decreased during sustained fasting due to lipolysis through reactive oxygen species (ROS) generation in adipocytes. Furthermore, the free FA released from lipolysis were shuttled to BAT through the activation of CD36 and catabolized by lipoprotein lipase in KO mice during sustained fasting. Although the exact mechanism for the activation of the FA receptor enzyme is still unclear, we found that ROS generation in adipocytes mediated the shuttling of FA to BAT.

Conclusions

Taken together, our findings uncover the novel role of catalase in lipid metabolism and thermogenesis in BAT, which may be useful in understanding metabolic dysfunction.

Background

Adipose tissue consists of white adipose tissue (WAT), which stores energy in the form of triacylglycerol (TG), and brown adipose tissue (BAT) which dissipates energy in the form of heat. BAT differs from WAT as it contains small and multilocular lipid droplets (LDs) and abundant mitochondria containing uncoupling protein 1 (UCP1), which is used for dissipation of energy in the form of heat [1]. The main fuel for BAT thermogenesis is the oxidation of fatty acids (FA) derived from WAT through lipolysis or through VLDL and acyl carnitines produced in the liver [2]. During stress or nutrient-deprivation, TG are the major form of lipids in the adipose tissue, where lipolysis takes place through the oxidation of FA, and are mobilized into the liver, where it induces excessive β-oxidation for energy homeostasis [3–5]. The enzyme lipoprotein lipase (LPL) catalyzes the hydrolysis of TG to generate free fatty acids (FFAs). During fasting, lipolysis is induced in WAT through regulation of angiopoietin-like protein 4 (ANGPTL4), which galvanizes LPL activity and thereby directs circulating TG to the liver [6]. ANGPTLs, are a family of proteins structurally similar to angiopoietins, that play an important role in lipid metabolism, inflammation and
cancer [7]; Among ANGPTLs, ANGPTL3, ANGPTL4, and ANGPTL8 play major roles in the trafficking and metabolism of lipid by suppressing LPL activity [3, 8]. Although both ANGPTL3 and ANGPTL4 act as an important regulators of LPL-mediated FA uptake, their expression in tissues varies. ANGPTL3 is a circulating protein synthesized exclusively in the liver [8–10], whereas ANGPTL4 is synthesized in adipose tissue (AT) [11, 12]. Hence, to fuel BAT thermogenesis, the FA transporter enzyme and LPL must be activated.

It is well documented that reactive oxygen species (ROS) regulate lipolysis and BAT thermogenesis [13–16]. ROS are generated during metabolism mainly in the mitochondria, peroxisomes, and endoplasmic reticulum, where oxygen consumption is high [17, 18]. High levels of ROS exert toxic effects on cells, causing the accumulation of oxidative damage in diverse cellular locations [18]. However, the enzyme catalase in the peroxisomes limit this damaging effect by detoxifying $\text{H}_2\text{O}_2$ to either water or oxidizing other organic compounds that provide a major source of metabolic energy for cell survival. During β-oxidation, several $\text{H}_2\text{O}_2$ producing oxidases generated in peroxisomes are converted to hydroxyl radicals (•OH), making peroxisomal membranes vulnerable to lipid peroxidation [18]. Although catalase is important for ROS detoxification, its deficiency in mice and humans is phenotypically normal [19, 20]. Recently, we showed that catalase depletion during starvation causes selective pexophagy in the liver through ROS accumulation in vivo and in vitro [21, 22], while overexpression of catalase has been shown to be beneficial [23–26]. Although catalase is well described as an antioxidant enzyme, its role in maintaining energy homeostasis during starvation has not yet been elucidated.

In this study, we aimed to investigate the role of catalase in the contribution of energy metabolism under fasting conditions. Both wild-type (WT) and catalase-knockout (KO) mice were subjected to sustained fasting for 48 h to ensure sufficient generation of ROS. Our study showed that lipolysis was induced in KO mice through ROS generation, which deviates from FFAs to BAT thermogenesis during sustained fasting.

Results

1. Catalase deficiency markedly decreased the hepatic lipid accumulation during sustained fasting

Starvation produces a complex array of adaptive metabolic responses that activate β-oxidation in both peroxisomes and mitochondria. To investigate the metabolic response of peroxisomes to starvation, mice were fasted for different time intervals (0, 6, 12, 24, 36, and 48 h) and immunoblot analysis was performed using liver homogenates to examine the differential expression of peroxisome markers. Among the peroxisome proteins, only catalase was significantly increased in a time-dependent manner after fasting. The expression levels of other peroxisome markers, including ACOX1, DBP, PEX5, and PMP70, did not change during different time intervals of fasting (Fig. S1A). In addition, there were no significant changes in the protein expression levels of other antioxidants, such as SOD1 and SOD2.
Consistently, the expression level of catalase mRNA significantly increased during sustained fasting, whereas transcriptional activation of other genes, including peroxisomes and antioxidants, remained at a consistent level similar to the protein levels (Fig. S1B). Taken together, these data suggest that catalase might be a key player in liver metabolism during sustained fasting.

To determine whether catalase expression might have been affected by nutrient status, catalase-KO and WT mice were subjected to 48 h of starvation, represented as sustained fasting. Catalase deficiency was confirmed by immunoblot and immunofluorescence analyses of the liver tissues (Fig. S2A-B). It is well known that starvation induces the hydrolysis of TG stored in adipose tissues, releasing FFAs in the plasma. These are taken up by the liver for TG formation, β-oxidation, and VLDL secretion for energy supply [27, 28]. Biochemical analysis was performed to examine TG levels in the liver and serum after fasting for different time intervals (0, 12, 24, 36, and 48 h). Hepatic TG of WT mice started to increase at early hours of fasting, reached a peak at 24 h, and remained sustained until 48 h. However, it drastically decreased after fasting for 24 h and returned to the resting status level at 48 h in KO mice (Fig. 1A). Similarly, TG in the serum was markedly decreased in KO mice after 24 h of fasting, whereas it was sustained in WT mice (Fig. 1B). To evaluate lipid accumulation in the liver during sustained fasting, Oil Red O (ORO) staining was performed (Fig. 1C). ORO staining, shown as a red signal, indicated lipid accumulation in the liver of WT mice during sustained fasting, whereas it was clearly absent in KO mice (Fig. 1C). Consistently, immunoblot analysis for the lipid droplet-associated protein perilipin 2 (PLIN2) and the lipid biogenesis protein seipin [27–30], revealed that increased lipogenesis and accumulation of lipid droplets occurred only in the liver of WT mice, but not in KO mice (Fig. 1D). To address the metabolic response to sustained fasting, we analyzed the expression of acetyl coenzyme A carboxylase 1 (ACC1), which is involved in FA synthesis. The mRNA expression of ACC1, decreased after sustained fasting in both WT and KO mice; however, there was no significant difference between the two experimental groups (Fig. 1E). SREBP1C mRNA, which is involved in the regulation of glucose metabolism, decreased significantly after sustained fasting without significant difference between the two experimental groups (Fig. 1E). Expression of genes involved in FA catabolism, including SCD-1 and FAS, were significantly decreased after sustained fasting without any significant difference between the two experimental groups. We also measured β-hydroxybutyrate levels in the liver after sustained fasting for 48 h (Fig. 1F); the levels started to increase at 12 h, reached a maximum at approximately 24 h, and sustained for a further 48 h in WT mice. However, it drastically dropped to the basal level at 48 h in KO mice. To assess total FA, gas chromatography-mass spectrometry (GC-MS) was performed (Fig. 1G). Accumulation of many FA species, including both long chain and median chain FA, decreased significantly during sustained fasting in KO mice compared to WT mice (Fig. 1G). Taken together, these data suggest that catalase deficiency resulted in decreased lipid accumulation in the liver during sustained fasting.

2. Catalase deficiency during sustained fasting accelerated lipolysis from the white adipose tissue through ROS generation

As hepatic TG levels were diminished in KO mice during sustained fasting, we reasoned that TG consumption might be induced by the activation of FA oxidation (FAO) in the liver of fasted KO mice.
Therefore, anomalies or extreme activation of mitochondrial or peroxisomal FAO could be the prime reason for the decreased hepatic TG accumulation in fasted KO mice. However, we did not find any involvement of mitochondrial or peroxisomal FAO [22]. Mitochondrial FAO did not change in either WT or KO mice, whereas peroxisome was decreased through autophagic degradation in KO mice during sustained fasting [22], indicating that TG depletion in the liver is not related to mitochondrial and peroxisomal FAO.

Fasting is a unique stress condition that enhances lipid catabolism. Prolonged fasting is known to induce TG hydrolysis in adipose tissue, increasing the plasma concentration of FFAs and inducing hepatic TG accumulation. Hence, we measured serum FFAs levels and found that it decreased in KO mice during sustained fasting (Fig. S3). Therefore, we assumed that KO mice might consume FFAs released from adipocytes faster than WT mice during sustained fasting. To examine this possibility, WAT from WT and KO mice was isolated and weighed. As expected, fat masses of KO mice were drastically reduced during sustained fasting compared to WT mice (Fig. 2A-B). Morphological analysis of adipose tissue sections by hematoxylin and eosin (H&E) staining showed bright eosinophilic cytoplasm with abundant lipid droplets in fed WT mice, whereas that of fasted WT mice were small in size with shrinkage of adipocytes and scanty eosinophilic cytoplasm. Similarly, the adipose tissue in fed KO mice was homogenous and uninnucleate with nearly white-colored lipid droplets, while fasted KO mice showed a marked loss of adipocytes and an extended eosinophilic cytoplasm (Fig. 2C). Immunofluorescence staining of PLIN1, a lipid droplet-associated protein found in adipose tissue, had markedly decreased only in fasted KO mice (Fig. 2D). As fat mass in adipose tissue was drastically decreased, we hypothesized that induction of lipolysis in WAT could be one of the reasons for the depletion of fat mass observed during sustained fasting in KO mice. Thus, we measured the serum FFAs and glycerol levels at different time intervals (0, 12, 24, 36, and 48 h) in WT and KO mice. After 36 h of fasting, FFAs level increased in a time-dependent manner and then remained stable in WT mice, while the level had decreased slightly in KO mice after 36 h of fasting (Fig. 2E). Serum glycerol levels remained unaltered in both WT and KO mice after sustained fasting in a time-dependent manner (Fig. 2F). To confirm lipolysis in adipose tissue, immunoblot analysis of lipolytic proteins, including hormone-sensitive lipase (HSL) and adipose triglyceride lipase (ATGL) was performed. Phosphorylation of HSL and expression of ATGL were markedly increased in fasted KO mice, suggesting that lipolysis signaling in adipose tissue was over activated (Fig. 2G). To address the mechanistic evidence of how lipolysis occurs in KO mice, we reasoned that ROS might be a key player in inducing lipolysis, as previously shown [13]. Hence, the WAT of mice was homogenized and incubated with the dye 2’7’-dichlorofluorescein diacetate (DCFH-DA) to measure total ROS. As expected, ROS generation was significantly increased in both fed and fasted KO mice compared to WT mice. In addition, ROS level in fasted KO mice was significantly higher than in fed mice (Fig. S4). Together, these data indicate that catalase deficiency during sustained fasting resulted in a marked reduction in fat mass, which is mediated through lipolysis of WAT caused by ROS generation.

**3. Catalase deficiency during sustained fasting induces brown adipose tissue thermogenesis**
Lipolysis has been shown to provide a source of energy for brown fat thermogenesis [29, 30]. In addition, reduction of fat mass is accompanied by increased energy expenditure and intrascapular BAT (iBAT) thermogenesis [31]. As FFAs released from WAT were neither consumed nor accumulated in the liver in KO mice, we assumed these FFAs might be consumed by BAT for thermogenesis. To test BAT activation, BAT isolated from WT and KO mice was used to perform morphological analysis by H&E staining. BAT showed with multilocular lipid droplets and reduced lipid deposition in KO mice after sustained fasting (Fig. 3A). It is well known that BAT activates UCP1 in the mitochondrial membrane to release potential energy as heat [29]. Expression of uncoupling proteins, including UCP1 and phosphorylated p38, were markedly increased in BAT from KO mice after sustained fasting compared to WT mice (Fig. 3B).

Furthermore, expression of the mitochondrial marker Tom20 was increased only in fasted KO mice. Consistently, mRNA levels of UCP1 and PGC-1α were increased in fasted KO mice compared to WT mice (Fig. 3C-D). As thermogenesis protein was increased in KO mice, we measured the rectal temperature of mice. The rectal temperature was significantly higher in fasted KO mice than in WT mice after 24 h of fasting (Fig. 3E). It is well known that activation of BAT is regulated by the sympathetic nervous system via β3-adrenergic receptors (ADBR3) [32]. Q-PCR for ADBR3 mRNA from BAT was performed, which revealed that expression of ADBR3 was significantly increased in fasted KO mice compared to WT mice (Fig. 3F). To evaluate whether increased expression of mitochondrial proteins in fasted KO mice was associated with mitochondrial activity in BAT, the enzymatic activities of mitochondrial complex I (COX I) and complex IV (COX IV) were measured. The enzymatic activity of COX I and IV was significantly increased in BAT from KO mice after sustained fasting (Fig. 3G-H). We showed that ROS increased lipolysis in WAT and lipolysis-induced BAT thermogenesis. ROS may also be the key player in inducing BAT thermogenesis. To prove this, BAT was homogenized and incubated with DCFH-DA, and total ROS was measured. ROS generation was significantly increased in BAT of KO mice compared to WT mice on sustained fasting (Fig. 3I). These data suggest that FFAs released from excessive lipolysis of WAT may be consumed by the ROS-mediated thermogenic pathway of BAT in fasted KO mice.

4. Catalase deficiency during sustained fasting facilitates the shuttling of FFA to the BAT but not in the liver through FA transporter

Fasting enhances lipid catabolism by hydrolyzing TG stored in adipocytes, which augments FFAs and glycerol in the liver to supply lipids to other tissues. We found that catalase deficiency during sustained fasting diminished hepatic lipid accumulation and augmented lipolysis to fuel the energy demand. Although lipolysis was induced, the liver did not obtain sufficient lipids during sustained fasting in KO mice. Thus, we hypothesized that the release of FFAs from fat tissues during sustained fasting might be shuttled to BAT rather than the liver in KO mice.

During fasting, triglyceride-rich lipoprotein (TRL) transports lipids to the bloodstream, where lipids are catabolized by the action of LPL. Hence, through LPL and transmembrane receptor cluster of differentiation 36 (CD36), other tissues such as liver, muscle, and brown adipocytes get energy during nutrient deprivation [33, 34]. Recently, it was reported that during lipolysis, FFAs are delivered by TRL to the FA receptor enzyme CD36 and catabolized by LPL, that promotes sufficient energy needed for BAT.
activation [34]. Hence, BAT utilizes FA that are liberated from lipolysis as a source that activates UCP1 for thermogenesis [35, 36]. BAT activation inhibits ANGPTL4, an important regulator of LPL-mediated FA uptake into BAT. Shuttling of ANGPTL4 enhances LPL activity to uptake FA derived from circulating TRL to BAT activation [11]. Hence to test the activity of the FA receptor, immunoblot analysis was performed to check FA shuttling protein in BAT of mice fasted for 48 h. The protein level of ANGPTL4 was markedly decreased whereas CD36 was markedly increased in the BAT of KO mice during sustained fasting (Fig. 4A). On the contrary, LPL activity significantly increased after 36 h of fasting in the BAT of KO mice compared to that of WT mice (Fig. 4B). Consistent with LPL activity, mRNA levels of CD36 and LPL were also significantly increased in the BAT of KO mice compared to WT mice (Fig. 4C-D). FA taken up by CD36 and catabolized by LPL in BAT could be a major fuel source to activate UCP1 in the mitochondrial membrane. Herein, we observed that the mRNA level of UCP1 continuously increased after 12 h of sustained fasting in KO mice compared to WT mice (Fig. 4E). In contrast, immunoblot analysis with ANGPTL3 protein in the liver showed a marked increase whereas CD36 level was slightly decreased in expression up to 48 h during sustained fasting in KO mice compared to WT mice (Fig. 4F). The lipolytic activity of LPL decreased significantly during sustained fasting in the liver of KO mice (Fig. 4G). Furthermore, mRNA levels of LPL and CD36 were significantly decreased in KO mice during sustained fasting compared to WT mice (Fig. 4H-I). Together, these data indicate that catalase deficiency during sustained fasting facilitates the shuttling of FFAs to BAT through the FA receptor, CD36, and were catabolized by LPL.

5. Both ROS generation and lipolysis by isoproterenol is attenuated by N-acetylcysteine in catalase deficient adipocytes

To confirm that the induction of lipolysis strictly occurred from WAT and not due to any other reason including hormonal changes in vivo, we analyzed the autonomous induction of lipolysis from primary inguinal white adipose tissue (iWAT). To mimic the effects of fasting and lipolysis in vitro, isoproterenol was added to differentiated iWAT of WT and KO mice. Isoproterenol treatment significantly increased the green fluorescence signals derived from DCFH-DA staining, reflecting increased ROS generation only in KO mice (Fig. 5A-B). The fluorescent signal in isoproterenol-treated iWAT of KO mice was inhibited after the addition of N-acetylcysteine (NAC), confirming the effect of NAC on ROS generation. We were further interested in the role of ROS in isoproterenol-induced lipolysis. As expected, treatment of differentiated iWAT with isoproterenol resulted in lipolysis, observed as fragmented smaller lipid droplets and decreased fluorescent intensity of PLIN1 compared to WT (Fig. 5C). However, features of lipolysis were more obvious in differentiated iWAT from KO mice than those from WT mice (Fig. 5C-D). We also confirmed that pre-treatment with NAC partially recovered lipolytic features, including the fluorescence intensity of PLIN1 and lipid content, of differentiated iWAT from WT and KO mice (Fig. 5C-D). In addition, the hydrolysis of cellular lipids induced by isoproterenol in differentiated iWAT of KO mice was more severe than that in WT mice, which was significantly inhibited by the addition of NAC. To determine whether NAC mitigated lipolysis in iWAT, we measured the levels of glycerol and FFAs (Fig. 5E-F). These levels significantly increased on treatment with isoproterenol, which was previously significantly attenuated by NAC in differentiated iWAT of both WT and KO mice. Consistently, the expression of lipolytic proteins,
including phospho-HSL and ATGL, decreased on addition of NAC to catalase-KO adipocytes treated with isoproterenol (Fig. 5G). Taken together, these data suggest that the antioxidant NAC prevents lipolysis induced by ROS in catalase-KO primary white adipocytes.

6. UCP1 activation in brown adipocytes induced by ROS is suppressed by N-acetylcysteine in catalase deficient adipocytes

To assess the effect of ROS generated during fasting on BAT thermogenesis, differentiated BAT was treated with isoproterenol, and ROS generation was measured by DCFH-DA staining in the presence or absence of NAC (Fig. 6A). The results showed that treatment with isoproterenol increased the green fluorescence signals derived from DCFH-DA staining, reflecting an increase in ROS generation from differentiated BAT of WT and KO mice (Fig. 6A-B). However, there was a marked increase in the green fluorescence signals of DCFH-DA in differentiated BAT of KO mice compared to that of WT mice, which was inhibited by the addition of NAC. Next, proteins involved in uncoupling and mitochondrial enzymes were analyzed by immunoblot analysis and Q-PCR. Phosphorylated p38, an upstream target of UCP1, and UCP1 protein were increased in differentiated BAT treated with isoproterenol compared to the control (Fig. 6C). The band intensities of phosphorylated p38, UCP1, and PGC-1α were dominant in isoproterenol-treated differentiated BAT from KO mice compared to those of WT mice. As expected, co-treatment with NAC diminished the expression of phosphorylated p38, UCP1, and PGC-1α in isoproterenol-treated differentiated BAT of both WT and KO mice. The mRNA expression of UCP1 was also increased in isoproterenol-treated differentiated BAT of both WT and KO mice, which was also diminished by the addition of NAC. Next, we measured the transcriptional activation of genes involved in mitochondrial biogenesis and β-oxidation, including PGC1a and PPARα, in differentiated BAT treated with isoproterenol (Fig. 6D-F). Treatment with isoproterenol predominantly increased the mRNA expression of PGC-1α and PPARα in differentiated BAT of KO mice, which was also clearly diminished by NAC. Together, these data suggest that treatment with fasting mimic isoproterenol results in ROS generation, which further activates the signaling pathway of WAT lipolysis and uncoupling events of BAT in KO mice. Furthermore, antioxidant NAC successfully prevented ROS-mediated lipolysis of WAT and uncoupling of BAT under fasting mimic conditions.

**Discussion**

During fasting, TG in adipose tissue is hydrolyzed to FFAs and transferred to the liver, where they are utilized for energy production and stored in lipid droplets. In contrast, we found that hepatic TG accumulation was reduced in fasted KO mice. We speculated that lipolysis in fasted KO mice induces enhancement of β-oxidation, which promotes H₂O₂ generating oxidase in peroxisomes. The H₂O₂ generated may cause a decrease in lipid accumulation in hepatocytes. Our speculation is supported by a previous report that ROS is generated in catalase-deficient mice and in nutrient deprived cells [21, 22]. Additionally, the cellular redox state affects the activity of several enzymes involved in lipid metabolism in the liver [37].
Lipolysis induces the release of FFAs from white adipocytes and triggers BAT activation via UCP1 activation for thermogenesis [34–36]. To generate heat, BAT uses a large amount of TG to perform efficient mechanisms for continuous replacement of intracellular lipid storage. We showed that catalase deficiency during sustained fasting induces BAT thermogenesis and hence requires a surplus amount of lipid to fuel thermogenesis. Therefore, the fatty acid receptor enzyme CD36 is active in BAT, and continuously takes up the FA delivered by circulating TRL and catabolized by LPL [11, 38]. Taken together, we believe that FFAs released from white adipocytes are taken up by BAT through the activation of CD36. Meanwhile, activation of ANGPTL3, a negative regulator of LPL in hepatocytes, reduced fatty acid receptor enzymes, resulting in the liver not getting sufficient FA delivered by circulating TRL. This observation caused a decrease in hepatic lipid accumulation during sustained fasting in KO mice. Recently, the fatty acid receptor enzyme CD36 was shown to be involved in the degradation of lipids in the liver [39], which overlaps with our data that a decrease in fatty acid receptor enzymes lessened lipid accumulation in the liver. Next, we investigated how ANGPTL3 in the liver and ANGPTL4 in BAT regulate fatty acid receptor enzymes in circulating TRL during fasting. Although the exact mechanism remains to be elucidated, we hypothesize that ROS generation in KO mice might regulate ANGPTLs in the bloodstream. Our assumption is consistent with a previous report that ROS inhibits ANGPTL4 expression in adipocytes during hyperoxia [40, 41]. Hence, ROS generation in BAT might be a key player in the inhibition of ANGPTL4, which activates LPL to trigger BAT thermogenesis. Two members of the angiopoietin-like family, ANGPTL3 and ANGPTL4, modulate lipoprotein metabolism with differences in key mechanisms. In addition to tissue-specific localization, ANGPTL3 in the liver and ANGPTL4 in adipocytes, their protein structure and mode of inhibition of LPL activity are also different [42, 43]. Furthermore, we also showed that fat mass was significantly decreased in fasted KO mice, which might be due to enhanced lipolysis. Reduced fat mass is also accompanied by the activation of BAT thermogenesis [31]. Indeed, our data suggest that lipolysis is activated in fasted KO mice and that the FFAs released from WAT were unavailable for delivery to the liver for TG synthesis because BAT consumed them for thermogenesis.

It is well known that UCP1 is responsible for non-shivering thermogenesis in BAT. However, the factor responsible for the production of heat by UCP1 in BAT is unknown. Many studies in mice and humans have shown that cold stimulation induces non-shivering thermogenesis in BAT. Moreover, liberation of FA from WAT by lipolysis is responsible for BAT thermogenesis during fasting [30, 35, 36]. As discussed above, FA released by lipolysis can be taken up by BAT for thermogenesis. Hence, fasting could also be a possible reason for BAT activation [36]. Other reports also suggest that BAT thermogenesis is activated by fasting, such as intermittent or every other day fasting, besides cold stress, to alleviate obesity [44, 45]. We also showed that ROS generation is responsible for lipolysis and BAT thermogenesis, concurrent with previous reports suggesting that BAT activation is associated with ROS generation [13–16]. Consistently, our study suggests that enhanced ROS generation induces UCP1 for thermogenesis in catalase-KO brown adipocytes.

Peroxisomes are abundant and play a vital role in the adipose tissue [46–49]. In adipocytes, they are closely associated with lipid droplets and play a key role during the differentiation of pre-adipocytes to...
mature adipocytes, suggesting that de novo peroxisome formation and transcriptional regulation of thermogenesis are interrelated [50]. We showed that the peroxisome number is decreased in the liver [22], whereas it is increased in BAT in KO mice during sustained fasting (Fig. S5). We argued that the drastic appearance of peroxisomes might be specific to the tissue. In the liver due to sustained fasting, we can speculate that excessive ROS is induced for meeting energy demands, which might increase peroxisome oxidases to induce autophagic degradation of peroxisomes as a self-defense mechanism in catalase-deficient mice. In contrast, peroxisomes are critical for BAT thermogenesis, as reported earlier [50].

Collectively, our study proposes that controlling catalase activity would be a valuable approach for maintaining homeostatic balance in lipid metabolism.

Materials And Methods

Animal treatments

Homozygous KO mice were kindly provided by Dr. Ye-Shih Ho (Institute of Environmental Health Sciences and Department of Biochemistry and Molecular Biology, Wayne State University, USA). Mice were interbred and experiment were designed as described previously [22]. Mice were maintained in accordance with the standard protocol approved by the Animal Care and Use Committee at Gwangju Institute of Science and Technology, Korea.

Isolation, culture and differentiation of primary adipocytes

Both iBAT, isolated from newborn to 7 days old mice, and iWAT, isolated from 7–8 weeks old mice were minced and subjected to collagenase digestion (2 mg of collagenase in 2 mL of isolation buffer containing 0.123 M NaCl, 5 mM KCl, 1.3 mM CaCl₂, 5 mM glucose, 100 mM HEPES, and 4% BSA) for 1 h at 37°C in a shaker. The cell suspension was filtered through a 100 µm filter. The filtrate was centrifuged at 600 ×g for 5 min. Pellets were resuspended in 2 mL SVF growth medium (DMEM consisting of 25 mM glucose, 20% FBS, 20 mM Hepes, 100 units/mL penicillin/streptomycin) and seeded on 6-, 12-, and 24 well plates in a humidified atmosphere of 5% CO₂. After 2 days post confluence (designated day 0), differentiation was induced by addition of differentiation media (DMEM containing 0.5 mM IBMX, 0.5 µM dexamethasone, 20 nM insulin) and incubated in a humidified atmosphere of 5% CO₂. At day 2, the medium was replaced with maintenance medium (DMEM medium containing 20 nM insulin and 0.5 µM rosiglitazone). This medium was renewed every 48 h until the adipocytes fully differentiated.

Lipolysis induction in adipocytes

After cell differentiation, the medium was replaced with lipolytic medium (DMEM medium containing 1 g/L glucose, 5% FBS, 2% BSA) and supplemented with or without 10 µM isoproterenol (I6504; Sigma-Aldrich) and incubated in a humidified atmosphere containing 5% CO₂. The lipolytic experiments were carried out for 12 h. Cultures were frequently observed with a phase contrast microscope. After 12 h of
incubation, serum from the cells was extracted. FFAs (Cayman #0488858) and glycerol (Biovision #K630-100) were measured according to the manufacturer’s protocol.

**Measurement of hepatic TG levels**

Serum TG levels were measured using a 7600 Clinical Analyzer (Hitachi). To measure hepatic TG, 50 mg of liver tissue was homogenized, and TG was measured using a Triglyceride Colorimetric Assay kit (Cayman Chemical) according to the manufacturer’s protocol.

**Histological analysis**

For histological analysis, tissues were fixed in 10% formalin solution, embedded in paraffin, and sectioned. H&E staining were performed as described previously [22].

For immunofluorescence (IF) staining of tissue and primary adipocytes, snap-frozen tissues were sectioned using a cryostat and then defrosted at room temperature for 1 h. The primary adipocytes and tissue sections were fixed in 4% paraformaldehyde (HT5014; Sigma-Aldrich) at room temperature for 20 min, washed thrice in PBS, incubated with 0.2% Triton X-100 for 5 min, and rinsed in PBS. The sections were blocked with 5% goat serum with 0.1% Triton X-100, whereas adipocytes were blocked with 3% bovine serum albumin (BSA) for 1 h at room temperature. Then, both were incubated with the target protein antibody (1:200) with their respective blocking solution overnight at 4°C. The next day, the tissues and adipocytes were again washed thrice with PBS, stained with Alexa Fluor-conjugated secondary antibody in blocking solution at room temperature in the dark for 1 h, washed twice with PBS, and incubated with 10 µM DAPI in PBS at room temperature for 10 min. Images were acquired and analyzed using an Olympus FluoView 1000 confocal laser scanning system. The slides were scored in a blinded manner.

**Oil Red O (ORO) staining**

Cryosectioned liver samples and primary adipocytes were fixed with paraformaldehyde for 20 min and washed twice with distilled water. ORO solution (01391; Sigma-Aldrich) was added to the samples for 1 h at room temperature. The sections were washed with distilled water and counterstained with hematoxylin solution for 1 min whereas adipocytes were washed with PBS and incubated with 10 µM DAPI in PBS at room temperature for 10 min. Images were acquired and analyzed using a light microscope for tissue and fluorescence microscope for adipocytes. The slides were scored in a blinded manner.

**Western blot analysis**

To determine the expression levels of target protein, western blot from whole cell analysis using tissues sample and primary adipocytes were performed as described previously [21, 22]. The antibodies used and their sources are listed in Supplementary Table S1.

**Measurement of total FAs**
Liver tissue (~ 50 mg) from the experimental mice was homogenized using a TissueLyzer (Qiagen). Heneicosanoic acid as an internal standard was added to the samples, and FA were extracted by the Folch method [51]. For FA analysis, base hydrolysis using KOH was performed, followed by neutralization with HCl. Methyl esterification was performed with BCl3-MeOH at 60°C for 30 min, and the FA content was analyzed by gas chromatography-mass spectrometry (GC-MS) analysis using 7890A/5975A (Agilent). All standards, including the internal standard for calibration, were purchased from Avanti Polar Lipids and Sigma-Aldrich.

**Quantitative reverse transcription-polymerase chain reaction (qPCR)**

qPCR from tissues lysates and primary adipocytes were analysed as described earlier [22]. The primer sequences are listed in Supplementary Table S2.

**Mitochondrial complex activity**

Liver and BAT were homogenized with buffer provided in the Complex I Enzyme Activity Microplate Assay Kit (ab109721, Abcam Biotech, UK) and Complex IV Rodent Enzyme Activity Microplate Assay Kit (ab109911, Abcam Biotech). Mitochondrial complex I and IV activities were measured according to the manufacturer's instructions.

**Determination of LPL activity**

Liver and BAT tissues were homogenized in ice-cold PBS. The centrifuged supematant was collected and the LPL activity was measured using the LPL activity assay kit (Biovision, # K72-100).

**Determination of ROS production**

ROS production from tissue lysates and adipocytes were performed as previously described [21, 22].

**Statistical analysis**

All values are presented as mean ± standard deviation (SD). One-way analysis of variance was used to compare means, and P-values < 0.05 were considered statistically significant.

**Declarations**

**Author Contributions**

RKD performed conceptualization, analysis, investigations, methodology development, visualization and writing original draft and validation of experimental results. JNL performed supervision, investigation, methodology development, writing, review and editing. YM performed analysis and investigation, methodology development, conceptualization, validation of experimental results. CP contributed to project administration, supervision, resource and software management. SKC contributed to project
administration, supervision review and editing. YSH contributed to resource management. 
RKP contributed to funding acquisition, conceptualization, visualization and supervision.

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Availability of data and materials:
All data generated and/or analyzed in this study are available from the corresponding author on reasonable request.

Ethics approval and consent to participate:
All procedures were approved by the Animal Care and Use Committee at Gwangju Institute of Science and Technology, Korea.

Consent for publication:
Not applicable.

Competing interests:
The authors have no competing interest to declare

Abbreviations
ANGPTLs: Angiopoietin-like proteins; BAT: Brown Adipose Tissue; CD36: Cluster of differentiation 36; FA: Fatty Acids; FAO: Fatty Acid Oxidation; FFAs: Free Fatty Acids; KO: Knock Out; LPL: Lipoprotein Lipase; ROS: Reactive Oxygen Species; TG: Triglyceride; UCP1: uncoupling protein 1; VLCFA: Very Long Chain Fatty Acid; WAT: White Adipose Tissue; WT: Wild Type.

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**Figures**
Figure 1

Catalase deficiency decreases the accumulation of lipid in liver during sustained fasting. Liver samples from mice fasted at the indicated time were homogenized, and TG levels were analyzed from (A) liver and (B) serum (n = 6). (C) Liver tissues from mice cryosectioned and stained with ORO. Scale bar represents 20μm. (D) Immunoblot analysis of liver tissue. Protein expression using whole-cell lysates with the indicated antibodies (n = 3). (E) qPCR analysis of FA metabolism-related genes from the liver of mice fed
and fasted for 48 h. (F) β-hydroxybutyrate was analyzed from the serum of mice fasted for indicated time. (G) GC-MS for total FA of long Chain and VLCFA from liver samples (n = 3). Values represent mean ± SD (n = 3, 4). *P < 0.001 WT 48 h fasting versus KO 48 h of fasting.

Figure 2

Catalase deficiency during sustained fasting decreases the fat mass that augmented lipolysis in WAT. (A) Images of WAT from mice (n = 6). (B) Analytical weight of WAT in gram (g) (C) Representative H&E staining of WAT from mice. (D) Representative fluorescence images of WAT, fixed and immunostained with anti-PLIN1 (green) and DAPI (blue). Scale bar represents 20μm. (E) FFA and (F) glycerol level from mice serum fasted for indicated time (n = 6). (G) Proteins were extracted from WAT and immunoblot analysis was performed using whole-cell lysates with the indicated antibodies (n = 3). Values represent mean ± SD (n = 3, 4). *P < 0.05 WT 48 h fasting versus KO 48 h of fasting.
Figure 3

Catalase deficiency during sustained fasting increased BAT thermogenesis through mitochondrial activation (A) Representative H&E staining of BAT from mice. Scale bar represents 20μm. (B) Proteins were extracted from BAT of WT and KO mice and immunoblot analysis was performed using whole-cell lysates with the indicated antibodies indicated (n = 3). α-Tubulin was used as a loading control. qPCR analysis of (C) UCP1 and (D) PGC-1α genes in BAT from mice. (E) Rectal body temperature was measured from mice fasted for indicated time (n = 6). (F) qPCR analysis of ADBR3. (G-H) Activities of mitochondrial complexes I and IV from BAT of mice. Details are described in Materials and Methods. (I) Total ROS was measured from BAT. Values represent mean ± SD (n = 3, 4). *P < 0.05 WT 48 h fasting versus KO 48 h of fasting.
Figure 4

Catalase deficiency during sustained fasting facilitates the shuttling of free fatty acid to BAT through fatty acid receptor enzymes (A) Proteins were extracted from BAT of mice fasted at indicated time and immunoblot analysis was performed using whole-cell lysates with indicated antibody (n = 2). (B) Isolated BAT from fasted mice at indicated time were homogenized and LPL activity was measured. qPCR analysis of (C) CD36, (D) LPL and (E) UCP1 in the BAT of mice fasted for indicated time. (F) Proteins were extracted from liver of mice fasted at indicated time and immunoblot analysis was performed using whole cell lysates with indicated antibodies (n = 2). (G) Isolated liver from fasted mice at indicated time were homogenized and LPL activity was measured. qPCR analysis of (H) CD36 and (I) LPL genes in liver of mice fasted for indicated time. Values represent mean ± SD (n = 3, 4). *P < 0.05 WT 48 h fasting versus KO 48 h of fasting; ##P < 0.001 WT 48 h fasting versus KO 48 h of fasting.
**Figure 5**

ROS generation induced lipolysis was suppressed by N-Acetylcyesteine in catalase deficient adipocytes (A) Representative fluorescence images of primary white adipocytes treated as indicated and fluorescence were measured with DCFH-DA and examined under a fluorescence microscope. Scale bar represents 100 μm. (B) Percentage of cells showing green fluorescence (corresponding to DCFH-DA) and fluorescence intensity of adipocytes. ##P < 0.001 WT ISO vs KO ISO; #P < 0.001 KO ISO vs KO ISO+NAC. (C) Representative fluorescence images primary white adipocytes, fixed and immunostained with anti-PLIN1 (green), ORO (red), and DAPI (blue). Scale bar represents 5 μm. (D) The quantification of ORO positive cells in white adipocytes from C. *P < 0.05 WT nor vs WT ISO; WT ISO vs KO ISO; KO ISO vs KO ISO+NAC. (E) Serum glycerol; **P < 0.01 WT nor vs WT ISO; WT ISO vs KO ISO; KO ISO vs KO ISO+NAC and (F) FFA level in primary white adipocytes. *P < 0.05 WT nor vs WT ISO; WT ISO vs KO ISO; KO ISO vs KO ISO+NAC. (G) Proteins were extracted from treated adipocytes and immunoblot analysis was performed using whole-cell lysates with the indicated antibodies. Data represent mean ± SD of three independent experiments.
Figure 6

UCP1 activation in brown adipocytes induced by ROS was suppressed by N-Acetylcysteine in catalase deficient adipocytes (A) Representative fluorescence images primary brown adipocytes treated as indicated and fluorescence were measured with DCFH-DA and examined under a fluorescence microscope. Scale bar represents 100 μm. (B) Percentage of cells showing green fluorescence (corresponding to DCFH-DA) and fluorescence intensity of adipocytes. *P < 0.05 WT ISO vs KO ISO; **P < 0.001 KO ISO vs KO ISO+NAC (C) Proteins were extracted from treated adipocytes and immunoblot analysis was performed using whole cell lysates with the indicated antibodies. qPCR analysis of (D) UCP1: *P < 0.01 WT nor vs WT ISO; WT ISO vs KO ISO; KO ISO vs KO ISO+NAC. (E) PGC-1α: **P < 0.05 WT ISO vs KO ISO; KO ISO vs KO ISO+NAC and (F) PPARα in isolated brown adipocytes. Data represent mean ± SD of three independent experiments.

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