

Ribosomal protein–Mdm2–p53 pathway coordinates nutrient stress with lipid metabolism by regulating MCD and promoting fatty acid oxidation

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The tumor suppressor p53 has recently been shown to regulate energy metabolism through multiple mechanisms. However, the *in vivo* signaling pathways related to p53-mediated metabolic regulation remain largely uncharacterized. By using mice bearing a single amino acid substitution at cysteine residue 305 of mouse double minute 2 (*Mdm2*^{C305F}), which renders Mdm2 deficient in binding ribosomal proteins (RPs) RPL11 and RPL5, we show that the RP–Mdm2–p53 signaling pathway is critical for sensing nutrient deprivation and maintaining liver lipid homeostasis. Although the *Mdm2*^{C305F} mutation does not significantly affect growth and development in mice, this mutation promotes fat accumulation under normal feeding conditions and hepatosteatosis under acute fasting conditions. We show that nutrient deprivation inhibits rRNA biosynthesis, increases RP–Mdm2 interaction, and induces p53-mediated transactivation of malonyl-CoA decarboxylase (MCD), which catalyzes the degradation of malonyl-CoA to acetyl-CoA, thus modulating lipid partitioning. Fasted *Mdm2*^{C305F} mice demonstrate attenuated MCD induction and enhanced malonyl-CoA accumulation in addition to decreased oxidative respiration and increased fatty acid accumulation in the liver. Thus, the RP–Mdm2–p53 pathway appears to function as an endogenous sensor responsible for stimulating fatty acid oxidation in response to nutrient depletion.

The dynamic process of cell growth and division is under constant surveillance. As one of the primary “gatekeepers” of the cell, p53 plays a major role in sensing and responding to a variety of internal and external stressors to maintain cellular homeostasis. In addition to its conventional roles in promoting cell cycle arrest, senescence, and apoptosis, p53 has recently been shown to regulate metabolism through the transcriptional activation of genes involved in glucose transport, glycolysis, oxidative phosphorylation (OXPHOS), and glutamine hydrolysis as well as in the activation of genes upstream of the mammalian target of rapamycin and autophagic pathways (reviewed in ref. 1). Virtually all cancers show metabolic changes that result in enhanced glucose consumption and elevated glycolytic activity known as the Warburg effect (2). Because cells continuously undergo metabolic perturbations as a result of constantly changing physiological and environmental cues such as daily feeding/fasting cycles, p53 may act in this context as a metabolic stress regulator altering cellular metabolic programs under nonlethal or “low-stress” conditions.

Recent studies have shown that inhibition of ribosomal biogenesis can activate p53 through the ribosomal protein (RP)-mediated suppression of mouse double minute 2 (*Mdm2*) in the RP–Mdm2–p53 stress response pathway (3). Detailed analysis revealed that several RPs require the *Mdm2* central zinc finger motif for efficient *Mdm2* binding. Interestingly, cancer-associated

MDM2 mutations have been reported to affect the central zinc finger motif (4, 5) and can specifically disrupt RP binding (6). To study the physiological functions of the RP–Mdm2–p53 pathway, we generated mice carrying a single human cancer-associated phenylalanine substitution at cysteine residue 305, which is structurally critical for the central zinc finger of *Mdm2* (*Mdm2*^{C305F}). The C305F mutation specifically disrupts *Mdm2* binding to RPL11 and RPL5 (6) and increases the rate at which *Mdm2*^{C305F} mice develop oncogenic c-Myc–induced lymphomas (7).

Here, we study the metabolic alterations of *Mdm2*^{C305F} mice in the context of acute dietary restriction and report that nutrient depletion inhibits ribosome biogenesis and induces RP–Mdm2 interaction. As a result, p53 becomes activated and induces malonyl-CoA decarboxylase (MCD), an enzyme that stimulates malonyl-CoA turnover and enhances carnitine palmitoyl-transferase (CPT) activity, thereby augmenting mitochondrial fatty acid uptake and oxidative respiration. To our knowledge, our results demonstrate for the first time in an animal model that the RP–Mdm2–p53 signaling pathway is critically involved in lipid homeostasis by coordinating the appropriate physiological response to nutrient deprivation.

Significance

Although progress has been made in the characterization of p53 in regulating metabolism, very little is known about the signaling pathways involved in this regulation in response to stress *in vivo*. Here we show that p53 controls hepatic fatty acid oxidation in mice in response to fasting. Disruption of ribosome protein (RP)-mouse double minute (*Mdm2*) binding in *Mdm2*^{C305F} mice results in fasting-induced hepatosteatosis. A full-dosage of p53 and an intact RP–Mdm2–p53 pathway are required for the induction of malonyl coA decarboxylase (MCD), a critical regulator of fatty acid oxidation. Thus, the RP–Mdm2–p53 pathway functions as a key regulator of hepatic lipid homeostasis in response to nutrient deprivation stress, a function that has implications in organismal survival and tumor suppression.

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Results

***Mdm2*^{C305F} Mice Have Altered Body Composition and Basal Metabolism.** General characterization of the *Mdm2*^{C305F} mice showed that they are viable to adulthood, fertile, and exhibit no clinical or pathological symptoms. The *Mdm2*^{C305F} mice gained weight at a similar rate to their WT counterparts when maintained on a normal diet (Fig. S1 *A* and *B*). Furthermore, the *Mdm2*^{C305F} mice showed a similar daily food intake to WT mice (Fig. S1*C*). Intriguingly, although the overall body weight was similar to the WT mice, body composition analysis revealed that the *Mdm2*^{C305F} mice maintain a higher percentage of body fat (Fig. 1*A*) and a correspondingly lower percentage of lean mass (Fig. 1*B*) than age-matched WT mice, indicating that the *Mdm2*^{C305F} mice may harbor a predisposition to obesity. To determine the basal metabolic rate, we analyzed energy expenditure in male mice fed a normal diet. During dark and light phases, O₂ consumption (Fig. 1*C*), CO₂ production (Fig. 1*D*), and heat generation (Fig. S1*D*) were all reduced in the *Mdm2*^{C305F} mice relative to WT littermates. Because basal

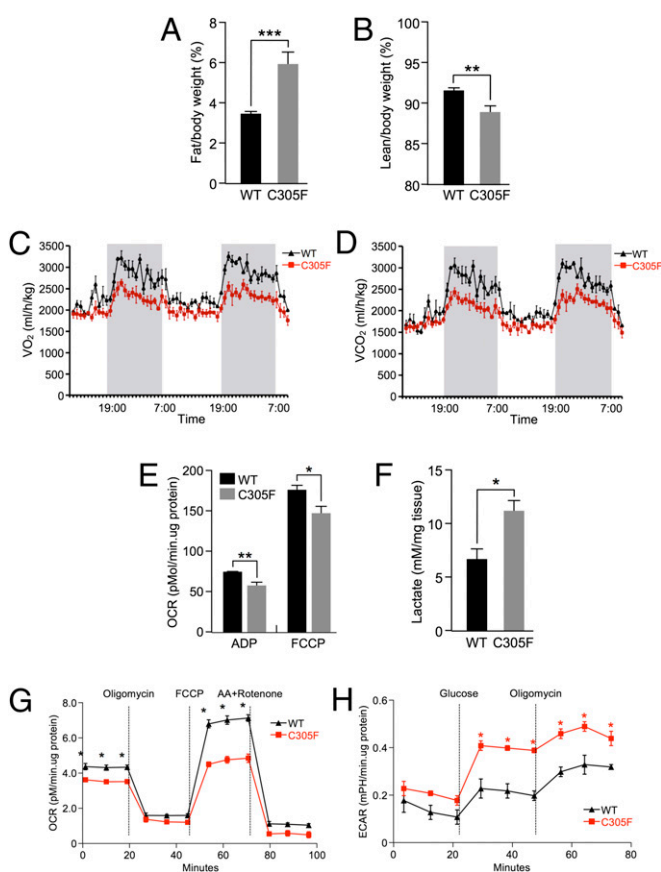


Fig. 1. *Mdm2*^{C305F} mice exhibit altered body composition and basal metabolism under unstressed conditions. Percentage of fat (*A*) and lean mass (*B*) was determined in 15-wk-old WT ($n = 36$) and *Mdm2*^{C305F} ($n = 19$) mice fed a normal chow diet by echo-MRI (fat percentage, $P = 0.0002$; lean mass percentage, $P = 0.003$). (*C* and *D*) Indirect calorimetry of WT (black line) and *Mdm2*^{C305F} (red line) mice fed a chow diet ad libitum. (*C*) VO₂, (*D*) carbon dioxide release (i.e., VCO₂). The white background indicates light hours, and the shaded background indicates dark hours. (*E*) OCR in the liver mitochondria of WT and *Mdm2*^{C305F} mice. (*F*) Lactate level in the livers of WT and *Mdm2*^{C305F} mice ($n = 3$ per genotype). (*G*) Linear graph showing OCR in WT and *Mdm2*^{C305F} MEFs in the presence of oligomycin (1 μ M), FCCP (0.5 μ M), antimycin A (2 μ M), and rotenone (1 μ M). (*H*) Linear graph showing the ECAR in WT and *Mdm2*^{C305F} MEFs in the presence of glucose (4.5 g/L) and oligomycin (1 μ M). Error bars denote SEM.

metabolic rate reflects mitochondrial activity, we directly tested mitochondrial OXPHOS [measured as oxygen consumption rate (OCR)] and glycolysis [measured as extracellular acidification rate (ECAR)] in liver homogenates. Basal oxygen consumption, which was measured in the presence of succinate and ADP, and the maximal oxygen consumption, which was measured in the presence of the mitochondrial uncoupler carbonyl cyanide 4-(trifluoromethoxy)-phenylhydrazone (FCCP), were significantly reduced in the *Mdm2*^{C305F} mice (Fig. 1*E*). In contrast, lactate levels were higher in *Mdm2*^{C305F} mouse livers (Fig. 1*F*), indicating that the reduced OXPHOS was compensated by enhanced glycolytic energy production in *Mdm2*^{C305F} mice. Simultaneously measuring respiration and glycolysis in mouse embryonic fibroblasts (MEFs) by using a Seahorse analyzer showed similar results whereby *Mdm2*^{C305F} MEFs have lower OXPHOS activity and higher glycolysis activity (Fig. 1*G* and *H*). The observed decreased energy expenditure, increased body fat gain, and impaired mitochondrial OXPHOS in the *Mdm2*^{C305F} mice suggest that the RP–Mdm2–p53 pathway plays an important role in energy homeostasis under unstressed normal feeding conditions.

***Mdm2*^{C305F} Mice Lose More Body Weight upon Fasting.** To further understand the role of the RP–Mdm2–p53 pathway in response to stress, we analyzed the *Mdm2*^{C305F} mice when subjected to the dietary stress of acute fasting. Previous experiments have shown that, in laboratory animals, fasting induces a significant increase in spontaneous ambulatory activity and voluntary exercise, an effect referred to as “fasting-induced foraging activity” (8). Consistent with this observation, WT mice showed increased in-cage foraging activity upon fasting, whereas, in contrast, the *Mdm2*^{C305F} mice showed no significant increase in foraging activity in response to fasting (Fig. S2). Interestingly, despite the decreased ambulatory activity of the *Mdm2*^{C305F} mice, the *Mdm2*^{C305F} mice lost significantly more body weight than their WT counterparts (Fig. 2*A* and *B*). To further investigate the difference in body weight loss, we analyzed the body composition of the mice by quantitative magnetic resonance spectroscopy. Our results show that, although the *Mdm2*^{C305F} mice lost more overall bodyweight, these mice were relatively resistant to fasting-induced fat loss (Fig. 2*C*), as they demonstrated a preference for the consumption of lean mass for their energy supply (Fig. 2*D*). To determine whether the inhibition of fat breakdown is caused by reduced lipolysis, we examined the epididymal fat pads of WT and *Mdm2*^{C305F} mice. Consistent with higher levels of body fat, the *Mdm2*^{C305F} mice showed 30% larger epididymal fat pads than the WT mice (Fig. 2*E*); however, fasting resulted in a similar rate of decrease in the epididymal fat pad size in the *Mdm2*^{C305F} and WT mice, suggesting that lipolysis occurs at a normal rate in the mutant mice (Fig. 2*E*).

Fasted *Mdm2*^{C305F} Mice Develop Hepatosteatosis. To gain insight into the mechanisms leading to decreased body fat loss in fasted *Mdm2*^{C305F} mice, we examined the livers because the liver is one of the primary organs involved in energy metabolism during fasting. Under ad libitum feeding conditions, the size and appearance of the *Mdm2*^{C305F} mouse livers were similar to the livers of WT mice (Fig. 3*A* and *C*). However, following fasting, whereas livers from the WT mice appeared normal in color and gross appearance (Fig. 3*B*), livers from *Mdm2*^{C305F} mice were paler and displayed a mottled appearance (Fig. 3*D*) indicative of increased fat content. Oil red O-stained liver sections showed a small amount of lipid droplets in the livers of WT and *Mdm2*^{C305F} mice under fed conditions (Fig. 3*E*, *Top*). After a 24-h period of fasting, a marked accumulation of lipid droplets in the WT and *Mdm2*^{C305F} livers was observed (Fig. 3*E*, *Middle*), which is characteristic of the temporary fasting-induced accumulation of lipids in the liver (9, 10). After a 48-h period of

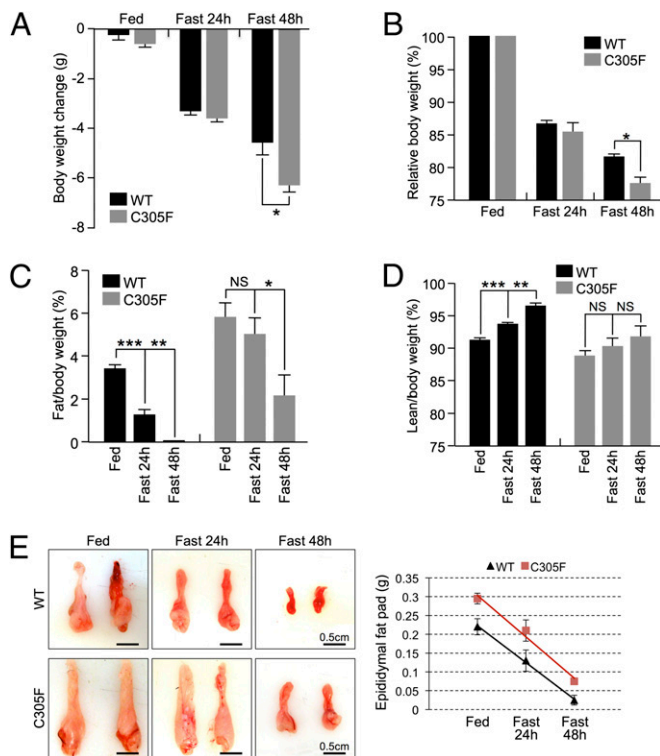


Fig. 2. Fasted *Mdm2^{C305F}* mice show accelerated loss of lean mass. (A–D) WT and *Mdm2^{C305F}* mice were fed a chow diet ad libitum (Fed) or were fasted for 24 or 48 h ($n = 5$ per genotype). Body composition was determined by echo-MRI. (A) Effect of fasting on body weight change. (B) Fasted body weight normalized to body weight before fasting. (C) Percentage of fat in WT and *Mdm2^{C305F}* mice under the indicated nutrient conditions. (D) Percentage of lean mass in WT and *Mdm2^{C305F}* mice under the indicated nutrient conditions. (E) Changes in the size (Left) and weight (Right) of the epididymal fat pads from WT and *Mdm2^{C305F}* mice under the indicated nutrient conditions. Error bars denote the SEM.

fasting, the number and size of lipid droplets in the WT mouse livers returned to the basal level, whereas lipid droplet accumulation continued in the *Mdm2^{C305F}* mouse livers (Fig. 3E, Bottom). Consistent with the lipid droplet accumulation in *Mdm2^{C305F}* mouse livers, we observed a significant increase in liver triacylglyceride (TAG) levels in fasted *Mdm2^{C305F}* mouse livers (Fig. 3F). These data indicate that *Mdm2^{C305F}* mouse livers have an attenuated ability to metabolize fatty acids. The excess TAG accumulation in the *Mdm2^{C305F}* mouse liver is consistent with hepatosteatosis, a consequence of metabolic syndrome (11). Collectively, these data suggest that the RP–Mdm2–p53 pathway is involved in maintaining hepatic lipid homeostasis.

Fasting-Induced Fatty Acid Oxidation Is Impaired in *Mdm2^{C305F}* Mice.

To further understand the metabolic alterations in the fasted mice, an integrated metabolomics study was conducted based on high-resolution NMR (¹H NMR) spectroscopy. When fed ad libitum, spectra based on liver lysates obtained from WT and *Mdm2^{C305F}* mice largely overlapped (Fig. 4A, filled symbols). Fasting resulted in a metabolic shift in the WT and *Mdm2^{C305F}* mice; however, the WT and the *Mdm2^{C305F}* metabolite spectra were notably different, producing distinct principal component analysis (PCA) plots (Fig. 4A, open symbols). These results suggest that fasting triggers unique global metabolic alterations in WT and *Mdm2^{C305F}* mice.

Upon further analysis of the metabolic profiles, fasted WT and *Mdm2^{C305F}* mouse livers differed significantly in the levels

of key individual metabolites. Specifically, β -hydroxybutyrate (β -HB), a ketone body derived from fatty acid oxidation (FAO; β -oxidation), accumulated in WT but not *Mdm2^{C305F}* mouse livers, whereas glucose, which is derived mostly from the digestion of lean tissue and muscle during prolonged starvation, was more abundant in the *Mdm2^{C305F}* mouse livers (Fig. 4B). Additionally, fasted *Mdm2^{C305F}* mice demonstrated high levels of muscle autophagy (Fig. S3A) and increased expression of glucose-6-phosphatase (Fig. S3B). These results are consistent with the observation that *Mdm2^{C305F}* mice tend to lose more muscle mass during nutrient deprivation (Fig. 2).

β -HB accumulation is a direct consequence of FAO during prolonged fasting and can serve as a hepatic and serum biomarker of starvation (12). Ketogenesis, a process that produces β -HB during FAO (13), was fully induced in fasted WT mice as indicated by high levels of serum β -HB (Fig. 4C). In contrast, ketogenesis was reduced in fasted *Mdm2^{C305F}* mice (Fig. 4C), indicative of attenuated FAO. Notably, even under fed conditions *Mdm2^{C305F}* mice generated less β -HB than WT mice, suggesting that the RP–Mdm2–p53 pathway may play a role in fatty acid catabolism during daily feeding/fasting cycles.

To directly measure fatty acid metabolism, we analyzed long-chain FAO (LC-FAO) in mouse liver homogenates using [¹⁻¹⁴C] palmitate (a 16-carbon saturated fatty acid) as a substrate. As shown in Fig. 4D, fasting resulted in a significant increase in the rate of maximal LC-FAO in WT but not in *Mdm2^{C305F}* mouse livers. Accordingly, fasted WT mice showed enhanced

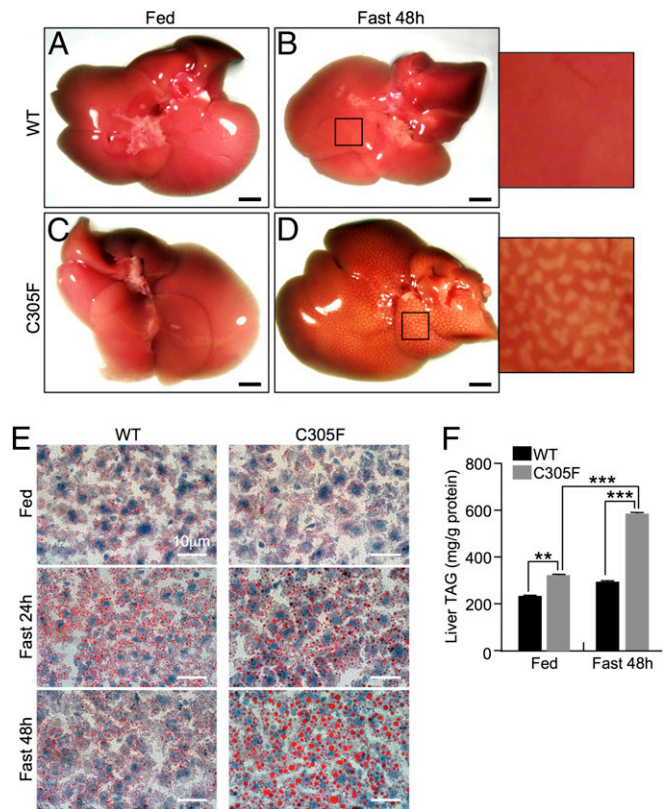


Fig. 3. Fasted *Mdm2^{C305F}* mice develop hepatosteatosis. (A–D) WT and *Mdm2^{C305F}* mice were fed a normal chow diet or were fasted for 48 h. The mice were killed, and the livers were collected. (Scale bars: 2.5 mm.) (E) Hepatic lipid droplets were visualized by oil red O staining on livers obtained from mice fed a normal chow diet or fasted for the indicated time. (F) Elevated TAG in *Mdm2^{C305F}* mouse livers upon fasting ($n = 3$). Error bars denote SEM.

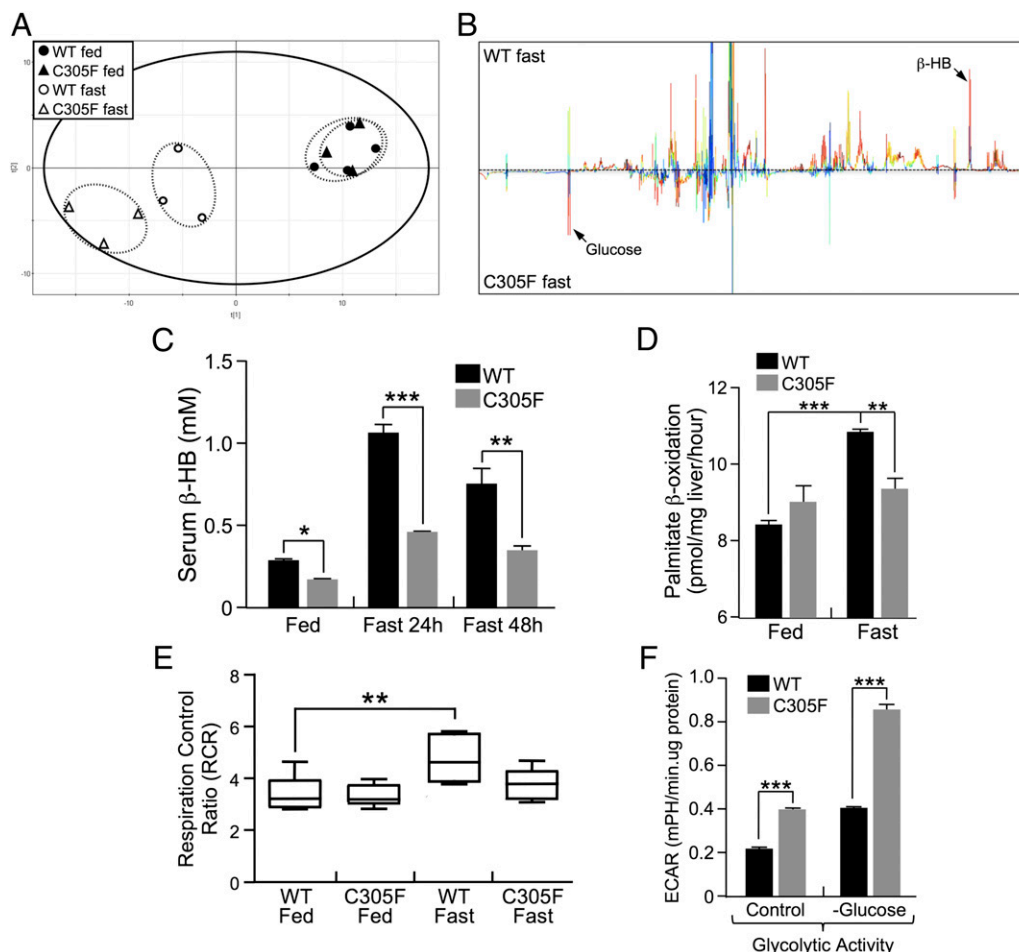


Fig. 4. $Mdm2^{C305F}$ mice demonstrate reduced FAO and mitochondrial dysfunction. (A) WT and $Mdm2^{C305F}$ mice were fed a chow diet ad libitum (Fed) or were fasted (Fast) for 48 h. Metabolomic analysis was performed on WT and $Mdm2^{C305F}$ mouse livers by H1-NMR. NMR spectra were acquired and were subjected to unsupervised PCA (WT, $n = 4$; $Mdm2^{C305F}$, $n = 3$). (B) WT and $Mdm2^{C305F}$ mice were treated as indicated in A. The major metabolic perturbations associated with the cluster PCA plots were identified from orthogonal partial least-squares loading coefficient plots. The identified metabolites are shown as individual peaks. β -HB and glucose are indicated with black arrows. (C) Serum β -HB was determined in WT and $Mdm2^{C305F}$ mice fed a normal chow diet ad libitum or fasted for 24 or 48 h (WT, $n = 10$; $Mdm2^{C305F}$, $n = 8$). (D) Fatty acid β -oxidation was examined in liver homogenates from WT and $Mdm2^{C305F}$ mice fed a normal chow diet ad libitum or fasted for 24 h ($n = 6$). The total β -oxidation activity was defined as the rate of [14 C]palmitate conversion to CO_2 and acid-soluble metabolites. (E) RCR of liver mitochondria isolated from WT and $Mdm2^{C305F}$ mice fed a normal chow diet ad libitum or fasted overnight (WT fed, $n = 9$; $Mdm2^{C305F}$ fed, $n = 9$; WT fast, $n = 5$; $Mdm2^{C305F}$ fast, $n = 5$). RCR was determined based on the ratio of the OCR in the presence of FCCP and the OCR in the presence of oligomycin. (F) WT and $Mdm2^{C305F}$ MEFs were cultured in normal DMEM or glucose-free DMEM supplemented with 10% FBS for 24 h, and then the cells were seeded into Seahorse XF24 plates (6×10^4 per well), and the ECAR was recorded upon sequential injection of glucose (4.5 g/L) and oligomycin (1 μ M). The maximal glycolytic activity was determined by the addition of oligomycin. Error bars denote the SEM. *, **, and *** are used to indicate statistical significance corresponding to P values < 0.05 , 0.01 , and 0.001 , respectively. NS indicates no significant difference.

mitochondrial OXPHOS relative to $Mdm2^{C305F}$ mice (Fig. 4E), and fasted $Mdm2^{C305F}$ MEFs demonstrated higher glycolytic activity (Fig. 4F). Altogether, our data support a model whereby the RP–Mdm2–p53 pathway regulates global metabolic homeostasis during nutrient deprivation in part through the modulation of hepatic FAO.

Nutrient Depletion Inhibits rRNA Synthesis, Induces RP–Mdm2 Binding, and Activates p53. Previous studies have demonstrated that the cells grown in serum-depleted medium show increased RP–Mdm2 binding and p53 activation (14). To investigate whether nutrient depletion inhibits ribosome biosynthesis in WT and $Mdm2^{C305F}$ MEFs, we examined nascent rRNA levels. MEFs were pulse-labeled with [3 H]methionine, after which total cellular RNA was isolated and probed for the amount of radioactivity, which was used as a surrogate marker for newly synthesized rRNA precursors. Upon serum depletion or glucose starvation, rRNA synthesis was reduced to $\sim 20\%$ and 40% ,

respectively (Fig. 5A). The dynamics and extent of inhibition of rRNA synthesis were similar in WT and $Mdm2^{C305F}$ MEFs, indicating that the C305F mutation does not affect rRNA synthesis. Next, we examined the RNA/DNA ratio in mouse livers, an assay that has been used to measure rRNA synthesis in fasted rats (15). Consistent with the MEF cell data, the RNA/DNA ratio in WT and $Mdm2^{C305F}$ mouse livers decreased similarly during the first 12 h of fasting (Fig. 5B), indicating that nutrient deprivation rapidly and profoundly affects rRNA synthesis. Next, we examined RPL11–Mdm2 binding and p53 activation in nutrient-deprived MEFs. Serum starvation and glucose deprivation augmented RPL11–Mdm2 binding in WT but not $Mdm2^{C305F}$ MEFs (Fig. 5C and D). As a result, p53 was stabilized and activated to a greater degree in WT MEFs compared with $Mdm2^{C305F}$ MEFs (Fig. 5E and F). Consistent with the increase in p53 activity, SCO2, a known p53 metabolic target (16), was induced in WT but not in $Mdm2^{C305F}$ MEFs. We have consistently observed that the p53 protein level is higher in

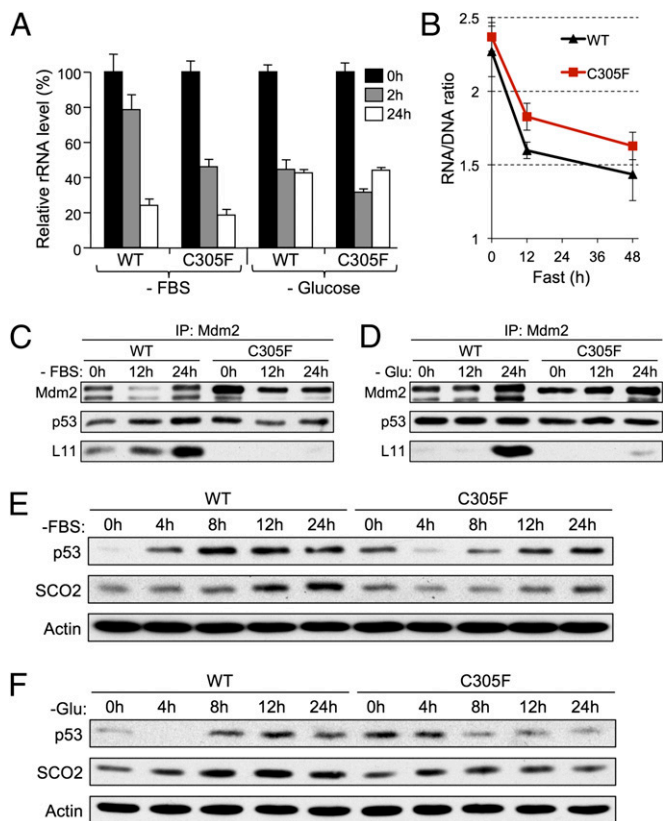


Fig. 5. Nutrient depletion inhibits rRNA synthesis, induces L11-Mdm2 binding, and activates p53. (A) WT and *Mdm2*^{C305F} MEFs were pulse-labeled with [methyl-³H]methionine, and then the total RNA was isolated. The radioactivity of the RNA was measured by using a scintillation counter. The bar graph represents the average of three experiments performed in duplicate. (B) Ratio of total liver RNA to DNA during fasting. Total RNA and DNA were extracted simultaneously with TRIzol reagent from 200 mg of liver pieces from WT and *Mdm2*^{C305F} mice under the indicated treatment conditions. (C and D) WT and *Mdm2*^{C305F} MEFs were treated with (C) serum starvation (-FBS) or (D) glucose starvation (-Glu) for the indicated times. Cell lysates were then immunoprecipitated with anti-Mdm2 antibody (2A10) and immunoblotted for Mdm2, p53, and L11. (E and F) WT and *Mdm2*^{C305F} MEFs were treated as in C and D for the indicated times. Cell lysates were analyzed by immunoblotting for total p53 and SCO2. Error bars denote the SEM.

untreated *Mdm2*^{C305F} MEFs than in WT MEFs, which was reported in one of our previous studies (7); the nature of the elevated basal level of p53 in the mutant cells remains unclear. Nonetheless, our results show that nutrient deprivation in WT and *Mdm2*^{C305F} MEFs results in the inhibition of rRNA biosynthesis, which results in RPL11-Mdm2 binding and activation of p53 in WT but not *Mdm2*^{C305F} cells.

Nutrient Depletion Induces RP-Mdm2-p53 Pathway-Dependent Activation of MCD. We reasoned that the phenotype displayed by the *Mdm2*^{C305F} mice is likely a result of altered p53 function. By using an inducible p53ER^{TAM} MEF cell system, we performed microarray analysis for p53 target genes. One robust candidate identified by microarray analysis was MCD (Fig. S44). MCD catalyzes the conversion of malonyl-CoA, a key fatty acid precursor and a FAO blocker, into acetyl-CoA and carbon dioxide. A rate-limiting step to LC-FAO in the liver is the transfer of fatty acids from the cytosol into the mitochondria by CPT1 α , an outer mitochondrial membrane enzyme that regulates the entry and subsequent oxidation of fatty acids by the mitochondria (17). Importantly, CPT1 α is allosterically inhibited by malonyl-CoA (13, 18). Thus, MCD functions as a central

regulator of LC-FAO by regulating the intracellular levels of malonyl-CoA to control the balance between lipid and glucose metabolism. To confirm the microarray results and to test the ability of p53 to directly activate MCD, a ChIP assay was performed to probe several putative p53 response elements present in the MCD promoter region (Fig. 6A). Of the three putative p53 response elements tested, RE2 and RE3 were robustly bound by p53, suggesting a direct regulatory mechanism. Furthermore, MCD transcript expression was induced in a p53-dependent manner in MEFs treated with nutlin-3 (Fig. 6B) or nutrient deprivation (Fig. 6C). Notably, in *Mdm2*^{C305F} MEFs, MCD was induced by nutlin-3 to a similar extent relative to WT MEFs. However, upon nutrient deprivation, *Mdm2*^{C305F} MEFs failed to induce MCD expression, indicating that the C305F mutation specifically affects nutrient stress signaling. We observed a slight induction of MCD in p53-null MEFs upon nutrient deprivation (Fig. 6C), as well as a slight increase in p53-null MEFs treated with nutlin-3 (Fig. 6B), suggesting that MCD can be partially induced in a p53-independent manner. Furthermore, MCD protein expression was greatly induced in WT MEFs but only slightly induced in *Mdm2*^{C305F} MEFs under conditions that inhibit ribosome biosynthesis including actinomycin D (Fig. 6D) and nutrient depletion (Fig. 6E and F). Consistent with a p53-dependent effect, MCD induction correlated with levels of total p53 (Fig. 5E and F) and S-18 phospho-p53 (Fig. 6D-F). We observed a similar p53-dependent induction of MCD in the p53 isogenic human cancer cell line HCT116 (Fig. S4B).

To determine the effect of nutrient deprivation on MCD expression in mice, young male mice were fasted, after which livers were collected and examined for MCD mRNA and protein expression. MCD mRNA (Fig. 6G) and protein (Fig. 6H-I) expression was induced in the livers of fasted WT mice but not in *Mdm2*^{C305F} mice. Furthermore, MCD protein levels peaked around 12 h of fasting, which is a characteristic expression pattern for genes that are involved in FAO (19). MCD catalyzes the degradation of malonyl-CoA, thereby activating CPT1 α and increasing liver LC-FAO (20). Consistent with the pattern of MCD induction in fasted mice, we observed a significant decrease of malonyl-CoA (Fig. 6J) and an increase in CPT1 α activity (Fig. 6K) in WT but not *Mdm2*^{C305F} mouse livers. Altogether, our data indicate that the RP-Mdm2-p53 pathway regulates nutrient depletion stress-stimulated FAO at least partially through the p53-dependent induction of MCD. The resulting induction of MCD functions to decrease intracellular malonyl-CoA levels, thereby increasing CPT1 α activity.

Reduction of p53 Dosage Mimics Hepatosteatotic Phenotype Observed in *Mdm2*^{C305F} Mice. To confirm that the hepatosteatotic phenotype observed in fasted *Mdm2*^{C305F} mice is a p53-dependent phenomenon, *p53*^{+/-} and *p53*^{-/-} mice were fasted, after which the livers were harvested and analyzed for lipid droplet accumulation. As shown in Fig. 7A, livers from fasted *p53*^{+/-} and *p53*^{-/-} mice showed a mottled appearance reminiscent of the appearance of livers in fasted *Mdm2*^{C305F} mice. Oil red O staining of liver sections obtained from fasted *p53*^{+/-} and *p53*^{-/-} mice also resembled fasted *Mdm2*^{C305F} mouse livers with respect to the accumulation of lipid droplets (Fig. 7B). To determine whether there is an additive effect between the *Mdm2*^{C305F} mutation and p53 deletion, we bred *Mdm2*^{C305F} mice into *p53*^{+/-} and *p53*^{-/-} backgrounds to examine the effect of p53 deletion on fasting-induced hepatosteatosis. Fasted *Mdm2*^{C305F}; *p53*^{+/-} and *Mdm2*^{C305F}; *p53*^{-/-} mice exhibited a mottled appearance (Fig. 7C) and lipid droplet accumulation (Fig. 7D) that was similar to the fasted *Mdm2*^{C305F} mice, suggesting that the observed hepatosteatotic phenotype from fasted *Mdm2*^{C305F} mice is p53-dependent. This observation was further corroborated based on p53 target gene expression in mouse liver

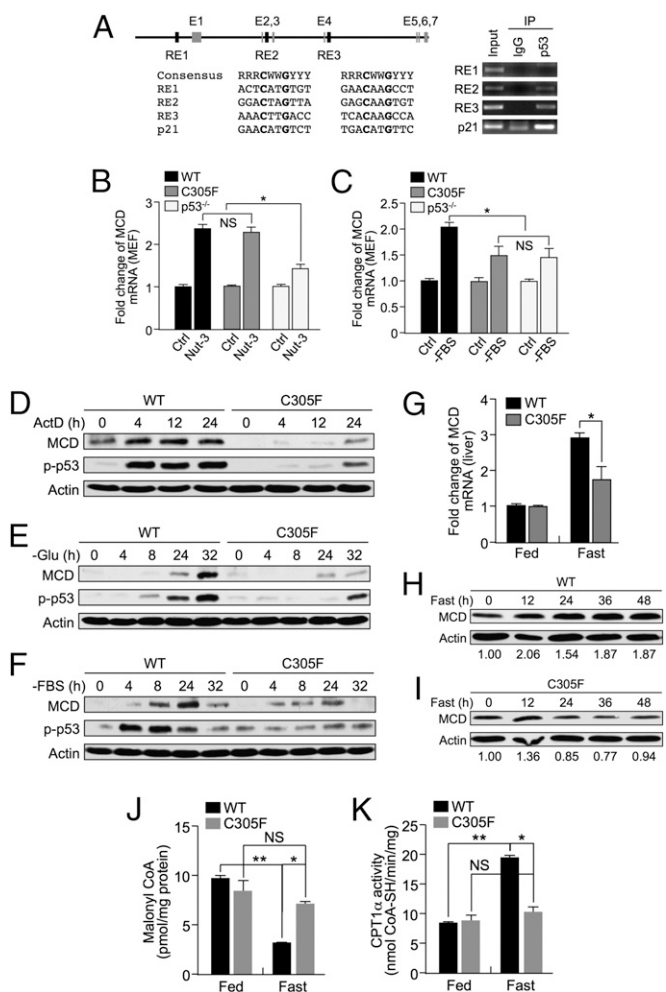


Fig. 6. Nutrient depletion induces RP-Mdm2-p53 pathway-dependent activation of MCD and augments CPT1 activity. (A) p53 binds to the *Mcd* gene promoter. (Left) Location and sequence of putative p53 response elements (RE1, RE2, and RE3). (Right) p53 binds to RE2 and RE3 as determined by a ChIP assay. The p53 response element found within the p21 promoter was included as a positive control. (B and C) Quantitative real-time-PCR (qRT-PCR) analysis of MCD from WT, *Mdm2*^{C305F}, and p53^{-/-} MEFs treated with (B) nutlin-3 (10 μ M) for 12 h or (C) serum starvation for 6 h. (D–F) Ribosomal biosynthesis stress activates p53 and induces MCD in MEF cells. WT and *Mdm2*^{C305F} MEFs were treated with (D) 5 nM actinomycin D, (E) glucose starvation (-Glu), or (F) serum starvation (-FBS) for the indicated time, and the total cell lysates were analyzed by immunoblotting. (G) qRT-PCR analysis of MCD mRNA expression in livers of WT and *Mdm2*^{C305F} mice fed a chow diet ad libitum (Fed) or fasted (Fast) for 12 h ($n = 3$ for each treatment). (H and I) WT and *Mdm2*^{C305F} mice were fasted for the indicated amount of time. Immunoblotting shows the MCD expression level in liver homogenates. Quantitative relative expression of MCD was normalized to β -actin. (J) Malonyl-CoA levels in livers isolated from WT and *Mdm2*^{C305F} mice ($n = 3$) fed a normal chow diet or fasted for 12 h were determined by HPLC and MS. (K) WT and *Mdm2*^{C305F} mice were fed a normal chow diet or were fasted for 12 h ($n = 3$). CPT1 α activity from liver homogenates was then measured spectrophotometrically. Error bars denote the SEM. *, **, and *** are used to indicate statistical significance corresponding to P values < 0.05 , 0.01 , and 0.001 , respectively. NS indicates no significant difference.

extracts. We observed a slight but noticeable increase in liver lipid staining in *Mdm2*^{C305F};p53^{+/-} and *Mdm2*^{C305F};p53^{-/-} livers compared with p53^{+/-} and p53^{-/-} livers, suggesting that there is a possible p53-independent function for the *Mdm2*^{C305F} mutation in the regulation of liver lipid metabolism. Transcript levels from the p53 target genes *Mdm2*, *Sco2*, and *Mcd* were significantly induced in

fasted WT mouse livers but were only slightly induced in livers from p53^{+/-}, p53^{-/-}, *Mdm2*^{C305F}, *Mdm2*^{C305F};p53^{+/-}, and *Mdm2*^{C305F};p53^{-/-} mice (Fig. 7 E–G). Importantly, the degree by which the gene expression of *Mdm2*, *Sco2*, and *Mcd* was reduced was similar among all five mutant mice when fasted, suggesting that reducing the p53 dosage by half is sufficient to attenuate much of the effect observed from fasting-induced p53 activation.

Discussion

Housekeeping Function for the RP-Mdm2-p53 Pathway in Energy Homeostasis. The results presented here demonstrate that, in mice, nutrient restriction elicits a p53 response that is mediated by the RP-Mdm2 interaction. Moreover, the RP-Mdm2 interaction is essential for regulating hepatic metabolism for efficient energy utilization. Starvation-mediated p53 induction requires an intact RP-Mdm2-p53 pathway, as demonstrated by a lack of p53 activity following nutrient deprivation in the RP-binding deficient *Mdm2*^{C305F} mutant. As a transcription factor, p53 serves as a point of convergence responsible for integrating a variety of upstream stress signals, including DNA damage, oncogene overexpression, hypoxic insults, and the inhibition of ribosome biosynthesis. Much attention has been devoted to dissecting how these upstream signals trigger p53 activation and how p53 modulates the expression of downstream genes to induce growth arrest, senescence, and apoptosis. Recent studies have pointed to the regulation of energy metabolism as a new theme for p53 function and have provided explanations to many previously puzzling observations. One such observation is that p53 is found almost throughout the animal kingdom, including in many ancient organisms predating the appearance of cancer (21). A plausible idea for the widespread presence of p53 in animals could be that p53 initially evolved as a regulator of energy homeostasis and later evolved to assume a role as a general stress sensor. In the present study, *Mdm2*^{C305F} mice are indistinguishable from WT mice in terms of growth and development; however, even under normal unstressed conditions, *Mdm2*^{C305F} mice clearly show altered basal metabolism (Fig. 1). We have found that p53^{-/-} MEFs accumulate more lipid droplets than WT MEFs when cultured in regular medium (Fig. S5A). Furthermore, when fed with normal chow diet, the livers from p53^{-/-} mice contain more lipid droplets than the livers from WT mice (Fig. S5B). Whether p53 acts as a stress sensor under day-to-day “unstressed” conditions depends on how “stress” is defined. We speculate that periodic feeding and fasting cycles in mice even fed ad libitum could generate small fluctuations in the rate of ribosome biosynthesis, resulting in the initiation of a signal that is relayed through the RP-Mdm2 interaction to p53. Oscillations in p53 activity can contribute to fine-tuning the metabolic response according to the energy requirements of the organism; fasting represents an extreme case of the normal energy fluctuation that occurs between meals. In the sense of responding to the daily feeding/fasting cycle, p53 can be viewed as a housekeeping gene responsible for maintaining the energy balance to favor fitness and survival.

RP-Mdm2-p53 Pathway Controls FAO Under Nutrient Shortage Stress. p53 regulates metabolism through a number of pathways; however, the complexity of p53-dependent metabolic regulation remains incompletely understood (22). Several studies have shown that p53 can reduce overall glycolytic function through mechanisms including the suppression of glucose transporters GLUT1/4 (23), the inhibition of phosphoglycerate mutase activity (24), and the induction of TIGAR (25). A recent study has shown that, in response to nutrient stress, p53 up-regulates FAO by inducing the expression of Lpin (26). To complement the function of inhibiting glycolysis, p53 increases OXPHOS through

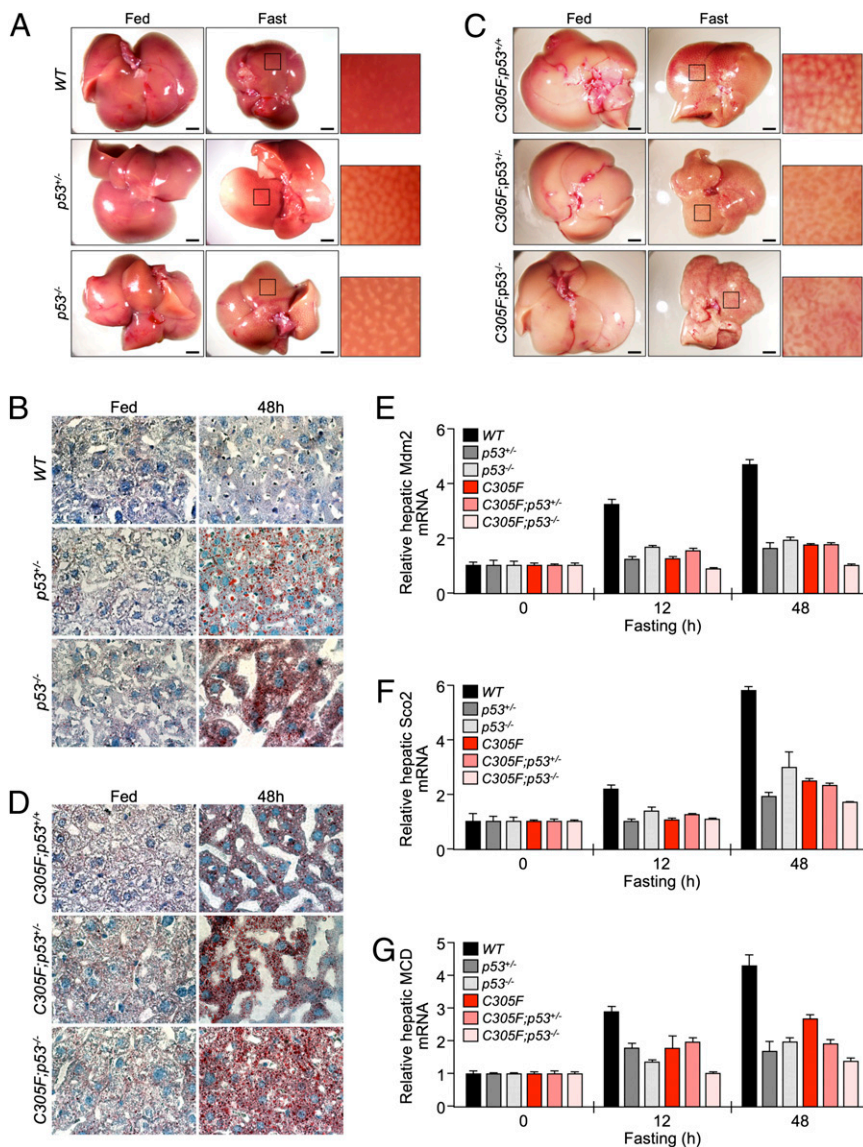


Fig. 7. Reduction of p53 dosage mimics hepatosteatotic phenotype observed in fasted *Mdm2*^{C305F} mice. (A and C) Representative images of livers harvested from mice with the indicated genotype fed a normal chow diet or fasted for 48 h. (Scale bars: 2.5 mm.) (B and D) Hepatic lipid droplets were visualized by oil red O staining on liver sections obtained from mice with the indicated genotypes fed a normal chow diet or fasted for the indicated amount of time. (E–G) qRT-PCR analysis of hepatic Mdm2 (E), SCO2 (F), and MCD (G) mRNA expression in mice with indicated genotypes fed a chow diet *ad libitum* (Fed) or fasted (Fast) for the indicated amount of time ($n = 3$ for each treatment). Error bars denote the SEM.

the transactivation of SCO2, an essential component of mitochondrial complex IV (16), and glutaminase 2, which promotes ATP production and suppresses oxidants (27, 28). Mitochondrial FAO plays a critical role in generating ATP when the glucose supply is low. Given the overall protective function of p53 in an organismal context, it is not surprising that p53 promotes FAO during glucose starvation (29). By using the *Mdm2*^{C305F} mouse model, we have shown that the RP–Mdm2–p53 pathway plays a central role in sensing nutrient depletion stress and in modulating energy output. Based on our results, we propose a model in which changes in nutrient availability impinge on ribosome biosynthesis, leading to an increase in RP–Mdm2 interaction and subsequent activation of p53. By inducing the expression of MCD, a key enzyme for malonyl-CoA turnover and CPT1 α activation, p53 promotes long-chain fatty acid uptake by the mitochondria. Together with the induction of SCO2, p53 promotes FAO and maintains metabolic homeostasis during nutrient shortage stress (Fig. 8).

The present study focuses on defining how nutrient deprivation influences cellular responses through the RP–Mdm2–p53 pathway. However, overnutrition represents another type of nutrient stress, and whether the RP–Mdm2–p53 pathway plays a role during nutrient overload is an interesting question for future studies. Given the emergence of obesity as a worldwide epidemic, it seems equally imperative to determine the cellular stress responses associated with nutrient overabundance. Future research focused on clarifying the role of the RP–Mdm2–p53 pathway as a metabolic stress response pathway and additional roles that this pathway might play in the p53-dependent modulation of nutrient partitioning and energy production provides an exciting opportunity to expand upon the myriad roles of p53 in regulating cellular homeostasis.

Experimental Procedures

Animals and Treatments. WT and *Mdm2*^{C305F} mice were bred and maintained in standard cages on a 12:12 h light/dark cycle with *ad libitum* access to regular

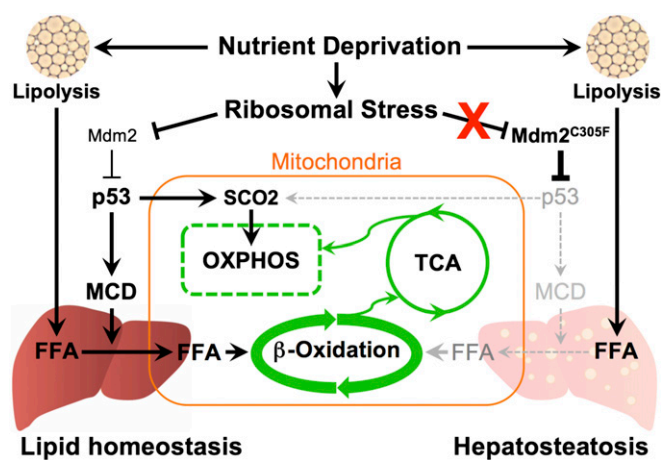


Fig. 8. RP–Mdm2–p53 pathway forms a crucial link between nutrient deprivation and cellular energy homeostasis. Nutrient deprivation induces lipolysis in adipocytes generating free fatty acids (FFAs), which enter the liver to serve as a direct energy source or as a precursor for ketone body formation. In normal mouse liver, nutrient depletion causes ribosomal stress by inhibiting rRNA biosynthesis, which in turn inhibits Mdm2 (indicated by the smaller letters) and activates p53 to induce MCD expression. MCD facilitates the transport of FFAs into the mitochondrial matrix through CPT1 α to be oxidized to acetyl-CoA by β -oxidation. Acetyl-CoA is then oxidized via the TCA cycle to produce energy and “reducing equivalents” (NADH and FADH₂), which are used for ATP synthesis by OXPHOS. p53 also induces the expression of SCO2, a crucial component of the electron transport chain, to promote OXPHOS. Together, the action of the RP–Mdm2–p53 pathway maintains liver lipid homeostasis during the fasting state. However, in *Mdm2*^{C305F} mouse livers, ribosomal stress in the form of nutrient deprivation cannot activate p53 (grayed out), which results in the accumulation of FFAs in the liver and hepatosteatosis.

chow and water. All mice were handled in strict accordance with our protocol (10-045) as approved by the institutional animal care and use committee at the University of North Carolina Animal Care Facility. Body weight was measured weekly. All mice used for experiments were between 9 and 11 wk of age unless otherwise specified. For the fasting studies, mice were housed individually with ad libitum access to water for 12 h (7:00 PM to 7:00 AM), 24 h, or 48 h.

Metabolic Analysis. Body water, body fat, and lean mass were analyzed by using an EchoMRI-100 quantitative magnetic resonance whole-body composition analyzer (Echo Medical Systems) according to the instructions from the manufacturer.

For metabolic studies, mice were placed individually in the metabolic chambers of an Oxymax system (Columbus Instruments) for 120 h, and readings were taken after a 24-h acclimation period. Physical activity, food consumption, OCR (VO₂), carbon dioxide production rate (VCO₂), and respiratory exchange ratio were obtained. VO₂, VCO₂, and respiratory exchange ratio were measured every 26 min during a 48-h period before or after fasting and were normalized to the body weight. Activity was measured on x, y, and z axes by counting interruptions in the infrared beams.

Hepatic TAG and β -HB (Stanbio) levels were measured by using colorimetric commercial kits according to the manufacturer’s instructions. Malonyl-CoA content was assessed from whole liver extracts. After separation, purification, and elution, lipid metabolite extracts were separated by HPLC, and individual and total lipid species were analyzed by liquid chromatography and tandem MS.

Metabolomic Analysis. Freshly isolated mouse livers were used for metabolomic analysis. *SI Experimental Procedures* provides a more detailed protocol.

Mitochondrial Activity and Glycolytic Flux. OCR and ECAR were evaluated by using a Seahorse XF24 Analyzer (Seahorse Bioscience). For MEFs, 6×10^4 cells were seeded 24 h before the assay. The basal level of OCR and ECAR were measured in XF assay medium containing 4.5 g/L glucose followed by the addition of oligomycin (1 μ M), which allowed for the assessment of the maximal glycolytic capacity (i.e., ECAR) or by the sequential addition of oligomycin (1 μ M), FCCP (0.5 μ M; maximal OXPHOS), and antimycin A (2 μ M)

plus rotenone (1 μ M) to determine the OCR. For animal experiments, mitochondria were isolated from mouse livers, and 7.5 μ g of mitochondria was transferred into each well of an XF24 plate immediately after extraction. Then, the OCR and ECAR were measured at basal levels in the presence of succinate (5 mM) as well as after the sequential addition of ADP (100 μ M), oligomycin (1 μ M), FCCP (0.5 μ M), and antimycin A (2 μ M). Mitochondrial respiration control ratio (RCR) was evaluated based on the OCR-FCCP/OCR-oligomycin ratio.

Oil Red O Staining. Mouse liver tissues were fixed in 10% (vol/vol) neutral formalin for 24 h and then were submerged in 20% sucrose for 2–3 d. The liver pieces were then frozen in optimal cutting temperature compound, and 5- μ m-thick tissue sections were cut. Oil red O staining was conducted according to the standard protocol used in the Histology Research Core Facility at the University of North Carolina (UNC).

Fatty Acid β -Oxidation in Liver. Freshly isolated liver was homogenized with a motor-driven Teflon pestle and glass mortar in ice-cold SET buffer (250 mM sucrose, 1 mM EDTA, 10 mM Tris-HCl, and 2 mM ATP, pH 7.4). Two milligrams of each liver homogenate was incubated with BSA-conjugated palmitate (200 μ M sodium palmitate plus 8 μ M [1-¹⁴C]palmitate at a final concentration of 0.1 μ Ci per reaction mixture) in a mixture containing 125 mM sucrose, 24 mM K₂HPO₄, 200 mM KCl, 2.5 mM MgCl₂, 2.5 mM L-carnitine, 0.25 mM malic acid, 20 mM Tris-HCl, 2.5 mM DTT, 0.25 mM NAD⁺, 4 mM ATP, and 0.125 mM CoA (pH 7.4) at 37 $^{\circ}$ C for 30 min in a customized two-well system: one well contained the reaction mixture with the liver homogenates, and the adjoining well contained 1 N NaOH. The reaction was terminated by adding 70% perchloric acid to the assay well, and then the plate was incubated for 1 h to drive the CO₂ into the NaOH-containing well. Radioactivity of acid-soluble metabolites in the supernatant of the reaction mixture and CO₂ was determined by liquid scintillation. FAO was quantified by using the following formula: [(dpm – BL)/SA]/[gram of liver (wet weight) \times time (in hours) of reaction mixture incubation], where dpm stands for disintegrations per minute, BL represents the disintegrations per minute of the blank wells, and SA is the fatty acid-specific radioactivity.

Determination of CPT1 Activity. The activity of CPT1 was measured spectrophotometrically by using a modified version of a previously described method (30). In brief, a 50-mg liver piece was homogenized in a buffer (0.25 M sucrose, 1 mM EDTA, and 10 mM Tris-HCl, pH 7.5) supplemented with protease inhibitor and PMSF, followed by centrifugation at 300 \times g at 4 $^{\circ}$ C for 10 min, after which the supernatant was further purified by a second centrifugation at 12,000 \times g at 4 $^{\circ}$ C for 15 min to pellet the mitochondria. Fifty microliters of the precleared supernatant was mixed with 50 μ L of Tris-5,5'-dithio-bis-(2-nitrobenzoic acid) buffer [116 mM Tris, 2.5 mM EDTA, 2 mM Tris-5,5'-dithio-bis-(2-nitrobenzoic acid), and 0.2% Triton X-100, pH 8.0] with or without malonyl-CoA (10 μ M) in individual wells of a 96-well plate. The mixture was preincubated for 5 min at room temperature, and then 50 μ L of 1 mM palmitoyl-CoA was added to the wells. The reaction was initiated by adding 5 μ L of L-carnitine solution (1.2 mM dissolved in 1 M Tris, pH 8.0), and the release of coenzyme A (CoA-SH) from palmitoyl-CoA was followed spectrophotometrically at 412 nm for 180 s by using a SpectraMax M5 Multi-Mode Microplate Reader (Molecular Devices). A CoA-SH (Sigma-Aldrich) standard curve (1–200 nmol per well) was generated. The activity of CPT1, defined as the amount of nmol of CoA-SH released per minute per milligram protein, was calculated by subtracting CPT2 activity (with malonyl-CoA) from the total CPT activity (without malonyl-CoA).

Microarray Analysis. Microarray analysis was performed as previously described (31). Briefly, total RNA was isolated from MEFs that were untreated or were treated with 4-hydroxytamoxifen for 24 h by using an RNeasy kit (Qiagen). Total RNA was amplified and labeled, and microarray hybridization was performed at the UNC genomics and bioinformatics core facility. Genes that were significantly up- or down-regulated were identified by using significance analysis of microarrays.

ChIP Assay. A ChIP assay was performed by using the QuikChIP kit (IMGENEX) according to the manufacturer’s suggested protocol. Briefly, MEFs were seeded in 10-cm dishes at a density of 1.5×10^4 cells per square centimeter. Twelve to twenty-four hours after plating, cells were cross-linked with formaldehyde (1%) at 37 $^{\circ}$ C for 10 min and incubated with glycine (0.125 M) for 5 min. Cells were washed with ice-cold PBS solution and lysed in SDS lysis buffer. Cell lysates were sonicated four times for 10 s each time with a Branson Sonifier. Insoluble material was removed by centrifugation at 12,000 \times g at 4 $^{\circ}$ C for 10 min. Chromatin extracts were precleared with

salmon sperm DNA/protein A/G agarose and then were incubated with a goat polyclonal antibody specific for p53 (FL393; Santa Cruz) or IgG control (Life Technologies) overnight at 4 °C. Reactions were then incubated for 1 h with salmon sperm DNA/protein A/G agarose beads. The beads were then washed several times, and the chromatin complex was eluted with the provided elution buffer. The eluates were further incubated at 65 °C overnight in the presence of 200 mM of NaCl to reverse the crosslinks followed by incubation with an RNase A solution and then a proteinase K solution. The DNA was purified by phenol/chloroform extraction. PCR was then performed in the presence of 5% DMSO. The primers used for the PCR amplification step were as follows: RE1 forward, 5'-CTGGACTGGCTGCTCTGTCC-3'; RE1 reverse, 5'-CTCTGCCTCGCTTCCGCTC-3'; RE2 forward, 5'-GTGCACTCCAGGAAGCCCTAG-3'; RE2 reverse, 5'-CTCTGCTCGGCTTCC-3'; RE3 forward, 5'-CCTTACAGGCCAAGCAAACACTTGAG-3'; RE3 reverse, 5'-GTGATAGCAGGACTCCCGAATAC-3'; p53 (p21^{Cip1}) forward, 5'-GAGACCAGCAGCAAAATCC-3'; p53 (p21^{Cip1}) reverse, 5'-CAGCCCCACCTCTCAATC-3'.

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Statistical Analysis. Data were analyzed by using Graph Pad 5.0 software. Repeated-measures ANOVA was performed to evaluate the differences in O₂ consumption, CO₂ production and heat generation between WT and *Mdm2*^{C305F} mice (SAS 9.3; SAS Institute). *P* values < 0.05 were considered statistically significant.

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