Up-regulation of FOXO1 and reduced inflammation by β-hydroxybutyric acid are essential diet restriction benefits against liver injury

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Liver ischemia and reperfusion injury (IRI) is a major challenge in liver surgery. Diet restriction reduces liver damage by increasing stress resistance; however, the underlying molecular mechanisms remain unclear. We investigated the preventive effect of 12-h fasting on mouse liver IRI. Partial warm hepatic IRI model in wild-type male C57BL/6 mice was used. The control ischemia and reperfusion (IR) group of mice was given food and water ad libitum, while the fasting IR group was given water but not food for 12 h before ischemic insult. In 12-h fasting mice, serum liver-derived enzyme level and tissue damages due to IR were strongly suppressed. Serum β-hydroxybutyric acid (BHB) was significantly raised before ischemia and during reperfusion. Up-regulated BHB induced an increment in the expression of FOXO1 transcription factor by raising the level of acetylated histone. Antioxidative enzy-Name: ume heme oxygenase 1 (HO-1), a target gene of FOXO1, then increased. Autophagy activity was also enhanced. Serum high-mobility group box 1 was remarkably lowered by the 12-h fasting, and activation of NF-κB and NLRP3 inflammasome was suppressed. Consequently, inflammatory cytokine production and liver injury were reduced. Exogenous BHB administration or histone deacetylase inhibitor administration into the control fed mice ameliorated liver IRI, while FOXO1 inhibitor administration to the 12-h fasting group exacerbated liver IRI. The 12-h fasting exerted beneficial effects on the prevention of liver IRI by increasing BHB, thus up-regulating FOXO1 and HO-1, and by reducing the inflammatory responses and apoptotic cell death via the down-regulation of NF-κB and NLRP3 inflammasome.

Significance

Long-term fasting for more than 24 h has been reported to ameliorate liver ischemia and reperfusion injury (IRI). However, it is difficult to apply long-term fasting for human preoperative management, and the underlying molecular mechanisms remain unclear. Our present research demonstrates that 12-h fasting markedly ameliorates liver IRI. The 12-h fasting induces up-regulation of FOXO1 by raising the level of acetylated histone and β-hydroxybutyric acid, followed by up-regulation of antioxidant enzymes and autophagy activity, and improves liver IRI through the reduction of the inflammation and apoptotic cell death. Perioperative administration of β-hydroxybutyric acid or histone deacetylase inhibitor may have beneficial effects by avoiding liver injury at liver surgery. This is an insight on the prevention of liver IRI.
Diet restriction also exerts a protective effect against IRI in several organs (19). Starvation for 48 h to 72 h reduced liver IRI by up-regulating antioxidant enzymes or autophagy (20, 21). In addition, fasting for 1 d, but not 2 d or 3 d, can prevent mouse liver IRI via the Sirtuin1-mediated down-regulation of circulating HMGB1 (22). We report here that 12-h fasting can remarkably suppress mouse liver IRI. This study aimed to clarify the effect of the 12-h fasting against liver IRI and the underlying molecular mechanisms.

Results
Preoperative 12-h Fasting Protects Against Ischemia and Reperfusion-Triggered Hepatocellular Damage. After ischemia and reperfusion (IR), serum liver-derived enzyme, alanine aminotransferase (sALT), levels markedly increased in the control fed mice. However, sALT remained significantly lower in the 12-h fasting group (Fig. 1 A and SI Appendix, Fig. S1B), indicating that 12-h fasting can ameliorate IR-induced liver damage. The livers of the 12-h fasted mice exhibited clear reduction in the hepatocellular necrosis caused by IR (Fig. 1 B, 1). Moreover, Suzuki’s score was significantly lowered for the 12-h fasting group than for the control fed mice (Fig. 1 B, 2). During liver IR, leukocytes infiltrate into the liver tissues and produce inflammatory cytokines, which lead to hepatocyte apoptosis and necrosis. The numbers of T cells (CD3), neutrophils (Ly-6G), and macrophages (CD68) infiltrating into the liver tissue after IR treatment were significantly lesser in the 12-h fasted mice (SI Appendix, Fig. S2). Terminal deoxynucleotidyl transferase-mediated dUTP nick end labeling (TUNEL)-positive cells induced by IR were significantly reduced in the liver of the 12-h fasting group compared with those of control fed mice (Fig. 1 C, 7 and 2), indicating that the 12-h fast clearly reduced liver cell injury caused by IR. Moreover, the 12-h fasting suppressed the expression of cleaved caspase-3, while it significantly increased the expression of antiapoptotic protein B cell lymphoma 2 (Bcl-2) in the liver (Fig. 1 D and E).

Preoperative 12-h Fasting Suppressed the Secretion of Proinflammatory Cytokines and Damage-Associated Molecular Patterns Induced by IR. Damage-associated molecular patterns (DAMPs) strongly influence the induction of inflammation (23, 24). HMGB1, a DAMP, is released into the extracellular space from the necrotic cells and stimulates immune cells (8, 23). Serum HMGB1 level increased during ischemia and was rapidly boosted following reperfusion in the controls. In contrast, the serum HMGB1 level was significantly suppressed during ischemia and reperfusion period in the 12-h fasting group (Fig. 2 A). HMGB1 activates the inflammatory cells in the liver through TLR-4 and induces the production of proinflammatory cytokines that are crucial for the development of liver IRI (25). In the inflammatory process of hepatic IRI, inflammatory cytokines, such as interleukin-6 (IL-6), tumor necrosis factor α (TNFα), and IFNγ are released from the macrophages and lymphocytes (25). Serum levels of IL-6, IL-18, TNFα, IFNγ, and IL-1β rapidly increased during reperfusion, and returned to the baseline level at 12 h after reperfusion, while the serum levels of all of these proinflammatory cytokines were

![Fig. 1. Hepatocellular damage induced by IR in the controls and the 12-h fasting mice. (A) The sALT levels were measured. Means and SD are shown (n = 8 mice per group; two-way ANOVA; P < 0.0001, Bonferroni’s posttest: ***P < 0.001 vs. control mice). (B-1) Liver histology (H & E staining) after IR and sham-operated (magnification 200x; in box, 400x). (B-2) Suzuki’s histological score after IR. Means and SD are shown (n = 8 mice per group; ***P < 0.001). (C-1) TUNEL-assisted detection of hepatic apoptosis after IR (magnification 200x; in box, 400x). (C-2) Quantification of TUNEL positive cells. Means and SD are shown (n = 8 mice per group; ***P < 0.001). (D) Western blot-assisted expression of cleaved caspase-3, Bcl-2 at 6 h of reperfusion. (E) Quantification of Western blot bands shown in D. Means and SD are shown (*P < 0.05).]

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significantly suppressed in the 12-h fasting group (Fig. 2B). Furthermore, the gene expressions of these proinflammatory cytokines in the liver tissue were also significantly suppressed in the 12-h fasting group compared with the control fed mice (Fig. 2C). These results clearly indicate that pretreatment with 12-h fasting can ameliorate liver IRI via suppression of HMGB1 and inflammatory cytokines induced by IR.

**Suppression of NF-κB and NLRP3 Induced by 12-h Fasting Before Treatment.** NF-κB is a protein complex that controls cytokine production and cell survival (26), also playing a key role in regulating the immune response against inflammation (27). NF-κB heterodimer comprises p65 and p50 proteins. In an inactivated state, NF-κB locates in the cytosol complexed with the inhibitory protein IκBα as the nonphosphorylated form. The amounts of phosphorylated IκBα (p-IκBα) reflect the activation state of NF-κB (26, 28). The p-IκBα expression at 1 h and more after reperfusion was significantly lower, while the amounts of cytosol NF-κB inversely increased in the 12-h fasting group compared with that in the control fed mice (Fig. 3A), indicating that the 12-h fasting clearly suppressed NF-κB activation in relation to the reduction in serum HMGB1. NLRP3 inflammasome also contributes to the early inflammatory phase and regulates inflammatory response (29). The NLRP3 expression was significantly decreased in the 12-h fasting group at early phase of IR (1 h of reperfusion) compared with that in the control fed mice (Fig. 3B). The reduced

Fig. 2. Preoperative fasting suppressed the release of HMGB1 and proinflammatory cytokines. (A) Serum HMGB1 levels were measured. Means and SD are shown (n = 7 mice per group; two-way ANOVA: P < 0.0001; Bonferroni’s posttest: **P < 0.01, ***P < 0.001 vs. control mice). (B) Serum levels of proinflammatory cytokines (IL-6, TNFα, IL-1β, IL-18, and IFNγ) were measured. Means and SD are shown (n = 8 mice per group; two-way ANOVA; IL-6, P < 0.0001; TNFα, P < 0.0001; IFNγ, P < 0.001; IL-1β, P < 0.0001; IL-18, P < 0.001; Bonferroni’s posttest: **P < 0.01, ***P < 0.001 vs. control mice). (C) Quantitative PCR detection of proinflammatory cytokines (IL-6, TNFα, IL-1β, and IFNγ) was performed at 3 h of reperfusion after 60 min of ischemia. Data were normalized to HPRT gene expression. Means and SD are shown (sham, n = 4; IR 3 h, n = 8 mice per group; *P < 0.05; **P < 0.01; ***P < 0.001).

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Fig. 3. Preoperative fasting protected liver from injury through the up-regulation of FOXO1 and suppressed NF-κB and NLRP3 signaling induced by IR. (A) Western blot-assisted analyses of the total and phosphorylated protein levels of FOXO1, IκBα and acetylated histone-3, histone-3, FOXO3a, HO-1, LC3B, and NLRP3 before ischemia and during reperfusion. β-actin was used as the internal control. (B) Representative of immunohistochemistry of FOXO1 (dark brown) at 6 h of reperfusion (magnification 400×). (C) Representative of immunofluorescence staining of FOXO1 (green) and DAPI (blue) at 6 h of reperfusion (magnification 400×). (D) Representative immunohistochemistry of 4-HNE (dark brown) at 6 h of reperfusion (magnification 200×; in box, 400×). (E) The sALT levels were measured for mice pretreated with FOXO1 inhibitor and the 12-h fasting mice pretreated with DMSO only. Means and SD are shown (n = 8 mice per group; **P < 0.01). (F) Serum levels of proinflammatory cytokines (IL-6, TNFα, IL-1β, and IFNγ) were measured at 6 h of reperfusion. Means and SD are shown (sham, n = 4; IR 6 h, n = 8 mice per group; *P < 0.05; **P < 0.01; ***P < 0.001). (G) Western blot-assisted expression of HO-1 and LC3B at 6 h of reperfusion.
expression of nuclear NF-κB and the reduction of NLRP3 from the early phase of reperfusion in the mice pretreated with the 12-h fasting may strongly suppress inflammatory cytokines, resulting in IRI amelioration.

Preoperative 12-h Fasting Protects Liver from Oxidative Stress through the Up-Regulation of FOXO1. FOXO transcription factor is an important target of the glycolytic pathway (30) that regulates gluconeogenesis and β-oxidation of fatty acids in the liver (31). Furthermore, FOXO1 is involved in the transcriptional regulation of antioxidant enzymes and in autophagy regulation (32, 33). FOXO1/3 expression in the liver of the 12-h fasted mice was higher than that in the control fed mice before ischemia, particularly during reperfusion. It was increased from the beginning of reperfusion and peaked at the later phase (3 h to 6 h), while the FOXO1/3 levels in the liver of the controls remained lower during reperfusion. The antioxidative enzymes, heme oxygenase 1 (HO-1), a target gene of FOXO1 (32), also clearly increased in the 12-h fasting group before ischemia and during the later phase (3 h to 6 h) of reperfusion. However, such an increase was not observed in the control fed mice (Fig. 3A). Autophagosome membrane protein LC3B, the marker of autophagy and a target of FOXO1 (33), was up-regulated from the early phase of reperfusion in the liver of the 12-h fasting group but not in the control fed mice (Fig. 3A). Immunohistochemistry (Fig. 3B) and immunofluorescence staining (Fig. 3C) of the liver tissues of the 12-h fasting group revealed that most FOXO1 positive cells were localized in the parenchymal and endothelial cells. These positive cells were significantly higher in the 12-h fasting group than in the controls 6 h after reperfusion (Fig. 3 B and C).

In the development of liver IRI, ROS are generated and induce oxidation of local macromolecules. Lipid peroxidation is essential to assess oxidative stress, and 4-hydroxynonenal (4-HNE) in the liver is an indicator of lipid peroxidation (34). Immunohistochemistry staining of 4-HNE in the liver tissues revealed clear reduction of lipid oxidation in the 12-h fasted mice (Fig. 3D), suggesting that the 12-h fasting could ameliorate oxidative stress in liver caused by IRI.

The i.p. administration of FOXO1 inhibitor (AS1842856) significantly exacerbated liver IRI in case of the 12-h fasting group (Fig. 3E and SI Appendix, Fig. S1C). Serum levels of IL-1γ, IL-6, and TNFα were also increased upon administration of FOXO1 inhibitor at 6 h after reperfusion (Fig. 3F). The up-regulated expressions of antioxidative enzyme, HO-1, and autophagosome protein, LC3B, which are induced by 12-h fasting, were suppressed by the FOXO1 inhibitor at 6 h after reperfusion (Fig. 3G). Conversely, the effect of FOXO1 inhibitor against liver IRI was not observed in the control fed mice (Fig. 3E). These results indicate that FOXO1 plays a preventive role against liver cell damage caused by IR via the up-regulation of antioxidative responses and autophagy.

Fig. 4. BHB controls the benefits of 12-h fasting. (A) Serum BHB levels were measured in control fed mice and 12-h fasting mice. Means and SD are shown (n = 8 mice per group; two-way ANOVA: P < 0.0001; Bonferroni’s posttest: ***P < 0.001 vs. control fed mice). (B) After the administration of BHB or PBS into the fed mice, serum BHB levels were measured. Means and SD are shown (n = 8 mice per group; two-way ANOVA: P < 0.0001; Bonferroni’s posttest: **P < 0.01, ***P < 0.001 vs. PBS-administered mice). (C) The sALT levels were measured. Means and SD are shown (n = 8 mice per group; ***P < 0.001). (D) After the administration of BHB or PBS, serum levels of proinflammatory cytokines (IL-6, TNFα, IL-1β, and IFNγ) were measured. Means and SD are shown (sham, n = 4; IR 6 h, n = 8 mice per group; *P < 0.05; **P < 0.01). (E-1) Liver histology (H & E staining) after IR (magnification 200×). (E-2) Suzuki’s histological score after IR. Means and SD are shown (n = 8 mice per group; **P < 0.01). (F) Western blot-assisted analyses of the FOXO1, HO-1 acetylated histone-3, and NLRP3.
β-Hydroxybutyric Acid Controls the Benefits of 12-h Fasting. In the animal body, fats are metabolized to acetocetate and β-hydroxybutyric acid (BHB) as sources of energy when the glycolytic pathway does not function (35). The serum BHB levels were significantly increased even before ischemia in the 12-h fasting group (Fig. 4A). The serum BHB levels slightly increased in the control fed mice 3 h to 12 h after reperfusion. However, the serum BHB level in the 12-h fasting group increased even after ischemia, and at 3 h of reperfusion. It was markedly up-regulated at 6 h of reperfusion. These results strongly suggest that the up-regulation of BHB may be a crucial event for preventing liver IRI in 12-h fasting mice.

To examine the effect of BHB on liver IRI, the control fed mice received i.p. administration of BHB 30 min before ischemia treatment (SI Appendix, Fig. S1E). After BHB administration, the serum BHB levels increased significantly and remained at higher levels during IR than that of the PBS-administered mice (Fig. 4B). The sALT levels and serum IL-1β levels were significantly lowered in the mice administered BHB at 6 h of reperfusion (Fig. 4C and D). The livers of the BHB-administered mice exhibited histological reduction in hepatocellular necrosis after IR treatment, and Suzuki’s score improved significantly in the BHB-administered mice compared with the PBS-administered mice (Fig. 4E, I and 2).

The expression of FOXO1 is up-regulated through the increment in acetylated histone (36). Thus, the FOXO1 expression is negatively regulated by histone deacetylase (HDAC). BHB has an endogenous HDAC inhibitory activity at a low concentration level (1 mM) (37). BHB induced histone acetylation at the promoter of oxidative stress resistance genes FOXO by inhibiting HDACs classes I and II (37, 38). Western blotting analyses showed that the expressions of FOXO1 and acetylated histone-3 markedly increased in the 12-h fasting group from the beginning of reperfusion (Fig. 3A). These results suggest that the FOXO1 expression in the liver was up-regulated owing to the increased amounts of acetylated histone induced by increased BHB activity; in turn, increased BHB was stimulated by the 12-h fasting. Exogenous BHB also induced acetylated histone-3, which resulted in higher expressions of FOXO1, followed by HDAC inhibition (40), at 16 h before IR treatment and just before IR treatment (SI Appendix, Fig. S1D). The sALT levels after reperfusion were significantly lowered in the HDACi administered fed mice (Fig. 3A). Serum levels of IL-6, TNFα, and IL-1β were also significantly suppressed in the HDACi administered fed mice (Fig. 3B). The expressions of FOXO1, HO-1, and LC3B were reversely up-regulated by the HDACi administration at 6 h of reperfusion (Fig. 3C).

These results indicate that the improvement in liver IRI induced by the 12-h fasting was partly owing to the enhancement of FOXO1 expression as well as the suppression of NLRP3, both of which were induced by the up-regulation of serum BHB, followed by the suppression of HDAC activity.

Addition of Glucose to the Drinking Water Reversed the Preventive Effects on Liver IRI of the 12-h Fasting Group. When glycolysis converts glucose to pyruvate, BHB is not produced in the animal body. We examined the effect of 10% glucose water on liver IRI in the 12-h fasting group (SI Appendix, Fig. S1B). The serum BHB level clearly increased during ischemia and reperfusion in the 12-h fasting group that was given only water but without glucose (Fig. 4A). However, in mice of the 12-h fasting group that was given 10% glucose water, the serum BHB level was strongly lowered (Fig. 4A). Amelioration of liver injury (measured by the elevation of sALT) was cancelled in mice fed with 10% glucose water (Fig. 6B). The i.p. administration of BHB inversely showed the protective effect against liver IRI in the mice of the 12-h fasting given water containing 10% glucose (Fig. 6B). Serum levels of IFNγ and TNFα at 6 h of reperfusion were significantly elevated in 12-h fasting mice given water containing 10% glucose (Fig. 6C). Increased expressions of FOXO1 and HO-1 in the 12-h fasting group at 6 h of reperfusion were markedly suppressed by the feeding with water containing 10% glucose (Fig. 6D).

Discussion
Long-term diet restriction without malnutrition improved stress resistance and extended lifespan (41); however, the underlying
are released from the infiltrated im-

Schematic summary of the present study.

Fig. 6. Addition of glucose to drinking water reversed the preventive effects of the 12-h fast on liver IRI. (A) Serum BHB levels were measured in 12-h fasting mice fed with water only or 10% glucose water. Means and SD are shown (n = 8 mice per group; two-way ANOVA; *P < 0.0001, Bonferroni’s posttest: ***P < 0.001 vs. 12-h fasting without glucose). (B) The ALT levels were measured in the control fed mice, 12-h fasting mice, and 12-h fasting mice fed with 10% glucose water. Means and SD are shown (n = 8 mice per group; *P < 0.05; **P < 0.01; ***P < 0.001). (C) Serum levels of proinflammatory cytokines (IL-6, TNFα, IL-1β, and IFNγ) were measured. Means and SD are shown (sham, n = 4; IR 6 h, n = 8 mice per group; *P < 0.05; **P < 0.01). (D) Western blot-assisted analyses of FOXO1 and HO-1.

Fig. 7. Schematic summary of the present study.
The NLRP3 inflammasome-mediated cell pyroptosis promotes HMGB1 secretion (45), and HMGB1 release is partly dependent on NLRP3 inflammasome (46). The NLRP3 expression in the 12-h fasting mice was significantly lowered before the ischemic phase (Fig. 3A), suggesting that the down-regulation of the NLRP3 inflammasome may induce suppression of HMGB1 at the early phase of IR in the 12-h fasting group. Thus, the enhanced expression of FOXO1 and HO-1, with reduced expression of NLRP3, may suppress HMGB1 release during ischemia through-out reperfusion in the 12-h fasting group.

It has been shown that the administration of BHB protected various organs from IRI (47, 48); however, the underlying molecular mechanisms are unclear. Recently, BHB has been reported to exert an inhibitory activity against NLRP3 inflammasome and prevent NLRP3-mediated inflammatory diseases (39). Activated NLRP3 inflammasome induces IL-1β and IL-18, and innate immune responses (49, 50), and promotes the maturation and secretion of proinflammatory cytokines from immune cells (28). Serum BHB levels were higher in the 12-h fasting group even before ischemia and during reperfusion (3 h to 12 h) (Fig. 4A). In contrast, the liver expressions of NLRP3 were suppressed during ischemia (before reperfusion) and at 1 h after reperfusion (Fig. 3A). Serum IL-1β and IL-18 levels were significantly lower during reperfusion (3 h to 6 h) in the 12-h fasting group (Fig. 2B). Thus, the enhanced expression of BHB may reduce inflammation by suppressing NLRP3 inflammasome. Exogenous BHB also suppressed the expression of NLRP3 at 1 h of reperfusion (Fig. 4F), and serum IL-1β levels at 6 h after reperfusion were also significantly suppressed (Fig. 4D). These results indicate a crucial role of BHB in suppressing liver inflammation and ameliorating liver IRI. BHB also causes the reduction of serum HMGB1 through the suppression of NLRP3 in the 12-h fasting group.

The serum BHB is known to increase during prolonged exercise or starvation (6 mM to 8 mM) and diabetic ketoacidosis (>25 mM) (51). BHB is known to display an endogenous HDAC inhibitory activity from a low concentration level (1 mM) (37). The expression of FOXO1 is reportedly up-regulated through the increment in acetylated histone (36, 38). The serum BHB concentrations in the 12-h fasting mice were 0.8 mM to 1.7 mM during reperfusion, much higher than in the control fed mice (Fig. 4A). Expression of acetylated histone-3 were up-regulated in the 12-h fasting group (Fig. 3A). These data indicate that the up-regulation of FOXO1 was induced through the increment of acetylated histone-3, which was accelerated by increased BHB. Exogenously administered BHB or HDACi also induced acetylated histone-3, leading to the up-regulation of FOXO1 and HO-1 (Fig. 4F and 5C). Thus, BHB-mediated enhancement of FOXO1 and then HO-1 are crucial events for the amelioration of IRI in the 12-h fasting group.

Autophagy is an intracellular self-digesting pathway responsible for removing long-lived proteins, damaged organelles, and malformed proteins. Autophagy exerts a protective effect against cell apoptosis (52). The inhibition of autophagy reportedly exacerbates liver IRI (52–54). Autophagy is regulated by FOXO1 and HO-1 (33). The expression of LC3B, an autophagosome membrane marker, was up-regulated in the 12-h fasting group (Fig. 3A). The up-regulation of autophagy induced by 12-h fasting through the increased expression of FOXO1 may also play an important role in suppression of liver IRI.

FOXO1 is regulated not only through acetylation but also through phosphorylation (55). The activation of FOXO1 is reported to be primarily regulated through phosphorylation by the insulin–Phosphoinositide 3-kinase (PI3K)–Akt pathway (55). Activated Akt (p-Akt) inhibits the activity of FOXO1 transcription factors via phosphorylation, resulting in nuclear exclusion and ubiquitination (56). Serum insulin levels were lowered before and during ischemia in the 12-h fasting group, suggesting that the serum insulin level was slightly affected before and during ischemia, due to 12-h fasting. The insulin level clearly increased at 3 h of reperfusion; however, no significant difference was observed between the 12-h fasting group and the control fed group (SI Appendix, Fig. S3A). The expression of p-Akt was significantly increased at 1 h of reperfusion in the control fed mice; however, this was not observed in the 12-h fasting group (SI Appendix, Fig. S3C). Although p-Akt was increased after reperfusion, the expression of phosphorylated FOXO1 (p-FOXO1) remained unchanged (Fig. 3A). In sum, these results suggest that the up-regulation of FOXO1 observed in the 12-h fasted mice may not be directly regulated by the insulin–PI3K–Akt pathway.

The activity of FOXO1 is reportedly regulated by the AMP-activated protein kinase (AMPK) pathway in response to nutrient deprivation (57). Activated AMPK (p-AMPKα) directly phosphorylates FOXO1 at six regulatory residues and enhances FOXO1 transcriptional activity (57). There was no difference in tissue adenosine triphosphate (ATP) levels in both the control fed group and the 12-h fasting group before ischemia and even during reperfusion. However, at 6 h of reperfusion, the ATP level was significantly increased in the 12-h fasting group (SI Appendix, Fig. S3B). Although tissue ATP levels were sustained, the expressions of p-AMPKα were significantly higher in the 12-h fasting group during the early phase of reperfusion (~3 h); however, there was no change at 6 h of reperfusion (SI Appendix, Fig. S3C). These results suggest the possibility that the levels of tissue ATP and p-AMPKα may have contributed to the amelioration of IRI in the 12-h fasting group via the up-regulation of FOXO1 activity.

Deacetylation also occurs in FOXO1 proteins. Sirtuin1 deacetylates FOXO1 and enhances FOXO1 transcriptional activity (58). The expression of Sirtuin1 was increased during ischemia in both the control fed group and the 12-h fasting group. Although Sirtuin1 was significantly higher in the 12-h fasted group at 3 h of reperfusion than in the control fed group (SI Appendix, Fig. S3C), the increment in Sirtuin1 occurred later than the up-regulation of FOXO1. Sirtuin1 might affect the increment in FOXO1 as well as HO-1 during the late phase of reperfusion and plays a role in the fasting-induced amelioration of IRI.

The present study showed that the preventive effect of fasting against liver IRI could be induced more rapidly by 12-h fasting. Higher level of serum BHB was quickly and efficiently induced by the 12-h fast, and the enhanced BHB increased FOXO1. Antioxidative enzyme HO-1 and autophagy activity were also enhanced through FOXO1 up-regulation. Furthermore, BHB inhibited the NLRP3 inflammasome activity during early reperfusion. HMGB1 release was suppressed by the 12-h fasting at the early stage of IR insult, leading to NF-κB inactivation. Taken together, the 12-h fasting helped suppress inflammatory responses by enhancing the BHB expression, followed by the up-regulation of acetylated histone-3 and the activation of FOXO1 and HO-1, together with the consequence of the reduced expression of HMGB1 and the inactivation of NF-κB and NLRP3. These changes resulted in a protective effect against hepatocyte apoptosis and necrosis caused by IRI.

Materials and Methods

Animals. Male C57BL/6 mice (8 wk, 22 g to 25 g weight) were purchased from Shimizu Laboratory Supplies 7 d before operation. Mice were housed (four mice per cage) in individually ventilated cages (TECHNIPLAST S.p.A.), kept under constant environmental conditions with a 12-h light–dark cycle (light 8:00 AM to 8:00 PM), and maintained under specific pathogen-free conditions. All mice were bred with standard rodent breeding chow (CA-1, CLEA Japan, Inc.) and sterile water ad libitum unless otherwise indicated, and
received human care as per the Guide for Care and Use of Laboratory Animals (National Institute of Health Publication, eighth edition, 2011).

Reagents. DL-BHB was purchased from Sigma-Aldrich Co. LLC., AS1842856 (FOXO1 inhibitor) (59) was purchased from Merck Millipore, and Trichostatin A (HDAC inhibitor) (40) was purchased from Tokyo Chemical Industry Co., Ltd.

Liver IRI Model. An established mouse model of partial warm hepatic IRI was used (10–13, 15, 18, 60). Surgical manipulation was performed from 8:30 AM. The mice were anesthetized under isoflurane (2 to 2.5%) and injected with heparin (100 U/kg). An atrumatic clip was used to interrupt the artery and portal venous supply and bile duct to the left and middle liver lobes. This method prevents mesenteric venous congestion by permitting portal de-compression through the right and caudate lobes. After 60 min of ischemia, the clamp was removed, and reperfusion was initiated. The mice were killed at different time points (SI Appendix, Fig. S1A). Liver samples were immediately dissected, mounted in optimal cutting temperature embedding compound, frozen at −80 °C, fixed overnight in 10% formaldehyde or frozen in liquid nitrogen, and reserved at −80 °C until extraction. Shaker-operated mice underwent the same procedure without vascular occlusion.

The mice were divided into groups (SI Appendix, Fig. S1B). The control fed group was provided food and sterile water ad libitum, while the fasting group was deprived of food but given free access to water for 12 h before the IR treatment. In the cases where the fasting group was fed 10% glucose water, sterile water was exchanged with 10% glucose water at the start of fasting.

The mice received i.p. administration of AS1842856 (FOXO1 inhibitor, 20 mg/kg) in 40 μL of dimethyl sulfoxide (DMSO) or 40 μL DMSO alone at 36 h and 12 h before ischemia (SI Appendix, Fig. S1C). The dose and usage of AS1842856 were determined based on a previous report (61). The control fed mice received i.p. administration of Trichostatin A (HDAC inhibitor, 1 mg/kg) in 0.2 mL of 20% DMSO in PBS or 0.2 mL of 20% DMSO in PBS alone 16 h before ischemia and just before ischemia (SI Appendix, Fig. S1D). The dose and route of administration of Trichostatin A were determined based on a previous report (62). The control fed mice received i.p. administration of BHB (10 mmol/kg) in 0.5 mL of PBS or 0.5 mL of PBS alone 30 min before the IR treatment. In the cases where the fasting group was fed 10% glucose water, sterile water was exchanged with 10% glucose water at the start of fasting.

Analysis of Blood Samples. The SALT level, an indicator of hepatocellular injury, was measured using a standard spectrophotometric method with an automated clinical analyzer (JA-58; JEOL Ltd.). Serum BHB was measured using an enzymatic method (ORIENTAL YEAST Co., Ltd.). ELISA. The serum HMGB1 was quantified with a HMGB1 ELISA Kit II (Shino-Test). Serum insulin was measured using Ultra-sensitive Mouse/Rat Insulin ELISA. Serum cytokines were measured using the Luminex multiplex cytokine analysis kit (Bio-plex Biotechnology, Inc.). Tissue ATP concentrations were determined based on a previous report (47) and by using the luciferase method with Luminescent ATP Detection Assay (Abcam).

Measurement of Tissue ATP Concentrations. Liver tissue samples were homogenized in 10 volumes of 0.25 mol/L sucrose in 10 mmol/L Heps-NaOH buffer (pH 7.4). The extracts were cleared via centrifugation, and amounts of protein were estimated using the BCA protein assay kit (Pierce Biotechnology, Inc.). Tissue ATP concentrations were determined using the luciferin–luciferase method with Luminescent ATP Detection Assay Kit (Abcam).

Western Blotting Assay. Proteins extracted from the liver (20 μg per sample) were subjected to sodium dodecyl sulphate (SDS) polyacrylamide gel electrophoresis and transferred to the polyvinyliden fluureide membrane (Bio-Rad). Thereafter, the membrane was incubated in blocking buffer [5% skim milk in Tris-buffered saline with polyoxymethylene sorbitan monolaureate (TBS-T)] for 1 h at room temperature. After blocking, the membrane was incubated in conjugated primary antibody (SI Appendix, Table S2) in the dilution buffer (2.5% skim milk in TBS-T) with overnight agitation at 4 °C. Then, the membrane was incubated in horseradish peroxidase-linked anti-rabbit IgG antibody (1:1000 dilution) in the dilution buffer with gentle agitation for 1 h at room temperature. Enhanced Chemi Luminescence (ECL prime; Amersham) and Lumino image analyzer (Image Quant LAS 4000; GE Healthcare) were used to detect each molecule. The intensity of the bands was quantified using Image J software (National Institutes of Health).

Apoptosis Assay. Apoptosis in 5-μm-thick liver paraffin sections was detected with TUNEL performed with the In Situ Apoptosis Detection Kit (TAKARA BIO) as per the manufacturer's protocol. Positive cells were counted blindly at 10 high power field (HPF)/section (magnification 400×).

Quantitative Reverse-Transcription PCR. Total RNA was extracted from liver tissue using the NucleoSpin RNA (Takara Bio). cDNA was prepared using the RT reagent kit (Takara Bio). Quantitative reverse-transcription PCR was performed using the StepOnePlusTM Real-Time PCR System (Life Technologies). Primers used to amplify specific genes were listed in SI Appendix, Table S3. Target gene expression was calculated by the ratio to the housekeeping gene hypoxanthine phosphoribosyltransferase (HPRT).

Statistical Analyses. Data are expressed as mean ± SD values. Differences between the experimental groups were analyzed using the Student's t test; two-way repeated-measures analysis of variance (ANOVA) with Bonferroni's postest was used to assess time-dependent changes and differences between the groups at each time point. P values of <0.05 were considered statistically significant.

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