

PKA phosphorylation couples hepatic inositol-requiring enzyme 1 α to glucagon signaling in glucose metabolism

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The endoplasmic reticulum (ER)-resident protein kinase/endoribonuclease inositol-requiring enzyme 1 (IRE1) is activated through transautophosphorylation in response to protein folding overload in the ER lumen and maintains ER homeostasis by triggering a key branch of the unfolded protein response. Here we show that mammalian IRE1 α in liver cells is also phosphorylated by a kinase other than itself in response to metabolic stimuli. Glucagon-stimulated protein kinase PKA, which in turn phosphorylated IRE1 α at Ser⁷²⁴, a highly conserved site within the kinase activation domain. Blocking Ser⁷²⁴ phosphorylation impaired the ability of IRE1 α to augment the up-regulation by glucagon signaling of the expression of gluconeogenic genes. Moreover, hepatic IRE1 α was highly phosphorylated at Ser⁷²⁴ by PKA in mice with obesity, and silencing hepatic IRE1 α markedly reduced hyperglycemia and glucose intolerance. Hence, these results suggest that IRE1 α integrates signals from both the ER lumen and the cytoplasm in the liver and is coupled to the glucagon signaling in the regulation of glucose metabolism.

endoplasmic reticulum stress | G protein-coupled receptor | metabolic disease

Homeostasis in the endoplasmic reticulum (ER) is central to the proper function and survival of eukaryotic cells. Accumulation of unfolded/misfolded proteins in the ER lumen triggers the unfolded protein response (UPR), which reduces ER stress by enhancing the ER's capacity to manage the workload of protein folding (1). The mammalian UPR is executed through three canonical signaling branches mediated by the ER-localized transmembrane proteins, inositol-requiring enzyme 1 (IRE1), PKR-like endoplasmic reticulum kinase (PERK), and activating transcription factor 6 (ATF6). Conserved from yeast to humans, IRE1 is the most ancient sensor of ER stress and possesses both protein Ser/Thr kinase and endoribonuclease (RNase) activities (1–5). Under ER stress conditions, IRE1 is activated through dimerization and transautophosphorylation (6, 7). Activation of IRE1 α results in the nonconventional splicing of X-box binding protein 1 (*Xbp1*) mRNA to generate an active form of this transcription factor that induces a major transcriptional program of the UPR (8–10). Interestingly, chemical manipulation of the activation mode of IRE1 α has been shown to generate alternate RNase outputs (11), suggesting complex structural features of this molecule upon autophosphorylation within its kinase domain to activate its RNase activity. Recently reported crystal structures of the cytoplasmic kinase/endoribonuclease region of yeast Ire1, as well as the dephosphorylated human IRE1 α , revealed a dimeric orientation (12, 13) or oligomeric arrangements (14) for transautophosphorylation, implying that this kinase is unlikely to be accessible for phosphorylation by other kinases than itself (14) and the XBP1-specific RNase activity is dependent on autophosphorylation (12). However, it remains largely unknown if mechanisms aside from auto-

phosphorylation are operative in activating IRE1, particularly under physiological or pathophysiological conditions.

The ER is thought to be a critical nutrient-sensing organelle that is functionally linked with metabolic responses (15). Emerging evidence has shown that in mammals, the UPR pathways are associated with changes in metabolic cues (15–19), as well as with obesity-associated metabolic stress conditions (20–23). Given our previous finding that glucose stimulation of IRE1 α phosphorylation in pancreatic β -cells represents a glucose-sensing event (18), we considered the possibility that induction of IRE1 α phosphorylation in response to metabolic stimuli could reflect a general phenomenon of physiological activation of this UPR sensor. Here we show that IRE1 α can integrate metabolic signals through phosphorylation by PKA of the G protein-coupled receptor (GPCR) pathway and is coupled to glucose metabolism in liver cells, thereby contributing to disruption of glucose homeostasis under obesity-associated metabolic ER stress.

Results

IRE1 α Is Metabolically Activated by Glucagon in the Liver. To test if mammalian IRE1 α is able to sense changes in metabolic cues, we first examined the phosphorylation status of liver IRE1 α in mice under fasted or fed states. In parallel with elevated phosphorylation of cAMP response element-binding protein (CREB), phosphorylation of IRE1 α at Ser⁷²⁴, a highly conserved positive regulatory site (Ser⁸⁴¹ in yeast Ire1) within its activation segment of the kinase domain (12–14), markedly increased upon fasting as detected by a phospho-site-specific antibody (Fig. 1A). In contrast, fasting did not enhance the phosphorylation of eukaryotic translation initiation factor 2 α (eIF2 α), the downstream target of the PERK pathway of the UPR (1). Furthermore, glucagon, the principal hormone that regulates hepatic glucose production in the fasted state, robustly augmented the phosphorylation of IRE1 α but not eIF2 α in livers of mice (Fig. 1B), as well as in

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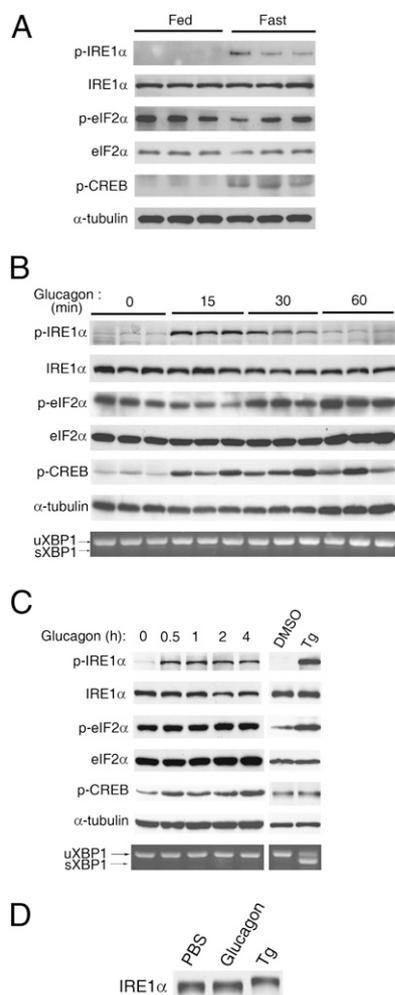


Fig. 1. Glucagon stimulates the phosphorylation of hepatic IRE1 α . (A) Fasting induced hepatic IRE1 α phosphorylation. Male C57BL/6 mice were ad libitum fed or subjected to a 6-h fast. Liver extracts were analyzed by immunoblotting using the indicated antibodies. Results are shown for three individual mice in the fed or fasted state, representative of two independent experiments. (B and C) Glucagon stimulated IRE1 α phosphorylation in vivo and in primary hepatocytes. (B) Mice were treated for the indicated time intervals ($n = 3$ per group) by intraperitoneal injection of glucagon (100 μ g/kg body weight). (C) Mouse primary hepatocytes were treated with 100 nM glucagon as indicated, or with dimethyl sulfoxide (DMSO) or thapsigargin (Tg, 1 μ M) for 1 h. Protein extracts were analyzed by immunoblotting, and spliced (s) and unspliced (u) *Xbp1* mRNA transcripts were measured by RT-PCR. Results are representative of at least three independent experiments. (D) Glucagon induced a distinct phosphorylation state of IRE1 α from that caused by ER stress. Primary hepatocytes were treated with PBS, 100 nM glucagon, or 1 μ M Tg for 1 h. Immunoblotting was performed for band-shift analysis using the IRE1 α antibody. Results are representative of three independent experiments.

isolated primary hepatocytes (Fig. 1C), suggesting that glucagon selectively affected the IRE1 α branch of the UPR pathways. Notably, glucagon-induced IRE1 α phosphorylation at Ser⁷²⁴ did not increase the splicing of *Xbp1* mRNA (Fig. 1B and C), contrasting that observed upon treatment with thapsigargin, a chemical inducer of ER stress (Fig. 1C). Although up-regulating the mRNA expression of two gluconeogenic genes, *G6pase* and *Pepck* (SI Appendix, Figs. S1A and S2A), glucagon did not increase the mRNA abundance of two UPR genes, *Chop* and *Bip* (SI Appendix, Figs. S1B and S2A). On the other hand, thapsigargin-induced phosphorylation of IRE1 α (Fig. 1C) was accompanied by increased mRNA abundance of *Chop* and *Bip* but

not that of *G6pase* and *Pepck* (SI Appendix, Fig. S2B). As an internal control, *actin* mRNA expression was not affected in hepatocytes when exposed to either glucagon or thapsigargin (SI Appendix, Fig. S2C). These results indicate that glucagon-induced acute phosphorylation of IRE1 α represents a metabolic activation event, which is distinct from that induced by typical ER stress. In keeping with this idea, glucagon-stimulated phosphorylation did not cause a detectable shift of the IRE1 α protein when examined by immunoblot analysis (Fig. 1D), in contrast to the thapsigargin-induced shift that most likely resulted from multiple-site autophosphorylation of IRE1 α upon ER stress (13, 14). These data thus raise the possibility that mechanisms aside from transautophosphorylation are operative in initiating the metabolic activation of the IRE1 α pathway.

Next, we investigated whether IRE1 α acts as a critical component in mediating glucagon's metabolic effects. In comparison with primary liver cells infected with a scrambled control adenovirus (Ad-shCON), knockdown of the expression of IRE1 α with Ad-shIRE1 α -#1 or -#2, which expressed one of two shRNAs directed against IRE1 α , significantly compromised the stimulation by glucagon of the mRNA expression of *G6pase* and *Pepck* without affecting the phosphorylation of CREB (Fig. 2A). Moreover, transcriptomic analysis (SI Appendix, Fig. S3A) revealed a remarkable blunting effect of IRE1 α knockdown on glucagon-regulated gene-expression programs in liver cells, as indicated by a Pearson Correlation Coefficient (PCC) of -0.64 (Fig. 2B). Among genes that were significantly up-regulated or down-regulated by glucagon induction, $\sim 37\%$ or $\sim 43\%$ were oppositely affected by IRE1 α knockdown (Fig. 2B), which were enriched by ~ 9.5 - or ~ 11.5 -fold over all array-probed genes ($P < 2.2E-16$, Fisher's exact test) (SI Appendix, Fig. S3B). Parametric analysis of gene set enrichment (PAGE) also showed a profound influence of IRE1 α deficiency upon glucagon-regulated cellular pathways and biological processes (SI Appendix, Fig. S3C), including those associated with carbohydrate, lipid, and amino acid metabolism. Notably, among transcription factors involved in the regulation of lipid metabolism and enzymes related to triglyceride synthesis, which were found to be up-regulated under ER stress in hepatocytes from mice with liver-specific deletion of IRE1 α (24), IRE1 α knockdown under glucagon-stimulated conditions did not cause prominent changes in the expression of *Clebp β* , *Clebp δ* , or *Ppar γ* , but resulted in increased expression of *Scd1*, *Dgat1/2*, and *Acc1* (SI Appendix, Fig. S3D). Importantly, knockdown of hepatic IRE1 α markedly decreased the glucose production capacity (Fig. 2C) of hepatocytes isolated from Ad-shIRE1 α -#2-infected mice in the absence or presence of stimulation with forskolin, a chemical activator of protein kinase A (PKA) of the glucagon signaling pathway. Suppression of hepatic IRE1 α in mice also led to a significant attenuation of the hyperglycemic response to glucagon (Fig. 2D). Together, these results demonstrate that the IRE1 α pathway of the UPR in the liver is responsive to glucagon stimulation and is functionally implicated in mediating glucagon's regulatory actions in glucose metabolism.

PKA Is Required for Glucagon-Induced Phosphorylation of IRE1 α .

Glucagon triggers a cascade of signal transduction events through activation of its seven-transmembrane GPCR. Stimulation of adenylylate cyclase activity through the stimulatory G protein (G_{α_s}) leads to increased cAMP production, thereby activating PKA for CREB phosphorylation (25). Glucagon can also increase the intracellular calcium in a phospholipase C (PLC)-dependent manner (26). To test the idea that a protein kinase other than IRE1 α itself may link glucagon receptor signaling with the observed stimulation of IRE1 α phosphorylation, we examined whether PKA is involved. Interestingly, pharmacological PKA inhibitor H89, but not PLC inhibitor U73122, blunted glucagon-stimulated phosphorylation of IRE1 α as well as CREB in primary hepatocytes (Fig. 3A). In contrast, H89 showed no such inhibitory effect on thapsigargin-induced phosphorylation of IRE1 α (SI Appendix, Fig. S4A). Epinephrine, another agonist of G_{α_s} -coupled receptor and regulator of hepatic glucose metabolism, could also stimulate the phosphorylation of

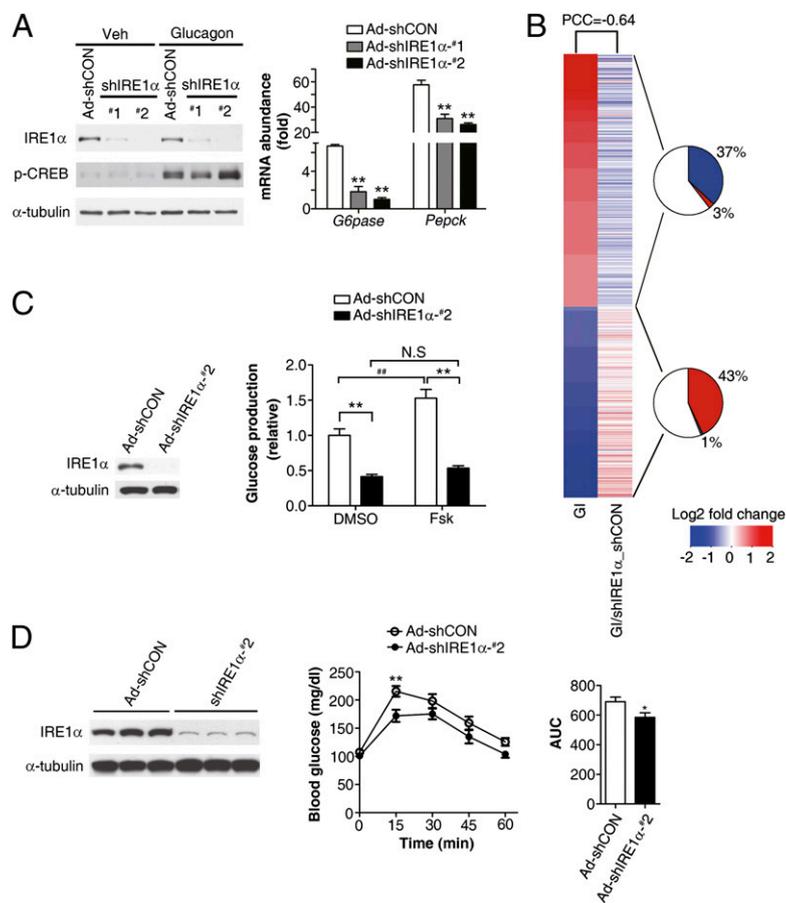


Fig. 2. IRE1 α mediates glucagon's metabolic effects in liver cells. (A) IRE1 α knockdown reduced glucagon-induced expression of gluconeogenic genes. Primary hepatocytes, infected for 72 h with the indicated adenoviruses, were incubated with 100 nM glucagon for 4 h. Immunoblotting was performed for IRE1 α expression analysis. Expression of *Pepck* and *G6pase* was analyzed by real-time RT-PCR using *actin* as an internal control, shown as averaged fold-induction by glucagon \pm SEM ($n = 3$ independent experiments). $^{***}P < 0.01$ compared with Ad-CON by one-way ANOVA. (B) Impact of IRE1 α knockdown on glucagon-induced transcriptomic changes in hepatocytes. Microarray analysis was performed as described in *SI Appendix, Fig. S3*. Heat maps represent averaged fold changes of gene expression from three independent experiments, showing differentially expressed genes caused by glucagon induction (GI) as aligned with changes of these genes as a result of IRE1 α knockdown in the presence of glucagon stimulation (GI/shIRE1 α -shCON). PCC value was calculated between the fold-changes in the two heat maps. The pie charts indicate the percentages of glucagon-up-regulated or -down-regulated genes that were oppositely changed or unaltered by IRE1 α knockdown. (C and D) IRE1 α deficiency reduced glucose production from liver cells and attenuated the hyperglycemic response to glucagon in vivo. Mice were infected with Ad-shCON or Ad-shIRE1 α -2 through tail vein injection. (C) Primary hepatocytes were isolated after 5 d of infection and glucose production was measured in the absence or presence of 10 μ M forskolin (Fsk). Data are presented as the mean \pm SEM ($n = 3$ independent experiments). $^{***}P < 0.01$, $^{##}P < 0.01$ by two-way ANOVA. (D) Glucagon challenge test. Mice infected for 10 d were injected intraperitoneally with glucagon (150 μ g/kg) after a 15-h fast. Blood glucose was measured at the indicated time points after glucagon injection. The areas under the curve (AUC) were shown as the mean \pm SEM ($n = 10$ –11/group). $^{*}P < 0.05$, $^{***}P < 0.01$ by two-way ANOVA or *t* test.

IRE1 α , which was likewise attenuated by H89 (Fig. 3B). Arguing for a direct connection of cAMP-dependent PKA activation with IRE1 α phosphorylation, both forskolin (an activator of adenylate cyclase) and Br-cAMP (a cAMP analog) were able to increase the phosphorylation of IRE1 α in liver cells (*SI Appendix, Fig. S4B*). Ruling out possible nonspecific effects of H89 (27), adenovirus-mediated knockdown of hepatic PKA expression with Ad-shPKA abolished glucagon-stimulated but not thapsigargin-induced phosphorylation of IRE1 α (Fig. 3C). These results clearly demonstrate that PKA is required for the phosphorylation of IRE1 α evoked by the GPCR-cAMP pathway. To verify further that glucagon-induced phosphorylation of IRE1 α is independent of its capability of transautophosphorylation, we used a Lys⁵⁹⁹→Ala⁵⁹⁹ (K599A) substitution kinase-dead mutant of human IRE1 α that was unable to autophosphorylate spontaneously (18) when adenovirally overexpressed in primary hepatocytes, as analyzed using the phospho-site-specific antibody that could not detect the Ser⁷²⁴→Ala⁷²⁴ (S724A) mutant of IRE1 α (Fig. 3D). Overexpressed S724A mutant failed to be autophosphorylated at this site located within the structurally disordered central activation segment of IRE1 α (12), and blocking its phosphorylation may cause disruption of IRE1 α 's functional outputs (17, 18). Consistently, glucagon increased the phosphorylation at Ser⁷²⁴ of IRE1 α -K599A, which did not alter its inactivity for splicing *Xbp1* mRNA (Fig. 3D). In contrast, thapsigargin was unable to increase overtly the phosphorylation of IRE1 α -K599A (*SI Appendix, Fig. S5A*) through activation of the endogenous IRE1 α . Because the IRE1 α -K599A mutant could be transphosphorylated at a detectable level by the overexpressed EGFP-IRE1 α -WT fusion protein that was robustly autophosphorylated (*SI Appendix, Fig. S5B*), Ser⁷²⁴ within the IRE1 α -K599A mutant might be more preferably phosphorylated by PKA than by the endogenous IRE1 α . In addition, H89 attenuated glucagon-induced phosphorylation of IRE1 α -K599A at

Ser⁷²⁴, but not that of IRE1 α -WT resulting from forced autophosphorylation (Fig. 3D). Notably, the S724A mutant retained its ability to splice *Xbp1* mRNA, suggesting that phosphorylation at this single site may not be essential for activating its RNase activity. Given that Ser⁷²⁴ is positioned within a consensus target sequence (RHS) of PKA (28), we reasoned that IRE1 α may be a direct phosphorylation substrate of PKA. Indeed, purified recombinant PKA (*SI Appendix, Fig. S6A*) but not PKC ϵ (*SI Appendix, Fig. S6B*) increased the phosphorylation at Ser⁷²⁴ of immunoprecipitated Flag-tagged IRE1 α -WT or IRE1 α -K599A from hepatocytes (Fig. 3E), as well as a purified cytoplasmic fragment of IRE1 α (Fig. 3F), both of which could be inhibited by H89. Thus, these data revealed a previously unanticipated PKA-dependent mechanism by which hepatic IRE1 α phosphorylation is metabolically linked to the GPCR pathway.

Blocking Phosphorylation at Ser⁷²⁴ Affects the Metabolic Output of IRE1 α . We then sought to assess the functional significance of PKA-directed phosphorylation at Ser⁷²⁴ of hepatic IRE1 α . First, band-shift analysis by immunoblotting showed a considerably less extent of shift of the adenovirally overexpressed IRE1 α -S724A mutant protein than that of IRE1 α -WT (Fig. 4A), whereas glucagon-induced phosphorylation did not cause a detectable shift of the autophosphorylation-deficient mutant IRE1 α -K599A (Fig. 4A). This finding indicates that blocking Ser⁷²⁴ phosphorylation might impede IRE1 α from being fully activated through transautophosphorylation. Next, we examined the effects of deficiency in Ser⁷²⁴ phosphorylation upon IRE1 α -evoked changes of the hepatic transcriptome. Genome-wide transcriptional profiling analysis (*SI Appendix, Fig. S7A*) showed that S724A mutation significantly attenuated IRE1 α -induced changes of gene-expression profiles, as indicated by a PCC of -0.51 (Fig. 4B). For genes significantly up-regulated or down-regulated by IRE1 α , $\sim 21\%$ or $\sim 13\%$ were oppositely influenced by S724A mutation (Fig. 4B),

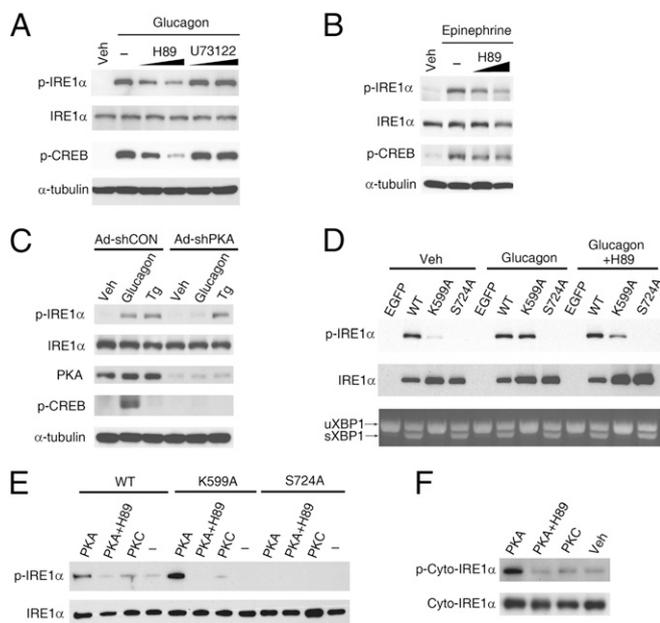


Fig. 3. PKA is directly responsible for phosphorylating IRE1 α . (A and B) Inhibition of PKA decreased glucagon- or epinephrine-induced phosphorylation of IRE1 α . (A) Primary hepatocytes precultured for 30 min with PKA inhibitor H89 (at 5 and 10 μ M) or PLC inhibitor U73122 (at 5 and 15 μ M) were treated with 100 nM glucagon or PBS/DMSO (Veh) for 1 h. (B) Hepatocytes precultured with 5 and 10 μ M H89 were stimulated with 10 μ M epinephrine for 10 min. Immunoblotting was performed with the indicated antibodies. (C) PKA knockdown blunted glucagon-stimulated IRE1 α phosphorylation. Hepatocytes infected for 72 h with adenoviruses Ad-shCON or Ad-shPKA were subsequently incubated for 1 h with 100 nM glucagon or 1 μ M Tg. (D) Glucagon-induced phosphorylation of IRE1 α was independent of its capability of autophosphorylation. Hepatocytes were infected with adenoviruses expressing EGFP, Flag-tagged human wild-type (WT) IRE1 α or IRE1 α -K599A and IRE1 α -S724A mutants. Cells with or without preincubation for 30 min in 10 μ M H89 were then treated with 100 nM glucagon for 1 h. Phosphorylation of IRE1 α and splicing of *Xbp1* mRNA were analyzed. (E and F) PKA directly phosphorylated IRE1 α in vitro. (E) Hepatocytes were infected for 48 h with adenoviruses expressing the indicated forms of Flag-tagged IRE1 α . IRE1 α proteins were immunoprecipitated with Flag antibody and subsequently incubated with purified mouse PKA or human PKC ϵ at 30 $^{\circ}$ C for 1 h. (F) Purified recombinant protein of the cytoplasmic portion of human IRE1 α was likewise incubated with PKA or PKC. Phosphorylation of IRE1 α was analyzed by immunoblotting. All results are shown as representative of three (A–E) or two (F) independent experiments.

which were enriched by \sim 5.3- or \sim 9.0-fold, respectively, relative to all genes ($P < 2.2E-16$, Fisher's exact test) (SI Appendix, Fig. S7B). PAGE analysis also revealed a large impact of S724A mutation upon IRE1 α -elicited alterations in metabolism-related cellular pathways (SI Appendix, Fig. S7C), including gluconeogenesis. Next, we verified the importance of Ser⁷²⁴ phosphorylation in IRE1 α regulation of gluconeogenic genes by quantitative RT-PCR. Under glucagon-stimulated conditions, S724A mutation significantly blunted—and K599A mutation completely abolished—the ability of IRE1 α to augment the mRNA abundance of *G6pase* and *Pepck* (Fig. 4C). These data suggest that phosphorylation of hepatic IRE1 α at Ser⁷²⁴, which is also targeted by PKA, dictates the functional output of IRE1 α , and may constitute a key step in glucagon regulation of the gluconeogenic program. As genetic ablation of hepatic IRE1 α or XBP1 has been shown to affect lipid metabolism (24, 29, 30), we wondered whether XBP1 can mediate IRE1 α 's effect on the up-regulation of gluconeogenic genes. Surprisingly, adenoviral overexpression in hepatocytes of the spliced form of XBP1 (SI Appendix, Fig. S8A) did not enhance, but rather suppressed, the expression of *G6pase* and *Pepck* upon glucagon stimulation (SI Appendix, Fig.

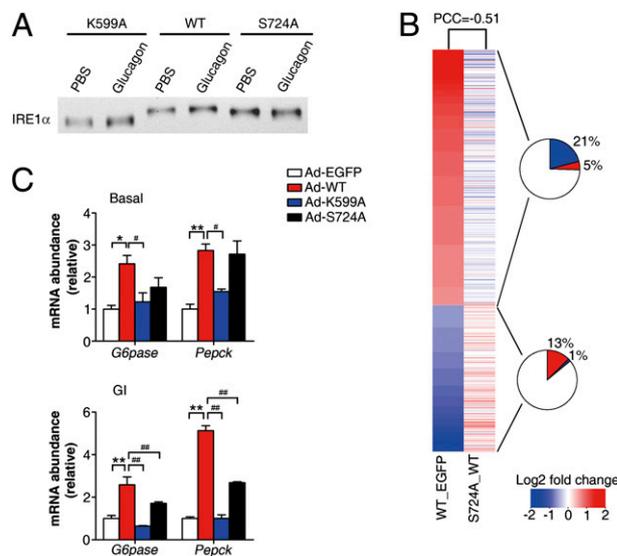


Fig. 4. Phosphorylation at Ser⁷²⁴ dictates the functional output of IRE1 α . (A) Blocking Ser⁷²⁴ phosphorylation resulted in altered autophosphorylation status of IRE1 α . Primary hepatocytes infected with adenoviruses expressing the indicated forms of Flag-tagged human IRE1 α were treated with PBS or 100 nM glucagon for 1 h. Immunoblotting was performed using IRE1 α antibody for band-shift analysis. (B) Impact of disruption of Ser⁷²⁴ phosphorylation on IRE1 α -evoked transcriptomic changes in hepatocytes. Microarray analysis was performed as described in SI Appendix, Fig. S7. Heat maps represent averaged fold-changes of gene expression from three independent experiments, showing differentially expressed genes upon IRE1 α -WT overexpression (WT versus EGFP control) as aligned with changes of these genes caused by S724A mutation (S724A versus WT). PCC value was calculated, and the pie charts indicate the percentages of IRE1 α up-regulated or down-regulated genes, which were oppositely changed or unaltered by S724A mutation. (C) S724A mutation impaired IRE1 α -WT up-regulation of gluconeogenic genes. Hepatocytes infected with the indicated adenoviruses were treated with PBS or 100 nM glucagon for 1 h. The mRNA abundance of *Pepck* and *G6pase* was determined by real-time RT-PCR using *actin* as an internal control. Data are shown as the mean \pm SEM after normalization to the Ad-EGFP control. * P or # $P < 0.05$, ** P or ## $P < 0.01$ by one-way ANOVA.

S8B), and markedly increased the expression of *Bip*. This finding is consistent with a recent study documenting that modest hepatic overexpression of the spliced form of XBP1 protein in mouse models suppressed the gluconeogenic program and improved glucose metabolism (31). Moreover, adenovirus-mediated knockdown of XBP1 expression in hepatocytes (SI Appendix, Fig. S8C) significantly reduced the expression of the XBP1 target gene *Erdj4* upon ER stress, but did not affect glucagon-induced expression of *G6pase* and *Pepck* (SI Appendix, Fig. S8D). Consistent with the reported findings that sXBP1 levels were increased in a pancreatic β -cell line upon prolonged exposure (14 h) to forskolin (32), chronic treatment of hepatocytes with glucagon for 16 h also increased *Xbp1* mRNA splicing (SI Appendix, Fig. S9A), without further influencing the expression of *G6pase* and *Pepck* (SI Appendix, Fig. S9B). These results thus argue for a lesser role for sXBP1 in glucagon up-regulation of gluconeogenic genes, further suggesting that IRE1 α most likely contributes, in an XBP1-independent manner, to glucagon-elicited metabolic regulatory programs in the liver.

Highly Phosphorylated Hepatic IRE1 α Is a Hyperglycemic Driver Under Metabolic ER Stress. To gain insights into the physiological significance of PKA-mediated crosstalk between IRE1 α and the glucagon signaling pathway, we determined the phosphorylation status of hepatic IRE1 α from leptin receptor-deficient *db/db* mice (18), which displayed hyperglycemia and hyperglucagonemia in the fasted state (SI Appendix, Fig. S10A). In comparison with wild-

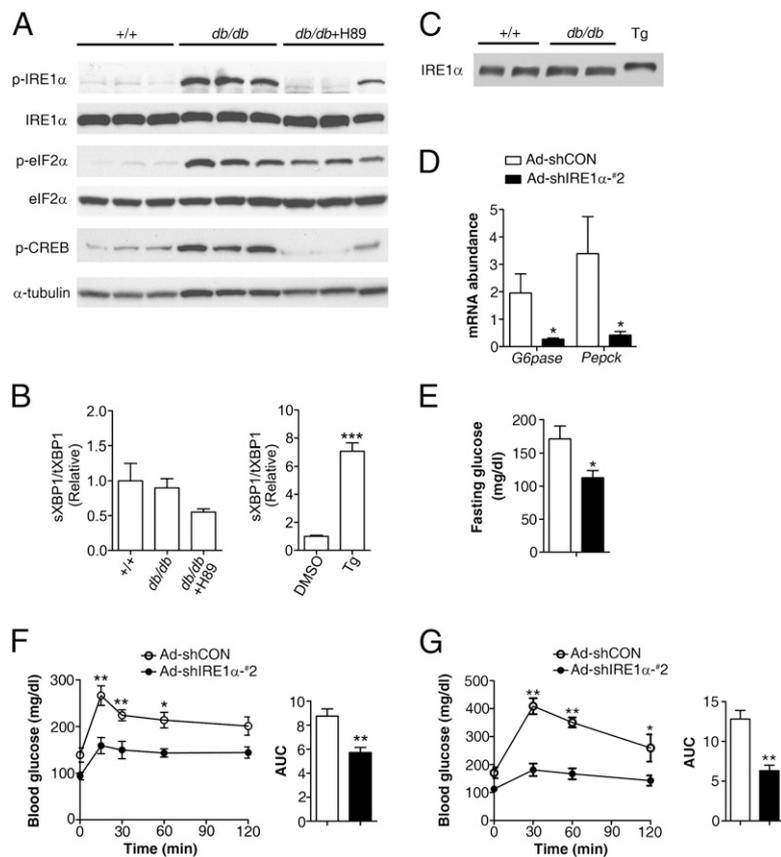


Fig. 5. PKA-dependent hyperactivation of hepatic IRE1 α contributes to obesity-associated disruption of glucose metabolism. (A–C) Highly increased IRE1 α phosphorylation in the livers of *db/db* mice is PKA-dependent and distinct from that induced by ER stress. Male *db/db* mice were treated for 2 h with PBS or H89 (5 mg/kg body weight) through intraperitoneal injection. For ER stress control, primary hepatocytes were treated with 1 μ M Tg or DMSO for 1 h. (A) Liver extracts from *db/db* mice and their wild-type littermates (+/+) were analyzed by immunoblotting with the indicated antibodies. Shown are representative results for three individual mice from each group ($n = 4$ –5 per group). (B) Abundance of the spliced (s) and total (t) *Xbp1* mRNA was determined by real time RT-PCR. (C) Band-shift immunoblot analysis of hepatic IRE1 α from wild-type or *db/db* mice and from Tg-treated hepatocytes. (D–G) Knockdown of hepatic IRE1 α improved glucose metabolism in *db/db* mice. Male *db/db* mice were infected with Ad-shCON or Ad-shIRE1 α -#2 ($n = 5$ /group). (D) At 21 d after infection, the mRNA of liver *G6pase* and *Pepck* was analyzed by real-time RT-PCR after a 6-h fast, using *actin* as an internal control. (E) Glucose was measured after a 6-h fast from mice infected for 18 d. (F and G) Pyruvate tolerance test and glucose tolerance test. Mice infected for 15 or 18 d were fasted for 6 h before intraperitoneal injection with 2 g/kg pyruvate (F) or 1.5 g/kg glucagon (G). Blood glucose was measured at the indicated time points, and the AUCs are shown as the mean \pm SEM ($n = 5$ /group). * $P < 0.05$, ** $P < 0.01$ by two-way ANOVA or t test.

type (+/+) littermates, *db/db* animals showed markedly increased phosphorylation of hepatic IRE1 α , CREB, and eIF2 α (Fig. 5A), suggesting simultaneous hyperactivation of PKA and PERK or PKR kinases under obesity-associated metabolic ER stress (20, 33, 34). Surprisingly, this hyperactivation state of IRE1 α in *db/db* livers, as detected by prominently elevated Ser⁷²⁴ phosphorylation, was not associated with increased *Xbp1* mRNA splicing (Fig. 5B); nor did this increased phosphorylation cause a shift of IRE1 α protein as that observed upon thapsigargin treatment (Fig. 5C). Supporting the idea that increased hepatic PKA activity could largely account for the increased phosphorylation of IRE1 α in *db/db* mice, treatment by H89 substantially decreased the phosphorylation of IRE1 α as well as CREB but not that of eIF2 α (Fig. 5A and SI Appendix, Fig. S10B), accompanied by a considerable reduction of XBP1 splicing. This finding clearly demonstrates that the obesity-associated increase of hepatic IRE1 α phosphorylation indeed primarily stemmed from dysregulation of PKA activity (33), rather than from classical ER stress resulting from increased workload of protein folding. Conceivably, in the face of obesity, this PKA-dependent hyperactivation of IRE1 α may arise largely from hyperglucagonemia, which plays a critical part in type 2 diabetes (26, 35). Next, we examined whether the highly phosphorylated IRE1 α in the liver is an essential contributor to hyperglycemia. Compared with the Ad-shCON-infected control group, adenovirus-mediated knockdown of hepatic IRE1 α by Ad-shIRE1 α -#2 (SI Appendix, Fig. S11A) markedly decreased the mRNA abundance of *G6pase* and *Pepck* (Fig. 5D), as well as that of the spliced form of XBP1 (SI Appendix, Fig. S11B), ruling out the involvement of XBP1 in the observed repression of gluconeogenic genes. Moreover, knockdown of hepatic IRE1 α expression substantially normalized fasting hyperglycemia in *db/db* mice (Fig. 5E) without significantly affecting their body weight (SI Appendix, Fig. S11C). Additionally, animals with hepatic IRE1 α suppression not only exhibited significantly reduced hyperglycemic responses to pyruvate (Fig. 5F) or glucagon (SI Appendix, Fig. S11D), but also showed markedly improved glucose tolerance

(Fig. 5G). These results further demonstrate that obesity-associated, PKA-dependent hyperactivation enables hepatic IRE1 α to act as a critical driver in disrupting glucose homeostasis, suggesting that targeted inhibition of hepatic IRE1 α may bear therapeutic benefits against hyperglycemia.

Discussion

Taken together, our findings establish a PKA-dependent mechanism that links the UPR sensor IRE1 α to the GPCR signaling pathway in the liver under both physiological and pathophysiological states. As depicted in our proposed model (SI Appendix, Fig. S12), upon stimulation by metabolic signals like the GPCR-agonist glucagon, PKA can directly phosphorylate IRE1 α at Ser⁷²⁴, a regulatory site that governs its function. This PKA-mediated phosphorylation represents a hitherto unanticipated cytoplasmic event leading to the metabolic activation of IRE1 α , which is distinct from that initiated through autophosphorylation in response to the ER luminal events during ER stress (i.e., accumulation of unfolded/misfolded proteins). Metabolic activation by the glucagon pathway of hepatic IRE1 α can exert a range of metabolic effects, including regulation of the gluconeogenic program in an XBP1-independent manner.

Currently, it remains an open question how IRE1 α , upon phosphorylation by PKA from the cytoplasm, can exert its effect on glucagon-regulated metabolic programs in the liver. Because we observed a lack of involvement of XBP1 splicing in IRE1 α -dependent up-regulation of gluconeogenic genes, it is possible that the acute activation mode of IRE1 α evoked by the glucagon-PKA pathway may be separated from activation of its XBP1-specific RNase activity. On the other hand, under conditions of chronic glucagon exposure, it is unclear whether PKA-dependent IRE1 α phosphorylation can promote the activation of its RNase activity for XBP1 splicing, and the precise metabolic action of the endogenous sXBP1 in this setting also remains elusive. However, given the multitude of downstream signaling events that can be engaged following IRE1 α activation (36), it is not

unreasonable to speculate that RNase-independent mechanisms may exist to execute its metabolic actions in response to glucagon stimulation.

It is interesting to note that another UPR sensor ATF6, upon activation by classic ER stress, was reported to interact with CRT2 and disturb its ability to coactivate CREB for up-regulating gluconeogenic genes (16). In addition, hepatic overexpression of the sXBP1 protein, the molecule targeted by IRE1 α during typical ER stress, was shown to repress the expression of gluconeogenic genes and improve glucose metabolism in the insulin resistant *ob/ob* mice (31). Whereas a recent study showed that alterations of lipid composition in the ER contributed to increased phosphorylation of IRE1 α in the obese livers (37), our results apparently unmasked a distinct PKA-dependent activation mode of IRE1 α that acted, without involving the splicing of *Xbp1* mRNA, as a critical driver of hyperglycemia in *db/db* mice. Thus, it seems that dysregulated metabolic signals leading to an increase in the PKA activity in the cytoplasm, as well as aberrant lipid and calcium metabolism in the ER, are both responsible for the metabolic hyperactivation of IRE1 α under obesity-associated ER stress conditions. Nonetheless, given the broad roles of the GPCR-PKA pathway in a myriad of biological processes, modulation of the IRE1 α pathway in this context may offer valuable therapeutic leads against many human diseases including diabetes.

Materials and Methods

Animal Studies. Male wild-type C57BL/6J mice were purchased from Shanghai Laboratory Animal Co. Ltd., and male C57BL/6J *db/db* mice and their wild-type littermates were from Model Animal Research Center, Nanjing University. Animals were housed in laboratory cages at a temperature of 23 \pm 3 $^{\circ}$ C and a humidity of 35 \pm 5% under a 12-h dark/light cycle with free access to standard chow (Shanghai Laboratory Animals Co.) and water in accredited animal facilities. All animals were killed under anesthetic conditions or

by cervical dislocation. Livers were snap-frozen in liquid nitrogen immediately after resection and stored at -80° C. All experimental protocols were approved by the Institutional Animal Care and Use Committee at the Institute for Nutritional Sciences, Shanghai Institutes for Biological Sciences, Chinese Academy of Sciences. Adenovirus administration, glucagon challenge test, and glucose or pyruvate tolerance test were carried out as described in the *SI Appendix, Materials and Methods*.

Isolation of Primary Hepatocytes and Adenovirus Infection. Primary hepatocytes were isolated from male C57BL/6J mice at 8 to 12 wk of age. Hepatocytes were infected for 48 or 72 h with adenoviruses at a multiplicity of infection of 40 and were treated with the desired reagents before protein extraction for Western immunoblot analysis. Generation of recombinant adenoviruses was as described in the *SI Appendix, Materials and Methods*.

Detailed materials and experimental procedures for immunoblotting, RT-PCR, IRE1 α phosphorylation analyses, glucose production assays, and microarray analyses were all described in *SI Appendix, Materials and Methods*.

Statistical Analysis. All data are presented as the mean \pm SEM. Statistical analysis was performed using unpaired two-tailed *t* test, and one-way or two-way ANOVA followed by Bonferroni's post test with GraphPad Prism 5.0. *P* < 0.05 was considered to be statistically significant.

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