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Genetic Deficiency of Glycogen Synthase Kinase-3β Corrects Diabetes in Mouse Models of Insulin Resistance

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Despite treatment with agents that enhance β-cell function and insulin action, reduction in β-cell mass is relentless in patients with insulin resistance and type 2 diabetes mellitus. Insulin resistance is characterized by impaired signaling through the insulin/insulin receptor/insulin receptor substrate/PI-3K/Akt pathway, leading to elevation of negatively regulated substrates such as glycogen synthase kinase-3β (Gsk-3β). When elevated, this enzyme has antiproliferative and proapoptotic properties. In these studies, we designed experiments to determine the contribution of Gsk-3β to regulation of β-cell mass in two mouse models of insulin resistance. Mice lacking one allele of the insulin receptor (Ir+/−) exhibit insulin resistance and a doubling of β-cell mass. Crossing these mice with those having haploinsufficiency for Gsk-3β (Gsk-3β+/−) reduced insulin resistance by augmenting whole-body glucose disposal, and significantly reduced β-cell mass. In the second model, mice missing two alleles of the insulin receptor substrate 2 (Ir2s−/−), like the Ir+/− mice, are insulin resistant, but develop profound β-cell loss, resulting in early diabetes. We found that islets from these mice had a 4-fold elevation of Gsk-3β activity associated with a marked reduction of β-cell proliferation and increased apoptosis. Ir2s−/− mice crossed with Gsk-3β+/− mice preserved β-cell mass by reversing the negative effects on proliferation and apoptosis, preventing onset of diabetes. Previous studies had shown that islets of Ir2s−/− mice had increased cyclin-dependent kinase inhibitor p27kip1 that was limiting for β-cell replication, and reduced Pdx1 levels associated with increased cell death. Preservation of β-cell mass in Gsk-3β+/− Ir2s−/− mice was accompanied by suppressed p27kip1 levels and increased Pdx1 levels. To separate peripheral versus β-cell-specific effects of reduction of Gsk3β activity on preservation of β-cell mass, mice homozygous for a floxed Gsk-3β allele (Gsk-3β−/−) were then crossed with rat insulin promoter-Cre (RIP-Cre) mice to produce β-cell-specific knockout of Gsk-3β (βGsk-3β−/−). Like Gsk-3β−/− mice, βGsk-3β−/− mice also prevented the diabetes of the Ir2s−/− mice. The results of these studies now define a new, negatively regulated substrate of the insulin signaling pathway specifically within β-cells that when elevated, can impair replication and increase apoptosis, resulting in loss of β-cells and diabetes. These results thus form the rationale for developing agents to inhibit this enzyme in obese insulin-resistant individuals to preserve β-cells and prevent diabetes onset.

Introduction

Despite treatment with agents that enhance β-cell function and insulin action, reduction in β-cell mass is relentless in type 2 diabetes (T2DM) [1–4]. Why β-cells fail in some individuals is a central issue in diabetes research today. The molecular mechanisms enabling β-cell adaptation to insulin resistance are being discovered primarily in animal models [5–7]. Important genetic models have focused on the requirement for insulin signaling through β-cell insulin/insulin-like growth factor 1 (IGF1) receptors (reviewed in [8,9]). Whereas mice with total-body deficiency for insulin receptor substrate 1 (Ir1−/−) have insulin resistance and significant expansion of β-cell mass, insulin receptor substrate 2-deficient mice (Ir2−/−) have insulin resistance yet develop postnatal β-cell loss and severe diabetes (reviewed in [10]). In this model and in others, the primary of PI-3K/Akt activity in expansion and postnatal maintenance of β-cell mass was apparent (reviewed in [11,12]). The remarkable ability of β-cell mass to expand via enhanced proliferation and reduced apoptosis was demonstrated in transgenic mice expressing constitutively active Akt in β-cells [13,14], illustrating the potential importance of this pathway for expanding β-cells in patients and perhaps resisting the apoptosis that accompanies long-standing diabetes. Knowing that increased expression of Akt in β-cells leads to marked expansion, these results have focused interest on the role of two negatively regulated Akt substrates, FoxO1 and Gsk-3β, each known to regulate carbohydrate and lipid metabolism in insulin target tissues while also exhibiting antiproliferative.

Abbreviations: Gsk, glycogen synthase kinase; Ir, insulin receptor; Ir2, insulin receptor substrate; S.E.M., standard error of the mean; TUNEL, terminal deoxynucleotidyl transferase-mediated dUTP-biotin nick-end labeling; WT, wild type.

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Author Summary

Diabetes is often characterized by a failure of insulin production by pancreatic β-cells to properly regulate glucose homeostasis. Insulin resistance can lead to β-cell failure, and our studies have focused on elucidating the mechanisms involved in this postnatal failure. In this study, we evaluated a new, negatively regulated enzyme of the insulin signaling pathway, glycogen synthase kinase 3 (Gsk-3), specifically within insulin-producing pancreatic β-cells. When this enzyme is elevated, it can impair replication and increase cell death, resulting in loss of insulin-producing cells and diabetes. Gsk-3 is also known to regulate cell death and proliferation in neurons. We assessed the role of Gsk-3 on glucose homeostasis in two different mouse models of insulin resistance. We demonstrated that genetically reducing the levels of Gsk-3β in the insulin-resistant mouse improved glucose homeostasis. In another model in which severe insulin resistance is associated with destruction of β-cells, reducing Gsk-3β not only preserved β-cells by increasing proliferation and reducing cell death, but it also corrected diabetes. Controlling activity of Gsk-3 could lead to new hopes for maintaining or improving β-cell number and prevention of diabetes.

Glycogen synthase kinase-3 (Gsk-3) was originally identified and proapoptotic properties when expressed at high levels [15,16]. There is substantial evidence, mostly from over-expression of a constitutively nuclear FoxO1, that FoxO1 has detrimental effects on β-cell proliferation and survival [17]. On the other hand, there is little known about the effects of expression of Gsk-3β on β-cell proliferation and/or survival.

Glycogen synthase kinase-3 (Gsk-3) was originally identified as a serine/threonine kinase that inactivates glycogen synthase [18]. Early studies showed that insulin inhibits Gsk-3β activity through PI-3K/Akt-induced phosphorylation promoting glycogen synthesis and glucose disposal [19–22]. Later, the enzyme was shown to affect many cellular processes, including transcription, translation, cell cycle regulation, and apoptosis [16,23–27]. Mammals express two isoforms, Gsk-3α and Gsk-3β, which share similar kinase domains but differ considerably in their termini. Inactivation of Gsk-3β appears to be the major route by which insulin activates glycogen synthesis [22,28], and recent studies have demonstrated that elimination of Gsk-3β is more effective at promoting neuronal survival than is elimination of Gsk-3α [29].

Gsk-3 activity has been shown to be increased in peripheral tissues in diabetic animals and patients [30–32], and diabetes was reversed in obese diabetic mice treated with Gsk-3 inhibitors [33–35]. Because inhibitors have differing degrees of kinase specificity, Gsk-3–deficient genetic models were created. Disruption of the Gsk-3β gene in mice results in embryonic lethality [23], yet mice with loss of one allele (Gsk-3β/+) are viable and express reduced levels of protein and enzymatic activity [23]. Although Gsk-3β/+ mice have been little studied, they have been shown to have behavioral effects similar to lithium-treated mice, suggesting that Gsk-3β is the main determinant of Gsk-3 activity in the nervous system [36]. The availability of these Gsk-3β/+ mice provided the opportunity to assess the role of Gsk-3β on insulin sensitivity and pancreatic β-cell function.

Mice haploinsufficient for the insulin receptor (Ir+/−) have insulin resistance with expanded islet β-cell mass and hyperinsulinemia [37]. Crossing Ir+/− mice with mice haploinsufficient for FoxO1 (Foxo1+/−) improved insulin sensitivity and reduced islet mass [17]. In the current study, we hypothesized that in Ir+/− mice, increased Gsk-3β as well as Foxo1 activity could be contributing to the insulin-resistant phenotype. We crossed Ir+/− mice with mice lacking one allele of Gsk-3β (Gsk-3β+/−) and found that Gsk-3β+/−Ir+/− mice, like Foxo1+/−Ir+/− mice, also had improved insulin sensitivity and reduced β-cell mass. Next, we investigated a mouse that is missing two alleles of the insulin receptor substrate 2 (Irs2+/−), that is also insulin resistant, but develops profound β-cell destruction resulting in marked diabetes [38]. Although crossing Irs2+/− mice with Foxo1+/− mice increased β-cell mass and proliferation [39], suggesting that increased β-cell FoxO1 activity was contributing to β-cell loss in Irs2+/− mice, we found that Gsk-3β activity in islets of Irs2+/− mice was also markedly elevated. We determined that Gsk-3β+/−Irs2+/− mice had reduced, but persistent, insulin resistance, yet do not develop diabetes, as a result of maintaining islet β-cell mass. Preservation of β-cell mass in Gsk-3β+/−Irs2+/− mice appeared to be due to accelerated proliferation and decreased apoptosis of β-cells. Reduction of Gsk-3β, like reduction of FoxO1, results in preservation of β-cell mass and rescues the diabetes in this model. The results of these studies now define a new, negatively regulated substrate of the insulin signaling pathway specifically within β-cells that when elevated, can impair replication and increase apoptosis, resulting in loss of β-cells and diabetes.

Results

Gsk-3β Deficiency (Gsk-3β+/−) Promotes Insulin Sensitivity and Reduces the Hyperinsulinemia of Insulin Receptor–Deficient Mice (Ir+/−)

To determine whether Gsk-3β is a downstream contributor to the insulin resistance of insulin receptor–deficient mice, Gsk-3β+/− mice were crossed with mice missing one allele of the insulin receptor (Ir+/−), previously shown to have insulin resistance and elevation of insulin levels in adult animals [37]. Fasting and fed glucose and insulin levels were assessed in Gsk-3β+/−, Ir+/−, and compound heterozygous (Gsk-3β+/−Ir+/−) mice and compared to levels in wild type (WT) at 26 wk of age (Figure 1A). Both fasting and fed insulin levels were significantly reduced in mice lacking one allele of Gsk-3β, thus indicating that genetic deficiency of Gsk-3β activity improves insulin sensitivity. In Ir+/− mice, both fasting and fed insulin levels were higher than in WT mice, consistent with previous reports [37]. In compound heterozygous Gsk-3β+/−Ir+/− mice, the serum insulin values were significantly decreased relative to those in Ir+/− mice in both the fasting and the fed state, although the values were significantly elevated relative to that in Gsk-3β+/− mice (p < 0.05), suggesting that Gsk-3β+/−Ir+/− mice are still insulin resistant. Similar differences in glucose and insulin values were observed in mice at 8–10 wk of age (Figure S1A and S1B). Gsk-3β+/−Ir+/− mice were found to exhibit improved glucose tolerance relative to that in Ir+/− mice (Figure S1C). The results of these experiments thus indicate that (1) endogenous Gsk-3β activity contributes to ambient insulin sensitivity, and (2) that Gsk-3β activity is a downstream mediator of the insulin resistance of the Ir+/− mice.

Gsk-3β+/−Ir+/− Mice Exhibit Enhanced Peripheral Insulin-Mediated Glucose Disposal and Reduced β-Cell Mass Relative to Ir+/− Mice

To further characterize the apparent improvement of insulin sensitivity, hyperinsulinemic-euglycemic clamps were...
performed. The rates of glucose disposal and glucose infusion were increased in Gsk-3β+/−/Ir+/− mice relative to those in Ir+/−/C0 mice, confirming enhanced insulin sensitivity (Figure 1C and 1D). Hepatic glucose production did not appear to differ between Ir+/−/C0 mice and Gsk-3β+/−/Ir+/−/C0 mice (Figure 1E), suggesting that the beneficial effects of genetic reduction of Gsk-3β on carbohydrate metabolism were a result of enhanced effects on peripheral insulin-mediated glucose disposal. These results are consistent with those of Patel et al. (S. Patel, B. W. Doble, K. MacAulay, E. M. Sinclair, D. J. Drucker, and J. R. Woodgett, unpublished data) in which tissue-specific knockout of Gsk-3β in skeletal muscle improved insulin sensitivity, whereas elimination of the gene in liver had no apparent effect on carbohydrate metabolism.

Pancreatic sections with insulin staining of each of the four genotypes are shown in Figures 2A–2D. Although there were no differences in pancreatic areas (unpublished data), the β-cell mass was increased in Ir+/−/C0 mice as previously noted [37], and reduced in Gsk-3β+/−/Ir+/−/C0 mice as assessed by pancreatic morphometry (Figure 2E). In conclusion, mice missing one allele of Gsk-3β when crossed with Ir+/−/C0 mice had reduced hyperinsulinemia associated with reduced β-cell mass.

Gsk-3β Is Activated in Islets of the Irs2-Deficient (Irs2−/−) Mice

Whereas the Ir+/−/C0 mice have peripheral insulin resistance, Irs2-deficient mice (Irs2−/−) have both peripheral insulin resistance as well as impaired insulin signaling, as measured
by reduced Akt activity in islets [40]. Because Akt is a negative regulator of Gsk-3β activity and Gsk-3β is a known regulator of both proliferation and apoptosis [16], we hypothesized that increased Gsk-3β activity could also contribute to the reduced β-cell mass of Irs2Δ−/− mice. Islets from Irs2Δ−/− mice were examined at 6 wk of age and shown to have decreased phosphorylation at serine 473 of Akt and phosphorylation at serine 9 of Gsk-3β in Irs2Δ−/− compared with WT mice (Figure 3A). Additionally, islets from Irs2Δ−/− mice were found to have a 4-fold elevation of phosphorylated glycogen synthase, a substrate of Gsk-3β and a measure of its increased activity (Figure 3B).

**Gsk-3β Haplodeficiency in Irs2Δ−/− Mice Corrects Diabetes**

Because of the antiproliferative and proapoptotic effects of Gsk-3β activity in other tissues [16], finding increased Gsk-3β activity in islets from Irs2Δ−/− mice was consistent with the possibility that this may contribute to the decreased β-cell mass and function of these mice. We therefore crossed mice haploinsufficient for Gsk-3β with Irs2Δ−/− mice to determine whether it would have beneficial effects on preserving β-cell mass and prevent diabetes. We generated double knockout mice (Gsk-3βΔ+/−Irs2Δ−/−) by interbreeding Gsk-3βΔ+/− and Irs2Δ−/− mice. The Irs2Δ−/− mice gained weight until about 9 wk, when body weight plateaued and began to decline. In contrast, the Gsk-3βΔ+/−Irs2Δ−/− mice continued to increase body weight, indistinguishable from that of the WT or Gsk-3βΔ−/− mice at 12 wk (Figure 3C) and at 24 wk (Figure S2A).

Fed glucose levels were determined at 6, 8, 10, and 12 wk of age (Figure 3D). Irs2Δ−/− mice developed a progressive increase with severe hyperglycemia, mean glucose >500 mg/dl, at 12 wk of age, confirming previous observations [38]. In contrast, blood glucose concentrations in Gsk-3βΔ+/−Irs2Δ−/− mice were significantly reduced relative to that in Irs2Δ−/− mice, although plasma insulin did not differ (Figure 3E). Fasting glucose levels at 6 wk of age did not differ, but did at 8 wk (Table S1). To interpret the basis for plasma insulin levels, insulin sensitivity was examined in Irs2Δ−/− and Gsk-3βΔ+/−Irs2Δ−/− mice relative to that in WT mice by insulin tolerance testing in 6-wk-old mice (Figure 3F). Interestingly, the Gsk-3βΔ+/−Irs2Δ−/− mice maintained insulin resistance relative to that in WT mice, suggesting that the beneficial effects of genetic deficiency of Gsk-3β on restoration of glucose homeostasis is not solely due to altered insulin sensitivity.

**β-Cell Mass Is Preserved in Gsk-3βΔ+/−Irs2Δ−/− Mice**

Islet morphology in mice of each genotype was assessed at 8 wk of age. Immunostaining for insulin and glucagon on pancreatic sections are shown in Figure 4A–4F. The insulin immunoreactive area in Irs2Δ−/− mice was severely reduced (Figure 4B and 4E) relative to that in Gsk-3βΔ−/− mice, consistent with previous reports [38,39]. In contrast, there appeared to be preservation of β-cell mass in the Gsk-3βΔ+/−Irs2Δ−/− mice (Figure 4C and 4F). As quantified in Figure 4G, islet mass of Irs2Δ−/− mice was reduced to 30% of WT and Gsk-3βΔ−/− mice, whereas the mass of Gsk-3βΔ+/−Irs2Δ−/− mice did not differ from that of the control mice. Although the α-cell mass was not
determined, the reduction in β- to α-cell ratio (Figure 4H) was consistent with the reduction in β-cell mass.

**β-Cell Proliferation Is Maintained and Apoptosis Reduced in Islets of*** \(Gsk-3β^+/-\) ***Irs2^−/−*** **Mice**

The progressive loss of β-cell mass in ***Irs2^−/−*** mice has been shown to be associated with reduced proliferation and increased apoptosis [38–41]. Preservation of β-cell mass in the ***Gsk-3β^+/-Irs2^−/−*** mice relative to that in the ***Irs2^−/−*** mice could be due to increased proliferation or reduced apoptosis or both. We measured Ki67-positive cells in β-cells to assess proliferation in ***Irs2^−/−*** and ***Gsk-3β^+/-Irs2^−/−*** mice, as shown in Figure 5A. The percent of Ki67-positive β-cells in ***Irs2^−/−*** mice was markedly reduced compared to that in WT and ***Gsk-3β^+/-*** mice at 8 wk of age (Figure 5B). Remarkably, the percentage of Ki67-positive cells in ***Gsk-3β^+/-Irs2^−/−*** mice was over four...
times greater than in \( Irs2^{-/-} \) mice, and even increased approximately 2-fold relative to that in WT and \( Gsk-3\beta^{-/-} \). We next assessed apoptosis by terminal deoxynucleotidyl transferase-mediated dUTP-biotin nick-end labeling (TUNEL) staining in pancreatic \( \beta \)-cells at 8 wk of age (Figure S8). The number of TUNEL-positive \( \beta \)-cells was markedly increased in \( Irs2^{-/-} \) mice compared to that in WT and \( Gsk-3\beta^{-/-} \) mice (Figure 5C). Whereas TUNEL positivity was significantly increased in \( Gsk-3\beta^{-/-}/Irs2^{-/-} \) mice relative to that in WT and \( Gsk-3\beta^{-/-} \) mice, importantly there was approximately a 60% reduction in TUNEL-positive cells in \( Gsk-3\beta^{-/-}/Irs2^{-/-} \) mice compared to \( Irs2^{-/-} \) mice. These observations indicate that the preservation of \( \beta \)-cell mass in \( Gsk-3\beta^{-/-}/Irs2^{-/-} \) mice was associated with reversal of decreased proliferation and increased apoptosis of \( Irs2^{-/-} \) mice.

\textbf{Pdx1 Levels Are Maintained and p27kip1 Levels Are Reduced in Islets from Gsk-3\beta^{-/-}/Irs2^{-/-} Mice}

To examine possible molecular mechanisms for the preservation of islet \( \beta \)-cell mass in \( Gsk-3\beta^{-/-}/Irs2^{-/-} \) mice, islets from WT, \( Irs2^{-/-} \), and \( Gsk-3\beta^{-/-}/Irs2^{-/-} \) mice at 6 wk of age were examined by western blot analysis to assess insulin signaling upstream and downstream of Gsk-3\( \beta \). Phosphorylation of Akt was equally reduced in \( Irs2^{-/-} \) and \( Gsk-3\beta^{-/-}/Irs2^{-/-} \) mice (Figure 6A), indicating that haploinsufficiency for Gsk-3\( \beta \) did not alter the insulin signaling pathway through Akt. Interestingly, total Gsk-3 activity remained reduced in \( Gsk-3\beta^{-/-}/Irs2^{-/-} \), indicating that neither the intact Gsk-3\( \beta \) allele nor the Gsk-3\( \alpha \) alleles compensated to increase Gsk-3 activity. \( Gsk-3\beta^{-/-}/Irs2^{-/-} \) mice had no change in Irs1 expression levels relative to those in \( Irs2^{-/-} \) mice (Figure 6B). Loss of \( \beta \)-cells in \( Irs2^{-/-} \) mice was shown to be associated with decreased islet Pdx1 protein [39,42], whereas transgenic expression of Pdx1 rescued the diabetic phenotype [43]. Associated with preservation of \( \beta \)-cell mass, \( Gsk-3\beta^{-/-}/Irs2^{-/-} \) mice had preservation of Pdx1 levels relative to that in \( Irs2^{-/-} \) mice by western blot analysis (Figure 6B) and nuclear localization by immunocytochemistry (Figure 6C).

The \( Irs2^{-/-} \) mice have been shown to have increased levels

![Figure 4. Morphologic Phenotypes of Irs2^{-/-} and Gsk-3\beta^{-/-}/Irs2^{-/-} Mice](image-url)
and the loss of represented as a percentage of the total number of insulin-positive increased p27kip1 protein, and found that the increased could only be detected in some of the nuclei of the islet cells (Figure 6D, right), with a staining that appears to be weaker than observed in islets, consistent with the increased proliferation observed in these mice.

β-Cell–Specific Gsk-3β Deficiency in Islets of the 

To separate peripheral versus β-cell–specific effects of reduction of Gsk-3β activity on preservation of β-cell mass, mice homozygous for a floxed Gsk-3β allele were then crossed with rat insulin 2 promoter-Cre (RIP-Cre) mice to produce β-cell–specific knockout of Gsk-3β. As shown in Figure 7A, expression of islet Gsk-3β was reduced more than 80%. βGsk-3β/Irs2 mice also maintained relatively normal plasma glucose compared to the severe hyperglycemia of the Irs2 mice for the 20 wk of observation (Figures 7B). Additionally, the hyperinsulinemia of the Irs2 mice was maintained in βGsk-3β/Irs2 mice (Figure 7C).

Gsk-3 Activity Stabilizes p27kip1 in Mouse Insulinoma Cells and in Primary Mouse Islets

To further examine effects of reduction in Gsk-3 activity on protein stability of p27kip1, mouse insulinoma cells were pre-treated with lithium to inhibit Gsk-3 activity, and protein levels were assayed 4 h after addition of cyclohexamide to inhibit new protein synthesis. Lithium treatment reduced Gsk-3 activity and markedly reduced levels of p27kip1 compared to cells treated with NaCl as an osmotic control (Figure 8A).

To confirm the physiological significance of the effects of Gsk-3 activity on p27kip1 levels, primary mouse islets were examined. Islets were isolated from WT mice and incubated for 4 h in serum-free medium, followed by 3 h with no addition, or with the addition of either IGF-1 to activate the insulin signaling pathway, with lithium, or with both, or islet protein lysates blotted for p27kip1 levels (Figure 8B). Addition of IGF-1 and lithium resulted in reduced p27kip1 levels, with maximum reduction with addition of both. These results provide physiological support for the conclusion that Gsk-3 activity stabilizes p27kip1 levels in pancreatic islets.

Discussion

The current study offers specific genetic approaches to assess the role of Gsk-3β in control of β-cell mass in insulin-resistant diabetic models, and as a consequence, several novel observations were made. Loss of one allele of Gsk-3β in WT mice promotes insulin sensitivity and in Irs2 mice reduces insulin resistance and improves glucose tolerance by enhancing glucose disposal. Severely insulin-resistant Irs2 mice were found to have elevated islet Gsk-3 activity associated with severe reduction of β-cell proliferation and elevated apoptosis. Loss of one allele of Gsk-3β in Irs2 mice reversed these findings, preserving β-cell mass and preventing diabetes. Additionally, Pdx1 levels were depressed and p27kip1 levels were increased in islets of Irs2 mice, and they were also reversed by loss of one allele of Gsk-3β. β-cell–specific deficiency of Gsk-3β reversed the diabetes of the Irs2 mice, indicating the importance of Gsk-3β in islet β-cells. Finally, in vitro studies demonstrated
that Gsk-3 activity stabilizes p27kip1 levels, suggesting a mechanism for impairment of proliferation. The results of these studies thus indicate that in insulin-resistant animals, Gsk-3β impairs replication and enhances cell death, leading to postnatal β-cell loss and diabetes.

FoxO1 and Gsk-3β are both negatively regulated targeted proteins of the insulin/PI-3K/Akt signaling pathway. Previous studies showed that Irs2−/− mice crossed with Foxo1+/−/C0 mice resulted in partial correction of fed plasma glucose, β-cell mass, and proliferation [39], along with improved Pdx1 expression. In the current study, we found that islets of Irs2−/− mice had increased Gsk-3 activity (Figure 3A and 3B), now demonstrating that both Gsk-3 and FoxO1 significantly contribute to the impaired proliferation and increased apoptosis in Irs2−/− mice.

What are the mechanisms that could account for postnatal loss of β-cell mass in the insulin-resistant models? Evidence to date suggests that FoxO1 is contributing through impaired proliferation and enhanced apoptosis via transcriptional mechanisms, as it has been shown to repress Pdx1 transcription in insulinoma cells [39]. In non-β-cells FoxO1 has also been shown to increase p27kip1 expression [45]. The results of the current studies suggest a novel mechanism for regulation of Pdx1 and p27kip1 levels in insulin-resistant β-cells through Gsk-3β activity. In non-β-cells, the half-life (t1/2) of p27kip1 protein was 12 h in the absence of growth factors, and 20 min when growth factors were restored, or when cells were treated with Gsk-3β inhibitors [44]. Gsk-3β phosphorylated and stabilized p27kip1 whereas Gsk-3β inhibitors targeted p27kip1 for proteosomal degradation. These results suggest a possible mechanism by which Gsk-3β activity might regulate cell proliferation in β-cells through altered p27kip1 stability. There is substantial evidence that this mechanism is operational in β-cells. First, p27kip1 levels were increased in islets from Irs2−/− mice (Figure 6B and 6D, and [40]), and levels were reduced with elimination of one allele of Gsk-3β.

**Figure 6.** Effects of Inhibition of Gsk-3β Activity on Pdx1 and p27kip1 Expression in Islets from Irs2−/− Mice

Islets were isolated from 6 wk-old WT, Gsk-3β+/−/Irs2−/−, and Irs2−/− mice. (A) Western blot analysis with anti-total Gsk-3, anti-phospho glycogen synthase, anti-phospho Akt (S473), anti-total Akt, and β-actin. Representative results of four independent experiments are presented. Densitometry of phosphorylation of Akt (S473) was measured and normalized over total Akt. Mean values ± S.E.M. are summarized on the graph. A single asterisk (*) indicates p < 0.05. (B) Western blot analysis with anti-total Gsk-3, anti-Irs2, anti-Irs1, anti-Pdx1, anti-p27kip1, and anti-β-actin. Representative results of four independent experiments are presented. Densitometry of Pdx1 and p27kip1 was measured and normalized over β-actin. Mean protein levels ± S.E.M. are summarized on the graph. A single asterisk (*) indicates p < 0.05; double asterisks (**) indicate p < 0.01. (C) Pancreatic sections from 8-wk-old mice of the indicated genotypes at 8 wk were stained with antibodies to Pdx1 (green) and insulin (red). Scale bars represent 50 μm (top) and 10 μm (bottom). (D) Immunostaining for p27kip1 in pancreatic islets of 8-wk-old WT, Irs2−/−, and Gsk-3β+/−/Irs2−/− mice. Hematoxylin counterstaining reveals nucleus in dark blue (and acinar cells in light blue); p27kip1 staining is in red. Scale bars represent 50 μm.

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**Gsk-3β Deficiency Rescues Diabetes**
The Irs2<sup>−/−</sup> mice have severe impairment of the insulin signaling PI-3K/Akt pathway, and rapidly lose β-cell mass. Elimination of one allele of Gsk-3β in Irs2<sup>−/−</sup> mice preserves β-cell mass and, for the most part, maintains glucose homeostasis, yet Gsk-3β is only one of many substrates regulated by the insulin signaling pathway. For example, Irs2<sup>−/−</sup> mice have increased Fox01 [39], and perhaps decreased S6K levels, along with alterations in other Akt substrates. Although the Gsk-3β<sup>+/−</sup>Irs2<sup>−/−</sup> mice maintain apparently normal β-cell mass, they are still insulin resistant, and therefore not fully functional; they would be anticipated to have expanded β-cell mass as shown in the Irs2<sup>−/−</sup> mice (Figure 2E). Thus Gsk-3β is only one protein among many necessary for fully functional β-cells.

Elimination of one allele of Gsk-3β in insulin-resistant Irs<sup>+/−</sup> mice enhanced insulin sensitivity by augmenting peripheral insulin-mediated glucose disposal, independent of effects on hepatic glucose output (Figure 1C–1E). These results are consistent with those in which tissue-specific knockout of Gsk-3β in skeletal muscle enhanced insulin sensitivity (S. Patel, B. W. Doble, K. MacAulay, E. M. Sinclair, D. J. Drucker, and J. R. Woodgett, unpublished data). Could enhanced peripheral insulin sensitivity by loss of one allele of Gsk-3β account for the preservation of β-cell mass in the Irs2<sup>−/−</sup> mice? The results with conditional knockout of the Gsk-3β gene in β-cells indicate the importance of this protein in the β-cell with impaired insulin signaling and that under these circumstances, Gsk-3β is not playing a major role. Although the exact contribution of Gsk-3β to β-cell function has yet to be determined, the results of Patel et al. (S. Patel, B. W. Doble, K. MacAulay, E. M. Sinclair, D. J. Drucker, and J. R. Woodgett, unpublished data) and MacAulay et al. [47] emphasize the isoform and tissue-selective effects of the two mammalian Gsk-3s in skeletal muscle and liver.

The results of these studies now define a new, negatively regulated substrate of the insulin signaling pathway specifically within β-cells that when elevated, can impair replication and increase apoptosis, resulting in postnatal loss of β-cells and diabetes. These results thus form the rationale for developing agents to inhibit this enzyme in obese insulin-resistant individuals to preserve β-cells and prevent diabetes onset.

**Materials and Methods**

Animal production and phenotypic analysis. Generation and genotyping of Gsk-3β<sup>+/−</sup>, Irs2<sup>−/−</sup>, and Irs2<sup>−/−</sup> mice have been described [23,37,38,48]. We maintained Irs2<sup>−/−</sup> mice on the C57BL/6J background and the Irs2<sup>−/−</sup> mice on a mixed C57BL/6J × 129Sv background, and crossed them with Gsk-3β<sup>+/−</sup> mice on the C57BL/6J × 129Sv back ground to obtain Gsk-3β<sup>−/−</sup>Irs2<sup>−/−</sup> and Gsk-3β<sup>−/−</sup>Irs2<sup>−/−</sup> mice on a mixed C57BL/6J × 129Sv background. Double-heterozygote F<sub>1</sub> offspring were intercrossed (Gsk-3β<sup>+/−</sup>Irs2<sup>−/−</sup>) to obtain Gsk-3β<sup>−/−</sup>Irs2<sup>−/−</sup> mice. The Gsk-3β<sup>−/−</sup>Irs2<sup>−/−</sup> progeny was observed at the expected Mendelian frequency in both instances. WT control mice have been obtained from littermates of double-heterozygous breeding.

The generation of mice expressing a conditional allele of Gsk-3β will be described in further detail (S. Patel, B. W. Doble, K. MacAulay, E. M. Sinclair, D. J. Drucker, and J. R. Woodgett, unpublished data). In brief, R1 embryonic stem (ES) cells were electroporated with a modified Gsk-3β targeting vector whereby LoxP sites were introduced by PCR into the intronic region flanking exon 2 of Gsk-3β, and a neomycin resistance cassette was inserted and flanked by FLP recombinase target (FRT) sites.

ES cell clones that had undergone correct homologous recombination were identified by Southern blot and microinjected into C57BL/6J blastocysts. The resultant chimeric mice were crossed to C57BL/6J, and germine transmission of the Gsk-3β floxed allele was verified by PCR. Resultant interbreeding of these mice yielded Gsk-3β floxed mice that...
Figure 8. Effects of Inhibition of Gsk-3 Activity on Protein Stability of p27kip1

(A) MIN6 cells were pretreated with either 40 mM lithium chloride (LiCl) or 40 mM NaCl in DMEM with 15% FBS for 1 h and were cotreated with 25 \( \mu \)g/ml cyclohexamide and lithium or NaCl for 4 h. The lysates were subjected to western blot analysis with anti-p27kip1, phospho-glycogen synthase, total Gsk-3\( \beta \), and \( \alpha \)-tubulin. Densitometry of p27kip1 was measured and normalized over \( \alpha \)-tubulin.

(B) Islets isolated from 16-wk-old WT mice were deprived of serum for 4 h and were incubated with no addition, or with addition of 100 nM IGF-1, or 40 mM LiCl, or both for 3 h. Lysates were then prepared from islets and were subjected to western blot analysis with anti-p27kip1, anti-phospho glycogen synthase, anti-total Gsk-3\( \beta \), and \( \beta \)-actin. Representative results of three independent experiments are presented. Densitometry of p27kip1 was measured and normalized over \( \beta \)-actin. Mean protein levels ± S.E.M. are summarized on the graph. A single asterisk (*) indicates \( p < 0.05 \); double asterisks (**) indicate \( p < 0.01 \).

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were viable, healthy, born with the expected Mendelian frequency, and expressed Gsk-3β at levels indistinguishable from WT animals.

Pancreatic β-cell-specific Gsk-3β knockout mice (βGSK-3β−/−) were generated by breeding Gsk-3β floxed mice with mice that express the Cre recombinase gene under the control of the promoter of the rat insulin 2 gene [49]. To obtain βGsk-3β−/−/Irs2−/− mice, F1 offspring (Gsk-3βfloxed/Irs2floxed) were intercrossed. The offspring were then further intercrossed with F2 offspring (βGsk-3β−/−/Irs2−/−) to obtain βGsk-3β−/−/Irs2−/− mice.

Blood glucose as well as serum insulin concentrations were determined as previously described [13]. For the glucose tolerance test, mice were subjected to an overnight fast followed by intraperitoneal glucose injection (2 g/kg). Blood samples were collected at 0, 30, 60, and 120 min after the injection. For the insulin tolerance tests, mice were subjected to a 4-h fast followed by intraperitoneal human regular insulin injection (0.5 U/kg). Blood samples were collected at 0, 30, 60, and 120 min after the injection. After sacrificing the mice, the pancreas was removed and enzymatic digestion was carried out with male mice. This project was approved by the Animal Ethics Committee of Washington University School of Medicine.

Hyperinsulinenic-euglycemic clamps. Clamp experiments were essentially performed as previously described [50,51]. Double-lumen catheters were placed and 3-[3H] glucose was infused to steady state. The 3-[3H] glucose infusion was continued during the clamp, with labeled glucose infused at the 25% of cold glucose infusion rate to maintain blood glucose concentration at 120 mg/dl for at least 90 min. The 3-[3H] glucose infusion was then stopped, and blood samples were collected at 30, 60, and 120 min after the injection. After sacrificing the mice, the pancreas was removed, enzymatic digestion was carried out, and the isolated islets were counted. Adjacent nonoverlapping fields were analyzed to determine the percentage of TUNEL-positive cells among insulin-positive cells.

Immunoblot analysis of the pancreatic islets. Islets were isolated and fixed from 8-wk-old Irs2−/−, Gsk-3β−/−/Irs2−/−, and Gsk-3β−/− mice. Isolated pancreatic islets were fixed overnight in 3.7% formaldehyde at room temperature. Tissue was then processed for paraffin embedding, and 5-μm sections were obtained. The sections were immunostained with antibodies to insulin (Dako), glucagon (Sigma Aldrich), Ki67 (Zymed Laboratories/Invitrogen), pAktSer473, and p-AktThr308 (Cell Signaling). Total Akt, total Gsk-3β, and total β-actin (Sigma-Aldrich) were also detected. Adjacent sections were stained for insulin and analyzed with NIH Image 1.38 software. Scale bar represents 50 μm.

Figure S3. Fasting glucose in mice, either WT (n = 10) or Gsk-3β−/−/Irs2−/− (n = 13). (A) Body weight in mice, either WT (n = 10) or Gsk-3β−/−/Irs2−/− (n = 13). (B) Fasting and fed insulin levels (8–10 wk of age). (C) Intraperitoneal glucose tolerance tests were performed on overnight-fasted male Irs2−/− mice at 8 wk of age and Gsk-3β−/−/Irs2−/− mice at 8 wk of age after an i.p. injection of 2 g/kg (body weight; n = 6). Glucose levels were assessed at the indicated time intervals. Results are presented as the mean ± S.E.M. in the graph. A single asterisk (*) indicates p < 0.05. Fasting blood glucose concentration at 6 and 8 wk of age in Either WT, Gsk-3β−/−/Irs2−/−, or Gsk-3β−/−/Irs2−/− Mice. (A) Body weight in mice, either WT (n = 10) or Gsk-3β−/−/Irs2−/− (n = 13). (B) Fasting glucose in mice, either WT (n = 10) or Gsk-3β−/−/Irs2−/− (n = 13). (C) Intraperitoneal glucose tolerance tests were performed on overnight-fasted male Irs2−/− mice at 8 wk of age and Gsk-3β−/−/Irs2−/− mice at 8 wk of age after an i.p. injection of 2 g/kg (body weight; n = 6). Glucose levels were assessed at the indicated time intervals. Results are presented as the mean ± S.E.M. in the graph. A single asterisk (*) indicates p < 0.05. Fasting blood glucose concentration at 6 and 8 wk of age in Either WT, Gsk-3β−/−/Irs2−/−, or Gsk-3β−/−/Irs2−/− Mice.

Table S1. Fasting Blood Glucose Concentration at 6 and 8 wk of Age in Either WT, Gsk-3β−/−/Irs2−/−, or Gsk-3β−/−/Irs2−/− Mice.

<table>
<thead>
<tr>
<th>Variable</th>
<th>WT (n = 10)</th>
<th>Gsk-3β−/−/Irs2−/− (n = 13)</th>
<th>n</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fasting glucose</td>
<td>6 wk</td>
<td>8.6 ± 0.6</td>
<td>8.4 ± 0.6</td>
<td>0.03</td>
</tr>
<tr>
<td>Fasting glucose</td>
<td>8 wk</td>
<td>8.9 ± 0.7</td>
<td>9.0 ± 0.7</td>
<td>0.74</td>
</tr>
</tbody>
</table>

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Author contributions. KT conceived, designed, and performed the experiments, analyzed the data, and wrote the paper. ZL and SCM performed experiments and analyzed data. SP and BWD performed experiments and contributed reagents/materials/analysis tools. LL and CMW performed experiments. CCM conceived and designed the experiments and wrote the paper. MPW and JRW contributed reagents/materials/analysis tools. FBM conceived and designed the experiments, MAP conceived, designed, and performed the experiments, and supervised the writing of the paper.

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