Pcif1 modulates Pdx1 protein stability and pancreatic β cell function and survival in mice

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The homeodomain transcription factor pancreatic duodenal homeobox 1 (Pdx1) is a major mediator of insulin transcription and a key regulator of the β cell phenotype. Heterozygous mutations in PDX1 are associated with the development of diabetes in humans. Understanding how Pdx1 expression levels are controlled is therefore of intense interest in the study and treatment of diabetes. Pdx1 C terminus–interacting factor-1 (Pcif1, also known as SPOP) is a nuclear protein that inhibits Pdx1 transactivation. Here, we show that Pcif1 targets Pdx1 for ubiquitination and proteasomal degradation. Silencing of Pcif1 increased Pdx1 protein levels in cultured mouse β cells, and Pcif1 heterozygosity normalized Pdx1 protein levels in Pdx1+/- mouse islets, thereby increasing expression of key Pdx1 transcriptional targets. Remarkably, Pcif1 heterozygosity improved glucose homeostasis and β cell function and normalized β cell mass in Pdx1+/- mice by modulating β cell survival. These findings indicate that in adult mouse β cells, Pcif1 limits Pdx1 protein accumulation and thus the expression of insulin and other gene targets important in the maintenance of β cell mass and function. They also provide evidence that targeting the turnover of a pancreatic transcription factor in vivo can improve glucose homeostasis.

Introduction
The homeodomain transcription factor pancreatic duodenal homeobox 1 (Pdx1) is required for pancreas development in mice and in humans (1–4). Pdx1 is required for the specification of pancreatic progenitors after which its expression becomes restricted primarily to the insulin-producing β cell, where it plays critical roles in insulin gene transcription and insulin secretion as well as β cell survival (reviewed in ref. 5). Tight regulation of Pdx1 protein levels is necessary for Pdx1 to play its developmental and adult physiological roles. Genetic studies in which Pdx1 is conditionally inactivated in mice suggest that Pdx1 gene dosage is critical both for development of the endocrine and exocrine pancreas and for the maintenance of adult β cells (6–8). Heterozygous loss of Pdx1 resulting in approximately 30% reduction in protein levels is sufficient to induce impaired glucose tolerance and insulin secretion in mice, and heterozygous mutations are associated with the development of diabetes in humans (9–11). Furthermore, Pdx1 haploinsufficiency limits the β cell compensatory mechanisms that occur in response to genetic and diet-induced insulin resistance (12–14).

Much of the regulation of Pdx1 dosage occurs at the transcriptional level. Upstream regulatory elements have been described that temporally and spatially control Pdx1 expression and are required for outgrowth and differentiation of pancreatic progenitors (8, 15–20); however, posttranslational regulation of Pdx1 via phosphorylation has also been proposed to have diverse functional effects, including regulation of Pdx1 transcriptional and protein stability (13, 21–25). We previously described Pdx1 C terminus–interacting factor 1 (Pcif1, also known as SPOP), a broad complex, tramtrack, bric-a-brac (BTB) domain–containing protein that interacts directly with the C terminus of Pdx1 (26, 27). BTB-domain containing proteins have recently been identified as substrate-specific adaptors for ubiquitin ligase complexes that contain the scaffolding protein Cul3 (28, 29). We demonstrate here that the Pcif1-Cul3 complex targets Pdx1 protein for ubiquitination and proteasomal degradation. Further, we derived and characterized mice with a mutant allele of Pcif1 (Pcif1<sup>−/−</sup>) and found that reduction of Pcif1 gene dosage improves glucose tolerance, normalizes β cell mass, and rescues β cell apoptosis rates in Pdx1+/- mice while normalizing Pdx1 protein levels and the expression of Pdx1 transcriptional targets.

Results
Pcif1 targets Pdx1 for ubiquitination by a Cul3-based E3 ubiquitin ligase. BTB domain–containing proteins have been identified as substrate-specific adaptors for ubiquitin ligase complexes that include the scaffold protein Cul3 (30). The Drosophila ortholog of Pcif1 targets cubitus interruptus for ubiquitination, while the human ortholog SPOP has been linked to multiple targets, including the polycomb protein Bmi1 (31, 32). To determine whether a similar mechanism underlies Pcif1-mediated downregulation of Pdx1 transactivation, we utilized heterologous HEK293T cells overexpressing Pdx1 with or without coexpression of Pcif1 and Cul3. In cells overexpressing Pdx1, treatment with the proteasome inhibitor MG-132 induced a slight increase in Pdx1 protein levels, suggesting that Pdx1 protein is proteasomally degraded in this system (Figure 1A). Addition of Pcif1 and Cul3 resulted in a dramatic decrease in Pdx1 protein accumulation that was partially reversed by proteasome inhibition. Notably, the ability of Cul3 to decrease Pdx1 accumulation was attenuated by alanine substitutions for key leucine and glutamic acid residues required for interactions between Cul3 and BTB domain–containing proteins (Cul3L52AE55A) (30, 33) (Figure 1A).

To determine whether ubiquitination of Pdx1 is promoted by Pcif1 and Cul3, we employed an in vivo ubiquitination assay in HEK293T cells. Cells expressing Myc-tagged ubiquitin were sub-
jected to immunoprecipitation with anti-Myc or isotype-matched control IgG and the products immunoblotted for Pdx1. Ubiquitination of Pdx1 was detectable as a ladder of high-molecular-weight bands that were immunoprecipitated in cells that expressed Pcif1 and Pdx1, and ubiquitination was further increased by the addition of Cul3 (Figure 1B and Supplemental Figure 1; supplemental material available online with this article; doi:10.1172/JCI40440DS1). Cul3L52AE55A supported a minimal level of ubiquitination, consistent with the proposed role of Cul3 interaction with Pcif1 in promoting Pdx1 ubiquitination.

We previously determined that the TRAF domain of Pcif1 is required for the physical and functional interaction with Pdx1 in vitro (27). To determine whether the interaction between Pcif1 and Pdx1 is required for Pdx1 ubiquitination, we replaced full-length Pcif1 with Pcif1 lacking the TRAF domain (Pcif1ΔTRAF) in the in vivo ubiquitination system and found that the TRAF domain was required to promote Pdx1 ubiquitination (Figure 1C). Further, deletion of a conserved 28-amino-acid motif in the C terminus of Pdx1 (aa 210–238) that is required for the physical and functional interaction with Pcif1 (26, 27) abolished Pcif1-mediated ubiquitination of Pdx1 (Figure 1D). Taken together, these data strongly suggest that physical interactions among Cul3, Pcif1, and Pdx1 are required to promote Pdx1 ubiquitination.

Pcif1 regulates Pdx1 protein accumulation in Min6 β cells. The experiments described above are in agreement with those of Bunce et al, who demonstrated Pcif1-mediated ubiquitination of Pdx1 in an overexpression system (34). To determine whether endogenous Pcif1 regulates Pdx1 protein levels in β cells, we next examined the impact of Pcif1 loss of function on Pdx1 protein accumulation in Min6 mouse insulinoma cells. shRNA-mediated silencing of Pcif1 resulted in a greater than 50% reduction in Pcif1 transcript (Figure 2A). Compared with cells transfected with a nontargeting shRNA, endogenous Pdx1 protein levels were substantially increased in cells after Pcif1 silencing (Figure 2B). Pdx1 protein levels are critical for maintaining β cell function and mediate these effects by regulating the transcription of β cell genes such as MafA and the glucose transporter Glut2 (35, 36). Downregulation of Pcif1 in Min6 cells was sufficient to induce a greater than 2-fold upregulation of MafA and 1.5-fold upregulation of Glut2 transcript (Figure 2C). Insulin mRNA was not regulated under these conditions, likely due to its long half-life. Downregulation of Pcif1 in Min6 cells also extended the half-life of endogenous Pdx1, dramatically increasing protein stability over the course of a 6-hour treatment with cycloheximide (Figure 2D). Treatment of Cul3-overexpressing Min6 cells with the proteasome inhibitor LLnL induced the accumulation of high-molecular-weight ubiquitinated proteins (Figure 2E), including bands detected by Pdx1 Western blot that may represent polyubiquitinated Pdx1. Accumulation of this species was abolished upon Pcif1 knockdown (Figure 2E, left panel). Probing Pdx1 immunoprecipitates from LLnL-treated Min6 cells with an anti-ubiquitin antibody revealed the presence of a high-molecular-weight (~98 kDa) band that was substantially diminished upon Pcif1 knockdown (Supplemental Figure 2). These data suggest that Pcif1-mediated regulation of endogenous Pdx1 protein levels directly impacts Pdx1 protein ubiquitination, accumulation, and stability and the expression of key Pdx1 target genes.

Derivation of a Pcif1 gene trap allele. To investigate the role of Pcif1 in vivo, we generated mice from an ES cell clone containing

![Figure 1](http://www.jci.org)
a gene trap insertion in the first intron of the Pcif1 locus. Mice heterozygous for the gene trap allele (Pcif1+/−) were viable and fertile. In heterozygous crosses, Pcif1+/− embryos harvested at E18.5 were found at Mendelian ratios (n = 372); however, no viable Pcif1−/− mice were found after postnatal day 1, indicating that homozygous loss of Pcif1 results in postnatal lethality. At E18.5, Pcif1−/− embryos displayed normal body and pancreas weight and normal blood sugar (Supplemental Figure 3).

The gene trap insertion cassette contains a strong splice acceptor sequence followed by the βGeo reporter and ends with a polyadenylation signal. The transcript generated by the gene trap should prevent splicing between exons 1 and 2, thereby blocking the generation of the normal Pcif1 transcript. Using primers that span exons 1 and 2, Pcif1 transcript levels were reduced by greater than 50% in E18.5 pancreas and lung from heterozygous animals and undetectable in Pcif1−/− tissues (Supplemental Figure 4, A and B). Similar reductions in Pcif1 protein were observed in Western blots of embryonic kidney lysates (Supplemental Figure 4C). No Pcif1 transcript was detected in Pcif1−/− tissues using primers spanning exons 7 and 8 (data not shown), indicating that the gene trap efficiently prevents the generation of a protein-coding Pcif1 transcript.

Transcript analysis indicated that Pcif1 is expressed during embryonic pancreas development (data not shown). To determine whether Pcif1 modulates Pdx1 protein accumulation during development, we assessed staining intensity in sections from E16.5 and E18.5 embryos using an anti-Pdx1 antibody at a dilution that enabled semiquantitative evaluation of Pdx1 protein levels. Pdx1 staining intensity was markedly increased in all pancreatic lineages in Pcif1−/− mice at these stages (Figure 2F and data not shown). Further, mice homozygous for the Pcif1 gene trap allele demonstrated an approximately 3-fold increase in relative β cell area at E18.5 (Figure 2G), consistent with previously established roles of Pdx1 protein level in regulating endocrine progenitor specification and lineage allocation during pancreatic development (7, 8, 37, 38). There was no significant change in β cell area in the Pcif1+/− mice at this stage of development (Figure 2E).

Increased Pdx1 protein levels in Pcif1-deficient Min6 β cells and embryonic pancreas. (A–C) Min6 cells nucleofected with control nontargeting shRNA (shNT) or shPcif1. (A) Pcif1 transcript measured by QPCR and normalized to Hprt. n = 3, **P < 0.01. (B) Western blot probed for Pdx1 and tubulin (loading control). (C) Transcript levels of Pdx1 transcriptional targets MafA and Glut2. n = 3, **P < 0.01. (D) Min6 cells expressing shNT or shPcif1, treated with the translation inhibitor cyclohexamide (CHX) and harvested at the time points indicated. Western blots are probed for Pdx1 and Ran (loading control). (E) Min6 cells overexpressing Cul3 and nontargeting shRNA or shPcif1 were treated with vehicle or 20 μM LLnL for 8 hours. Western blots are probed for Pdx1 (left) and ubiquitin (right). (F) Fluorescence staining of E16.5 embryos with dilute Pdx1 antibody. Original magnification ×10. (G) β Cell area of E18.5 embryos. n = 6, **P < 0.01 compared with Pcif1+/−.
antly (Figure 3C). Further, the normalization of Pdx1 protein levels in Pcif1+/–Pdx1+/– islets was associated with normalization of pre-
mRNA level for insulin1, a key Pdx1 transcriptional target (Figure 3D). These data demonstrate that Pcif1 regulates the accumula-
tion of Pdx1 protein in the adult β cell and may thus modulate the actions of Pdx1 as a critical regulator of glucose homeostasis.

Pcif1 heterozygosity normalizes β cell mass and improves β cell survival in Pdx1+/– mice. Pdx1+/– mice were previously reported to exhibit an age-related decline in β cell mass due to impaired β cell survival (39). Examination of islet architecture in Pdx1+/– pancreata revealed the previously described “mixed islet” phenotype, with glucagon-positive α cells scattered throughout the core of the islet; this defect was not normalized in Pcif1+/–Pdx1+/– mice (Figure 4A). We quantified β cell mass by determining the relative area occupied by insulin staining of pancreatic sections and observed a marked decrease in the cross-sectional area stained by insulin in Pdx1+/– pancreas compared with all other genotypes. Upon quantification, Pcif1+/+ mice were found to have β cell mass similar to that of wild-type littermates, whereas Pdx1+/– mice displayed a greater than 50% decrease in β cell mass compared with wild-type and Pcif1+/+ mice (Figure 4B). Notably, Pcif1 heterozygosity normalized the reduced β cell mass of Pdx1+/– mice. This was accompanied by a reduction in the rate of apoptotic β cells in Pcif1+/–Pdx1+/– mice as compared with Pdx1+/– mice (Figure 5A).

To determine whether the rescue of β cell area in Pcif1+/–Pdx1+/– was associated with changes in islet size, we determined the distribution of islet size in all genotypes. Although average islet size was not different among genotypes (data not shown), Pdx1+/– mice had a dramatic increase in the number of small (101–250 μm²) islets and a corresponding decrease in the percentage of medium-sized (1,001–2,500 μm²) and large (>25,000 μm²) islets (Figure 4C). Notably, each of these differences in islet size distribution was normalized in Pcif1+/–Pdx1+/– mice (Figure 4C).

In Pcif1+/– mice, the finding of increased β cell apoptosis in the setting of normal β cell mass (Figure 4B and Figure 5A) suggested that β cell replication may also be influenced by Pcif1 gene dosage. Indeed, Pcif1+/– mice displayed an increase in the rate of β cell replication compared with wild-type littermates; however, the rates of BrdU incorporation were not different among Pcif1+/+, Pdx1+/–, and Pcif1+/–Pdx1+/– mice (Figure 5B). The finding of increased β cell replication in Pcif1+/– mice was confirmed by quantification of Ki67 and phospho–histone H3–positive β cells (Supplemental Figure 5, A and B). We also observed a modest but significant decrease in average β cell size in Pdx1+/– mice that was not altered in the context of Pcif1 heterozygosity (Figure 5C). Taken together, these data indicate that the normalization of β cell mass observed in Pcif1+/–Pdx1+/– mice is due primarily to a reduction in the rate of apoptosis.

To determine whether the effect of Pcif1 heterozygosity on β cell turnover was age dependent, we assessed the β cell replication rate of each genotype in 5-week-old mice. Notably, we observed no differences in BrdU incorporation among genotypes at this age (Supplemental Figure 6). In these adolescent animals, pancreas size and β cell apoptosis rates as assessed by TUNEL staining were extremely low; precluding rigorous evaluation; preliminary analysis did not reveal differences based on genotype (data not shown). Taken together, these data suggest that the effect of Pcif1 heterozygosity on β cell replication rates is age dependent and that Pcif1 deficiency improves the age-related decline in β cell mass previously observed in Pdx1+/– mice (39).

Pcif1 deficiency reduces ER stress–associated apoptosis in Pdx1-deficient β cells. Pdx1 deficiency was recently shown to result in increased susceptibility to ER stress–induced apoptosis in β cells (14). To determine whether improved ER homeostasis contributed to the improvement in β cell survival in Pcif1+/–Pdx1+/– mice, we measured mRNA levels of the ER chaperone protein Bip in islets isolated from each genotype. As recently described (14), we observed an increase in Bip mRNA levels in Pdx1+/– islets compared with those of wild-type littermates (Figure 6A). Bip mRNA levels were normalized in Pcif1+/–Pdx1+/– compared with Pdx1+/– islets, suggesting that improved ER homeostasis may underlie the improved β cell survival in transheterozygous animals. In support of this finding, knockdown of endogenous Pdx1 protein in Min6 cells resulted in increased cleavage of the ER-resident caspase-12, while simultane-
ous knockdown of Pci1 restored Pdx1 protein to control levels and prevented caspase-12 cleavage (Figure 6B).

Pci1 heterozygosity improves glucose homeostasis in Pdx1+/− mice. To determine whether the observed normalization of β cell mass in Pci1+/−Pdx1+/− mice is associated with improved β cell function, we assessed glucose tolerance and insulin secretion. Male and female Pci1+/− mice displayed normal glucose tolerance upon intraperitoneal glucose tolerance testing, and Pdx1+/− mice were glucose intolerant as previously described (Figures 7, A and B). The glucose intolerance phenotype of Pdx1+/− mice was significantly improved in both male and female Pci1+/−Pdx1+/− transheterozygous mice. All genotypes displayed similar insulin sensitivity, determined by insulin tolerance testing (Figures 7, C and D), suggesting that the observed alterations in glucose tolerance were due to differences in insulin secretion; therefore, acute glucose-stimulated insulin secretion was assessed in vivo. As expected, Pdx1+/− mice displayed a blunted insulin secretory response to glucose. Notably, the defect in Pdx1+/− mice was attenuated, though not fully rescued, in Pci1+/−Pdx1+/− mice (insulin AUC: Pdx1+/− vs. Pci1+/−Pdx1+/−, 373 ± 44 vs. 595 ± 58 pM min; P < 0.05) (Figure 7E).

To determine whether Pci1 heterozygosity directly influences islet function, we further analyzed insulin secretion in isolated islets. Wild-type and Pci1+/− islets displayed a significant glucose stimulation of insulin secretion in static incubations (Figure 7F). Although Pdx1+/− islets were previously reported to display normal insulin secretion ex vivo, we observed a blunted response.

**Figure 4**

Pci1 heterozygosity normalizes β cell mass in Pdx1+/− mice. (A) Representative images of insulin- and glucagon-stained adult pancreas sections. Original magnification ×20. (B) Quantification of β cell mass. n = 7–10 mice per group; *P < 0.05 compared with wild-type. (C) Islet size distribution. n = 7–9 mice per group; *P < 0.05, **P < 0.01 compared with wild-type.

**Figure 5**

Normalized β cell mass is mediated by improved survival in Pci1+/−Pdx1+/− mice. (A) Apoptosis measured by quantifying insulin- and TUNEL-positive cells. (B) β Cell replication measured by quantifying insulin and BrdU double positive cells. (C) β Cell size calculated by dividing insulin-positive islet area by the number of DAPI-stained nuclei counted within that area. All morphometric analyses performed on pancreas sections from 16-week-old male mice. n = 5–8 per group; *P < 0.05, **P < 0.01 compared with wild-type.
after static incubation (1.76 ± 0.26-fold over basal secretion; P = NS). Glucose-stimulated insulin secretion was restored in Pclf1+/−Pdx1+/− islets (2.76 ± 0.4-fold over basal secretion, P = 0.07) (Figure 7F). These data suggest that Pclf1 haploinsufficiency improves glucose responsiveness in Pdx1+/− β cells.

Pclf1 regulation of Pdx1 protein levels is regulated by glucose. The increase in Pdx1 protein level that followed shRNA-mediated Pclf1 silencing in Min6 cells (Figure 2B and Figure 8A) was observed in cells cultured in low-glucose-containing medium (5.5 mM). Although Min6 cells chronically maintained in high glucose exhibit levels of Pclf1 similar to those maintained in low glucose (Supplemental Figure 7), Pclf1 silencing in Min6 cells cultured in high-glucose medium (25 mM) did not significantly alter endogenous Pdx1 protein levels (Figure 8A), suggesting that Pclf1 modulation of Pdx1 protein accumulation is regulated by glucose. In addition, the level of Pdx1 protein was increased in high-glucose–compared with low-glucose–cultured cells, in agreement with recent findings by Humphrey et al. demonstrating that Pdx1 protein is stabilized in Min6 cells and primary islets cultured under high-glucose conditions (25). Similarly, we found that in the HEK293T overexpression system, the previously observed effect of Pclf1 and Cul3 on Pdx1 protein accumulation (Figure 1A) was amplified in cells cultured in 5.5 mM glucose as compared with standard (25 mM glucose) conditions (Figure 8B). In agreement with these findings, overexpressed Pdx1 more readily communoprecipitated with Flag-tagged Pclf1 in cells cultured in low glucose (Figure 8C), suggesting that decreased interaction between Pclf1 and Pdx1 in high-glucose conditions may underlie the increase in Pdx1 protein accumulation in those cells.

In primary mouse islets, overnight culture in high glucose (30 mM) resulted in an approximately 50% decrease in Pclf1 mRNA, which corresponded to a posttranscriptional increase in Pdx1 protein (Supplemental Figure 8, A and B). This increase in Pdx1 protein was associated with a greater than 10-fold increase in insulin mRNA level (Supplemental Figure 8C), suggesting that acute glucose regulation of Pclf1 transcription could contribute to the posttranscriptional increase in Pdx1 protein levels and increased expression of Pdx1 transcriptional targets. Taken together, these data suggest that glucose modulation of Pdx1 protein accumulation involves multiple mechanisms including regulation of levels of and interaction with Pclf1.

Discussion

Here, we demonstrate that Pclf1, a BTB domain protein partner of Pdx1, directly contributes to the accumulation of Pdx1 protein in the adult β cell and thus to the actions of Pdx1 as a critical regulator of glucose homeostasis. Our results demonstrate that Pclf1 and Cul3 cooperatively target Pdx1 for ubiquitination and proteasomal degradation and that Pclf1 deficiency increases Pdx1 protein levels and stability in Min6 cells and in the developing pancreas and adult islets in vivo. Normalization of Pdx1 protein levels in Pclf1−/−Pdx1+/− transheterozygous mice was associated with improved β cell function and mass, leading to improved glucose tolerance and insulin secretion in Pdx1+/− mice. The improvement in β cell mass was due to an attenuation of the survival defect caused by Pdx1 deficiency and associated with an improvement in ER homeostasis. These results provide genetic evidence of posttranscriptional regulation of Pdx1 as a mediator of glucose homeostasis and of β cell survival in vivo.

Pclf1 heterozygosity completely rescued β cell mass and islet insulin transcript levels in Pdx1+/− mice but mediated only a partial rescue of insulin secretion and thus glucose tolerance. Further, the defect in islet architecture observed in Pdx1+/− mice was not normalized. These discrepancies may reflect different thresholds of Pdx1 required for optimal activation of Pdx1 transcriptional targets involved in islet development, insulin secretion, and β cell growth and survival. Our analysis of Pclf1−/− embryos demonstrated that although Pclf1 is not required for pancreatic organogenesis, it does influence the formation of developing β cells; however, the spatial and temporal sequence of Pclf1 expression in the pancreas has not yet been fully elucidated, and thus the relative importance of Pclf1 in regulation of the multiple roles of Pdx1 in the developing pancreas merits further investigation.

Although our findings suggest that the improved glucose homeostasis in Pclf1−/−/Pdx1+/− mice is due to rescue of islet Pdx1 protein levels and thus improved β cell mass and function, the data also suggest a complex role for Pclf1 in the maintenance of β cell mass. Pclf1−/− mice displayed normal β cell mass, but also balanced increases in both β cell replication and apoptosis rates. Furthermore, Pclf1 heterozygosity induced an increased rate of β cell replication that was not affected in the setting of Pdx1 heterozygosity, indicating roles of Pclf1 in β cell replication, and perhaps apoptosis, that may be independent of its regulation of Pdx1 protein accumulation. In addition, Pclf1 heterozygosity had no effect on β cell replication in younger (5-week-old) mice, pointing to a role in age-dependent replication rates. The human ortholog of Pclf1, SPOP, has been implicated in the ubiquitination of multiple targets, including the polycystic kidney group protein Bmi1 (32), recently implicated in pancreatic β cell proliferation via its control of the Ink4a/Arf locus (40, 41). SPOP has also been described to ubiquitinate the proapoptotic protein Daxx (42). Thus, the contribution of Pclf1 to the β cell–replicative and apoptotic rates may be due to regulation of Bmi1, Daxx, or other, as-yet-undiscovered ubiquitination targets.
Our findings indicate that in adult β cells, Pcif1 limits Pdx1 protein accumulation and thus the expression of insulin and other gene targets critical to the maintenance of β cell mass and function. The identification of this new Pdx1 regulatory mechanism and validation of its role in vivo suggest that the therapeutic targeting of Pcif1-Cul3–mediated turnover of Pdx1 could raise Pdx1 protein levels to improve β cell function and viability in the treatment of diabetes.

**Methods**

**Animals and physiological experiments.** Mouse ES cells containing a gene trap insertion in the first intron of Pcif1 were purchased from BayGenomics (clone Bgb118). Mice were derived by injection of ES cells into blastocysts by the University of Pennsylvania Transgenic and Chimeric Mouse Facility. Animals were housed in the animal care facility at the University of Pennsylvania and maintained on a 12-hour light/12-hour dark cycle with ad libitum access to food and water. All experiments described were performed on mice of a mixed 129/Sv × C57BL/6 background. For embryonic studies, the date of vaginal plug was considered E0.5. For glucose tolerance tests, animals were fasted for 16 hours before delivery of a 2-g/kg glucose bolus. Blood glucose was measured by handheld glucometers (Freestyle/One Touch). Serum was collected during the glucose tolerance test and insulin assessed by ELISA (Chemicon). For insulin tolerance tests, animals

**Figure 7**

Pcif1 heterozygosity improves glucose homeostasis and β cell function in Pdx1+/– mice. (A) Intraperitoneal glucose tolerance of male mice (7–9 weeks old), n = 6–9 per group; *P < 0.05, Pdx1+/– versus Pcif1+/–Pdx1+/– by ANOVA. (B) Intraperitoneal glucose tolerance of female mice (10–12 weeks old), n = 6–11 per group; P < 0.001, Pdx1+/– versus Pcif1+/–Pdx1+/– by ANOVA. (C) Insulin tolerance of male mice (8–9 weeks old), P = NS for all groups. (D) Insulin tolerance of female mice (11–13 weeks old), P = NS for all groups. (E) Acute glucose-stimulated insulin secretion calculated as AUC for 15 minutes after glucose bolus (12-week-old male mice), n = 7–12 per group. *P < 0.05, **P < 0.01. (F) Glucose-stimulated insulin secretion during static incubations of isolated islets from 6- to 8-week-old male mice. Insulin secretion values were normalized to islet insulin content and expressed as fold stimulation. n = 3–6 mice per group, *P < 0.05.

**Figure 8**

Posttranscriptional regulation of Pdx1 protein level by glucose. (A) Western blot of lysates from Min6 cells maintained in 5.5 or 25 mM glucose, expressing nontargeting shRNA or shPCIF1. Blots are probed for Pdx1 and Ran (loading control). (B) Western blot of lysates from HEK293T cells maintained in 5.5 or 25 mM glucose, expressing Pdx1 in the presence or absence of overexpressed Pcif1 and Cul3. Blots are probed for Pdx1 and Ran. (C) HEK293T cells maintained in 5.5 or 25 mM glucose, transfected with plasmids expressing Pdx1 and Flag-Pcif1, and subjected to immunoprecipitation with IgG or Pdx1 antisera. Western blots are probed for Pdx1 or Flag(Pcif1). Arrow indicates migration of Flag-Pcif1; asterisk indicates heavy chain IgG signal.
were fasted for 6 hours prior to an i.p. injection of 0.75 U/kg insulin. All animal procedures were approved by the University of Pennsylvania Institutional Animal Care and Use Committee.

Pancreas morphometry. BrDu was administered in the drinking water (1 g/l) for 1 week prior to sacrifice. Five- and 16-week-old male mice were anesthetized and the pancreas dissected rapidly, weighed, and fixed overnight in 4% paraformaldehyde. Paraffin sections were stained for insulin (Linco), BrDu (US Biologicals), and TUNEL (Chemicon Apoptag Peroxidase In Situ Kit). For β cell mass measurements, whole slide images of insulin-stained sections were captured on an Aperio slide scanner and the stained area quantified using Image-Pro software (Media Cybernetics). The percentage of insulin-stained area was then multiplied by pancreas weight to obtain an estimate of β cell mass. For replication measurements, an average of 1,119 and 1,463 β cells were counted per animal in the 5- and 16-week-old cohorts, respectively. For apoptosis measurements, an average of 2,398 β cells were counted per animal. For Ki67 and pHH3 confirmation of replication measurements, an average of 2,398 β cells were counted per animal in the 5- and 16-week-old mice by collagenase digestion followed by 3–4 rounds of hand picking. For RNA and protein analysis, samples were matched for islet purity by measuring amylase and insulin transcript levels. For static incubations, 5 islets were analyzed in triplicate from each animal. Islets were incubated in glucose-free KRBB buffer for 1 hour before the addition of 1.67 mM glucose. Samples were taken after 1 hour of incubation, and glucose was added to a final concentration of 16.7 mM. After 1 hour of stimulation, secretion samples were collected and islets were harvested in islet lysis buffer and analyzed for insulin content. Insulin secretion and content were measured by ELISA (Chemicon).

Cell culture and transfections. Min6 β cells were cultured in DMEM with 5.5 mM or 25 mM glucose (Invitrogen) supplemented with 10% FBS and 1% penicillin/streptomycin. Cells were transfected as described above. After 48 hours, cells were harvested in RIPA buffer and subjected to immunoprecipitation with mouse α-Myc–coupled agarose beads or mouse IgG–coupled control beads (Santa Cruz Biotechnology Inc.). To assess Pcf1–Pdx1 interaction, HEK293T cells were transfected as described above and subjected to immunoprecipitation with goat α-Pdx1 antibody or goat IgG (Santa Cruz Biotechnology Inc.) 48 hours later.

Western blots and antisera. Proteins were separated by SDS-PAGE and immunoblotted with the following antibodies: rabbit anit-Pcf1 (26), mouse anti-tubulin (Sigma-Aldrich), rabbit anti-Pdx1 (43), mouse anti-Flag epitope (Sigma-Aldrich), rabbit anti-HA epitope (Santa Cruz Biotechnology Inc.), rabbit anti-ubiquitin (Dako), rat anti–caspase-12 (Sigma-Aldrich), and mouse anti-Ran (BD Biosciences).

RNA isolation and transcript analysis. Embryonic tissues were harvested and stored in RNalater (Ambion), then homogenized in TRIzol (Invitrogen) and processed according to the manufacturer’s instructions. Islet RNA was extracted using the RNeasy Mini Kit (QIAGEN). All samples were reverse transcribed using SuperScript (Invitrogen) and oligo(dT) for priming. Transcript analysis was performed using semi-quantitative PCR and normalized to the Hprt transcript as an internal control. For isolated islets, samples were purity matched by comparing insulin and amylase transcripts.

Statistics. Data are presented as mean ± SEM. Differences between groups were compared by 2-tailed Student’s t tests and considered significant when P values were less than 0.05. Group measurements of glucose and insulin tolerance were compared by factorial ANOVA.

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