RNA m6A reader IMP2/IGF2BP2 promotes pancreatic β-cell proliferation and insulin secretion by enhancing PDX1 expression

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ABSTRACT

Background: Type 2 diabetes (T2D) is a common metabolic disease. Variants in human IGF2 mRNA binding protein 2 (IMP2/IGF2BP2) are associated with increased risk of T2D. IMP2 contributes to T2D susceptibility primarily through effects on insulin secretion. However, the underlying mechanism is not known.

Methods: To understand the role of IMP2 in insulin secretion and T2D pathophysiology, we generated Imp2 pancreatic β-cell specific knockout mice (Imp2KO) by recombining the Imp2lox allele with Cre recombinase driven by the rat insulin 2 promoter. We further characterized metabolic phenotypes of Imp2KO mice and assessed their β-cell functions.

Results: The deletion of IMP2 in pancreatic β-cells leads to reduced compensatory β-cell proliferation and function. Mechanically, IMP2 directly binds to Pdx1 mRNA and stimulates its translation in an m6A dependent manner. Moreover, IMP2 orchestrates IGF2-AKT-GSK3β-PDX1 signaling to stable PDX1 polypeptides. In human EndoC-bH1 cells, the over-expression of IMP2 is capable to enhance cell proliferation, PDX1 protein level and insulin secretion.

Conclusion: Our work therefore reveals IMP2 as a critical regulator of pancreatic β-cell proliferation and function; highlights the importance of posttranscriptional gene expression in T2D pathology.

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Keywords IMP2/IGF2BP2; T2D; Insulin secretion; m6A; Post-transcriptional gene expression regulation

1. INTRODUCTION

Human type 2 diabetes (T2D) is a common metabolic disease attributable to both environment and genetics [1]. In 2007, three independent genome-wide association studies (GWAS), WTCCC, Fusion, and DGI scans, reported a strong association between IMP2 genetic variations and T2D [2–4]. These GWAS each identified a cluster of single-nucleotide polymorphisms (SNPs) in the second intron of IMP2 gene as the basis for the T2D association [2–4]. To date, GWAS have identified ~100 SNPs in the second intron of the human IMP2 gene associated with increased risk for T2D [5–10]. IMP2 contributes to T2D susceptibility primarily through effects on insulin secretion [11–13]. Any strong evidence for IMP2 association with insulin sensitivity, a primary component of T2D susceptibility, is strikingly absent [11–13]. In line with these discoveries, IMP2 mRNA and protein are readily detected in human islets and pancreatic β cells [14] (microarray data from the Diabetes Genome Anatomy Project), and IMP2 transcript levels are significantly altered in the islets of T2D patients [15]. Notably, in 2019, two independent studies provided evidence that the association of the IMP2 intronic SNPs with T2D is affected through modification of the expression of IMP2 itself instead of neighboring genes [16,17]. These human findings have further confirmed that IMP2 is a T2D causal gene. IMP2 belongs to a small IMP protein family (IMP1-3) and is composed of six canonical RNA-binding domains: two RNA recognition motif (RRM) domains and four K homology (KH) domains [18]. IMPs have been recognized as “onco-fetal” proteins as they are highly expressed in development and carcinogenesis [19–24]. Distinct from IMP1 and IMP3, which are expressed at very low levels in most adult tissues, IMP2 expression is broadly maintained postnatally [25,26]. IMPs play key roles in post-transcriptional regulation of RNAs, in a cell typespecific manner, through ribonucleoprotein complex (RNP) [27]. They participate in the life cycle control of a wide variety of RNAs, including long noncoding RNAs and mRNAs [28–34]. The conserved IMP binding sites are enriched in “GGAC”, which is highly overlapped with N6-methyladenosine (m6A) motif [35–37]. IMPs preferentially recognize m6A-modified mRNAs and regulate their stability and translation [38]. Recent studies have suggested that the m6 A modification plays a critical role in the regulation of T2D [38,39]. For...
example, the increased expression of m6A methylation upregulates the IGF1 –AKT-PDX1 pathway in human β-cells, which ultimately inhibits cell-cycle arrest and protects insulin secretion [40]. Therefore, m6A modulators are potential therapeutic targets for maintaining glucose metabolism and insulin functions in T2D.

Nevertheless, nothing was known about the metabolic functions of IMP2 when it was first nominated as a human T2D-associated gene. To understand the role of IMP2 in T2D pathophysiology, we previously characterized global Imp2-deficient mice [26]. Imp2 null mice have decreased lean and fat body mass, increased energy expenditure, reduced insulin secretion coupled with improved insulin sensitivity, and glucose tolerance [26]. The complex metabolic effects of global Imp2 deficiency make it impossible to clarify Imp2’s function in each major metabolic tissue and to provide a foundation for patient care. Guided by human studies, we decided to further focus on the islet-specific function of IMP2 and its function in insulin secretion. Here, we report that IMP2 pancreatic β-cell-specific deletion mice (IMP2KO) have reduced compensatory β-cell proliferation and insulin secretion in response to the increased insulin demand of high-fat diet (HFD) feeding. We identified IMP2-binding mRNAs in islets, uncovered alterations of mRNAs and polypeptides from IMP2KO mice, confirmed the binding of IMP2 protein to Igf2, Pdx1 and many other islet transcripts, demonstrated that IMP2 regulates PDX1 mRNA translation in an m6A dependent manner, and showed that Imp2 regulates insulin secretion by orchestrating Igf2-Akt-Gsk3-Pdx1 signaling. Moreover, expression of IMP2 in human EndoC-βH1 cells promotes cell proliferation, PDX1 expression and Igf2-Akt-Gsk3 signaling. Based on the above evidence, we conclude that the ablation of Imp2 in mouse pancreatic β cells leads to impaired insulin secretion recapitulating human T2D traits.

2. RESULTS

2.1. Deletion of IMP2 in mouse pancreatic β-cells causes reduced compensatory β-cell proliferation and insulin secretion

IMP2 is expressed at moderate levels in most mouse tissues, including liver, muscle, fat, and brain [25]. In the pancreas, IMP2 is highly expressed in islets, particularly in insulin-producing β cells (Figure 1A). As the causal T2D risk alleles at the investigated T2D risk locus are correlated with reduced insulin secretion and lower islet IMP2 mRNA expression [16,17], we hypothesized that the decreased IMP2 activity in pancreatic β-cells contributes to the T2D pathophysiology. To test this hypothesis, we generated Imp2 pancreatic β-cell-specific knockout mice (IMP2KO) under RIP2 promoter (Supplementary Figure 1a), which showed specific reduced expression of Imp2 in isolated islets (Figure 1B). IMP2KO mice have normal body weight, body composition, physical activity, energy expenditure and serum lipid levels (Supplementary Table 1) as littermate controls. On normal calorie diet (NCD), HFD induced a 1.3-fold increase in islet mass of βIMP2KO mice (NCD: 1.84 ± 0.09 mg, HFD: 2.38 ± 0.07 mg), much less than a 3.0-fold increase in islet mass (NCD: 2.06 ± 0.12 mg, HFD: 6.23 ± 0.19 mg) in RIP2-Cre mice (Figure 1B). Although βIMP2KO mice have more small islets and less big islets (Figure 2C), the average islet diameter is not different due to the large variation (Figure 2D). This suggests that significantly decreased islet mass from HFD-fed βIMP2KO mice was primarily due to reduced β-cell number instead of cell size. As IMP2 is known to promote cell proliferation in various cell types, we next analyzed β-cell proliferation by BrdU incorporation and immunofluorescence staining of 6-week-old, HFD-fed mice. The number of BrdU+ β-cells from βIMP2KO mice was reduced 70% compared to age-matched RIP2-Cre controls (Figure 1E), indicating that the loss of Imp2 led to impaired compensatory β-cell proliferation. Next, we performed glucose-stimulated insulin secretion (GSIS) using isolated islets. Although the size-matched islets from βIMP2KO and RIP2-Cre mice secrete similar insulin levels upon low glucose (2.8 mM) stimulation, βIMP2KO islets released approximately half the amount of insulin as controls after high glucose (16.7 mM) stimulation (Figure 1F). Therefore, deleting Imp2 in mouse β-cells impairs high glucose-stimulated insulin secretion. Based on all data above, we conclude that mouse pancreatic β-cell-specific Imp2 is required for the full, compensatory β-cell proliferation and glucose stimulated insulin secretion in response to elevated insulin demand of HFD feeding.

2.2. Imp2 promotes insulin secretion through post-transcriptional regulation

To understand the molecular mechanism through which Imp2 contributes to β-cell proliferation and function, we conducted RNA-seq and Imp2 RNA immunoprecipitation sequencing (RIP-seq) using islets from 8-week-old mice fed on HFD. RNA-seq identified total 30,636 transcripts and the expression levels of 1,291 transcripts were altered significantly more than 2 folds (Supplementary 2a). Consistent with reduced β-cell proliferation observed in βIMP2KO mice, RNA sequencing analyses showed downregulation of key cell-cycle genes (Ccn1, Ccn2, Cdk4) and increased expression of the cyclin-dependent kinase inhibitor (p21). Enriched pathway analyses of differentially expressed genes (FDR < 0.10) revealed response to insulin, cell proliferation/cell cycle, and insulin secretion as the most commonly affected biological events in βIMP2KO islets (Figure 3A). To identify Imp2-binding RNAs, each IP-associated RNA was corrected for abundance in the total extract and divided by the corrected IP value from βIMP2KO islets. This nominated ~1,600 islet mRNAs as Imp2 binding partners (Supplementary Figure 2a–b) that are highly enriched for insulin secretion, RNA processing, and cell cycle (Figure 3B).

Since IMP2 plays a key role in post-transcriptional gene expression regulation, we next performed liquid chromatography tandem mass spectrometry (LC-MS/MS)-based proteomics using islets from the same cohort of mice for RNA-seq and RIP-Seq. The analysis resulted in the confident identification of 29,021 different tryptic peptides covering 3,365 proteins (≥2 unique peptide identifications per protein). As expected, three major islet hormones (insulin, glucagon, and somatostatin) were detected, as well as various β-cell enriched secretory products, ion channels, and transcription factors. In addition, significant proteome coverage of islet metabolic enzymes and cellular pathways was observed. Quantitative measurement of relative polypeptide abundances showed significant differences of a total of 1,252 proteins. Pathway analyses of differentially expressed proteins revealed a highly significant enrichment of the insulin secretion, cell proliferation/cell cycle, and IGF1-AKT pathway (Figure 3A). Although the latter was not evident in the transcriptome analysis, the impairment...
of insulin/IGF—AKT signaling is a well-known mechanism associated with T2D [40–42]. Guided by the results of genome-wide research, we further validated the abundances of critical RNAs and polypeptides for β cell function (Figure 3C–E). Consistent with RNA-Seq and proteomic studies, 50% of validated Imp2 targets did not show significant alternation at either mRNA or protein level. As expected, Igf2 mRNA was confirmed as an Imp2 binding partner and reduced in βIMP2KO islets (Figure 3F).

Figure 1: The pancreatic β-cell specific deletion of IMP2 impairs insulin secretion on HFD feeding. A. Immunostaining of insulin and IMP2 in 8-week-old mouse pancreas. Scale bar: 80 μm B. Relative IMP2 protein abundance in control (black) and βIMP2KO (red) mice (n = 4 pairs). C–F. 6-hour fast blood glucose, serum insulin, C-peptide and glucagon from 12-week-old male control (black) and βIMP2KO (red) mice (n = 6 pairs), G–H. Glucose tolerance test of 14-week-old, HFD-fed male control (black) and βIMP2KO (red) mice (n = 8 pairs). Data are means ± SD. Two-way ANOVA test. *P < 0.05; **P < 0.01.

Figure 2: IMP2 deletion reduces β-cell proliferation and function. A. The representative images of isolated islets from HFD-fed, 16-weeks-old control and βIMP2KO mice (n = 6 pairs). B. The islet mass from 16-weeks-old NCD (n = 7 pairs) and HFD fed mice (n = 7 pairs). C–D. The distribution of islet size and area from HFD-fed control (black) and βIMP2KO (red) mice (n = 6 pairs). E. The percentage of BrdU labeled β-cells from 6-week-old mice (n = 8 pairs). F. the glucose stimulated insulin secretion of isolated 16-week-old, HFD-fed control (black) and βIMP2KO (red) mice (n = 8 pairs). Data are means ± SD. Two-way ANOVA test. *P < 0.05. **P < 0.01.
Notably, validated as one of imp2 targeted mRNAs (Figure 3C), the Pdx1 transcriptional level is similar between β-imp2ko and control (Figure 3D). However, the Pdx1 polypeptides significantly decreased in β-imp2ko (Figure 3G), indicating imp2 regulates Pdx1 mRNA translation and/or Pdx1 polypeptide stability. In line with reduced Pdx1 protein levels, the mRNA abundances of major Pdx1 transcriptionally regulated genes were reduced by 12–54% (Figure 3H).

2.3. IMP2 directly binds PDX1 mRNA and promotes its translation in an m6A-dependent manner

As IMPs preferentially bind m6A-modified mRNAs [36], we next hypothesized that IMP2 regulates PDX1 mRNA translation through m6A recognition. We first confirmed the direct binding of IMP2 with PDX1 mRNAs by IMP2 RIP-qPCR (Figure 4A), and demonstrated m6A modification of PDX1 mRNAs through PDX1-specific m6A assays using human islets (Figure 4B). Notably, the PDX1 stop codon containing region which showed dramatic hypomethylation in T2D patients [40] has most abundant m6A modifications (Figure 4C) relative to other regions of PDX1 transcript (such as 5'UTR). As conserved IMP2 binding motifs significantly overlaps with m6A modification sites (Figure 4D), this locus is also enriched for IMP2 binding. To determine whether the stop codon-containing region is critical for IMP2-guided PDX1 mRNA translation, we inserted 200 nt PDX1 mRNA wild-type or mutant sequence (Supplementary Figure 3a) flanking stop codon into firefly luciferase reporters (Supplementary Figure 3b) into EndoC-βH1 cells [43] stably expressing Flag-IMP2 or Flag-GFP. IMP2 RIP-qPCR demonstrated a strong binding of IMP2 with wild-type PDX1 reporter but not with the mutant form (Figure 4F). Moreover, IMP2 dramatically enhanced luciferase activity, which was also dependent on the presence of wild-type m6A motifs surrounding the stop codon of PDX1 mRNA (Figure 4G) as the mutant m6A motif form abolished the IMP2-stimulated luciferase activity. We further confirmed the effect of IMP2 stimulated PDX1 mRNA translation in vitro using rabbit reticulocyte lysate as Flag-IMP2-induced a significant increase of PDX1 wild-type reporter activity in a dose-dependent manner (Figure 4H). This effect was greatly impaired by mutations in the m6A consensus sites/IMP2 binding sites in the PDX1 stop codon-containing region (Figure 4H). Taken together, these data demonstrate that IMP2 preferentially binds m6A-modified PDX1 mRNAs and promotes their translation.

2.4. IMP2 stabilizes PDX1 polypeptides by enhancing IGF2–AKT–GSK3 signaling

In addition to reduced PDX1 mRNA translation, the decreased Pdx1 proteins in β-imp2ko mice could also be attributed to post-translational regulation. Indeed, the impairment of IGF2-AKT signaling promotes PDX1 protein degradation and decreases β-cell mass and function [44,45]. The nominated reduced IGF1–AKT signaling by proteomics studies promoted us to further explore the post-translational mechanism. We previously published that, in addition to its ability to promote IGF2 production, IMP2 strongly stimulates IGF2 action [46]. In mouse muscles, the deficiency of IGF2 results in impaired IGF2 signaling through diminishing the phosphorylation of insulin/IGF1 receptors, reducing Akt1 activation and disinhibiting Gsk3β [30]. Therefore, we hypothesized that IMP2 regulates IGF2 signaling, in a similar way, to promote insulin secretion in pancreatic β cells. Islets from hfd-fed β-imp2ko and control mice were stimulated with exogenous IGF2. Immunoblots of total lnsR and lrs polypeptides showed no difference (Figure 5A–B); however, the phosphorylated forms of both proteins reduced by 35–40% (P < 0.05) (Figure 5A–B) in β-imp2ko islets. Similarly, the total Akt polypeptide levels are also not different. However, phos-Akts (Thr308 and Ser473) are both markedly reduced in β-imp2ko islets (Figure 5A–B). We next examined Akt-catalyzed inhibitory phosphorylation of Gsk3β. While Gsk3β (Ser21) phosphorylation is not dramatically different (Figure 5C–D), the corresponding phosphorylation of Gsk3β (Ser9) is reduced by ~50%, pointing to

![Figure 3: IMP2 promotes β-cell proliferation and function. A. GO analyses of differentially expressed islet proteins (red) and RNAs (blue). B. Functions of IMP2 binding RNAs. C–E. Confirmation of abundances of selected IMP2 binding mRNAs (c), total mRNAs (d) and proteins (e) from 8-week-old, HFD fed control and β-imp2ko animals (n = 5 pairs). F. The islet and serum IGF2 levels from 12-weeks-old HFD-fed mice (n = 7 pairs). G. Western blot of PDX1 protein from isolated islets of 12-weeks-old, HFD-fed mice (n = 3 pairs) H. the relative expression of PDX1 transcriptionally regulated genes. RNA was isolated from 12-weeks-old, HFD-fed control and β-imp2ko mice (n = 5 pairs). Data are means ± SD. T-test *P < 0.05;**P < 0.01.](image-url)
higher activity of Gsk3β in β-pancreatic H1 cells (Figure 5C–D). Most importantly, the Gsk3β-catalyzed Pdx1 degradation is increased more than 3-fold in β-pancreatic H1 islets over controls measured by the reduced half-life of Pdx1 protein in β-pancreatic H1 islets (Figure 5E). Taken together, the above data show that the IMP2 deficiency in mouse islets led to impaired insulin secretion through the downregulation of the IGF2–AKT–GSK3β signaling pathway.

2.5. Overexpression of IMP2 in EndoC-β-H1 cells promotes insulin secretion

We found that IMP2 is highly expressed in human islets but cannot be detected by immunoblotting in β-pancreatic H1 cells (Figure 6A), a commonly used human β-cell line. In addition, EndoC-β-H1 cells have much lower capability of insulin secretion compared to human islets [43]. We therefore hypothesized that the expression of IMP2 is able to promote insulin secretion in EndoC-β-H1 cells. Next, we transfected EndoC-β-H1 cells with Flag-IMP2 or Flag-GFP, and selected stable cells. EndoC-β-H1 cells stably expressing IMP2 showed accelerated cell proliferation (Figure 6B), a marker of β-cell function progressively declines over time, blood glucose rises. This study focuses on the pancreatic β-cell specific function of IMP2. We provide experimental evidence for the molecular mechanisms of IMP2 in β-cell biology which highlights the significance of post-transcriptional gene expression regulation in T2D pathophysiology.

IMP2 was first nominated as a human T2D-associated gene in 2007 without any experimental evidence supporting its role in metabolic regulations [2–4]. To understand the function of IMP2 in T2D pathogenesis and provide a foundation for patient care, we generated and studied the Imp2 global knockout mice [26]. The global Imp2-null mice exhibited a normal birth size and a body weight similar to that of control littermates until weaning [26]. However, they gain less weight due to impaired Igf2 and Pdx1 expression, stimulates cell proliferation, enhances IGF2 and PDX1 expression, promotes IGF2-AKT-GSK3β signaling, and increases insulin secretion. Collectively, our study revealed an unexpected function of Imp2 in pancreatic β-cells through promoting Pdx1 mRNA translation and enhancing Igf2–Akt–Gsk3β signaling, which impairs insulin secretion and β-cell proliferation upon the ablation of Imp2.

3. DISCUSSION

Human T2D is widely considered as a chronic and progressive disease without cure [47]. As insulin-producing β-cell function progressively declines over time, blood glucose rises. This study focuses on the pancreatic β-cell specific function of IMP2. We provide experimental evidence for the molecular mechanisms of IMP2 in β-cell biology which highlights the significance of post-transcriptional gene expression regulation in T2D pathophysiology.

IMP2 was first nominated as a human T2D-associated gene in 2007 without any experimental evidence supporting its role in metabolic regulations [2–4]. To understand the function of IMP2 in T2D pathogenesis and provide a foundation for patient care, we generated and studied the Imp2 global knockout mice [26]. The global Imp2-null mice exhibited a normal birth size and a body weight similar to that of control littermates until weaning [26]. However, they gain less weight due to impaired Igf2 and Pdx1 expression, stimulates cell proliferation, enhances IGF2 and PDX1 expression, promotes IGF2-AKT-GSK3β signaling, and increases insulin secretion. Collectively, our study revealed an unexpected function of Imp2 in pancreatic β-cells through promoting Pdx1 mRNA translation and enhancing Igf2–Akt–Gsk3β signaling, which impairs insulin secretion and β-cell proliferation upon the ablation of Imp2.
hepatic steatosis, superior insulin sensitivity, reduced insulin secretion) is relevant to its role in human T2D; it is also unclear which is a direct consequence of Imp2 deficiency in the target organ (adipocyte, liver, skeletal muscle, islet). Seeking to understand the tissue(s) in which Imp2 has its primary impact, we generated a series of mice lacking Imp2 in specific organs. The postnatal inactivation of Imp2 in skeletal muscle decreased accrual of muscle mass after weaning and reduced wheel-running activity [30]. However, their whole-body glucose...
metabolism and insulin sensitivity are unaltered [39], in contrast to the very pronounced improvements seen in global Imp2 deficiency [26]. The primary finding of elimination of the hepatocyte Imp2 is increased intrahepatic triglyceride deposition in HFD-fed mice, which developed slight hypertriglyceridemia and hyperglycemia by 30 weeks of age [31]. This indicates that the marked protection against fatty liver observed in global Imp2 deficiency mice is entirely due to their reduced adiposity. Although we have plausibly attributed the reduced insulin secretion of the global Imp2-null mice to their enhanced insulin sensitivity, the majority of human studies have identified impaired insulin secretion as the predominant defect of IMP2 T2D risk alleles [16,17]. In the current study, we showed that β-cell specific deletion of Imp2 in mice resulted in impaired compensatory β-cell proliferation and function due to the increased demand of insulin secretion by HFD. This study revealed that pancreatic β-cell intrinsic functions of Imp2 are completely obscured by the tremendous insulin sensitivity of the global Imp2-null mice caused by their dramatically reduced fat mass. While studying Imp2 murine models, we noticed that many phenotypes are diet-dependent. For example, the impaired insulin secretion and reduced β-cell proliferation in βIMP2KO mice only showed on HFD but not on NCD. This phenomenon is likely due to Imp2 functioning as a post-transcriptional gene regulator, which is critical for stress conditions [48], such as HFD feeding. As Imp2 is not critical under normal conditions, the Imp2 deficiency has less/no impact on whole-body metabolism on NCD.

The data collected from all Imp2 murine models has shown that GWAS signals can be moved forward into biomedical understanding by embracing all the technologies at our disposal. To date, no deleterious mutations encoding IMP2 have been found in human populations; this knowledge would be informative to the future T2D research. In line with our data collected from murine models, the recent human T2D research grouped Imp2 into a “proinsulin cluster” featured with decreased proinsulin and reduced β-cell function [49]. The blood Imp2 mRNA levels were positively associated with fasting insulin from people without diabetes [50] indicating that IMP2 may be involved in the regulation of β-cell function before onset of T2D. Despite these emerging data, it remained unclear what these findings brought to our greater understanding of human T2D. The caveat is that, despite a large number of variants discovered by GWAS, the total of associated variants explains only a small proportion (10%) of the heritability of T2D.

Recently, mRNA m6A methylation has been reported to contribute significantly to the pathogenesis of T2D and is a better mark to metabolic stress such as HFD feeding [52]. In this study, we demonstrated that Imp2 is required to promote Igf2–Akt–Gsk3β–Pdx1 signaling in response to HFD feeding. Taken together, we propose that reduced IMP2 expression in pancreatic β-cells contributes to the pathophysiology of human T2D. Therapeutic targeting of IMP2 in a β-cell-specific manner in combination with current therapeutic agents might be a new avenue to counter the decreased m6A levels in T2D islets and to promote β-cell proliferation and function.

4. METHODS

4.1. Animal studies

All animal procedures were approved by the Institutional Animal Care and Use Committee of Massachusetts General Hospital and were performed in accordance with the National Research Council guidelines for laboratory animal care. RIP2-Cre transgenic mice were purchased from Jackson Lab (Stock Number #003573), which has a 668 bp fragment of the rat insulin II promoter. βIMP2KO and RIP2-Cre control mice were maintained on C57BL/6J background in a specific pathogen-free facility with 12:12 light:dark cycle and fed irradiated chow (Prolab SP75 Isopro 3000; 5% crude fat; PMI nutrition international) or a HFD (D12492i; 60 kcal% fat; Research Diets, Inc.).

4.2. Glucose and insulin tolerance tests

For glucose tolerance tests, mice were fasted overnight. Twenty percent D-glucose (Sigma) (1 g/kg body weight) was administered by intraperitoneal injection. At 0, 30, 60, 90, and 120 minutes after administration, blood was collected by tail vein bleeding. Glucose levels were measured by a One Touch Ultra AlphaTrak2 glucometer (Zoetis, Parsippany, NJ). For insulin tolerance tests, HFD-fed mice were fasted for 6 h. Human insulin (Eli Lilly) at 0.75 U/kg was injected intraperitoneally. Blood was drawn from the tail vein at 0, 30, 60, 90, and 120 minutes after injection, and glucose levels were measured.

4.3. Serum analyses

Blood was collected into EDTA-coated tubes (Sarstedt, Newton, NC). Serum was separated by centrifugation at 4 °C, frozen in liquid nitrogen, and assayed at the Vanderbilt University Mouse Metabolic Phenotyping Center (MMPC). The blood Igf2 concentration was measured using a mouse Igf2 enzyme-linked immunosorbent assay (ELISA) kit (catalog number EK0381; Boster Biological Technology Co., Ltd., CA) according to the manufacturer’s instructions.

4.4. Mouse islet isolation and quantification

Mouse islet isolations were performed at MGH islet transplant core. Briefly, mice were anesthetized and their pancreases were infused with liberase (Roche) and dissected. Following incubation at 37 °C, the digested pancreases were washed, filtered, and run on a sucrose gradient. The islet cell counter (ICC) used for this study (Biorep Technologies) is a machine vision system that includes both hardware and software designed specifically for automated islet counting [53]. For each sample, a single high-resolution image is taken to assess the amount and purity of the islet cell sample. ICC uses a fully automated counting method implemented in LabVIEW software [Laboratory Virtual Instrumentation Engineering Workbench (LabVIEW); National Instruments, Austin, TX, USA]. Islet samples were prepared and stained with diphenylthiocarbazone. The software uses pixel coloration to separate islets from acinar tissue and a segmentation-based algorithm to delineate each individual islet and calculate its area. The IEQ (mass) contribution of each islet is then obtained using the 50-μm increment group classification and standard IEQ group values. Estimated islet
mass (total IEQ), number, purity, and size group distributions, as well as the high-resolution image, are saved in the final output as both Word document and Adobe pdf files.

For islet histomorphometry study, number of islet/mm² of pancreatic area, mean area of islet and number of β-cell nuclei/mm² of insulin-positive area were carried out for each group of animals.

4.5. Insulin secretion assay

After overnight culture in RPMI 1640 medium (11 mmol/L glucose and 10% FBS), triplicate samples of 10 equilibrated islets for each mouse placed in wells of a 24-well plate were sequentially incubated with 2.8 and 16.8 mmol/L glucose in Krebs–Ringer buffer (16 mmol/L HEPES and 0.1% BSA, pH 7.4) for 50 minutes. Supernatant fractions and cell lysates were frozen until assayed for insulin.

4.6. BrdU treatment

Mice were provided BrdU in drinking water (80 mg/ml) after weaning at 4-week-old and received intraperitoneal injections of BrdU (100 mg/kg body weight) every three days for two weeks. Mice were sacrificed at 6 weeks old and β-cell proliferation was assessed by co-immunostaining sections of each pancreas sample with BrdU, insulin, and DAPI. β-cell proliferation was assessed by analyzing images acquired from 30 islets per animal.

4.7. Pancreas immunomorphology and analyses

Mouse pancreas was collected and fixed in 3.7% formaldehyde at 4 °C overnight, followed by paraffin embedding. Four micron-thick slides were cut and subjected to immunostaining. Slides were heated in 10 mM sodium citrate, followed by blocking with donkey serum and incubated with various primary antibodies (Supplementary Table 2). The specific signal was detected using fluorescence-conjugated secondary antibodies.

4.8. RNA-seq

RNA from 8-week-old, HFD-fed mouse islets was extracted using a Qiagen RNase kit. RNA-seq libraries were constructed from poly[A]-selected RNA sequenced on an Illumina HiSeq2500 instrument, resulting in approximately 30 million reads per sample on average. STAR aligner was used to map sequencing reads to transcripts in the mouse mm10 reference genome. Read counts for individual transcripts were produced with HTSeq-count, followed by the estimation of expression values and detection of differentially expressed transcripts using EdgeR.

4.9. Proteomic studies

Purified islets from 8-week-old, HFD-fed mice were processed and analyzed through the Thermo Fisher Scientific Center for Multiplexed Proteomics at Harvard Medical School. Peptides derived from digestion using LysC and trypsin were labeled with Tandem Mass Tag 8-plex reagents and fractionated. Multiplexed quantitative mass spectrometry data were collected on an Orbitrap Fusion mass spectrometer operating in a MS3 mode using synchronous precursor selection for tryptic data were collected on an Orbitrap Fusion mass spectrometer.

4.10. IMP2 RIP-seq

Purified islets from 8-week-old, HFD-fed mice were extracted directly using a tissue lyser (Qiagen) in ice-cold lysis buffer (140 mM KCl, 1.5 mM MgCl₂, 20 mM Tris–HCl at pH 7.4, 0.5% Nonidet P-40, 0.5 mM dithiothreitol, 1 U/μl RNase inhibitor, one complete EDTA-free protease inhibitor cocktail tablet) for 10 min. The lysates were centrifuged for 10 min at 14,000 rpm, and the supernatant was transferred to a fresh 1.5-ml tube. Total protein was measured by a Bradford assay, and 5 mg of the cytoplasmic lysate protein was subjected to immunoprecipitation. Lysates were incubated with 500 μl of protein A Dynabeads magnetic beads precoated with IMP2 antibody and incubated for 6 hours at 4 °C with rotation. Dyna beads were extensively washed with lysis buffer five times and digested with DNase I and protease K. RNA was extracted with phenol-chloroform and precipitated with ethanol. RNA sequencing was performed to examine RNAs associated with cytoplasmic IMP2 in islets.

4.11. Protein assays

Proteins were extracted using a tissue homogenizer (Qiagen). Purified islets from 8-week-old, HFD-fed mice were homogenized in ice-cold buffer (20 mM Tris [pH 7.5], 2.7 mM KCl, 1 mM MgCl₂, 1% Triton X-100, 10% [wt/vol] glycerol, 1 mM EDTA, 1 mM dithiothreitol) supplemented with protease (Thermo Scientific) and a phosphatase inhibitor cocktails (Millipore). Samples were then centrifuged at 13,000 rpm for 10 min at 4 °C, and the supernatants were collected. The protein content of the supernatant was determined using a bicinchoninic acid (BCA) assay (Thermo Scientific). Aliquots of each extract containing 50 μg protein were loaded for Western blotting. Proteins were resolved on a 4–12% gradient SDS Bis-Tris gel (Invitrogen, Carlsbad, CA). Antibodies for immunoblotting are listed in Supplementary Table 2. For protein half-life assay, 250 mM cycloheximide was used to inhibit mRNA translation; protein half-life was estimated from immunoblot of extracts prepared before and at various times after cycloheximide addition.

4.12. Gene-specific m6A qPCR

m6A modifications of specific genes were determined using Magna MeRIP m6A Kit (Millipore, Billerica, MA). In short, 1 mg of total RNA was sheared to 100 nt, incubated with anti-m6A antibody (Synaptic Systems) or mouse IgG (Millipore) with RNase inhibitors at 4 °C. Methylated RNA was immunoprecipitated with beads, eluted by competition with free m6A, and recovered with an RNAeasy kit (Qiagen). Relative enrichment of m6A was calculated by normalizing to input.

4.13. Human samples

Human islet samples were obtained from the Pancreatic Islet Isolation Core of Boston Area Diabetes Endocrinology Research Center (BADRC). The detailed donor information is in Supplementary Table 3.

4.14. EndoC-βH1 cell culture

Culture flasks were coated with DMEM (glucose 4.5 g/L, Gibco) containing PS (1%, Gibco), fibronectin (2 μg/ml; Gibco), and extracellular matrix (1% vol/vol, Sigma) and incubated for 2 hours in 5% CO₂ at 37 °C before the cells were seeded. EndoC-βH1 cells were grown on Matrigel/fibronectin-coated culture flasks containing DMEM (glucose 1 g/L), BSA fraction V (2% wt/vol) (Roche), 2-mercaptoethanol (50 mM, Sigma), nicotinamide (10 mM, Sigma), transferrin (5.5 μg/ml Sigma), sodium selenite (6.7 ng/ml Sigma), PS (1%). For insulin secretion assays, EndoC-βH1 cells were starved for 2 h in Krebs ringer buffer (KR) containing NaCl (115 mM), NaHCO₃ (24 mM), KCl (5 mM), MgCl₂ (1 mM), CaCl₂ (1 mM), HEPES (10 mM), BSA (0.2% wt/vol), and 0.5 mM glucose. Static insulin secretion assays were then initiated by adding KRB containing 2.8 mM or 16.7 mM of glucose for 1 hour. Aliquots of supernatant were removed, centrifuged to discard dead
cells, and stored for analysis in ice-cold acid ethanol that was added to extract insulin content from cells. Insulin secretion and content were measured by the human insulin ELISA (Merodia) according to the manufacturer’s instructions. For stable cell lines, EndoC-βH1 cells were transduced with lentiviral particles. Puromycin (1 μg/ml) was added to the culture media to select cells stably expressing Flag-IMP2 or Flag-GFP. Transient transfections were performed by mixing cells with Lipofectamine according to the manufacturers’ instructions.

4.15. Dual luciferase reporter assay
Luciferase activity was examined by a dual luciferase reporter assay using the dual luciferase reporter assay kit (Promega) on a TD-20/20 luminometer (Turner Designs). To generate the PDX1 firefly luciferase reporter construct, DNA fragments of wild-type and mutant sequences were synthesized by Integrated DNA Technologies (Coralville, Iowa) and cloned into the pcDNA5/TO vector (Invitrogen). The EndoC-βH1 cells were transiently transfected with luciferase vector together with Renilla luciferase control (Promega). Luciferase activity was measured at 36 hours post-transfection.

4.16. Statistical analysis
Comparisons between the mean ± SD of two groups were calculated using two-way analysis of variance (ANOVA) or Student’s unpaired two-tailed t test. P values of 0.05 or less were considered statistically significant.

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CONFLICT OF INTEREST
All authors do not have conflict of interest.

APPENDIX A. SUPPLEMENTARY DATA
Supplementary data to this article can be found online at https://doi.org/10.1016/j.molmet.2021.101209.

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