Examining How the MAFB Transcription Factor Affects Islet β-Cell Function Postnatally

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The sustained expression of the MAFB transcription factor in human islet β-cells represents a distinct difference in mice. Moreover, mRNA expression of closely related and islet β-cell–enriched MAFA does not peak in humans until after 9 years of age. We show that the MAFA protein also is weakly produced within the juvenile human islet β-cell population and that *MafB* expression is postnatally restricted in mouse β-cells by de novo DNA methylation. To gain insight into how MAFB affects human β-cells, we developed a mouse model to ectopically express *MafB* in adult mouse β-cells using *MafA* transcriptional control sequences. Coexpression of MafB with MafA had no overt impact on mouse β-cells, suggesting that the human adult β-cell MAFA/MAFB heterodimer is functionally equivalent to the mouse MafA homodimer. However, MafB alone was unable to rescue the islet β-cell defects in a mouse mutant lacking MafA in β-cells. Of note, transgenic production of MafB in β-cells elevated tryptophan hydroxylase 1 mRNA production during pregnancy, which drives the serotonin biosynthesis critical for adaptive maternal β-cell responses. Together, these studies provide novel insight into the role of MAFB in human islet β-cells.

Type 2 diabetes is characterized by peripheral insulin resistance and impaired pancreatic α- and β-cell activity (1). Although many distinct genetic lesions appear to contribute to disease susceptibility, islet-enriched transcription factor mutations commonly are associated with a monogenic form of diabetes termed maturity-onset diabetes of the young (e.g., HNF1α [2], HNF1β [3], PDX1 [4], MAFA [5]). As a consequence of extensive mutational analysis of these and other islet-enriched transcription factors in mice, many were shown to play essential roles in islet cell development and/or function (6). However, striking differences exist in the expression pattern of a few of these key regulators between humans and mice. For example, *MAFA* mRNA is produced at lower levels in juvenile (<9 years of age) than in adult (>29 years of age) human islet β-cells (7), whereas expression peaks soon after birth in mice (8,9). In addition, the closely related *MAFB* gene is expressed in primate islet β-cells postnatally (10) but not in rodents (8,9).

Both MafA and MafB are made relatively late during mouse islet cell development compared with other islet-enriched transcription factors (11). Thus, MafB expression begins around embryonic day 10.5 (e10.5) in both insulin-positive (i.e., β-cell) and glucagon-positive (i.e., α-cell) progenitors, whereas MafA is first detected at e13.5 and only in insulin-positive cells (8,9). In contrast, most other islet-enriched transcription factors are produced much

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earlier during development (e.g., Pdx1 [e8.5 (12)]) and within a larger fraction of the adult islet cell population (e.g., α, β, δ, PP, Pax6 [13], and FoxA1/2 [14]). MafA expression persists in the mouse islet β-cell population postnatally, whereas MafB is restricted to α-cells (8,9). However, MafB is re-expressed in a small fraction of islet β-cells during pregnancy (15). Analysis of mice that lack *MafA* or *MafB* during pancreas development has demonstrated that the *MafA* mutant has the most significant phenotype ($MafA^{\Delta pancreas}$ [16]), which is manifested as defects in β-cell activity and islet cell architecture after birth. In contrast, there is no overt effect in $MafB^{\Delta pancreas}$ islet β-cells as a result of postnatal compensation by MafA, although plasma glucagon secretion levels from α-cells are reduced (10).

Of note, human *MAFA* mRNA is made at low levels in juvenile β-cells in relation to adult islets (7), and MAFB is expressed throughout the lifetime of these cells in nonhuman primates (NHPs) and humans (7,10,17). Here, we first show that the MAFA protein is found in relatively few juvenile and adolescent human islet β-cells in relation to MAFB and that DNA methylation within the 5′ flanking region of mouse *MafB* correlates with gene silencing in β-cells. The impact of MafB on adult islet β-cells was next examined in βMafBTg transgenic mice that sustain production of this transcription factor postnatally, thus mimicking the expression pattern in humans. Although little impact was observed on coexpression of MafB with endogenous MafA in islet β-cells, production of MafB alone was unable to rescue any of the islet deficiencies of $MafA^{\Delta pancreas}$ mice. These results suggest that the juvenile $MAFB_2$ homodimer and adult MAFA/MAFB heterodimer regulators could be involved in controlling age-dependent differences in human β-cell gene expression (7). Of note, maternal βMafBTg mice displayed increased β-cell proliferation and function compared with wild-type mice. These data illustrate the many distinct ways MAFB may regulate human islet β-cells.

RESEARCH DESIGN AND METHODS

Tissue Collection and Immunofluorescence

Human pancreata from healthy juvenile (<10 years of age) donors were received in partnership with the International Institute for the Advancement of Medicine and National Disease Research Interchange for histological analysis (Supplementary Table 1). The Vanderbilt University institutional review board declared that studies on deidentified human pancreatic specimens do not qualify as human subject research.

Human and mouse pancreata were fixed in 4% paraformaldehyde in PBS and embedded in either Tissue-Plus O.C.T. (Thermo Fisher Scientific) or paraffin wax. Immunofluorescent images were obtained using the Zeiss Axio Imager M2 widefield microscope with ApoTome. Antibodies are listed in Supplementary Data.

Islet Isolation, Cell Sorting, and DNA Methylation

Mouse islets were isolated using standard islet isolation conditions (18). Islet α- and β-cells were isolated as previously described (19). α-Cells were sorted by FACS to an average purity of 80–85%. For isolation of β-cells from neonatal (postnatal day 5) and adult (age 2.5 months) mice, islets from mouse INS-1 gene promoter-green fluorescent protein transgenic mice were sorted for green fluorescent protein by FACS to an average purity of 85–95%. Cadaveric human islets and mouse islets were processed for DNA extraction and bisulfite treatment as detailed previously (19). Primers and additional details are available upon request.

siRNA-Based Knockdown, RNA Isolation, cDNA Synthesis, and Real-time PCR

Knockdown of *Dnmt3a* in Min6 cells was performed by transfection with a pool of specific targeting siRNAs or scrambled controls (Dharmacon) using Lipofectamine 2000 (Invitrogen) every 2 days (average transfection efficiency ~80%). RNA was isolated 4 days posttransfection using the RNeasy Micro Kit (QIAGEN). cDNA was generated using Superscript III Reverse Transcriptase (Invitrogen) by the oligo(dT) priming method. Real-time PCR assays were performed using the LightCycler FastStart DNA Master PLUS SYBR Green kit (Roche) and a LightCycler PCR instrument (Roche). The expression levels were normalized to cyclophilin with previously published primers (19).

Mouse islet RNA was collected using the TRIzol reagent (Life Technologies) and a DNA-free RNA kit (Zymo Research). cDNAs were produced using the iScript cDNA Synthesis Kit (Bio-Rad), and real-time PCR was performed on a LightCycler 480 system (Roche) with primers listed in Supplementary Table 2. Real-time PCR results were analyzed using the ΔΔCt method; *Gapdh* was used to normalize the mouse islet data.

Chromatin Immunoprecipitation Assays

Chromatin immunoprecipitation (ChIP) experiments with purified α- and β-cells were carried out using the micro-ChIP protocol (20). Islets from 2-month-old mice were used for MafB (Bethyl Laboratories) and Dnmt3a (Novus Biologicals) ChIP. Approximately 500 islets per data point were cross-linked using 1% formaldehyde for 10 min at 37°C and quenched using glycine. Sonication was performed for 3 × 5 min using a Diagenode Bioruptor (Diagenode Diagnostics). Approximately 4 μg antibody was incubated at 4°C overnight. Protein A Dynabeads (Thermo Fisher Scientific) were treated overnight with BSA, herring sperm, and protease inhibitors before being added to the chromatin-antibody mixture for 4 h at 4°C. After washes, chromatin elution was performed with 1% SDS and 0.1 mol/L sodium bicarbonate for 15 min followed by a 5 mol/L NaCl overnight incubation at 65°C and then proteinase K digestion (1 μg) for 1 h at 45°C. The eluted DNA was purified using the QIAGEN PCR

Purification Kit. Real-time PCR enrichment of MafB binding was calculated using, as controls, IgG and the *albumin* gene, whereas IgG and the *actin* gene were used in the Dnmt3a ChIP (21,22).

Generation of βMafBTg Transgenic Mice

Bacterial artificial chromosome (BAC) recombineering (recombination-mediated genetic engineering) was used to exchange the *MafA* coding sequences in bMQ-159k16, a BAC clone of ~120 kb spanning the single exon *MafA* gene as well as the 5′ and 3′ flanking sequences, with *MafB* single exon sequences (Fig. 3*A*). Purified linear $MafA^{BAC}$-*MafB* DNA was injected into pronuclei from B6D2 donors and then implanted into pseudopregnant recipient mice. Thirty-one pups derived from microinjection were screened for the BAC transgene, with six $MafA^{BAC}$-*MafB*. βMafBTg founders bred with C57BL/67 females, and three independent mouse lines were characterized in depth. MafB was expressed in ~87% of islet β-cells in the D line, 70% in the A line, and 36% in the E line (Supplementary Fig. 1). The analysis presented here principally used the D-expressing line, although similar results were obtained with the A line. Mouse studies were performed in compliance with protocols approved by the Vanderbilt institutional animal care and use committee.

$MafA^{\Delta\beta}$ Generation, Intraperitoneal Glucose Tolerance Testing, Blood Serum, and Islet Hormone Measurements

$MafA^{\Delta\beta}$ mice were generated by crossing floxed *MafA* ($MafA^{fl/fl}$) (8) with rat *Ins2* enhancer/promoter-driven Cre (RIP-Cre) mice (23). $MafA^{\Delta\beta}$ mice were subsequently crossed with either the βMafBTg D or the βMafBTg A lines. Intraperitoneal glucose tolerance testing (IPGTT) was performed on adult nonpregnant and pregnant mice fasted for 6 h; tail blood glucose was measured before (0 min) and at 15, 30, 60, and 120 min after intraperitoneal injection of glucose (2 mg/g body weight) prepared in sterile PBS (20% w/v). Blood serum was collected through the tail vein after a 6-h fast. Insulin measurements were conducted in the Vanderbilt Hormone Assay & Analytical Services Core. Serotonin measurements were determined in the Vanderbilt University Neurochemistry Core.

Statistics

Significance was evaluated with the Student two-tailed *t* test for the quantitative PCR (qPCR), area under the curve (AUC), and β-cell proliferation tests. Statistical significance is denoted in the figure legends.

RESULTS

The MAFA Protein Is Expressed at Very Low Levels in Islet β-Cells From Juvenile Human Donors

Human and NHP islets are distinct from rodents in many ways, including glucose-stimulated insulin secretion (i.e., basal [5.6 mmol/L] insulin secretion is higher in primates, and stimulated secretion is lower [17,24,25]), islet cell composition (i.e., α-cells are a much larger fraction of the islet cell population in humans [26–28]), and islet cell architecture (27,28). In addition, *MAFB* mRNA and protein are only produced postnatally in NHP and human islet β-cells, and *MAFA* mRNA expression is very low in adult NHP (10) and human juvenile islets (7) but produced throughout the lifetime of a rodent β-cell (8,9). We now show that the MAFA protein is rarely detected in juvenile human β-cells (i.e., 3 months, 10 months, 19 months, 4 years, 5 years) in relation to MAFB as well as in a smaller fraction of cells in a 10-year-old donor (Fig. 1). This represents a clear distinction from the penetrant pattern of MAFA in rodent (~80% from birth [16]) and adult human (~60% [29]) islet β-cells. Of note, most other islet-enriched transcription factors appear to be at similar levels and cellular distribution within rodents and primates (10). These results not only highlight the possible importance of MAFB in regulating human β-cells but also suggest that coexpression of MAFA with MAFB imparts unique regulatory properties.

DNA Methylation Drives the Postnatal Silencing of *MafB* Expression in Mouse β-Cells

DNA methylation at CpG sites within gene promoters and enhancers is a fundamental epigenetic mark associated with transcriptional repression (30). Because mouse *MafB* expression in β-cells is no longer detectable 2 weeks after birth (8,9), we analyzed the *MafB* gene methylation status by bisulfite sequencing in FACS-purified cells in which the gene was either transcriptionally active (i.e., adult α- and neonatal β-cells) or inactive (adult β-cells). Of note, the region between −1,032 and −838 base pairs (bp) upstream from the *MafB* transcriptional start site was differentially methylated, with much lower CpG methylation in *MafB*-expressing than -nonexpressing cells (Fig. 2*A*). Thus, the methylation pattern within the *MafB* −1,032 to −838 bp region in expressing neonatal mouse β- and α-cells is more similar than to nonexpressing adult β-cells. However, these sequences, as well as those surrounding them, were hypomethylated in *MAFB*-expressing human islet α- and β-cells (i.e., −934 to −734 bp) (Fig. 2*A* and Supplementary Fig. 2), indicating that this region is involved in regulating species-specific expression.

The de novo DNA methyltransferase Dnmt3a was shown to bind to the −1,032 to −838 bp region in MafB-nonexpressing adult mouse β-cell nuclei (Fig. 2*B*). Of note, this enzyme also is involved in repression of functionally disallowed genes in maturing islet β-cells (e.g., hexokinase 1 and lactate dehydrogenase A [18]). Suppression of Dnmt3a by siRNA treatment in the mouse Min6 β-cell line also led to increased *MafB* expression (Fig. 2*C*), providing further support that this epigenetic regulator represses *MafB* in mouse islet β-cells in an age- and cell type–specific manner. Furthermore, the dynamic histone protein modification state within the *MafB* gene added evidence for gene silencing and Dnmt3a regulation, with only repressive chromatin marks enriched in adult β-cells

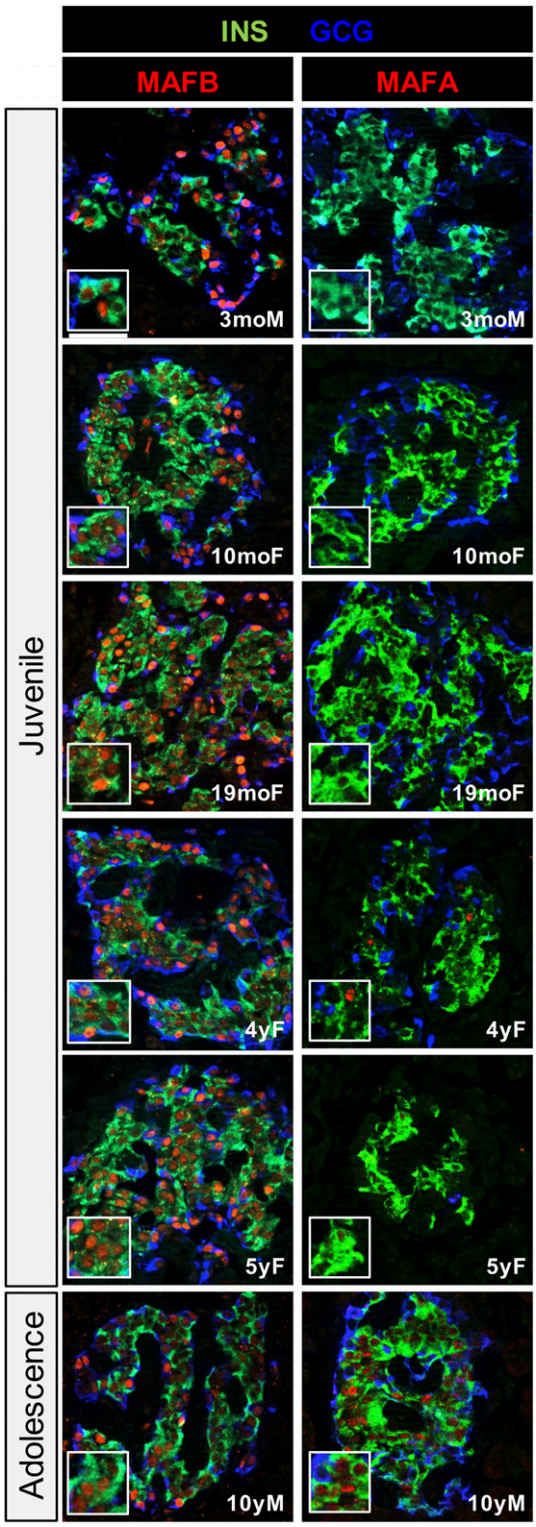

Figure 1—MAFA is expressed in a very small fraction of juvenile human islet β-cells relative to MAFB. Immunodetection of insulin (INS), glucagon (GCG), MAFB, and MAFA in 3-month-old male (3moM), 10-month-old female (10moF), 19-month-old female (19moF), 4-year-old female (4yF), 5-year-old female (5yF), and 10-year-old male (10yM) pancreatic tissue. Insets show nuclear expression of MAFB in early human pancreatic β-cells with MAFA expression principally absent. MAFA is expressed in a much greater fraction of adult human islet β-cells (~60% [17,29]). Scale bar = 50 μm.

(i.e., H3K27me2, H2Aub, H3K9me2, H3K9me3) and activating (H3K4me3, H3K9/14Ac) in neonatal β- and α-cells (Fig. 2*D*).

Development of a β-Cell–Specific MafB Overexpression Mouse Model (βMafBTg) to Study the Impact of Sustained MafB Production in β-Cells

A mouse *MafA* BAC spanning the 5′ flanking regions 1–6 transcription control sequences necessary for directing islet cell–specific expression (31) was used to transgenically drive *MafB* in ~86% (SD ±4.91) of adult islet β-cells (Fig. 3*A* and Supplementary Fig. 1). This is similar to the number of adult islet β-cells that normally express MAFA (8). Of note, the coding sequences for human and mouse MafB are 97% identical. Transgenically produced *MafB* mRNA was not observed until e15.5 and sustained in adult β-cells (Supplementary Fig. 3), a pattern characteristic of the endogenous *MafA* gene (8,9). βMafBTg mice were born in normal Mendelian ratios with no overt physical or physiological deficiencies (data not shown). Immunohistochemical analysis revealed that nuclear MafB was produced in a staining pattern similar to that of MafA within adult insulin-positive cells of βMafBTg mice (Fig. 3*B*). Islet architecture was indistinguishable between $MafA^{fl/fl}$ and βMafBTg mice, with core insulin-positive β-cells surrounded by glucagon-positive α-cells (Fig. 3*B*). There was a 2.5-fold increase in βMafBTg *MafB* mRNA levels in 2-month-old pancreatic islets compared with the $MafA^{fl/fl}$ control, with no change in *MafA* mRNA expression (Fig. 3*C*) and a staining intensity in β-cells similar to the endogenous MafB of α-cells (Fig. 3*B*).

Transcriptional regulation by the MafA and MafB proteins is achieved by homo- or heterodimer binding to large Maf-response elements (32–34). ChIP analysis demonstrated MafB-induced binding to MafA-regulated *Insulin* and *G6pc2* gene transcriptional control sequences in βMafBTg nuclei (Fig. 4*A*). This resulted in a roughly twofold elevation in *Insulin*, *G6pc2*, and *Ccnd2* mRNA levels, but not MafA-activated *Slc2a2* or *Slc30a8* expression (Fig. 4*B*). However, the induction of these insulin-signaling (i.e., *Insulin*, *G6pc2*) or cell proliferation (*Ccnd2*) genes did not affect the rate of glucose clearance in IPGTTs (Fig. 4*C*) or β-cell proliferation (Fig. 4*D*). These results suggest that the heterodimeric MAFA/MAFB regulator of human adult β-cells is essentially functionally equivalent to the mouse $MafA_2$ homodimer.

The $MafB_2$ Homodimer Is Not Functionally Equivalent to $MafA_2$ Homodimer in Mouse Islet β-Cells

The $MafB_2$ regulator not only is enriched in NHP (10) and human (7) islet β-cells (Fig. 1), but also is associated with immature β-cell function in mice (35). Therefore, we next analyzed whether production of MafB alone would affect the deficiencies associated with loss of islet β-cell MafA in $MafA^{\Delta\beta}$ mice, which includes abnormalities in islet architecture, impaired glucose tolerance, and reduced expression of factors important for insulin production and secretion (16).

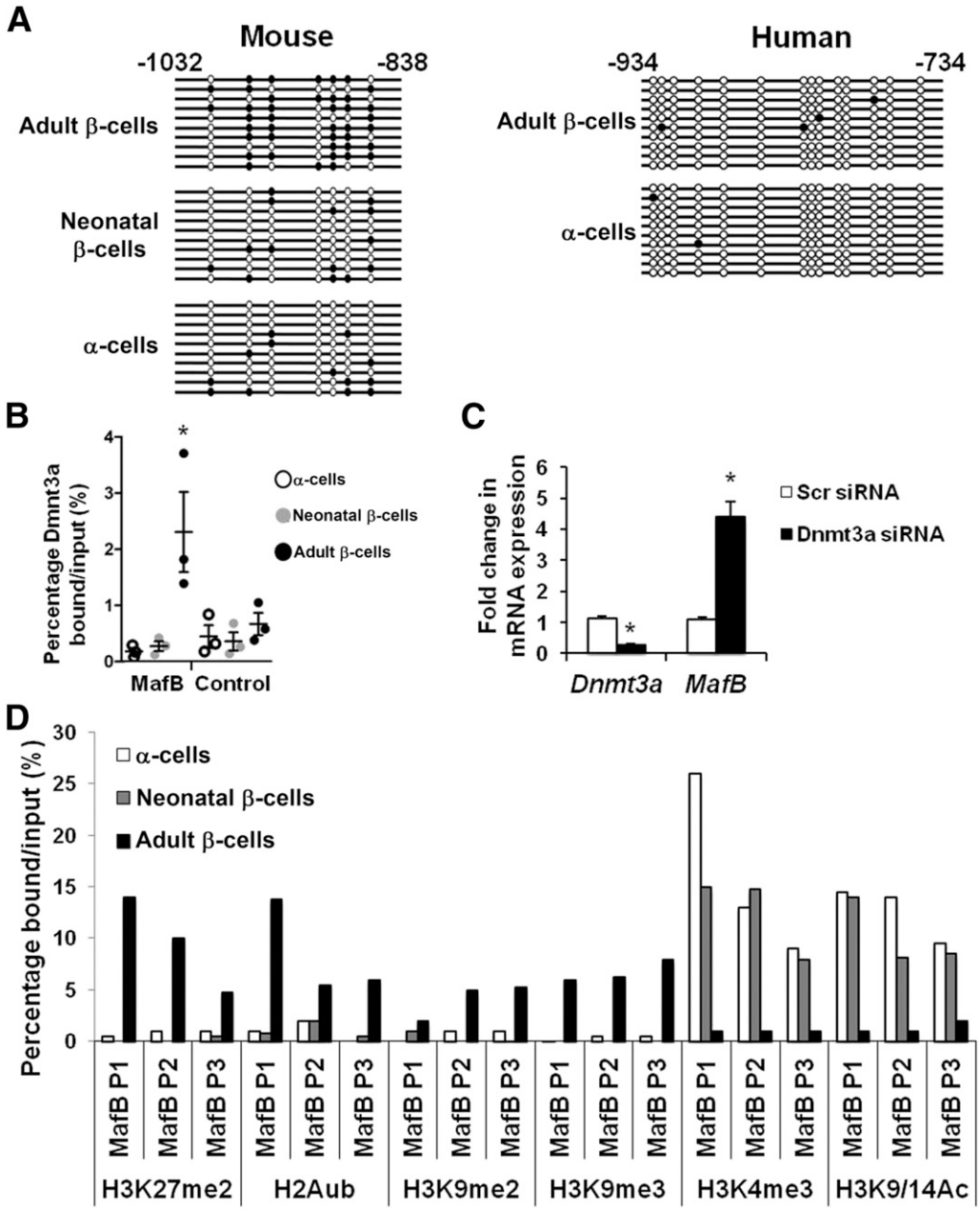

Figure 2— Epigenetic modification analysis reveals the inhibited state of the mouse *MafB* gene in β-cells. *A*: Bisulfite sequencing results of the −1,032 to −838 bp region of the mouse and analogous −934 to −734 bp region of the human *MAFB* gene in α- and β-cells. The only methylation site not conserved within this region is located in mouse at −789 CpG −788 bp and human −791 CC −789 bp. *B*: Dnmt3a binding to the −1,032 to −838 bp region in FACS-sorted mouse adult α-cells, neonatal α-cells, and adult β-cells. Error bars represent the average and SEM ($n = 3$). *C*: Fold change in *Dnmt3a* and *MafB* mRNA levels after scrambled (Scr) and Dnmt3a siRNA treatment in Min6 cells. *D*: H3K27me, H2Aub, H3K9me2, H3K9me3, H3K4me3, and H3K9/14Ac signals over the mouse MafB −1,032 to 838 bp region. P1, P2, and P3 represent independent experiments. *$P < 0.05$.

$MafA^{\Delta\beta}$ and βMafBTg mice were crossed to generate islet β-cells only expressing MafB. The resulting $MafA^{\Delta\beta}$;βMafBTg mice had the same abnormal islet cell architecture of the $MafA^{\Delta\beta}$ mutant marked by the presence of glucagon-positive α-cells throughout the islet core (Fig. 5*A*). In contrast, α-cells and other non–β-islet cell types were found on the islet periphery in control $MafA^{fl/fl}$ mice (Fig. 5*A* and data not shown). Moreover, no improvement was observed in the ability of $MafA^{\Delta\beta}$:βMafBTg mice to clear blood glucose in relation to βMafBTg and $MafA^{fl/fl}$ mice, as indicated by IPGTT (Fig. 5*B*). Not surprisingly, we found that MafB alone could not rescue the gene expression changes resulting from loss of MafA, including *Ins2*, *G6pc2*, *Slc30a8*, and *Slc2a2* (Fig. 5*C*). The only gene differentially expressed in our candidate gene survey was *Ccnd2*, which also was elevated in βMafBTg mice (Fig. 4*B*). Collectively, these results indicate that there will be differences in gene expression, and potentially function, in

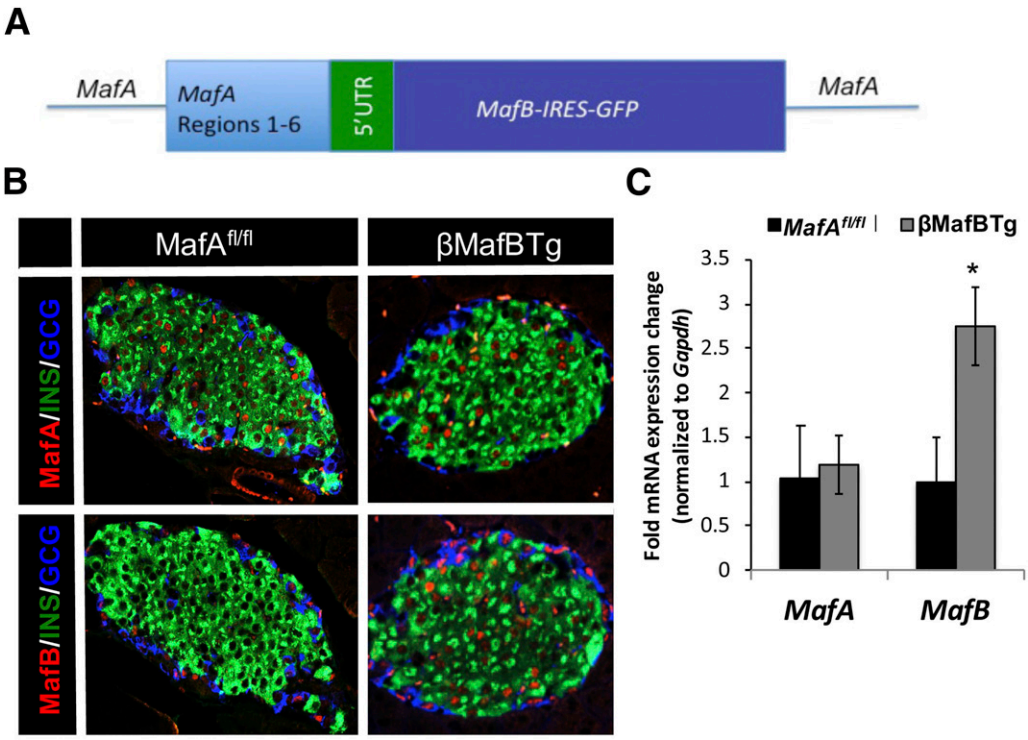

Figure 3—MafB is produced in a large fraction of adult islet β-cells in βMafBTg mice. *A*: The *MafB* transgene is driven by a mouse BAC spanning the region 1–6 control sequences required for β-cell type–specific expression in vivo (31). *B*: MafB is present in ∼85% of 2-month-old βMafBTg islet β-cells (insulin [INS]) and essentially only α-cells (glucagon [GCG]) in age-matched *MafA*$^{fl/fl}$ islets. *C*: *MafB* mRNA was increased by 2.5-fold, whereas *MafA* mRNA levels were unchanged in βMafBTg islets. qPCR signals were normalized to *Gapdh* expression (n = 5). *$P < 0.05$. UTR, untranslated region.

juvenile β-cells principally expressing $MAFB_2$ compared with adult (mouse) $MafA_2$ or (human) MAFA/MAFB β-cells.

MafB Expression Throughout the Islet β-Cell Population Enhances Serotonin Signaling During Pregnancy, Leading to Increased β-Cell Function and Proliferation

MafB is re-expressed in ∼25% of mouse β-cells during pregnancy (15,36), and is presumably present in even a greater percentage in humans. Increased peripheral tissue insulin resistance in pregnancy ensures proper glucose shuttling to the growing fetus and raises insulin demand (36). Thus, β-cells undergo a period of proliferation and increased insulin secretion to adapt to these changes, which occurs in response to serotonin signaling (37–39). We have shown that removal of MafB in maternal β-cells leads to a reduction in serotonin expression and gestational diabetes (36).

Thus, we asked whether broader MafB production within the islet β-cell population of βMafBTg mice is consequential during pregnancy. Indeed, βMafBTg mice had improved glucose tolerance at gestational day (GD) 14.5 compared with control pregnant *MafA*$^{fl/fl}$ dams (Fig. 6*A*). In addition, serum insulin levels in fasted mice were increased in βMafBTg animals (Fig. 6*B*). There was also a 2.4-fold increase in the number of insulin-positive cells labeled with the Ki67$^+$ proliferation marker in pregnant βMafBTg mice (Fig. 6*C*), suggesting that MafB directly affects serotonin signaling and, consequently, adaptive maternal β-cell function and mass expansion. Consistent with this proposal, there was also an increase in maternal βMafBTg serotonin staining (Fig. 7*A*) and islet serotonin levels (Fig. 7*B*) compared with *MafA*$^{fl/fl}$ controls.

To obtain mechanistic insight into how MafB influenced serotonin production, the expression of multiple genes involved in this signaling pathway was examined in GD15.5 βMafBTg and *MafA*$^{fl/fl}$ islets (Fig. 7*C*). No difference was observed in expression of either the prolactin receptor (*Prlr*), which lies upstream of MafB and serotonin production (36), or the serotonin-responsive receptors that mediate β-cell proliferation (i.e., *Htr2b* [39]) or function (*Htr3a* [39]). In fact, only *Tph1* (and not *Tph2* [38]), the rate-limiting enzyme in serotonin production from tryptophan in maternal β-cells (38), was significantly increased in βMafBTg islets (Fig. 7*C*). We propose that the high level of MafB expression within the primate β-cell population also affects the adaptive response of these cells during pregnancy (Fig. 7*D*).

DISCUSSION

Human pancreas tissue and cellular samples recently have become much more accessible for analyses because of the

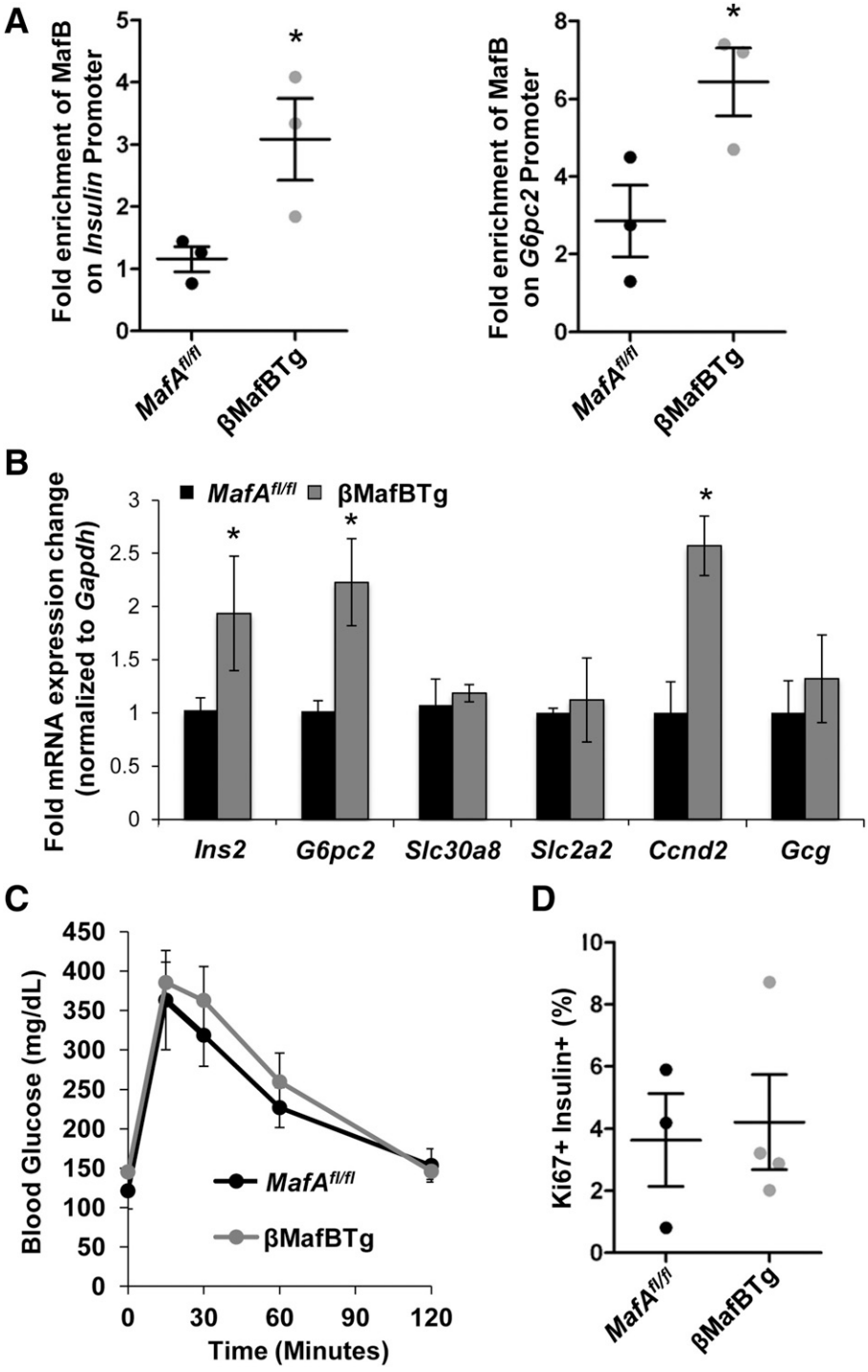

Figure 4—The MafA/MafB regulator has activity similar to that of $MafA_2$ in adult islet βMafBTg β-cells. *A*: ChIP analysis reveals that MafB binds to *Insulin* and *G6pc2* control sequences in 2-month-old βMafBTg islet nuclei. Fold enrichment was calculated relative to the *albumin* promoter (n = 3). *B*: MafB enhances the expression of *Insulin*, *G6pc2*, and *Ccnd2* in 2-month-old βMafBTg islets. qPCR signals were normalized to *Gapdh* (n = 8–10). *C*: βMafBTg and *MafA*$^{fl/fl}$ have the same fasting blood glucose levels and ability to clear a blood glucose bolus in IPGTT assays. *D*: The number of Ki67$^+$-proliferating islet β-cells between βMafBTg and *MafA*$^{fl/fl}$ mice were unchanged (n = 3–4). *$P < 0.05$.

establishment of a variety of collaborative human tissue repositories, including, for example, the Network of Pancreatic Organ Donors With Diabetes, Integrated Islet Distribution Program, and Human Pancreas Analysis Program. Consequently, we now recognize similarities and differences in pancreatic biology between our extensively analyzed rodent models and humans, with distinctions in islets ranging from aspects of architecture/cell

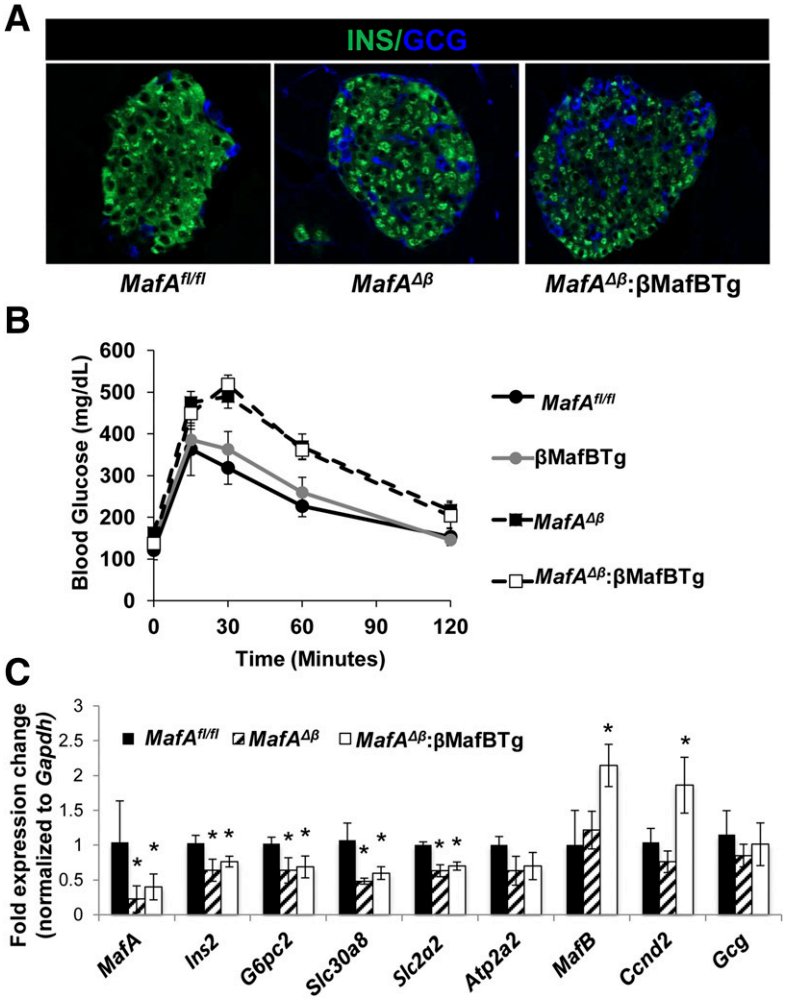

Figure 5—$MafB_2$ is not functionally equivalent to $MafA_2$ in *MafA*$^{\Delta\beta}$:βMafBTg mice. *A*: In contrast to 2-month-old control *MafA*$^{fl/fl}$ mice, the islet architectural core contains both α-cells (glucagon [GCG]) and β-cells (insulin [INS]) in *MafA*$^{\Delta\beta}$:βMafBTg and *MafA*$^{\Delta\beta}$ islets. *B*: Two-month-old *MafA*$^{\Delta\beta}$ and *MafA*$^{\Delta\beta}$:βMafBTg mice are glucose intolerant compared with *MafA*$^{fl/fl}$ and βMafBTg animals. *C*: Gene expression changes in *MafA*$^{\Delta\beta}$ islets are not corrected in *MafA*$^{\Delta\beta}$:βMafBTg mice. qPCR signals normalized to *Gapdh* expression (n = 5–8). *$P < 0.05$.

composition to physiological function (17,24,25,27,28). However, very few experimental systems currently exist to study human islet cell biology, so rodent models are still necessary for revealing the potential significance of these differences to physiology and disease. Here, we focus on the MAFA and MAFB transcription factors, essential regulators of postnatal mouse islet β-cell (i.e., MafA) and α-cell (MafB) activity whose expression pattern is distinctly different between rodents and primates. The current study leverages mouse genetics to understand the mechanistic basis of transcription factor regulation unique to human β-cells.

Distinct age-dependent differences in chromatin landscape, gene expression, and proliferative capacity between juvenile and adult human islet α- and β-cells have been shown (7). This includes epigenetic regulatory modifications of both the N-terminal tails of histone proteins and CpG dinucleotides in DNA. In addition, only adult human islets have enhanced basal, low (5.6 mmol/L) glucose-stimulated insulin secretion in perfusion assays (7), a property not observed in either juvenile human or rodent islets. Yet, what factors control this novel feature of human islets is still unclear. Of note, misexpression of the adult-induced SIX2 and SIX3 transcription factors was found to enhance human β-cell maturation (7). Islet β-cell–specific *MAFA* mRNA levels also are induced in adult human cells, and we now show that little to no cellular protein is produced within juvenile β-cells compared with MAFB (Fig. 1). A caveat to this observation is limitations in human donor availability, which precluded possible distinguishing variabilities among individuals, sex, and ethnicity.

The very low human juvenile MAFA protein levels is strikingly different from the high and penetrant expression pattern within the rodent embryonic, postnatal, and adult β-cell population (8,9). In fact, misexpression of MafA within MafB-enriched immature rat perinatal β-cells induces maturation marker expression and glucose-stimulated

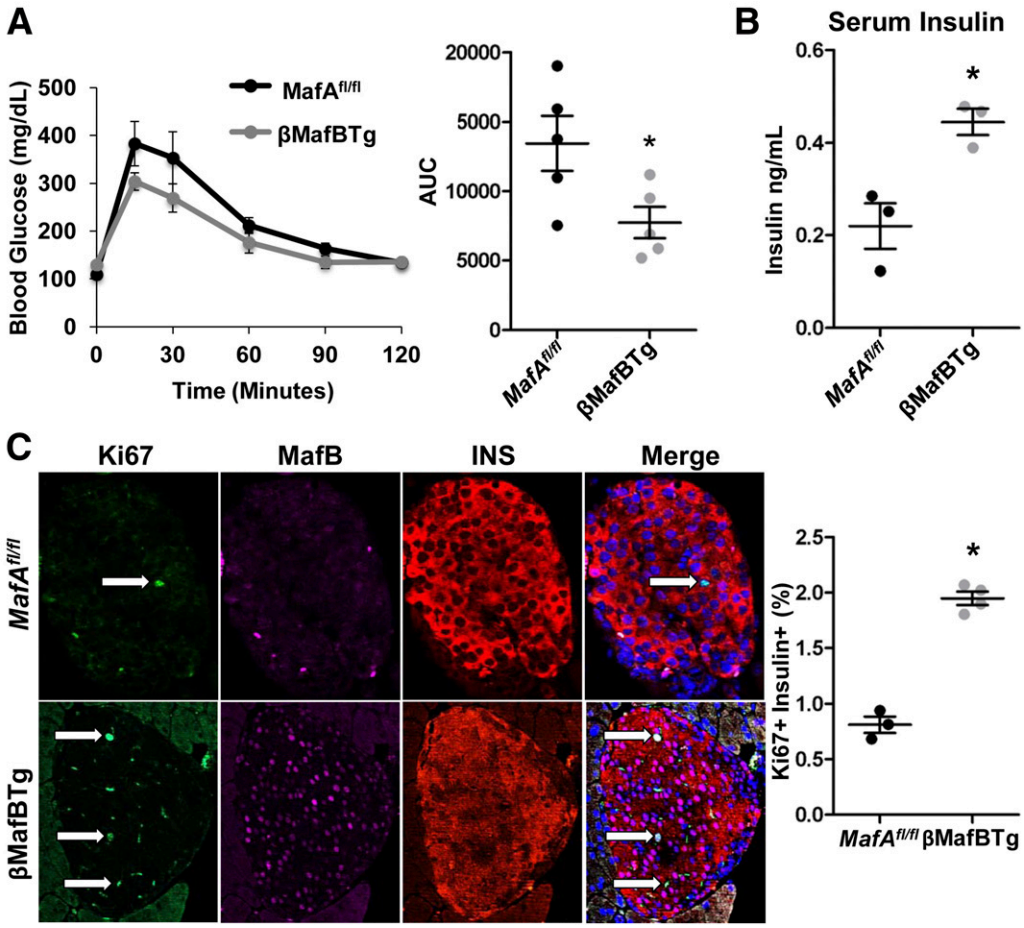

Figure 6—Glucose tolerance and β-cell proliferation levels are enhanced during pregnancy in βMafBTg mice. *A*: *MafA*$^{fl/fl}$ and βMafBTg pregnant dams (GD15.5) were subjected to IPGTT, and βMafBTg mice had improved glucose tolerance. AUC is shown for statistical analysis (n = 8–10). *B*: GD15.5 βMafBTg mice have increased serum insulin levels after a 6-h fast. *C*: βMafBTg islets have an increased number of Ki67$^+$ β-cell nuclei (arrows) compared with *MafA*$^{fl/fl}$ pregnant mice. Graph indicates the percentage of proliferating (Ki67$^+$) insulin-positive (INS$^+$) cells (n = 4). *$P < 0.05$.

insulin section (40). These results indicate that elevated MAFA affects adult islet β-cell gene expression and in vitro glucose-dependent insulin secretion. Of note, juvenile islets had a much lower insulin secretion level when exposed to basal 5.6 mmol/L glucose levels than adult human islets in perfusion assays (7). However, these juvenile and adult cell populations must be functionally similar in vivo because there are no overt age-dependent differences in homeostatic glucose levels. Consequently, we propose that juvenile MAFB-enriched human embryonic-derived β-like cells will have therapeutic utility similar to those expressing adult markers, as presently evaluated, for example, by MAFA production (41–44).

In stark contrast to rodents, MAFB is predominantly expressed in primate β-cells, with no apparent induction of MAFA in aging NHP β-cells (10). Methylation by Dnmt3a of CpG sequences between −1,032 and −838 bp was found to facilitate silencing of mouse *MafB* gene transcription in β-cells postnatally (Fig. 2), whereas this region was unmethylated in MafB-expressing mouse α-cells, newborn mouse β-cells, human β-cells, and human α-cells. It is possible that the absence of the mouse −789 CpG −788 bp site in the human *MAFB* gene prevents DNMT3A action within this otherwise highly conserved region. Of note, the loss of MafB in *MafB*$^{\Delta pancreas}$ mice during development has no consequence to islet β-cells because of compensation by MafA (10). Determining how MAFB influences formation of the β-like cells derived from human embryonic stem cells, a system used by others to analyze how key model system–defined islet-enriched transcription factors act during this important formative period (45,46), would be of great interest. Moreover, because some juvenile human β-cells do not appear to express MAFB, varying levels of this transcription factor may contribute to functional differences within the cell population. Such experimentation is predicted to reveal a unique role for MAFB in human β-cells.

βMafBTg animals were derived to gain insight into how MAFB could affect adult islet β-cells. Ectopic MafB appeared to be expressed first at the onset of embryonic *MafA* expression (Supplementary Fig. 3) and at levels comparable to islet α-cells (Fig. 3) and neonatal β-cells (data not shown). MafB also was bound to endogenous

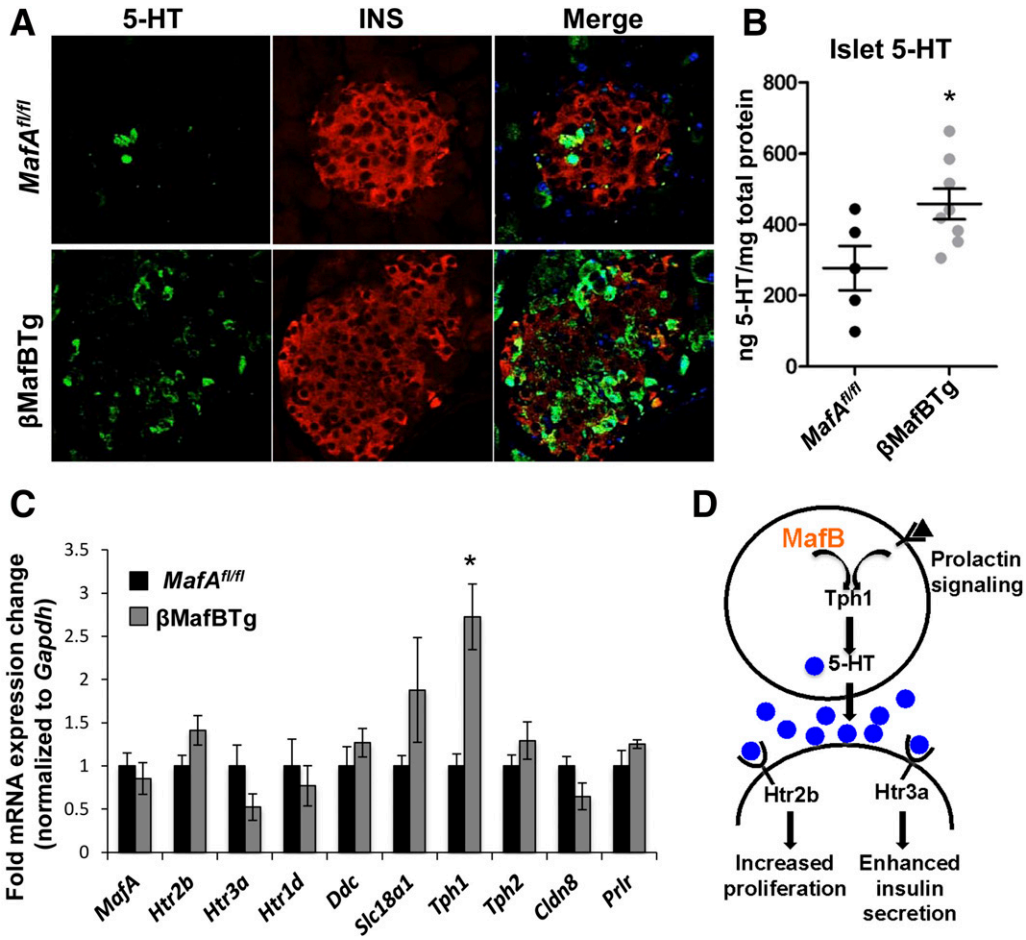

Figure 7—Increased Tph1 elevates serotonin signaling in maternal βMafBTg β-cells. *A* and *B*: βMafBTg islets have increased cellular serotonin (5-HT) staining (n = 3) and total islet 5-HT protein levels compared with *MafA*$^{fl/fl}$ islets during pregnancy (n = 5–8). *C*: Only *Tph1* transcript levels were increased at GD15.5 and not *MafA* or a variety of analyzed mediators of 5-HT signaling in βMafBTg islets (n = 4). *D*: Model illustrating that MafB activation of Tph1 elevates the 5-HT–mediated increase in β-cell mass and function. *$P < 0.05$.

MafA target gene sequences in islet βMafBTg β-cell nuclei (Fig. 4*A*). We conclude from these results that MafB was being produced in βMafBTg β-cells at functionally active levels. The lack of any overt physiological or molecular phenotype upon coexpression of MafA and MafB in βMafBTg β-cells likely means that the mouse $MafA_2$ homodimer is functionally equivalent to the adult human MAFA/MAFB heterodimer. However, $MAFB_2$ alone was unable to rescue any of the islet cell deficiencies of *MafA*$^{\Delta\beta}$ mice (Fig. 5). It is possible that unique and enabling sets of transcriptional coregulators are recruited by human $MAFB_2$ or MAFA/MAFB. If so, this is likely due to binding within the COOH-terminal sequences of these proteins, which have been shown in chimeric analysis to impart defining functional regulatory properties (47). MafA activation is known to be controlled by the histone acetyltransferase p300/CBP-factor (PCAF) coactivator (48), whereas both MAFA and MAFB recruit the Mll3 and Mll4 histone 3 lysine 4 methyltransferase complexes (22). Such $MAFB_2$- and MAFA/MAFB-recruited factors also could influence the functional properties of the four distinct adult human islet β-cell populations termed β1 (most active) to β4 (least active) (49).

MafB expression in βMafBTg islet β-cells led to increased serotonin signaling during pregnancy, which presumably resulted in enhanced cell proliferation and function (Fig. 7). Our results demonstrate that this entails activation of *Tph1* mRNA levels, which encodes the rate-limiting enzyme for serotonin synthesis from tryptophan in maternal β-cells. Of note, the mouse *Tph1* gene contains three MafB consensus binding sites at −9,184/−9,189, −8,184/−8,189, and −8,147/−8,152 bp that are near a region bound by Stat5 (−8,754/−8,723 bp), which mediates prolactin signaling in maternal β-cells (50). Thus, MafB may directly bind in conjunction with Stat5 to increase *Tph1* transcription, thereby increasing serotonin production and leading to greater islet β-cell mass and function that compensates for the insulin resistance induced by pregnancy. Preventing MafB induction in these cells leads to a form of gestational diabetes as a result of an inability to facilitate these processes (36). Although it is unclear whether increased MafB expression in βMafBTg mice

simulates the conditions in the human maternal islet β-cell population, these results raise the possibility that such an increase in MAFB levels would buffer the detrimental effects of obesity in the mother and her offspring (51).

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