Regulation of pancreatic islet gene expression in mouse islets by pregnancy

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Abstract

Pancreatic β cells adapt to pregnancy-induced insulin resistance by unclear mechanisms. This study sought to identify genes involved in β cell adaptation during pregnancy. To examine changes in global RNA expression during pregnancy, murine islets were isolated at a time point of increased β cell proliferation (E13.5), and RNA levels were determined by two different assays (global gene expression array and G-protein-coupled receptor (GPCR) array). Follow-up studies confirmed the findings for select genes. Differential expression of 110 genes was identified and follow-up studies confirmed the changes in select genes at both the RNA and protein level. Surfactant protein D (SP-D) mRNA and protein levels exhibited large increases, which

were confirmed in murine islets. Cytokine-induced expression of SP-D in islets was also demonstrated, suggesting a possible role as an anti-inflammatory molecule. Complementing these studies, an expression array was performed to define pregnancy-induced changes in expression of GPCRs that are known to impact islet cell function and proliferation. This assay, the results of which were confirmed using real-time reverse transcription-PCR assays, demonstrated that free fatty acid receptor 2 and cholecystokinin receptor A mRNA levels were increased at E13.5. This study has identified multiple novel targets that may be important for the adaptation of islets to pregnancy.

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Introduction

Pancreatic β cells within the islets of Langerhans play a critical role in the regulation of glucose homeostasis by sensing glucose levels and secreting insulin. Compensation in response to increased demands for insulin occurs through changes in both insulin synthesis and release, as well as changes in β cell mass. These changes observed in β cells play an important role in regulating glucose homeostasis during different physiological and pathological states such as the insulin resistant states of pregnancy and obesity (Butler *et al.* 2007). Studying states in which insulin needs change can provide insight into pathways mediating β cell adaptations.

In mice, β cell mass increases about twofold during pregnancy, with a maximum rate of β cell proliferation near day 14 post coitum. This response is dynamic, as β cell mass decreases in the *postpartum* period (Karnik *et al.* 2007). Although the activation of the prolactin receptor (PRLR) by prolactin and placental lactogens contributes to the regulation of β cell mass during pregnancy (Brelje *et al.* 1993), the mechanisms controlling the expansion of β cell mass during

pregnancy remain incompletely defined, and it is likely that additional signaling pathways contribute to or are directly involved in the PRLR-mediated proliferative response (Karnik *et al.* 2007, Sorenson & Brelje 2009).

This study was designed to begin to elucidate the molecular mechanisms responsible for pregnancy-induced changes in β cell mass. This was accomplished by comparing the results of whole genome RNA expression analysis using islets from pregnant mice on day E13.5 and control mice. We identified 110 genes that are differentially expressed in pregnant islets, including the gene encoding surfactant protein D (SP-D). Furthermore, the impact of pregnancy on the expression of G-protein-coupled receptors (GPCRs), a family of proteins involved in the regulation of β cell function and mass and effectors for a number of therapeutic agents (Vasavada *et al.* 1996, Holz & Chepurny 2005, Xiao *et al.* 2008), was examined in detail using a separate but complementary microarray approach. Together, the results of our studies suggest multiple novel targets that will facilitate further studies seeking to define molecular changes characteristic of the pregnancy-induced increase in β cell mass.

Materials and Methods

Islet isolation

C57BL/6 pregnant (day 13.5 of pregnancy) and control (age- and sex-matched) mice were purchased from Jackson Laboratories (Bar Harbor, ME, USA). Islets were isolated using standard protocols as described previously (Chen *et al.* 2006), and RNA was isolated using RNeasy (Qiagen). For RNA analyses, islets isolated from three mice were pooled for each sample to reduce interindividual variability in the RNA expression analyses. All procedures performed on animals were approved by the Northwestern University Animal Care and Use Committee.

Microarray analysis

RNA expression analysis was performed using the Illumina MouseWG-6 BeadChip, which provides coverage of over 45 200 genes and expressed sequence tags. Islet RNA from pregnant and control mice was labeled using a commercial kit (TargetAmp 1-Round Aminoallyl-aRNA Ki; Epicentre, Madison, WI, USA). To reduce changes related to interindividual variability, each sample of islet RNA was prepared using pooled islets from three pregnant or control mice. Labeled RNA was hybridized to microarrays. Raw signal intensities of each probe were obtained and processed as described previously (Du *et al.* 2008). Differentially expressed genes were identified using an ANOVA model with empirical Bayesian variance estimation. The problem of multiple comparisons was corrected using the false discovery rate (FDR). Genes were identified as being differentially expressed on the basis of a statistical significance (P value < 0.01), FDR $< 5\%$, and 1.5-fold change in expression level in pregnant compared with control islets. Functional classification was performed using the ingenuity pathway analysis (IPA) software, based on statistical significance (IPA pathway test, $P < 0.01$) and biological importance.

GPCR RNA array

A commercially available GPCR-specific 384 well TaqMan real-time reverse transcription (RT)-PCR microarray (Applied Biosystems, Carlsbad, CA, USA), which allows for the detection of mRNAs encoding 343 GPCRs (not including the odorant, olfactory, gustatory, and pheromone receptors), was used with RNA extracted as described above. cDNA was prepared using the high-capacity reverse transcriptase kit (Applied Biosystems) from the RNA that had contaminating DNA removed by DNAase I (Ambion, Applied Biosystems, Austin, TX, USA), and each port was loaded with cDNA (from 1 μ g RNA) and the TaqMan Gene Expression Master Mix (Applied Biosystems). The resulting plate was analyzed on the 7900HT ABI PRISM, and the data were analyzed using SDS 2.3 and RQ Manager 1.2 software provided by Applied Biosystems. The results for each

individual GPCR were examined, and if the sample had a calculated threshold < 0.1 , it was defined as undetectable (this generally indicated no discernible melt curve) and for data processing, the cycle time (C_t) was set at 40.0. A comparative C_t approach was utilized to quantify the relative levels of mRNA. C_t s were normalized to the geometric mean of three housekeeping (HK) genes. Each HK gene on the microarray was evaluated for stability by geNorm (Vandesompele *et al.* 2002), and the three most stable HK genes were used. This experiment was performed on two independent samples, where each sample consisted of islets pooled from three mice.

Real-time RT-PCR

RNA was isolated and cDNA was prepared as described above. cDNA samples were examined by real-time RT-PCR and were normalized to the geometric mean of the three most stable HK genes as determined by geNorm (Vandesompele *et al.* 2002). These were hypoxanthine-guanine phosphoribosyltransferase (*HPRT*), TATA-binding protein (*TBP*), and peptidylprolyl isomerase A (*PPIA*). Sequences of primers used in the different assays are listed in Supplementary Table 1, see section on supplementary data given at the end of this article. A comparative C_t analysis was utilized to determine fold change in RNA expression based on the following approach: fold change = $2^{(\Delta\Delta C_t)} = 2^{-(\Delta C_t \text{ sample} - \Delta C_t \text{ control})}$ (Fleige *et al.* 2006). For each real-time RT-PCR experiment, three independent replicates were used.

Western blot

Islets from the control or pregnant mice (day 13.5 of pregnancy) were isolated as described above, and proteins were extracted in RIPA cell lysis buffer containing a cocktail of protease inhibitors (Calbiochem, LaJolla, CA, USA). Primary antibodies for free fatty acid receptor 2 (FFAR2) and PRLR were from Santa Cruz (Santa Cruz, CA, USA) and for tryptophan hydroxylase 1 (TPH1) from Thermo Scientific (Rockford, IL, USA). Mouse monoclonal antibodies to SP-D (hyb-246) were prepared as described earlier (Sorensen *et al.* 2009). The protein content of the lysate was determined using the Coomassie blue protein assay. Twenty micrograms of protein were separated by 10% SDS-PAGE and transferred to a polyvinylidene fluoride membrane in a semidry apparatus. Western blot analyses were performed as previously described (El Muayed *et al.* 2010). For SP-D samples, both reduced and nonreduced protein samples were examined. For reduced samples for SP-D, protein preparations were prepared by mixing the protein samples 1:2 with Laemmli sample buffer with 5% 2-mercaptoethanol and heating at 100 °C for 1 min (nonreduced preparations were heated in sample buffer with the addition of iodoacetamide, final concentration of 90 mM) as previously described (Madsen *et al.* 2000, Sorensen *et al.* 2009). Each assay was repeated with two or three independent samples.

Figure 1 Global gene expression analysis of pancreatic islets at day 13.5 of pregnancy. (A) Heat map of differentially regulated pancreatic islet genes in pooled islets from either pregnant mice on day 13.5 (E1–E3) or control mice. E1, E2, C1, and C2 represent independent samples prepared from separate pools of pregnant (E1 and 2) and control (C1 and 2) islets. E3 and C3 are data obtained using the same RNA as E1/C1 respectively but prepared for microarray analysis independently. Scale bar: relative RNA expression (mean centered, standardized). (B) Graphical representation of the 110 differentially regulated genes from islets of pregnant mice compared with those of control mice using a functional classification scheme from ingenuity. Genes could be included in more than one category. The number of genes in each category is shown. *P* values for each category were determined by a category overrepresentation *P* test to examine the relative proportion of the genes in each category among the differentially regulated genes compared with the chip as a whole. The *P* values were as follows: cellular movement ($P=2.9 \times 10^{-3}$), molecular transport ($P=4.5 \times 10^{-4}$), cell-to-cell signaling and interaction ($P=4.5 \times 10^{-4}$), cell death ($P=9.3 \times 10^{-4}$), cellular growth and proliferation ($P=4.5 \times 10^{-4}$), inflammation ($P=1.5 \times 10^{-2}$), and metabolism ($P=3.4 \times 10^{-3}$).

Table 1 Global expression analysis of RNA isolated from pregnant mice (E13.5) compared with that from control mice (age- and sex-matched)

Genes	Description	Fold	P value
<i>Cell-to-cell signaling and interaction</i>			
<i>Sftpd</i>	Surfactant-associated protein D	12.8	<0.0005
<i>Tph1</i>	Tryptophan hydroxylase 1	6.1	<0.0005
<i>Tnfrsf11b</i>	Osteoprotegerin	3.7	<0.0005
<i>Vip</i>	Vasoactive intestinal polypeptide	2.9	0.002
<i>Tac1</i>	Tachykinin 1	2.6	<0.0005
<i>Prlr</i>	Prolactin receptor	2.2	<0.0005
<i>C3</i>	Complement component 3	2.1	<0.0005
<i>Cartpt</i>	CART prepropeptide	2.1	<0.0005
<i>Glycam1</i>	Glycosylation-dependent cell adhesion molecule 1	2.1	0.002
<i>Grp</i>	Gastrin-releasing peptide	2.1	0.001
<i>C1qa</i>	Complement component 1, q subcomponent, alpha polypeptide	1.7	<0.0005
<i>Ttyh1</i>	Tweety homolog 1 (<i>Drosophila</i>)	1.7	<0.0005
<i>Ffar2</i>	Free fatty acid receptor 2	1.6	<0.0005
<i>Ptpre</i>	Protein tyrosine phosphatase, receptor type, E	1.6	0.003
<i>Ntrk2</i>	Neurotrophic tyrosine kinase, receptor, type 2	1.5	<0.0005
<i>Lpl</i>	Lipoprotein lipase	-1.5	<0.0005
<i>Pcsk9</i>	Proprotein convertase subtilisin/kexin type 9	-1.5	0.0001
<i>Rap2a</i>	RAS-related protein 2a	-1.6	0.003
<i>Ibsp</i>	Integrin-binding sialoprotein	-1.7	0.002
<i>Spon2</i>	Spondin 2, extracellular matrix protein	-1.7	<0.0005
<i>Cellular growth and proliferation</i>			
<i>Sftpd</i>	Surfactant-associated protein D	12.8	<0.0005
<i>Cish</i>	Cytokine-inducible SH2-containing protein	7.5	<0.0005
<i>Tnfrsf11b</i>	Osteoprotegerin	3.7	<0.0005
<i>Vip</i>	Vasoactive intestinal polypeptide	2.9	0.002
<i>Nupr1</i>	Nuclear protein 1	2.9	0.001
<i>Ovol2</i>	Ovo-like 2 (<i>Drosophila</i>)	2.7	<0.0005
<i>Tac1</i>	Tachykinin 1	2.6	<0.0005
<i>Socs2</i>	Suppressor of cytokine signaling 2	2.6	<0.0005
<i>Igfbp-5</i>	Insulin-like growth factor-binding protein-5	2.5	<0.0005
<i>Hopx</i>	HOP homeobox	2.3	<0.0005
<i>Enpp2</i>	Ectonucleotide pyrophosphatase/phosphodiesterase 2	2.3	<0.0005
<i>Prlr</i>	Prolactin receptor	2.2	<0.0005
<i>C3</i>	Complement component 3	2.1	<0.0005
<i>Grp</i>	Gastrin-releasing peptide	2.1	0.001
<i>Gadd45α</i>	Growth arrest and DNA-damage-inducible 45 alpha	2.0	<0.0005
<i>Apobec1</i>	Apolipoprotein B editing complex 1	1.7	0.0003
<i>Dnmt3b</i>	DNA methyltransferase 3B	1.7	<0.0005
<i>Mcf2</i>	Multiple coagulation factor deficiency 2	1.7	0.0001
<i>Eif4ebp-1</i>	Eukaryotic translation initiation factor 4E-binding protein-1	1.6	0.003
<i>Cited2</i>	Cbp/p300-interacting transactivator, w/Glu/Asp carb.-terminaldom.2	1.6	0.0008
<i>Ptpre</i>	Protein tyrosine phosphatase, receptor type, E	1.6	0.003
<i>Fgl2</i>	Fibrinogen-like protein 2	1.6	<0.0005
<i>Fhit</i>	Fragile histidine triad gene	1.6	<0.0005
<i>Isg20</i>	Interferon-stimulated protein	1.6	0.0002
<i>Ntrk2</i>	Neurotrophic tyrosine kinase, receptor, type 2	1.5	<0.0005
<i>Defb1</i>	Defensin beta 1	-1.5	0.0002
<i>Tox</i>	Thymocyte selection-associated high-mobility group box	-1.5	<0.0005
<i>Fosb</i>	FBj osteosarcoma oncogene B	-1.5	<0.0005
<i>Tm4sf4</i>	Transmembrane 4 superfamily member 4	-1.7	<0.0005
<i>Molecular transport</i>			
<i>Sftpd</i>	Surfactant-associated protein D	12.8	<0.0005
<i>Tnfrsf11b</i>	Osteoprotegerin	3.7	<0.0005
<i>Vip</i>	Vasoactive intestinal polypeptide	2.9	0.002
<i>Tac1</i>	Tachykinin 1	2.6	<0.0005
<i>Igfbp-5</i>	Insulin-like growth factor-binding protein-5	2.5	<0.0005
<i>Enpp2</i>	Ectonucleotide pyrophosphatase/phosphodiesterase 2	2.3	<0.0005
<i>Prlr</i>	Prolactin receptor	2.2	<0.0005
<i>C3</i>	Complement component 3	2.1	<0.0005
<i>Grp</i>	Gastrin-releasing peptide	2.1	0.001

(continued)

Table 1 Continued

Genes	Description	Fold	P value
<i>Cartpt</i>	CART prepropeptide	2.1	<0.0005
<i>Slc6a8</i>	Solute carrier family 6 (neurotransmitter transporter, creatine), member 8	1.8	<0.0005
<i>Apobec1</i>	Apolipoprotein B editing complex 1	1.7	0.0003
<i>Hba-α1</i>	Hemoglobin alpha, adult chain 1	1.6	0.0008
<i>Hbb-β1</i>	Hemoglobin, beta adult major chain	1.6	<0.0005
<i>Lpl</i>	Lipoprotein lipase	-1.5	<0.0005
<i>Pcsk9</i>	Proprotein convertase subtilisin/kexin type 9	-1.5	0.0001
<i>Adh1</i>	Alcohol dehydrogenase 1 (class I)	-1.7	0.0002
<i>Ibsp</i>	Integrin-binding sialoprotein	-1.7	0.002
<i>Cell death</i>			
<i>Sftpd</i>	Surfactant-associated protein D	12.8	<0.0005
<i>Tnfrsf11b</i>	Osteoprotegerin	3.7	<0.0005
<i>Vip</i>	Vasoactive intestinal polypeptide	2.9	0.002
<i>Nupr1</i>	Nuclear protein 1	2.9	0.001
<i>Tac1</i>	Tachykinin 1	2.6	<0.0005
<i>Socs2</i>	Suppressor of cytokine signaling 2	2.6	<0.0005
<i>Igfbp-5</i>	Insulin-like growth factor-binding protein-5	2.5	<0.0005
<i>Prlr</i>	Prolactin receptor	2.2	<0.0005
<i>C3</i>	Complement component 3	2.1	<0.0005
<i>Gadd45α</i>	Growth arrest and DNA-damage-inducible 45 alpha	2.0	<0.0005
<i>Dnmt3b</i>	DNA methyltransferase 3B	1.7	<0.0005
<i>C1qa</i>	Complement component 1, q subcomponent, alpha polypeptide	1.7	<0.0005
<i>Fgl2</i>	Fibrinogen-like protein 2	1.6	<0.0005
<i>Fhit</i>	Fragile histidine triad gene	1.6	<0.0005
<i>Ptpre</i>	Protein tyrosine phosphatase, receptor type, E	1.6	0.003
<i>Ezr</i>	Ezrin	1.6	<0.0005
<i>Ern1</i>	Endoplasmic reticulum (ER) to nucleus signaling 1	1.6	0.0003
<i>Eif4ebp1</i>	Eukaryotic translation initiation factor 4E binding protein 1	1.6	0.003
<i>Ntrk2</i>	Neurotrophic tyrosine kinase, receptor, type 2	1.5	<0.0005
<i>Pcsk9</i>	Proprotein convertase subtilisin/kexin type 9	-1.5	0.0001
<i>Defb1</i>	Defensin beta 1	-1.5	0.0002
<i>Anpep</i>	Alanyl (membrane) aminopeptidase	-1.5	<0.0005
<i>Fosb</i>	FBJ osteosarcoma oncogene B	-1.5	<0.0005
<i>Cellular movement</i>			
<i>Sftpd</i>	Surfactant-associated protein D	12.8	<0.0005
<i>Tnfrsf11b</i>	Osteoprotegerin	3.7	<0.0005
<i>Vip</i>	Vasoactive intestinal polypeptide	2.9	0.002
<i>Tac1</i>	Tachykinin 1	2.6	<0.0005
<i>Socs2</i>	Suppressor of cytokine signaling 2	2.6	<0.0005
<i>Enpp2</i>	Ectonucleotide pyrophosphatase/phosphodiesterase 2	2.3	<0.0005
<i>Cartpt</i>	CART prepropeptide	2.1	<0.0005
<i>Grp</i>	Gastrin-releasing peptide	2.1	0.001
<i>C3</i>	Complement component 3	2.1	<0.0005
<i>Gadd45α</i>	Growth arrest and DNA-damage-inducible 45 alpha	2.0	<0.0005
<i>Dnmt3b</i>	DNA methyltransferase 3B	1.7	<0.0005
<i>Ezr</i>	Ezrin	1.6	<0.0005
<i>Fgl2</i>	Fibrinogen-like protein 2	1.6	<0.0005
<i>Fhit</i>	Fragile histidine triad gene	1.6	<0.0005
<i>Ntrk2</i>	Neurotrophic tyrosine kinase, receptor, type 2	1.5	<0.0005
<i>Defb1</i>	Defensin beta 1	-1.5	0.0002
<i>Anpep</i>	Alanyl (membrane) aminopeptidase	-1.5	<0.0005
<i>Rap2a</i>	RAS-related protein 2a	-1.6	0.003
<i>Metabolism</i>			
<i>Tph2</i>	Tryptophan hydroxylase 2	3.9	<0.0005
<i>Ivd</i>	Isovaleryl coenzyme A dehydrogenase	3.7	<0.0005
<i>Tnfrsf11b</i>	Osteoprotegerin	3.7	<0.0005
<i>Tac1</i>	Tachykinin 1	2.6	<0.0005
<i>Igfbp-5</i>	Insulin-like growth factor-binding protein-5	2.5	<0.0005
<i>Prlr</i>	Prolactin receptor	2.2	<0.0005

(continued)

Table 1 Continued

Genes	Description	Fold	P value
<i>Prlr</i>	Prolactin receptor	2.2	<0.0005
<i>Grp</i>	Gastrin-releasing peptide	2.1	0.001
<i>C3</i>	Complement component 3	2.1	<0.0005
<i>Cartpt</i>	CART prepropeptide	2.1	<0.0005
<i>Slc40a1</i>	Solute carrier family 40, member 1	2.1	<0.0005
<i>Grhl1</i>	Grainyhead-like 1 (<i>Drosophila</i>)	2.0	<0.0005
<i>Slc2a13</i>	Solute carrier family 2 (facilitated glucose transporter), 13	2.0	<0.0005
<i>Enox1</i>	Ecto-NOX disulfide-thiol exchanger 1	1.8	<0.0005
<i>Slc6a8</i>	Solute carrier family 6 (neurotransmitter transporter, creatine), member 8	1.8	<0.0005
<i>Cdh7</i>	Cadherin 7, type 2	1.7	<0.0005
<i>Matn2</i>	Matrilin 2	1.7	<0.0005
<i>Gcdh</i>	Glutaryl-coenzyme A dehydrogenase	1.6	<0.0005
<i>Fhit</i>	Fragile histidine triad gene	1.6	<0.0005
<i>Ptpre</i>	Protein tyrosine phosphatase, receptor type, E	1.6	0.003
<i>Slc39a11</i>	Solute carrier family 39 (metal ion transporter), member 11	1.6	<0.0005
<i>Uhrf2</i>	Ubiquitin-like, containing PHD and RING finger, 2	1.6	<0.0005
<i>Ntrk2</i>	Neurotrophic tyrosine kinase, receptor, type 2	1.5	<0.0005
<i>Atp8a1</i>	ATPase, aminophospholipid transporter	1.5	<0.0005
<i>Edem2</i>	ER degradation enhancer, mannosidase alpha-like 2	1.5	0.0003
<i>Tmem2</i>	Transmembrane protein 2	1.5	<0.0005
<i>Lpl</i>	Lipoprotein lipase	-1.5	<0.0005
<i>Pcsk9</i>	Proprotein convertase subtilisin/kexin type 9	-1.5	0.0001
<i>Neurog3</i>	Neurogenin 3	-1.5	<0.0005
<i>Tox</i>	Thymocyte selection-associated high-mobility group box	-1.5	<0.0005
<i>Ddc</i>	DOPA decarboxylase	-1.6	<0.0005
<i>Hmgn3</i>	High-mobility group nucleosomal-binding domain 3	-1.6	<0.0005
<i>Scg3</i>	Secretogranin III	-1.6	<0.0005
<i>Inflammation</i>			
<i>Sftpd</i>	Surfactant-associated protein D	12.8	<0.0005
<i>Tph2</i>	Tryptophan hydroxylase 2	3.9	<0.0005
<i>Tnfrsf11b</i>	Osteoprotegerin	3.7	<0.0005
<i>Chgb</i>	Chromogranin B	2.9	<0.0005
<i>Tac1</i>	Tachykinin 1	2.6	<0.0005
<i>Hopx</i>	HOP homeobox	2.3	<0.0005
<i>Enpp2</i>	Ectonucleotide pyrophosphatase/phosphodiesterase 2	2.3	<0.0005
<i>Prlr</i>	Prolactin receptor	2.2	<0.0005
<i>C3</i>	Complement component 3	2.1	<0.0005
<i>Grp</i>	Gastrin-releasing peptide	2.1	0.001
<i>Slc2a13</i>	Solute carrier family 2 (facilitated glucose transporter), 13	2.0	<0.0005
<i>Sgip1</i>	SH3-domain GRB2-like (endophilin) interacting protein 1	2.0	<0.0005
<i>Gadd45a</i>	Growth arrest and DNA-damage-inducible 45 alpha	2.0	<0.0005
<i>Enox1</i>	Ecto-NOX disulfide-thiol exchanger 1	1.8	<0.0005
<i>C1qa</i>	Complement component 1, q subcomponent, alpha polypeptide	1.7	<0.0005
<i>Matn2</i>	Matrilin 2	1.7	<0.0005
<i>Hba-α1</i>	Hemoglobin alpha, adult chain 1	1.6	0.0008
<i>Hbb-β1</i>	Hemoglobin, beta adult major chain	1.6	<0.0005
<i>Slc39a11</i>	Solute carrier family 39 (metal ion transporter), member 11	1.6	<0.0005
<i>Fhit</i>	Fragile histidine triad gene	1.6	<0.0005
<i>Ern1</i>	Endoplasmic reticulum (ER) to nucleus signaling 1	1.6	0.0003
<i>Atp8a1</i>	ATPase, aminophospholipid transporter	1.5	<0.0005
<i>Tspan14</i>	Tetraspanin 14	1.5	<0.0005
<i>Ntrk2</i>	Neurotrophic tyrosine kinase, receptor, type 2	1.5	<0.0005
<i>Edem2</i>	ER degradation enhancer, mannosidase alpha-like 2	1.5	0.0003
<i>Tox</i>	Thymocyte selection-associated high-mobility group box	-1.5	<0.0005
<i>Anpep</i>	Alanyl (membrane) aminopeptidase	-1.5	<0.0005
<i>Fosb</i>	FBJ osteosarcoma oncogene B	-1.5	<0.0005
<i>Acpl2</i>	Acid phosphatase-like 2	-1.6	<0.0005
<i>Sphkap</i>	SPHK1 interactor, AKAP domain containing	-1.6	0.0005
<i>Ddc</i>	DOPA decarboxylase	-1.6	<0.0005
<i>Aard</i>	Alanine and arginine-rich domain-containing protein	-1.7	<0.0005

(continued)

Table 1 Continued

Genes	Description	Fold	P value
<i>Miscellaneous</i>			
<i>Fmo1</i>	Flavin-containing monooxygenase 1	10.7	<0.0005
<i>B3galnt1</i>	UDP-beta 1, 3-galactosaminyltransferase, polypep-1	2.9	0.001
<i>EG244911</i>	Predicted gene, EG244911	2.8	<0.0005
<i>H21Rik</i>	RIKEN cDNA 64 305 50H21 gene	2.8	<0.0005
<i>Hsbp-1</i>	Heat shock factor-binding protein-1	2.7	0.0007
<i>Wipi1</i>	WD repeat domain, phosphoinositide interacting 1	2.6	<0.0005
<i>Znrf2</i>	Zinc and ring finger 2	2.4	0.0009
<i>Car15</i>	Carbonic anhydrase 15	2.4	0.003
<i>Chac1</i>	ChaC, cation transport regulator-like 1 (<i>Escherichia coli</i>)	2.2	0.002
<i>Aqp4</i>	Aquaporin 4	2.2	0.0009
<i>Txnrd2</i>	Thioredoxin reductase 2	2.0	<0.0005
<i>Fkbp-11</i>	FK506-binding protein-11	2.0	0.001
<i>Ehhadh</i>	Enoyl-Coenzyme A, hydratase/3-hydroxyacyl CoA dehydrogenase	1.9	<0.0005
<i>Igfals</i>	Insulin-like growth factor-binding protein, acid labile subunit	1.9	0.0005
<i>Rag1ap1</i>	Recombination activating gene 1 activating protein 1	1.9	0.02
<i>Aldh1l2</i>	Aldehyde dehydrogenase 1 family, member L2	1.9	0.002
<i>Pycr1</i>	Pyrroline-5-carboxylate reductase 1	1.8	0.001
<i>BC006662</i>	cDNA sequence BC006662	1.7	0.0005
<i>Cldn8</i>	Claudin 8	1.7	0.0005
<i>Gnβ11</i>	G-protein, beta polypeptide 1-like	1.7	0.0009
<i>Rnf182</i>	Ring finger protein 182	1.7	0.001
<i>Hapln4</i>	Hyaluronan and proteoglycan link protein 4	1.7	0.006
<i>Tmem28</i>	Transmembrane protein 28	1.7	0.006
<i>Lsm6</i>	LSM6 homolog, U6 small nuclear RNA associated	1.6	<0.0005
<i>Ugt1a7c</i>	UDP glucuronosyltransferase 1 family, polypeptide A7C	1.6	0.006
<i>Ugt1a6b</i>	UDP glucuronosyltransferase 1 family, polypeptide A6B	1.5	0.001
<i>Edem1</i>	ER degradation enhancer, mannosidase alpha-like 1	1.5	0.002
<i>Uck2</i>	Uridine-cytidine kinase 2	1.5	0.01
<i>D01Rik</i>	RIKEN cDNA 22 104 12D01 gene	-1.5	0.002
<i>H3f3b</i>	H3 histone, family 3B	-1.5	0.04
<i>Mlxip1</i>	MLX interacting protein-like	-1.5	0.001
<i>Slc30a8</i>	Solute carrier family 30 (zinc transporter), member 8	-1.5	0.0006
<i>Mdm1</i>	Transformed mouse 3T3 cell double minute 1	-1.5	<0.0005
<i>Slc29a4</i>	Solute carrier family 29 (nucleoside transporters), member 4	-1.6	0.001
<i>Nt5dc2</i>	5'-nucleotidase domain-containing 2	-1.7	0.02
<i>Sult1d1</i>	Sulfotransferase family 1D, member 1	-1.7	0.01
<i>Msx3</i>	Homeobox, msh-like 3	-1.8	0.04
<i>Cib-3</i>	Calcium and integrin-binding family member-3	-1.8	0.01
<i>CryβA2</i>	Crystallin, beta A2	-1.8	0.0009

^aShown are the 110 unique total transcripts that were identified, where each of these genes demonstrated at least 1.5-fold change, being statistically significant (P value < 0.01) with false discovery rate (P < 0.05). These 110 genes were categorized based on a classification scheme from ingenuity, and could be classified into more than one category. The miscellaneous category corresponds to genes that did not appear in any category shown here.

Cytokine treatment

For studies with murine islets, purified islets were incubated for 24 h in RPMI 1640 (ATCC, Manassas, VA, USA) supplemented with 1% nonheat inactivated FBS. The medium was supplemented with or without 5 ng/ml interleukin 1 β (IL1 β) or 10 ng/ml tumor necrosis factor- α (TNF- α , R&D Systems, Minneapolis, MN, USA), and RNA was isolated as described above. Three independent experiments were performed.

Statistical analysis

P values were calculated by Student's t -test (one tailed) with a significance level at P < 0.05 using Excel software. To

determine the statistical significance for a change in the real-time RT-PCR studies, the fold change was used relative to the control condition, which was set at 1.0. For TPH1, statistical analysis was performed using logarithmic transformed values.

Results

Global RNA expression analysis of mouse islets during pregnancy

Pregnancy-induced changes in islet gene expression were examined by RNA expression analysis using RNA prepared from islets isolated from pregnant (day 13.5) and control

(age- and sex-matched) mice. In all, 110 differentially expressed transcripts were identified (Fig. 1A), as defined by a statistically significant ($P<0.01$, FDR $<5\%$) >1.5 -fold change in expression. Table 1 lists the 110 differentially regulated transcripts using a classification scheme from ingenuity (see Materials and Methods). In Fig. 1B, these 110 differentially regulated genes are categorized into seven groups. An overrepresentation P test, which is a hypergeometric test and provides a representation of the enrichment of the genes in each category compared with the chip as a whole, demonstrated that the genes in each of the categories were overrepresented relative to the genes on the chip as a whole.

To confirm the results of the RNA expression analysis, real-time RT-PCR was performed on a select group of the differentially regulated genes (rationale for their selection is provided below). The gene encoding SP-D was examined as it demonstrated the greatest pregnancy-induced change in expression in the global expression array (12.8-fold increase in SP-D mRNA levels). This finding was verified by real-time RT-PCR (Fig. 2A) where the level of SP-D mRNA increased 22.4 ± 1.7 -fold (mean $\pm$ S.E.M., $n=3$, $P<0.05$) in pregnant compared with that in control islets. Real-time RT-PCR studies were performed on additional genes that demonstrated significant pregnancy-induced increases in mRNA levels and have been directly or indirectly implicated in islet function: osteoprotegerin (Schrader *et al.* 2007), TPH1 (Paulmann *et al.* 2009), TPH2 (Paulmann *et al.* 2009), cytokine-inducible SH2-containing protein (Dif *et al.* 2001), and isovaleryl-CoA dehydrogenase (MacDonald 2007). The pregnancy-induced changes in the expression were confirmed for each of these genes (Fig. 2A). In addition, the expression of two ligands for GPCRs, gastrin-releasing peptide (GRP), and vasoactive intestinal polypeptide (VIP) was increased; however, in follow-up real-time RT-PCR analyses, the changes in GRP and VIP mRNA levels, while trending in the same direction, did not reach statistical significance (Fig. 2A). In addition to the above mRNAs, the levels of mRNA encoding the PRLR, which has a well described role in β cells during pregnancy, as well as suppressor of cytokine signaling 2, which is known to attenuate PRLR signaling (Sutherland *et al.* 2007), were also increased by pregnancy in the global expression analysis and confirmed in real-time RT-PCR studies.

Among the mRNAs with a decreased level of expression in pregnant compared with that in control islets were mRNAs encoding zinc transporter-8 (ZnT8), neurogenin 3 (NGN3), and MLX interacting protein-like 1 (Mlxip1). ZnT8 and NGN3 are important for islet cell function and/or development (Gu *et al.* 2002, Lemaire *et al.* 2009), while the role of Mlxip1 in islets is unclear, although an association with NGN3 has been reported (Soyer *et al.* 2010). The pregnancy-induced changes in the expression for each of these genes were confirmed by real-time RT-PCR (Fig. 2B).

Figure 2C shows the results of western blot analyses that examined changes in the protein level of select genes in islets from pregnant mice compared with those in control mice.

Figure 2 Pregnancy induced changes in expression of select genes by real-time RT-PCR and western blot analysis. (A and B) Fold change in the level of select mRNAs in islets isolated from pregnant mice at day E13.5 compared with that in age- and sex-matched control mice. (A) Shows the results for genes with increased mRNA levels and (B) shows the results for genes with decreased mRNA levels in pregnant compared with the control islets. The change in the level of each of the transcripts in both (A) and (B) was statistically significant at $P<0.05$ (indicated by *), except for GRP ($P=0.13$) and VIP ($P=0.13$). Values are the mean fold change $\pm$ S.E.M. from three independently prepared samples (each individual sample was a pooled sample from three mice). Fold change was calculated by a comparative cycle time (C_t) approach (see Materials and Methods) where the housekeeping genes (*HPRT*, *TBP*, and *PPIA*) were determined by geNORM (Vandesompele *et al.* 2002) using an approach defined in section ‘Materials and Methods’. The house-keeping C_t s were used to derive the geometric means for the comparative C_t analyses. (C) Changes in the level of proteins encoded by three genes that exhibited pregnancy-induced changes in expression. Proteins were extracted from isolated islets from pregnant mice at day E13.5 and age- and sex-matched control mice. Representative immunoblots for prolactin receptor (PRLR), FFAR2, and TPH1 are shown. Shown are the results of a single experiment. Three independent experiments were performed. Each experiment was performed using islets prepared from a different control and pregnant mouse. After hybridization, the blots were stripped and re-probed with antibody against GAPDH.

The expression of each of the proteins was increased in pregnant compared with that in control islets, consistent with the pregnancy-induced change in the level of their mRNA.

A potential role of SP-D in pancreatic β cells

The transcript encoding SP-D exhibited the largest pregnancy-induced increase in expression in the global expression analysis, which was confirmed by the real-time RT-PCR studies. As the expression of SP-D in adult islets has not been observed before, we examined SP-D expression in islets during pregnancy. Using mouse monoclonal antibodies, SP-D was detected in the mouse lung tissue (Fig. 3A, lane 2, at ~ 40 – 45 kDa) but not in the brain (lane 1), consistent with the known pattern of SP-D expression (Motwani *et al.* 1995). The immunoreactive SP-D detected in mouse lung was consistent with monomeric SP-D, which is ~ 45 kDa (Sørensen *et al.* 2009). However, SP-D is commonly present in large complex structures (Sørensen *et al.* 2009). In unreduced preparations, the antibody typically recognizes proteins with a molecular weight of ~ 120 kDa and above, which likely represent trimers and other multimers. In our studies, using unreduced protein preparations, the anti-SP-D antibodies recognized proteins of ~ 120 and 180 kDa (see Fig. 3A, lane 3) consistent with previous reports (Sørensen *et al.* 2009). Interestingly, in murine islets, a single band of ~ 100 kDa was detected and in the nonreduced state, a band of ~ 180 kDa was observed (Fig. 3A, lanes 4 and 5). It has been previously described that differences in covalent binding patterns between SP-D monomers exist and can result in the formation of nonreducible dimers (Crouch *et al.* 1993, Guo *et al.* 2008, Matalon *et al.* 2009, Sørensen *et al.* 2009) that are detected by western blots. Therefore, this band at 100 kDa likely represents a dimer. To further explore this, we examined SP-D expression in the lung, pancreas, spleen, and islets (Fig. 3A, lanes 6–9). All tissues exhibited immunoreactive bands at both 45 and 100 kDa, although the relative intensities of the bands varied widely between tissues. These data are consistent with differences in SP-D complexes between the tissues. Finally, we compared the level of SP-D in islets from control and pregnant mice, and found increased expression in pregnant islets, consistent with the pregnancy-induced increase in the level of SP-D mRNA (Fig. 3B).

In addition to its role in surfactant biology, SP-D may have an anti-inflammatory role in the lung tissue (Sørensen *et al.* 2007). As pregnancy is characterized by increased levels of pro-inflammatory peptides (Richardson & Carpenter 2007), we hypothesized that SP-D mRNA levels may be regulated by pro-inflammatory molecules in islets. To address this possibility, the impact of treating isolated native murine islets for 48 h with IL1 β or TNF- α on SP-D expression was examined. As can be seen (Fig. 3C), IL1 β and TNF- α each increased SP-D mRNA levels in isolated murine islets.

Figure 3 Surfactant protein D (SP-D) expression in mouse islets. (A) Expression of SP-D in the lungs and islets. Proteins were extracted from mouse lung and islets. Samples from the brain (lane 1), lung (lane 2), mouse lung – not reduced (lane 3), islets (lane 4), islets – not reduced (lane 5), lung (lane 6), pancreas (lane 7), spleen (lane 8), and islets (lane 9) were either reduced or not reduced prior to separation by PAGE. The results are representative of the results of two to three independent experiments. (B) SP-D expression in pregnant islets. Protein extracts were prepared from islets isolated from pregnant mice at day E13.5 and age- and sex-matched control mice. Western blot analysis was performed as described in section ‘Materials and Methods.’ C, control; P, pregnant. After hybridization, the blots were stripped and re-probed with antibody against GAPDH. The findings are representative of the results of three independent experiments. (C) Effect of cytokines on SP-D mRNA levels in murine islets. RNA was prepared from murine islets treated for 48 h with 5 ng/ml IL1 β or 10 ng/ml TNF- α . The fold change in SP-D mRNA was determined by real-time RT-PCR, using the comparative method and the housekeeping gene, *18S*. Values represent the mean $\pm$ S.E.M. (three independent samples, $n=3$) and are the relative level of SP-D mRNA compared with the level in control islets not treated with cytokines. * $P<0.05$ compared with control mice.

Analysis of GPCR-specific expression in mouse islets during pregnancy

In the global RNA expression analysis, one GPCR, FFAR2, was differentially regulated by pregnancy (>1.5 -fold change in expression, $P<0.01$). As the genes for GPCRs tend to be underrepresented in the global RNA expression analyses, potentially leading to missed targets (Fredriksson & Schiöth 2005), further studies were performed to determine whether pregnancy induced changes in the expression of genes encoding members of this receptor family. To do this, a secondary approach, a real-time RT-PCR array, was used to examine the expression of transcripts encoding 343 different GPCRs. The expression of GPCRs in the mouse islets during pregnancy at E13.5 was compared with the expression in islets isolated from nonpregnant female mice. Of the 343 GPCRs probed in the real-time array, the mRNAs encoding 216 of these GPCRs were detectable in the islets, as defined by the

detection of a discernible melt curve in the control islets. Of the 343 GPCR transcripts that were examined in the real-time RT-PCR array, 92 were detected in the global RNA expression array. For this study, we focused on the 100 most highly expressed GPCR mRNAs in control islets (Table 2). Three of these GPCRs (FFAR2; GAL3R, galanin receptor 3; and CCKAR, cholecystokinin receptor A) exhibited a >2.5-fold change in the expression in pregnant compared with those in control islets. In the global RNA expression analysis, the mRNA encoding CCKAR was not detected. In contrast, FFAR2 mRNA exhibited a significant 1.5-fold increase in expression in the global RNA expression analysis and GAL3R mRNA was increased 1.3-fold ($P=0.19$). Follow-up real-time RT-PCR studies of these three GPCRs were performed (Fig. 4), and a significant pregnancy-induced increase in mRNA levels was observed for only FFAR2 and CCKAR ($P<0.05$).

Discussion

The global RNA expression analysis and the confirmatory real-time RT-PCR assays demonstrated a marked increase in the level of the mRNA encoding SP-D, a protein best known for its role in lung function (Botas *et al.* 1998). A clear role for SP-D in lung and surfactant homeostasis has been described (Sørensen *et al.* 2007), but expression of this gene in the adult islets has not been previously reported. However, a recent study identified SP-D as a marker of new β cell formation during the perinatal period (Aye *et al.* 2010). A global SP-D knockout has been created (Botas *et al.* 1998), although alterations in glucose metabolism and/or islet biology in these mice have not been reported. A possible metabolic role of SP-D is suggested by the observation that SP-D impacts systemic lipid levels and atherosclerosis (Sørensen *et al.* 2006b). Consistent with that is the observation that circulating SP-D levels are inversely associated with obesity in humans (Zhao *et al.* 2007). Furthermore, mice with a null mutation of the *SP-D* gene maintained on a normal diet exhibited more weight gain and increased adiposity compared with control mice (Sørensen *et al.* 2006a).

The novel observation that SP-D mRNA is expressed in β cells was confirmed by western blot and immunohistochemical analyses (see Fig. 3A and B). Through a combination of studies, it is apparent that SP-D in islets exists in a form that leads to nonreducible dimers. This is not surprising, considering the complex nature of this protein. First, SP-D ranges in size from 37 to 50 kDa in the monomeric form because of different glycosylation patterns (Leth-Larsen *et al.* 1999, Sørensen *et al.* 2009). Furthermore, SP-D trimers contain multiple disulfide bonds between cysteine residues as well as nondisulfide covalent bonds, and these nondisulfide covalent bonds result in dimers resistant to reduction (Crouch *et al.* 1993, Guo *et al.* 2008, Matalon *et al.* 2009, Sørensen *et al.* 2009). As the function of SP-D is dependent on these modifications (Leth-Larsen *et al.* 1999, Sørensen *et al.* 2009),

the modifications of SP-D in islets may have important implications for its function.

If SP-D indeed has a role in islet function, it could be hypothesized that, similar to endothelial cells where SP-D has an anti-inflammatory role (Snyder *et al.* 2008), it may function as an anti-inflammatory molecule in islets. As pregnancy is characterized by increased levels of circulating pro-inflammatory molecules (Richardson & Carpenter 2007), we examined the impact of the pro-inflammatory cytokines, IL1 β and TNF- α , on SP-D expression in islets. We demonstrated that not only SP-D mRNA is expressed in islets but also its expression is regulated by cytokines. This is consistent with the possibility that SP-D serves as an anti-inflammatory factor in islets during pregnancy, although future studies will be needed to more fully examine this possibility. As with another anti-inflammatory molecule, cytokine IL1 receptor antagonist, which protects mice fed with high-fat diets from hyperglycemia (Sauter *et al.* 2008), this type of molecule could have unique therapeutic potential.

Recent genome-wide association studies have reported association of a missense mutation in *SLC30A8*, the gene that encodes the ZnT8, with type 2 diabetes (Chimienti *et al.* 2006, Sladek *et al.* 2007). ZnT8 may also be an autoantigen in type 1 diabetes (Wenzlau *et al.* 2007). Expression of *SLC30A8* is largely localized to islets and, more specifically, β cells (Chimienti *et al.* 2006). ZnT8 is thought to play an important role in zinc transport into the insulin secretory granules (Nicolson *et al.* 2009). Thus, the decrease in ZnT8 mRNA levels in islets from pregnant compared with those in control mice was unexpected given the increased insulin secretory demands in pregnancy. However, pregnancy is a pro-inflammatory state, and recent observations (Egefjord *et al.* 2009, El Muayed *et al.* 2010) have shown that cytokines decrease the level of ZnT8 mRNA in islet cells. This may explain, in part, the pregnancy-induced decrease in ZnT8 mRNA levels.

In addition to the global RNA expression analysis, we examined genes encoding the GPCRs in greater detail. Multiple GPCRs are known to be expressed in islets (Winzell & Ahren 2007), but the impact of pregnancy on GPCR expression has not been previously examined. These receptors are well-known targets for multiple medicinal therapies. Moreover, given the important role of some GPCRs (e.g. the glucagon-like peptide 1 and parathyroid hormone-1 receptors) in the regulation of islet function and mass (De Leon *et al.* 2003), we explored the impact of pregnancy on the expression of mRNAs encoding GPCRs in islets using a real-time RT-PCR array. This array examined the level of mRNAs encoding 343 GPCRs in pregnant compared with that in control islets and was complementary to the global RNA expression analyses. Low levels of GPCR mRNA (Fredriksson & Schioth 2005) and the stringent requirements for statistical significance can lead to missed targets with the global RNA expression analysis. A recent study reported a role for serotonin in the regulation of β cell mass in pregnancy (Kim *et al.* 2010). The expression of mRNA encoding the serotonin

Table 2 Comparison of the 100 most highly expressed G-protein-coupled receptors (GPCRs) in the GPCR-specific real-time reverse transcriptase-PCR array compared with the global expression array for RNA from islets isolated from pregnant mice (E13.5) compared with control mice (age- and sex-matched)

	GPCR array		Global array	
	Relative expression ^a	Fold change ^a	Fold change ^a	<i>P</i> <value
GPCRs				
Glp1r	32 988	1.7	1.2	0.01
Sstr3	11 663	1.1	−1.1	0.03
Adra2a	9473	1.1	−1.0	0.59
Gabbr2	5441	2.1	ND	–
Gpr56	5441	1.4	−1.0	0.89
Gipr	5077	1.3	ND	–
Gpr116	5077	0.6	ND	–
Lphn1	5077	0.9	ND	–
Casr	4737	0.8	−1.0	0.29
Ffar1	4419	1.1	−1.1	0.14
Gpr108	4419	1.4	1.2	0.01
Gpr119	4123	2.0	ND	–
Vipr1	4123	0.7	1.0	0.52
Adora1	2720	0.7	−1.1	0.10
Gpr137b	2683	0.9	ND	–
Chrm3	2538	1.3	−1.0	0.40
Galr1	2368	1.0	−1.1	0.04
Gpr137	2210	1.2	ND	–
P2ry6	2062	1.6	1.0	0.56
Crcp	1924	0.8	−1.1	0.34
F2rl1	1795	0.7	−1.3	7.04×10^{-5}
P2ry5	1795	1.1	1.2	0.004
Ptger3	1360	0.9	ND	–
Bai3	1269	0.5	−1.0	0.31
Tm7sf3	1269	1.3	ND	–
Eltf1	1184	0.9	ND	–
Gabbr1	1184	0.9	ND	–
Gcgr	1184	1.2	−1.2	0.01
Gprc5c	1184	1.6	ND	–
Lgr4	1184	1.2	ND	–
Adcyap1r1	1105	0.7	ND	–
Fzd4	1105	0.9	1.0	0.58
Fzd3	1031	1.4	ND	–
Fzd6	1031	1.2	−1.1	0.01
Gpr125	1031	1.1	−1.0	0.12
Fzd7	962	1.0	−1.1	0.07
Taar1	962	1.0	−1.1	0.03
F2r	897	0.9	−1.1	0.07
Edg2	837	0.6	−1.1	0.06
Ednra	837	0.6	ND	–
Lphn3	837	0.8	1.0	0.54
Ccrl2	729	1.2	−1.0	0.40
Calclrl	680	1.1	ND	–
Cd97	680	1.2	ND	–
Gpr146	680	1.2	−1.0	0.10
Chrm4	635	0.6	ND	–
Fzd5	635	1.0	ND	–
Gpr120	635	1.1	−1.3	2.67×10^{-5}
Oxtr	635	1.6	ND	–
Celsr3	592	1.3	−1.2	0.04
Ednrb	592	1.4	1.0	0.53
Gprc5b	592	0.8	−1.1	0.01
Lphn2	592	1.1	ND	–
Fzd8	552	1.3	ND	–
Gpr85	552	0.8	−1.2	0.03
Cnr1	515	0.7	−1.0	0.58
Galr3	515	6.5	1.1	0.01
Gpr39	515	0.9	ND	–

(continued)

Table 2 Continued

	GPCR array		Global array	
	Relative expression ^a	Fold change ^a	Fold change ^a	P<value
P2ry1	515	0.8	ND	–
Gpr161	481	1.1	ND	–
Gpr75	449	0.7	ND	–
Gpr81	449	0.7	ND	–
Ffar2	419	2.8	1.6	3.96×10^{−5}
Mrgpre	364	1.8	ND	–
Ghsr	317	0.7	−1.1	0.06
Agtrl1	296	2.2	−1.0	0.66
Gpr41	296	1.5	ND	–
Npy1r	296	0.6	−1.0	0.93
Adora2b	258	0.7	−1.1	0.26
Gpr124	258	0.9	−1.0	0.20
Gpr63	258	0.8	ND	–
Cckar	240	11.1	ND	–
Adora2a	224	0.7	ND	–
Gpr19	224	1.1	1.1	0.03
Admr	209	1.2	ND	–
Avpr1b	209	0.9	ND	–
Cx3cr1	209	1.1	−1.0	0.51
Edg1	209	1.7	ND	–
Gprc5a	209	0.6	−1.0	0.82
Ebi2	195	0.8	ND	–
Gpr4	195	1.7	ND	–
Sstr1	195	1.0	ND	–
Celsr1	182	2.2	1.1	0.03
Cxcr3	182	1.3	ND	–
Gpr153	182	1.2	ND	–
Avpr1a	170	0.3	−1.0	0.61
Chrm1	170	0.5	ND	–
Crhr1	170	0.7	−1.0	0.13
Gprc6a	170	0.6	1.0	0.50
P2ry2	159	0.4	ND	–
Ccr2	148	1.1	ND	–
Gpr135	148	1.1	1.0	0.46
Tbxa2r	148	1.4	−1.1	0.01
Chrm2	138	0.6	ND	–
Emr1	138	1.6	−1.0	0.24
Fzd1	129	1.4	ND	–
Gpr109a	129	1.4	1.0	0.30
Gpr65	129	0.8	−1.0	0.28
Galr2	129	1.3	ND	–
Celsr2	129	0.6	ND	–

Indicated in bold are the G-protein-coupled receptors that demonstrated a fold change of >2.5.
^aShown are the 100 GPCRs with the greatest expression and their corresponding fold change in pregnant islets compared with those in control. Relative expression was calculated by $2^{C_T(\text{GPCR}) - C_T(\text{housekeeping gene})} \times 10^{-5}$. Fold change for the GPCR array and global array were calculated as outlined in section 'Materials and Methods'.

receptor, Htr2b, was increased at mid-gestation while that encoding Htr1d was increased at the end of gestation. In our studies, Htr2b was only weakly expressed in islets from both pregnant and control mice and a clear pregnancy-induced change in expression was not observed; Htr1d mRNA was detectable on day E13.5 but a pregnancy-induced change in the expression was not evident at that time point during pregnancy.

Our studies did exhibit a reproducible pregnancy-induced change in two GPCRs, FFAR2 and CCKAR. Of the two,

the role of CCKAR in islets has been more fully examined, with initial studies suggesting that CCKAR plays a role in mediating insulin secretion through CCK binding (Winzell & Ahren 2007). FFAR2, which exhibited a pregnancy-induced increase in mRNA and protein levels in islets, is a member of a recently identified family of GPCRs that are activated by free fatty acids. This family can be divided by the chain length (long versus short) of the free fatty acids that bind to the receptors. GPCRs that are specific for long-chain free fatty acids include GPR40 (FFAR1) and

Figure 4 Real-time RT-PCR analysis of GPCR mRNAs regulated by pregnancy in mouse islets at day 13.5 of pregnancy. The fold change in the level of mRNA encoding the following GPCRs, CCKAR, FFAR2, and GAL3R, in islets isolated from pregnant mice (day E13.5) compared with that in age- and sex-matched control mice was determined by real-time RT-PCR. Values are the mean $\pm$ S.E.M. from three independent experiments. The change in the level of each transcript was statistically significant at $P < 0.05$ (indicated by *), except for GAL3R ($P = 0.19$).

GPR120. In contrast, FFAR2 binds specifically to short-chain fatty acids (Kebede *et al.* 2009). Pregnancy-induced changes in the expression of the mRNAs encoding GPR40 and GPR120 were not observed. The role of FFAR2 in islets is unclear, although a previous study has demonstrated the expression of FFAR2 in islets (Regard *et al.* 2007). Moreover, evidence exists that FFAR2 is expressed in β cells, at least in pancreatic β cell lines (Kebede *et al.* 2009). Taken together, these data suggest that FFAR2 may be involved in the regulation of islet cell function or mass in pregnancy, although future studies will be required to define its role in islet cell biology.

During the preparation of this manuscript, a study exploring the global RNA expression analysis in islets from pregnant mice was reported (Rieck *et al.* 2009). These investigators identified 1907 genes that exhibited differential regulation compared with 110 in our study. This difference relates to the use of different analytic approaches. Using an approach similar to that of Rieck *et al.*, we also detected ~ 1500 differentially regulated genes. Of the 110 differentially regulated genes identified in our study, 37 were not shown to be differentially regulated (Rieck *et al.* 2009). A few of the notable exceptions were suppressor of cytokine signaling 2, *Ng2* and *ZnT8*, genes in which a pregnancy-induced change in expression was verified in our study. These differences may be explained, in part, by differences in the gene expression platforms used and the day of gestation studied, E13.5 vs E14.5.

To date, there are three genes that have been specifically identified as playing a role in gestational diabetes using mouse

models. The *PRLR*, as discussed below, has a clearly established role. The other two genes suggested to be involved in gestational diabetes are *FoxM1* and *menin*. The level of *FoxM1* mRNA was shown by Zhang *et al.* (2010) to be increased on E14.5. Our data show that *FoxM1* mRNA levels were increased on E13.5; however, this increase was not statistically significant. Zhang *et al.* (2010) also developed mice with pancreatic specific null mutations for *FoxM1* and found that these mice developed gestational diabetes. The other gene identified as having an important role in gestational diabetes was *menin* (Karnik *et al.* 2007). Karnik *et al.* demonstrated decreased *menin* mRNA levels in islets during pregnancy, while overexpression of *menin* led to hyperglycemia during pregnancy. Finally, expression of *menin* was shown to be regulated by prolactin (Karnik *et al.* 2007). In our study, a pregnancy-induced change in *menin* mRNA levels was not demonstrated, although only a single time point in pregnancy was examined.

Increased *PRLR* expression in islets during pregnancy was demonstrated over 10 years ago (Sorenson & Brelje 2009). In this report, we have confirmed those findings using the global RNA expression analysis and follow-up confirmation by real-time RT-PCR. Furthermore, a transgenic mouse model has confirmed the role of the *PRLR* in glucose homeostasis during pregnancy (Huang *et al.* 2009). Although prolactin signaling undoubtedly contributes to pregnancy-induced changes in β cell mass (Sorenson & Brelje 2009), other signaling pathways and downstream molecules are likely important in this process. Together, with the recently published study by Rieck *et al.* (2009), our study demonstrates multiple novel proteins that warrant further investigation for their role in pregnancy-induced changes in β cells. Many of these genes were not previously known to be regulated by pregnancy, and, in some cases, expression of the gene in islets had not been previously demonstrated. These data should lay the foundation upon which select genes can be more fully examined for their role in pregnancy-induced changes in pancreatic β cell function.

Supplementary data

This is linked to the online version of the paper at <http://dx.doi.org/10.1677/JOE-10-0298>.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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