The Notch Family Transcription Factor, RBPJ_K, Modulates Glucose Transporter and Ovarian Steroid Hormone Receptor Expression During Decidualization

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Abstract

During decidualization, endometrial stromal cells differentiate into a secretory phenotype to modulate the uterine micro-environment and promote embryo implantation. This highly metabolic process relies on ovarian steroid receptors and glucose transporters. Canonical Notch signaling is mediated by the transcription factor Recombination Signal Binding Protein for Immunoglobulin Kappa J Region (RBPJ). Loss of RBPJ in the mouse uterus (*Pgr^{crel}+Rbpj^{flox/flox}*; *Rbpj* c-KO) results in subfertility in part due to an abnormal uterine–embryonic axis during implantation and, as described herein, decidualization failure. Induced in vivo decidualization in *Rbpj* c-KO mice was impaired with the downregulation of decidual markers and decreased progesterone receptor (*Pgr*) signaling. Consistent with in vivo mouse data, *RBPJ* knockdown during in vitro Human uterine fibroblast (HuF) cell decidualization results in the reduced expression of decidual marker genes along with *PGR*. Expression of the glucose transporter, *SLC2A1*, was decreased in the *RBPJ*-silenced HuF cells, which corresponded to decreased *Slc2a1* in the secondary decidual zone of *Rbpj* c-KO mouse uteri. Exogenous administration of pyruvate, which bypasses the need for glucose, rescues *PRL* expression in *RBPJ*-deficient HuF cells. In summary, Notch signaling through *RBPJ* controls both ovarian steroid receptor *PGR* and glucose transporter *SLC2A1* expression during decidualization, and this dysregulation likely contributes to embryo implantation failure.

Keywords

decidualization, RBPJ, notch, progesterone receptor, glucose transport

Introduction

Receptivity of the uterus to an active blastocyst is initiated through both hormonal regulation by the corpus luteum and embryonic signals, which favor embryo attachment and trophoblast invasion.^{1–3} The process of invasion and embryonic cues trigger transformation of endometrial stromal fibroblasts into secretory, epithelioid-like decidual cells, termed decidualization.^{4–6} Decidualized stromal cells migrate toward and encapsulate the implanting embryo, where they simultaneously support and control trophoblast invasion.^{7–10} Clinical consequences of decidualization defects include recurrent pregnancy loss, preeclampsia, preterm labor, and intrauterine growth restriction.^{11,12} Implantation defects contribute to ~75% of failed human pregnancies, suggesting a need to identify mechanisms regulating the early events, including decidualization, during the establishment of a successful pregnancy.¹³

The evolutionarily conserved Notch signaling pathway mediates key developmental and homeostatic functions,

including cellular differentiation, survival, and apoptosis.^{14,15} Notch signaling consists of 4 heterodimeric transmembrane receptors (NOTCH1–4) whose 5 ligands (Delta-like1, 3, 4 and Jagged1, 2) are expressed on the cell membrane of adjacent cells, thereby initiating receptor–ligand interactions through a juxtacrine mechanism. Upon ligand binding, activated Notch receptors undergo an ADAM protease and gamma-secretase-mediated cleavage cascade that results in the intracellular

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release of the Notch intracellular domain (NICD), which is the active form of Notch.¹⁶ In canonical Notch signaling, the NICD translocates to the nucleus and binds to the Notch family transcription factor recombination signal binding protein-Jk (RBPJ), converting it from a basal transcriptional repressor to an activator of downstream target genes, such as the “Hairy enhancer of split” (HES) and Hes-related (HEY) transcription factor families.^{17–20}

During the implantation window, NOTCH1 expression is induced in endometrial stromal cells of the nonhuman primate and in women in response to the embryonic signal Chorionic Gonadotropin (CG), which promotes uterine receptivity and decidualization.^{1,21–24} We have shown that Notch signaling, specifically NOTCH1, is indispensable for the initiation of decidualization in both mice and women.^{23,25} Further, Notch receptors, ligands, and target genes are all reduced in eutopic endometrium of baboons and women with endometriosis, contributing to their disease-specific decidualization failure.²⁶ In mice, during implantation, Notch1 is initially induced in the stromal cells of the primary decidual zone and then subsequently increases in the secondary decidual zone.²⁵ Ablation of *Notch1* in the mouse uterus using a progesterone receptor (*Pgr*)-driven cre-expressing (*Pgr^{cre/+}Notch1^{fl/fl}*; *Notch1* c-KO) drastically inhibited their response to an *in vivo* artificial decidualization stimulus.²⁵ Decidualization failure in *Notch1* c-KO mice was associated with the downregulation of genes crucial for progesterone-induced stromal cell differentiation and pathways related to cell proliferation, cell cycle progression, and estrogen receptor α (*Esr1*) signaling.²⁵ *Notch1* c-KO mice produce reduced litter sizes in the first litter but, otherwise, exhibited normal fecundity,²⁵ suggesting compensation by other Notch receptors may have rescued fertility in these mice, since other Notch receptors have been implicated in decidualization.²⁷ Disruption of all potential Notch receptor signaling through ablation of the shared transcription factor, *Rbpj*, in the mouse uterus (*Pgr^{cre/+}Rbpj^{fl/fl}*; *Rbpj* c-KO) resulted in subfertility due to an abnormal embryonic–uterine axis and impaired decidual remodeling.²⁸ More recently, we identified secondary infertility in *Rbpj* c-KO mice that results from impaired postpartum uterine repair and enhanced inflammatory signaling.²⁹ While secondary infertility was also associated with impaired *in vivo* artificial decidualization response, it was unclear whether decidualization defects occur in nulliparous *Rbpj* c-KO mice contributing to their subfertility. In the current study, we sought to determine the role that RBPJ plays in decidual function beyond the initial implantation phase and its relevance in the setting of human decidualization.

Growing evidence supports the importance of endometrial glucose transporters during decidualization, and their dysregulation results in infertility.^{30–34} Facilitative glucose transporters (known as the GLUT or SLC2 family) regulate cellular glucose uptake and have distinct tissue-specific and temporal expression patterns.³⁵ SLC2A1 (GLUT1) is ubiquitously expressed in the uteri of both women and rodents.³⁰ Additionally, SLC2A1 is further induced during early gestation and decidualization of mouse and human endometrial stromal cells.^{32,36} Progesterone

administration increases SLC2A1 expression in stromal cells in both women and mice.^{31–33} Further, *Rbpj* mediates SLC2A1 expression in memory CD4⁺ T cells, which is essential for their survival in the mouse,³⁷ suggesting that this mechanism may be important for decidualization.

We hypothesized that Notch signaling through RBPJ mediates differentiation of endometrial stromal cells during the establishment of pregnancy. In the current study, we induced *in vivo* artificial decidualization in *Rbpj* c-KO and control mice (*Pgr^{cre/+}Rbpj^{fl/fl}*). *Rbpj* c-KO mice displayed reduced decidualization associated with altered *Pgr* signaling and downregulation of glucose transporter *Slc2a1* during decidualization. Our mouse studies were performed in parallel with *in vitro* decidualization of human uterine fibroblast (HuF) cells after siRNA silencing of *RBPJ*. Similar to *Rbpj* c-KO mice, the decidualization response was impaired when *RBPJ* was silenced and was also associated with reduced *Pgr* and *SLC2A1* expression. Finally, supplementation of pyruvate, bypassing the necessity for glucose, restored the decidualization response in *RBPJ*-silenced HuF cells. Through these translational studies, we have determined that RBPJ is essential for decidualization in both the mouse and the human, in part, through regulation of progesterone signaling and glucose transporter expression.

Materials and Methods

Mouse Models

All studies performed using animals were approved by the Institutional Animal Care and Use Committee of Michigan State University, East Lansing, Michigan. To avoid the embryonic lethality of a complete *Rbpj* knockout,³⁸ we generated a uterine-specific, conditional *Rbpj* knockout mouse. *Rbpj^{fl/fl}* mice contain LoxP sites flanking the sixth and seventh exons, corresponding to the DNA binding domain.³⁹ We crossed *Pgr^{cre/+}* mice, containing a knock-in of Cre recombinase in 1 copy of the *Pgr* gene while maintaining normal fecundity and *Pgr* function,^{40,41} with *Rbpj^{fl/fl}* (control) mice to generate *Pgr^{cre/+}Rbpj^{fl/fl}* (*Rbpj* c-KO) mice, in order to ablate *Rbpj* in *Pgr*-positive tissues as described previously.^{28,29}

In Vivo Artificial Decidualization Model

Induction of artificial decidualization was performed on 6-week-old *Rbpj* c-KO and control mice as described previously.²⁵ Mice were ovariectomized and then primed to mimic the hormonal milieu of early pregnancy with daily E2 (100 ng; Sigma-Aldrich, St. Louis, Missouri) for 3 days followed by 2 days of rest and then daily P4 (1 mg; Sigma-Aldrich) plus E2 (6.7 ng). Six hours following the third E2 plus P4 injection, a mechanical scratch of the antimesometrial luminal epithelium of the left uterine horn (S; stimulated) was performed using a blunted needle to induce decidualization. The unscratched right horn served as the hormonal control (US; unstimulated). Subsequently, daily injections of E2 and P4 were provided with sacrifice 3 or 5 days post-scratch ($n = 3\text{--}5$ mice/group). Uterine

tissues were collected, weighed, and either snap frozen in liquid nitrogen or fixed in 4% paraformaldehyde for paraffin embedding. Decidual response was calculated based on the ratio of uterine decidualized to control horn weight.

Immunohistochemistry

Paraformaldehyde 4% fixed paraffin-embedded tissues were sectioned to 6- μ m thickness and placed onto microscope slides (Thermo Fisher Scientific, Waltham, Massachusetts). For immunohistochemistry, each tissue section was dewaxed, rehydrated with a graded alcohol series, followed by heat-mediated antigen retrieval in citrate buffer (Antigen unmasking solution; Vector Laboratories, Burlingame, California) and 10% hydrogen peroxide treatment. Sections were blocked for 1 hour in 10% Normal Horse Serum (Vector Laboratories) in phosphate-buffered saline (PBS) incubated overnight at 4°C in one of the following primary antibodies: rabbit anti-PGR (1:500; DAKO, Carpinteria, California), mouse anti-SLC2A1 (1:100; Santa Cruz Biotechnology, Dallas, Texas). Next, sections were incubated in respective biotinylated secondary antibodies for anti-mouse IgG or anti-rabbit IgG (Vector Laboratories) followed by HRP-conjugated streptavidin. Detection for immunoreactivity was achieved using the DAB Substrate Kit (Vector Laboratories) followed by hematoxylin counterstaining. Staining intensity of each section was quantified by image analysis software *ImageJ* (NIH) resulting in a Digital HSCORE (D-HSCORE) of staining intensity as previously reported.⁴²

Isolation and Culture of Human Uterine Fibroblast Cells

Human uterine fibroblast cells were isolated from the decidua parietalis from term placentas.^{23,43} Tissues were procured after patient informed consent was obtained under a protocol approved by the Institutional Review Board at Michigan State University. Corresponding clinical information for each cell line used in this study are summarized in Supplemental Table S1. After trypsinization, cells were propagated in phenol-red-free RPMI 1640 (Thermo Fisher Scientific) supplemented with 10% Charcoal: Dextran Stripped-Fetal Bovine Serum (CDS-FBS). Each cell line obtained from a single placenta is considered a single sample for statistical analysis; therefore, at least 3 HuF cell lines plated in triplicate were used for each experiment. Further, cell lines beyond 6 passages were not used for experiments due to the potential for diminished decidualization response.

Transfection of RBPJ Small-Interfering RNA in HuF cells

Human uterine fibroblast cells were seeded in 6-well dishes at 1.5×10^5 cells per well in 10% CDS-FBS in RPMI. Once cells reached ~30% to 50% confluence, they were washed with sterile PBS then acclimated to serum and antibiotic-free Opti-MEM media (Thermo Fisher Scientific) for at least 30 minutes prior to transfection. Subsequently, nontargeting (Control; ON-TARGETplus Non-targeting pool; GE

Dharmacon, Lafayette, CO) and *RBPJ* small-interfering RNA (siRNAs; SMARTpool ON-TARGETplus RBPJ siRNA; GE Dharmacon, Lafayette, Colorado) were dissolved in Lipofectamine RNAiMAX (Thermo Fisher Scientific) diluted in Opti-MEM. This mixture was added to the cells to achieve a final concentration of 40 nmol/L of either control or RBPJ siRNA, and cells were incubated with the siRNA mixture for 5 hours. Finally, cells were washed with PBS and then incubated overnight in fresh RPMI with 10% CDS-FBS and antibiotics prior to any future experiments.

In Vitro Decidualization of HuF Cells

In vitro decidualization of HuF cells was performed through administration of an EPC cocktail (a combination of 36 nmol/L estradiol-17 β , 1 μ mol/L medroxyprogesterone acetate [MPA] and 0.5 mmol/L di-butyl-cyclic adenosine monophosphate [dbcAMP; dbcAMP; Sigma]) in 2% CDS-FBS in RPMI.^{23,43,44} Human uterine fibroblast cells were treated with EPC or vehicle (Veh) for 5 days with replacement of treatment and media every 2 days. For in vitro decidualization studies with glucose and pyruvate supplementation, 8 mmol/L D-Glucose or 16 mmol/L Sodium Pyruvate (Sigma) in RPMI, respectively, were used for dilution of EPC or vehicle. Glucose and pyruvate concentrations were chosen based on dose-response experiments using supra-physiologic concentrations.

RNA Isolation and Real Time-Quantitative PCR (RT-qPCR)

Total RNA was isolated from snap-frozen mouse tissues or cultured cells using TRIzol reagent (Invitrogen, Carlsbad, California) followed by reverse transcription to produce cDNA using the High Capacity cDNA Reverse Transcription Kit (Applied Biosystems, Foster City, California). Differential gene expression was determined using RT-qPCR performed with either TaqMan Universal Master Mix II (Applied Biosystems) or SYBR Green PCR Master Mix (Applied Biosystems) using the ViiA 7 qPCR System (Applied Biosystems). *RPL17* and *Rpl19* were used for normalization of cDNA from human and mouse samples, respectively. Primer sequences used for RT-qPCR are provided (Supplemental Table S2).

Protein Isolation and Western Blot Analysis

Total protein was isolated from HuF cells using RIPA lysis buffer, and the concentrations were calculated using the bicinchoninic acid (BCA) method (Thermo Fisher Scientific). Eight micrograms of total protein were separated on SDS/PAGE gels (Invitrogen) and then transferred to a PVDF membrane (Millipore, Burlington, Massachusetts). The membranes were then blocked and incubated overnight with primary antibodies: rabbit anti-PGR (1:2000; Thermo Fisher Scientific), mouse anti-SLC2A1 (1:1000; Santa Cruz Biotechnology), and rabbit anti- β -ACTIN (1:10,000; Sigma-Aldrich). After incubation with respective anti-rabbit IgG or anti-mouse

Figure 1. Decreased *in vivo* artificial decidualization response in *Rbpj* c-KO mice. Control and *Rbpj* c-KO mice ($n = 3-5/\text{group}$) were ovariectomized, hormonally primed and then mechanically induced to decidualize. (A) Decreased decidual response in *Rbpj* c-KO mice was present both 3 and 5 days following mechanical decidualization based on stimulated to unstimulated uterine horn weight. (B) Impaired decidualization was associated with reduced mRNA expression of decidual marker genes, *Bmp2* and *Wnt4*, in *Rbpj* c-KO mice on AD5. Data represented as mean $\pm$ standard error of the mean (SEM). $^{**}P < .01$; scale bar = 0.5 cm.

IgG secondary antibodies labeled with HRP (Vector Laboratories, Burlingame, CA), immune complexes were visualized using enhanced chemiluminescence (GE Life Sciences, Pittsburgh, PA).

Statistical Analysis

Variation between groups was determined using 2-way analysis of variance with repeated measures followed by an Holm-Sidak post hoc test to determine group-specific differences. Statistical significance was defined as $P < .05$. All statistical analyses were performed by GraphPad Prism 6.0 (GraphPad Software, San Diego, California).

Results

Recombination Signal Binding Protein- κ is Essential for Decidualization in the Mouse and Drives Progesterone Receptor Signaling

To induce *in vivo* decidualization, *Rbpj* c-KO and control mice were ovariectomized, hormonally primed to mimic early gestation and induced to decidualize via a mechanical scratch with collection as described previously.^{25,29} Decidualization was significantly reduced in *Rbpj* c-KO mice on both day 3 and day 5 of artificial decidualization (AD) based on stimulated (S) to US uterine horn weight ratio (Figure 1A). We chose to focus our studies on AD5 because this time point corresponded to day 9.5 postconception in natural mouse gestation, when initial

Figure 2. Downregulation of Pgr signaling in *Rbpj* c-KO mice during in vivo artificial decidualization. Ovarian steroid hormone receptor levels were investigated in *Rbpj* c-KO and control mice by RT-qPCR and immunohistochemistry. (A) *Pgr* mRNA was significantly increased in the decidual horn of control mice compared to the unstimulated horn, which was not the case in *Rbpj* c-KO mice ($n = 3$ -5/group). Additionally, *Pgr* target genes, *Hoxa10*, *Areg*, *Ptgs2* and *Cdk6* were significantly decreased in the decidual horn of *Rbpj* c-KO mice compared to control mice. (B) Reduced Pgr expression was localized to the decida of *Rbpj* c-KO mice. Data represented as Mean $\pm$ standard error of the mean (SEM). Comparisons were made using Two-Way ANOVA. * $P < .05$, ** $P < .01$; scale bar = 50 μ m. US indicates unstimulated, non-decidualized uterine horn, S, stimulated, decidualized uterine horn.

embryo loss begins in *Rbpj* c-KO mice potentially contributing to their subfertility.²⁸ Impaired decidualization in *Rbpj* c-KO mice on AD5 was associated with reduced uterine mRNA expression of decidual markers *Bmp2* and *Wnt4* (Figure 1B). Orchestrated ovarian steroid hormone receptor expression is necessary for promoting stromal survival and, ultimately, mediating the epithelial to stromal proliferation shift that occurs during implantation and decidualization.⁴⁵ Thus, expression patterns of Pgr and downstream signaling effectors were investigated in the uteri of artificially decidualized mice (Figure 2).

Pgr mRNA levels were significantly decreased in the stimulated uterine horns of *Rbpj* c-KO mice compared control mice (Figure 2A). Consistent with this finding, decreased levels of Pgr downstream effectors *Hoxa10* and its target *Cdk6*, *Areg*, and *Ptgs2* were decreased in the stimulated horn of *Rbpj* c-KO mice compared to the stimulated horn of control mice (Figure 2A). *Cdk6* plays an important role in mediating the *Hoxa10*-induced differentiation of stromal cells toward the decidual phenotype.⁴⁶ Immunostaining revealed reduced levels of Pgr protein in the stimulated horn of the *Rbpj* c-KO mouse

Figure 3. Reduced glucose transporter, *Slc2a1*, expression is associated with *Rbpj* loss during in vivo artificial decidualization. Glucose transporter mRNA and protein levels were investigated by RT-qPCR and immunohistochemistry, respectively, during artificial decidualization. (A) *Slc2a1* was significantly induced in the decidual horns of the control and *Rbpj* c-KO mice compared to control horns. *Slc2a1* mRNA expression was reduced in the control horn of *Rbpj* c-KO mice compared to control mice ($n = 3$ -5/group). (B) *Slc2a1* protein localization was significantly reduced in the decidual horn of *Rbpj* c-KO mice, particularly in the secondary decidual zone ($n = 4$ -5/group). Data represented as Mean $\pm$ standard error of the mean (SEM). Comparisons were made using Two-Way ANOVA for more than 2 groups or using a Student *t*-test for 2 groups. * $P < .05$, ** $P < .01$; Scale Bar = 100 μ m; US indicates unstimulated, non-decidualized uterine horn; S, stimulated, decidualized uterine horn; SDZ, secondary decidual zone; PDZ, primary decidual zone.

(Figure 2B). The Notch pathway regulates cell survival and differentiation,^{14,15} and failed initiation of stromal cell differentiation for decidualization was a consequence of reduced Pgr signaling in the absence of Notch signaling transcriptional factor *Rbpj*.

Endometrial Loss of *Rbpj* Reduces *Slc2a1* Induction During Decidualization

Glucose metabolism plays an important role in rapidly proliferating and differentiating cells, and there is mounting evidence that supports a role for endometrial glucose transporters during decidualization and embryo implantation.⁴⁶ We examined the mRNA levels of glucose transporter *Slc2a1* on AD5, which has been described extensively in the setting of natural pregnancy and in vitro decidualization of mouse endometrial stromal cells.^{32,47} However, to our knowledge, its expression has not been investigated in the setting of in vivo artificial decidualization. *Slc2a1* was up regulated in the stimulated horns of *Rbpj* c-KO and control mice (Figure 3A). There was no statistically significant difference between stimulated

horn mRNA levels of *Slc2a1*, although there was a trend toward decreased expression in *Rbpj* c-KO mice. Since glucose transporters are often transiently induced in a temporospatial manner, we performed immunostaining to determine both expression and localization of SLC2A1 in the mouse decidua on AD5. Consistent with the trend of decreased *Slc2a1* mRNA, *Slc2a1* protein levels were reduced in the decidual horn of *Rbpj* c-KO mice compared to control mice (Figure 3B). Interestingly, *Slc2a1* staining in control mice more specifically localized within the secondary decidual zone.

Recombination Signal Binding Protein- κ is Indispensable for Progesterone Receptor and Glucose Transporter Expression During Decidualization in HuF Cells

Human uterine fibroblast cells represent a proliferating population of undifferentiated stromal fibroblasts, which can be decidualized in vitro.^{23,43,44} To determine the role of RBPJ in human decidualization and the translational relevance of our work in the *Rbpj*-ablated mouse, HuF cells were decidualized for 5 days *in vitro* with a well-described hormone cocktail

Figure 4. Loss of *RBPJ* in HuF cells phenocopies the mouse during decidualization. Human uterine fibroblast cells were transfected with either control or *RBPJ* siRNA and subsequently treated with either vehicle or EPC [estradiol, medroxyprogesterone (MPA), and di-butylryl-cyclic adenosine monophosphate (dbcAMP)] to induce in vitro decidualization. Decidual markers, ovarian steroid hormone receptors and glucose transport molecules were investigated by RT-qPCR. (A) In response to EPC, mRNA expression of human decidual markers *PRL* and *IGFBP1* were significantly reduced with *RBPJ* siRNA knockdown compared to control siRNA treated cells. (B) Similar to the *Rbpj* c-KO mouse, *Pgr* expression was significantly reduced with loss of *RBPJ* during decidualization. (C) Glucose transporter, *SLC2A1*, mRNA expression was reduced in EPC-treated HuF cells with *RBPJ* knockdown. However, pyruvate transporter *SLC16A1* was unaffected by *RBPJ* loss with EPC treatment. (D) *PGR* A and B, along with *SLC2A1*, protein levels are significantly reduced with *RBPJ* siRNA knockdown during decidualization in HuF cells. Data represented as Mean $\pm$ SEM ($n = 3$ -4 HuF cell lines performed in triplicates); comparisons were made using Two-Way ANOVA. * $P < 0.05$.

consisting of estradiol, medroxyprogesterone acetate (MPA), and db-cAMP (EPC) following siRNA-mediated knockdown of *RBPJ*.²³ We confirmed that *RBPJ* expression was reduced in siRNA silenced HuF cells (Supplemental Figure S1A). Expression of human decidual markers *PRL* and *IGFBP1* were significantly reduced following *RBPJ* knockdown (Figure 4A). Similar to the *Rbpj* c-KO mouse, impaired decidualization was coupled with decreased *PGR* expression (Figure 4B). We investigated mRNA levels of glucose transporter *SLC2A1*, which was reduced in *Rbpj* c-KO mice. As expected, *RBPJ*

siRNA knockdown in decidualized HuF cells resulted in decreased mRNA expression of *SLC2A1* compared to control siRNA treatment (Figure 4C). Subsequently, we confirmed that gene expression levels correlated with protein levels by Western blot analysis. *PGR* A and B along with *SLC2A1* protein levels were significantly reduced with loss of *RBPJ* during decidualization in HuF cells (Figure 4D). Altogether, our findings indicate that similar to the mouse, *RBPJ* expression coordinates both *PGR* and glucose transporter expression in human endometrial fibroblasts, which are essential for decidualization.

Pyruvate Supplementation Rescues Decidualization Failure With RBPJ Loss in HuF Cells

Both in *Rbpj* c-KO mouse endometrium and isolated HuF cells where *RBPJ* was suppressed, decidualization was dramatically altered and was associated with decreased expression of glucose transporter *SLC2A1*. Therefore, we hypothesized that supplementation of additional glucose or pyruvate, which bypasses the necessity for glucose, might restore decidualization associated with loss of *RBPJ* in HuF cells. First, we confirmed that pyruvate transporter, *SLC16A1*, expression was unaffected by *RBPJ* loss (Figure 4C). Next, we supplemented the control and *RBPJ* knockdown HuF cells with additional glucose or pyruvate during in vitro decidualization of HuF cells. Dosages of glucose and pyruvate (8 mmol/L and 16 mmol/L, respectively) selected for this experiment were determined based on physiological dose–response experiments conducted previously in our laboratory and were ultimately chosen based on their ability to favor decidualization without negatively affecting cellular osmotic pressure. Remarkably, administration of pyruvate but not glucose restored expression of decidual marker *PRL* in *RBPJ* knockdown HuF cells during decidualization (Figure 5A). Additionally, *Pgr* mRNA levels were also restored through supplementation of glucose during decidualization with loss of *RBPJ* in HuF cells (Figure 5B).

Discussion

Notch is an arbiter of cell fate, and, based on our previous work, its expression regulates endometrial transformation during embryo implantation and more specifically, decidualization.^{23,26} In the current study, we demonstrate that ablation of *RBPJ* in the mouse uterus and in HuF cells reduces their capacity to differentiate toward the decidual phenotype. Contributors to impaired decidualization in both models included dysregulated *Pgr* expression and reduced glucose transporter *Slc2a1* expression. Additionally, supplemental pyruvate partially restores the expression of decidual markers in HuF Cells in which *RBPJ* was silenced during decidualization. Our findings support a previously undescribed contributor to subfertility in *Rbpj* c-KO mice, where implantation failure was shown to result from abnormal embryonic–uterine axis.²⁸ However, in the current study, we determined that *Rbpj* c-KO mice display a reduced in vivo response to artificial decidualization based on decidual horn wet weight and differentiation marker expression, which is independent of the presence of an embryo. Additionally, we determined a translational role for *RBPJ* in women, using an in vitro decidualization HuF cell model.

Similar to Notch1, *Rbpj* is dynamically expressed in a temporospatial manner in the mouse uterus during the establishment of pregnancy. Embryonic signals induce NOTCH1 expression, which include CG in primates or physical contact in mice.^{23–25} During implantation in mice, *Notch1* is initially expressed in the subepithelial stromal cells of the primary decidua, and as pregnancy progresses there is a shift of expression to the secondary decidua.²⁵ Consistent with the mouse,

Figure 5. Pyruvate supplementation rescues decidual marker gene and *PGR* expression in the setting of *RBPJ* loss. Human uterine fibroblast cells were transfected with either control or *RBPJ* siRNA and subsequently treated with either vehicle or EPC [estradiol, medroxyprogesterone (MPA), and di-butyl- γ -cyclic adenosine monophosphate (dbcAMP)] to induce in vitro decidualization with cell culture media supplementation of 8 mM Glucose, 16 mM Pyruvate, or neither. (A) Administration of pyruvate but not glucose restored expression of decidual marker *PRL* in EPC-treated *RBPJ* knockdown HuF cells. (B) Pyruvate supplementation increased *PGR* expression in decidualized *RBPJ* knockdown HuF cells compared to no glucose or glucose treatment. Data represented as Mean $\pm$ standard error of the mean (SEM) ($n = 3$ –4 HuF cell lines performed in triplicates). The effect of pyruvate in increasing *PRL* and *PGR* levels in *RBPJ* knockdown cells during decidualization has been highlighted with bars. Comparisons were made using Two-Way ANOVA. Significantly changed mRNA levels between two groups are represented by matching letters, where $P < .05$.

NOTCH1 initially increases in decidualizing HuF cells and subsequently becomes downregulated at the completion of decidualization.²³ Expression of *Notch1* is tightly regulated, where overactivation of Notch1 signaling impairs decidualization and causes infertility.⁴⁸ Similar to Notch1, *Rbpj* is initially induced in the primary decidual zone in the mouse and subsequently downregulated, at which point its expression is localized within the secondary decidual zone.²⁸ In the current study, *Slc2a1* staining was concentrated within cells of the

secondary decidual zone of control mice, which did not occur in the *Rbpj* c-KO mice. Overall, these findings suggest Slc2a1 expression is induced to a greater extent in stromal cells during the initiation of decidualization rather than upon completion of differentiation, which parallels the increase and the subsequent downregulation of Notch1 and Rbpj expression during decidualization.^{23,25,28}

During implantation, Pgr drives stromal proliferation and promotes transformation of endometrial fibroblasts toward a secretory phenotype while suppressing epithelial Esr1.⁴⁵ In part, Pgr signaling through *Hoxa10* promotes differentiation of stromal cells through induction of cell cycle regulators, such as Cdk6.^{46,49} In the *Rbpj* c-KO mice, broad downregulation of the Pgr signaling pathway was evident. A limitation to this study is the potential for haploinsufficiency of the Pgr gene in *Rbpj* c-KO mice, which could contribute to decreased Pgr expression. However, other studies in Pgr-cre mice have shown these mice display preserved Pgr expression.⁴⁰ Further, Pgr expression was decreased with *RBPJ* knockdown during decidualization of HuF cells, where haploinsufficiency does not occur.

Based on our prior and present work, a close relationship between Pgr and Notch signaling exists. P4-targets were decreased in *Notch1* c-KO mice during decidualization, including Cdk6, and Pgr mediates cleavage of Notch1 to its active form.^{23,25} Further, overactivation of Notch signaling indirectly suppresses Pgr expression via recruitment of DNA methyltransferases and hypermethylation of its promoter in an Rbpj-dependent manner.⁴⁸ Since Rbpj basally suppresses Notch target expression, its loss can mimic either downregulation or upregulation of Notch signaling.⁵⁰ Therefore, RBPJ mediates Notch receptor-independent functions, which may include regulation of Pgr signaling.

Glucose transporter SLC2A1 expression was downregulated during *in vivo* and *in vitro* decidualization in the absence of *RBPJ*. In women, stromal SLC2A1 expression is low in the proliferative phase and increases during the mid-secretory phase, corresponding to the period of uterine receptivity and increased progesterone levels.³¹ Further, inhibition of glucose transporters drastically reduced PRL secretion during *in vitro* decidualization of human endometrial stromal cells.³¹ In rat, endometrial stromal Slc2a1 expression pattern during early pregnancy mirrors that of Rbpj.^{28,47} Initially, Slc2a1 is induced in the stromal cells of the primary decidual zone and as implantation progresses, expression shifts to the secondary decidual zone.⁴⁷ In the current study, Slc2a1 staining was concentrated in the stromal cells of the secondary decidual zone on AD5 in control mice, which was reduced in the *Rbpj* c-KO mouse decidual horn. We confirmed decreased Slc2a1 expression in the mouse with loss of Rbpj occurs with *RBPJ* knockdown during decidualization in HuF cells. Based on our findings, we hypothesize that Rbpj acts upstream of Slc2a1 and that both are early mediators of proliferation and differentiation of stromal fibroblasts into decidual cells. After completion of differentiation during decidualization, both Notch1 and Rbpj are reduced in the primary decidual zone, which may

subsequently reduce Slc2a1 expression in the primary decidual zone.^{23,25}

In CD4⁺ memory T cells, Rbpj mediates survival through regulation of Slc2a1 expression and loss of Rbpj could be overcome by administration of pyruvate, which bypasses the requirement for glucose.³⁷ In the current study, we found that supplementation of pyruvate rescued expression of *PGR* and the decidual marker *PRL* associated with *RBPJ* knockdown in decidualizing HuF cells. Dysregulated glucose transporter expression represents an understudied contributor to infertility despite reduced levels of SLC2A1 in midsecretory biopsies of patients with idiopathic versus clear anatomical causes for infertility.³¹ A positive impact of pyruvate supplementation on embryo development in culture media used for *in vitro* fertilization patients has been previously reported.⁵¹ However, to our knowledge, the effect of supplementing pyruvate in IVF culture media during embryo transfer on implantation rates in the setting of impaired uterine receptivity has not been determined. We have shown that impaired decidualization can be restored with supplementation of low levels of pyruvate, through potentially bypassing the necessity for glucose. Future studies are indicated to determine whether endometrial receptivity can be restored in the setting of infertility through supplementation of pyruvate.

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Declaration of Conflicting Interests

The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

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Supplemental Material

Supplemental material is available for this article online.

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