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Source / Izvornik: **Cellular physiology and biochemistry**, 2017, 42, 2318 - 2329

Journal article, Published version

Rad u časopisu, Objavljena verzija rada (izdavačev PDF)

<https://doi.org/10.1159/000480004>

Permanent link / Trajna poveznica: <https://urn.nsk.hr/urn:nbn:hr:184:416742>

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Original Paper

Spatial and Temporal Analyses of FGF9 Expression During Early Pregnancy

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Fibroblast growth factor 9 (FGF9) • Fibroblast growth factor receptor (FGFR) • Implantation • Angiogenesis

Abstract

Background: Fibroblast growth factors (FGFs), in complex with their receptors (FGFRs), regulate a broad spectrum of biological functions including cellular proliferation, survival, migration, and differentiation. In human endometrial stromal cells, FGF9 is regulated with estrogen (E). **Methods/ Results:** First, we report that in uterus tissue of ovariectomized wild type mice, FGF9 is present in three isoforms and is regulated with E. Second, we found that during peri-implantation, *Fgf9* expression reached its peak at day 4.5 of pregnancy. Immunofluorescence analyses demonstrated overlapping FGF9 and COX2 expression surrounding the blastocyst attachment site. Next, we identified FGF9- and CD31-positive cells as a part of the microvessels; however, expression was localized to a distinct population of cells. Finally, our data showed synchronized, spatial expression of FGF9 on the luminal epithelium with FGFR2 present on the trophectoderm. **Conclusion:** Our data suggest that FGF9 is a crucial factor required to establish the appropriate microenvironment for successful implantation and the maintenance of pregnancy.

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Published by S. Karger AG, Basel**Introduction**

The steroid hormones estrogen (E) and progesterone (P), through their cognate receptors estrogen receptor (ER) and progesterone receptor (PR), prepare the uterus for pregnancy [1, 2]. The transcriptional activity of PR and ER leads to the synthesis of cytokines, growth factors, lipid mediators, and genes that control the dynamics of uterine transition to a receptive state and the establishment of pregnancy [2, 3]. In mice, the window of uterine receptivity corresponds to the P-mediated downregulation of ER activity in the uterine luminal epithelium (LE). At day 4 of pregnancy, in coordination with P, a short surge of E is nidatory for embryo attachment and implantation [1]. Suppression of ER-mediated proliferative activity by PR is mandatory for implantation and the establishment of pregnancy [4]. A recent study showed that the anti-proliferative activity of P in the uterine epithelium

is mediated by heart and neural crest derivatives expressed transcript 2 (HAND2). Specific deletion of *Hand2* in postnatal uteri [5] results in infertility and increased expression of several fibroblast growth factors (*Fgfs*).

The vertebrate FGF family consists of 22 heparin-binding polypeptide growth factors. Most FGFs are mitogens and stimulate various cellular functions including migration, proliferation, and differentiation [6, 7]. These activities are critical for a wide variety of physiological and pathological processes including embryonic development, angiogenesis, and tumorigenesis [8]. The biological function of FGFs is mediated by signaling through transmembrane tyrosine kinases receptors (FGFR) that are encoded by four distinct genes (*Fgfr1–Fgfr4*) [9]. In addition to the membrane localization of FGFRs, FGF induced cellular differentiation was found to be accompanied by the nuclear translocation of FGFRs [10].

In the uterus, components of the FGF signaling pathway can act as paracrine and/or autocrine mediators of epithelial-stromal interactions. Paracrine activation of FGFRs induces proliferation of the luminal epithelium in the uterus [11, 12]. Mice with uterine-specific *Hand2* deletions failed to suppress the expression of *Fgf1*, *Fgf2*, *Fgf9*, and *Fgf18* on day 3.5 of pregnancy [5]. A previous study demonstrated dynamic changes in FGF2 expression during peri-implantation [13]. In addition, FGF9 is a survival and mitogenic factor that is induced in endometrial cells by E and prostaglandins (PGs) [14, 15].

FGF9 is widely expressed in mouse embryos [16]. Ablation of *Fgf9* alleles leads to lethality at the neonatal stage mainly due to malformations of the lung, in addition to causing male-to-female sex reversal [17]. To date, the expression of FGF9 and its receptors during peri-implantation has not been investigated. In this study, we report the spatial and temporal analyses of FGF9 expression during early pregnancy. We first compared the expression of P-regulated FGFs at day 2.5, 4.5, and 7.5 of pregnancy, and found that only *Fgf9* and *Fgf2* expression changed significantly during peri-implantation. We found that the expression of *Fgf2* mRNA was highest at day 7.5 of pregnancy, whereas *Fgf9* expression peaked at day 4.5 of pregnancy when implantation takes place.

Materials and Methods

Animals and tissue collection

All experimental and surgical procedures complied with the Guide of Care and Use of Laboratory Animals and were approved by the ethical committee of the University of Rijeka Faculty of Medicine and Ministry of Agriculture. Mice (BALB/c) were housed in humidity (55 ± 10%) and temperature (22 ± 2 °C) controlled rooms with a 12-hour light-dark cycle with free access to food and water.

Wild-type females (10–12-weeks old) were housed with fertile males; the day that the vaginal plug was observed was considered day 0.5 of pregnancy. The animals were sacrificed by cervical dislocation on day 2.5, 4.5, and 7.5 of pregnancy. Implantation sites at day 4.5 of pregnancy were visualized by intravenous injection (0.1 ml/mouse) of Chicago blue B dye (Sigma Aldrich, Taufkirchen, Germany) solution (in 1% saline). A discrete blue band represented the implantation site. One uterine horn was fixed in 4% paraformaldehyde and the other was frozen at -80 °C for RNA or protein isolation. For reverse transcription-real-time-polymerase chain reaction experiments and western blot analysis, the conceptuses on day 7.5 of pregnancy were carefully dissected out of the implantation site. To analyze protein expression during implantation, implantation and non-implantation sites were separated.

Wild-type mice (BALB/c) were ovariectomized [18] (from 6–8-week old animals) and allowed to rest for 2 weeks before hormonal treatment. Mice (n = 3/treatment) were given daily subcutaneous injections of sesame oil (vehicle), sesame oil containing E (250 ng), and sesame oil containing E and P (EP) (250 ng / 2.5 mg) for 4 days. Uterus tissues were frozen at -80 °C for mRNA or protein isolation.

Immunofluorescence

Uterine sections (5 µm) were prepared from tissues that were fixed in 4% (wt/vol) paraformaldehyde and paraffin-embedded. The sections were deparaffinized, rehydrated in graded concentrations of ethanol, and subjected to heat-induced antigen retrieval (10 mM sodium citrate, pH 6.0). The slides were then

washed in PBS, pre-incubated in 1% bovine serum albumin (BSA, Affymetrix, Santa Clara, CA, USA) in PBS containing 0.001% NaN_3 for 1 hour at room temperature. The following primary antibodies were used: mouse monoclonal FGF9 (D-8) IgG₃ (1:100 dilution, Santa Cruz Biotechnology, Santa Cruz, CA, USA), rabbit monoclonal Cox2 (SP21) IgG (1:200 dilution, Thermo Scientific, Waltham, MA, USA), rabbit monoclonal CD31 (PECAM-1) (D8V9E) IgG (1:100 dilution, Cell Signaling, Leiden, Netherlands), rabbit polyclonal FGFR-3 (C-15) IgG (1:200 dilution, Santa Cruz Biotechnology, Santa Cruz, CA, USA), and rabbit polyclonal FGFR-2 (Bek, C-17) IgG (1:50 dilution, Santa Cruz Biotechnology, Santa Cruz, CA, USA). Primary antibodies were diluted in 1% BSA in PBS containing 0.001% NaN_3 and incubated with tissue sections overnight at 4 °C in a humid environment. To visualize immunocomplexes, Alexa donkey anti-rabbit 594 nm (1:500 dilution, Molecular Probes, Eugene, OR, USA) and Alexa goat anti-mouse IgG₃ (1:300 dilution, Molecular Probes, Eugene, OR, USA) were used. Secondary antibodies were diluted in blocking solution and incubated with tissue sections in the dark for 1 hour at room temperature in a humid environment. Nuclei were visualized with DAPI (Vector Laboratories, Burlingame, CA, USA). Images were captured using an Olympus (Tokyo, Japan) imaging system equipped with a DP71CCD camera, and CellF imaging software was used. Images were edited using Photoshop CS6 (Adobe, San Jose, CA, SAD).

RNA isolation and real-time RT-PCR analysis

Total RNA was isolated from uterine tissues (n = 3 per pool) using Trizol (Invitrogen, Carlsbad, CA, USA) according to the manufacturer's protocol. The RNA was reverse transcribed and gene levels were quantified by quantitative real time-PCR (qRT-PCR) using the ABI Prism 7300 detection system (Applied Biosystems, Foster City, CA, USA). This procedure was independently repeated three times. Oligonucleotide primers for mouse *Fgf1* (forward 5'-TTTATACGGCTCGCAGACAC-3'; reverse 5'-CGCTTACAGCTCCCGTTCTT-3'), *Fgf2* (forward 5'-GGCTGCTGGCTTCTAAGTGT-3'; reverse 5'-GTCCCGTTTTGGATCCGAGT-3'), *Fgf9* (forward 5'-GCAGTCACGGACTTGATCA-3'; reverse 5'-AATTCCAGAATGCCGAAGCG-3'), *Fgf18* (forward 5'-CCTGCACTTGCTGTGTTTA-3'; reverse 5'-GCTGCTCCGACTCACATCA-3') and *Cox2* (forward 5'-GGGCCATGGAGTGGACTTAAA-3'; reverse 5'-TGCAGGTTCTCAGGGATGTG-3') were obtained from Metabion (Steinkirschen, Germany). Relative mRNA levels were determined after normalization to mouse *Gapdh* expression [19].

Western blotting

Protein extracts were obtained from uterine tissues (n = 3) using RIPA buffer (Thermo Scientific, Waltham, MA, USA). The concentration was determined using a Bradford protein assay kit (BioRad, Hercules, CA, USA). Proteins were separated by 12% sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE), under reducing conditions, and then transferred onto a nitrocellulose membrane (BioRad, Hercules, CA, USA). The membrane was blocked with 5% BSA in tris-buffered saline (TBS) with 0.05% tween-20 (TBST) and incubated with rabbit anti-FGF9 IgG antibody (1:2000 dilution, Abcam, Cambridge, UK) followed by incubation with species-specific HRP-conjugated secondary antibodies. In accordance with the datasheet, this antibody recognizes three species of FGF9, specifically, 30, 29, and 25 kDa. Mouse anti- β -actin (1:160 000 dilution, Millipore, Billerica, MA, USA) was used as a control of protein loading. For detection, membranes were incubated with Amersham ECL Prime (GE Healthcare, Little Chalfont, UK), and scanned with an Alliance 4.0 imager (Uvitec, Cambridge, UK).

Statistical analysis

All data are reported as mean \pm S.E.M. and were analyzed by a Student's t test and one-way ANOVA followed by the Scheffe post-hoc method. $P < 0.05$ was considered significant. All statistical analyses were performed using STATISTICA 10 (StatSoft Inc, Tulsa, OK, SAD).

Results

Expression of Fgfs in the uterus during early pregnancy

Gene expression profiling on day 3.5 of pregnancy revealed elevated expression of mRNA corresponding to several FGF members in mice with uterine-specific Cre-mediated ablation of *Hand2* [5]. To further assess the role of FGFs, we determined the mRNA expression of

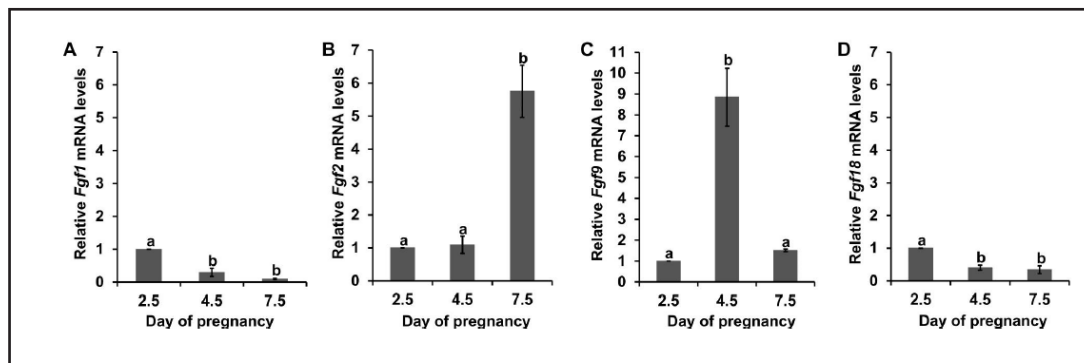


Fig. 1. Relative levels of *Fgf1* (A), *Fgf2* (B), *Fgf9* (C) and *Fgf18* (D) mRNA expression in uteri during early pregnancy. Mice were bred and pregnancy was indicated by the appearance of a vaginal plug (day 0.5 of pregnancy), after which animals were sacrificed at the indicated time points. Uterine horn tissue was removed and real time reverse-transcription PCR was performed for quantification of gene expression. Bars represent mean \pm SEM and those with different letters are significantly ($p < 0.05$) different based on an ANOVA followed by the Scheffe post-hoc method.

Fgf1, *Fgf2*, *Fgf9*, and *Fgf18* on days 2.5, 4.5, and 7.5 of pregnancy (Fig. 1). With pregnancy progression, we observed significant changes in the expression levels of mRNA encoding FGF2 and FGF9. *Fgf9* mRNA expression reached its peak at day 4.5 of pregnancy when embryo attachment induces the initiation of decidualization (Fig. 1C). With the development of the mesometrial decidua (basalis) at day 7.5 of pregnancy, a decline in *Fgf9* mRNA expression occurred (Fig. 1C). Whereas *Fgf9* mRNA decreased, *Fgf2* mRNA expression reached its maximum (Fig. 1B) at day 7.5 of pregnancy. These data strongly suggest that FGF9 and FGF2 have distinct but overlapping roles during peri-implantation; specifically, FGF9 dominates during implantation and primary decidualization, whereas FGF2 activity is involved in the control of secondary decidua zone formation.

FGF9 protein expression during peri-implantation

To the best of our knowledge, spatiotemporal expression of FGF9 during the peri-implantation period in mouse uteri has not been analyzed previously. Therefore, our next aim was to determine the localization of FGF9 on day 2.5, 4.5, and 7.5 of pregnancy. For this, we employed immunohistochemistry for the fluorescent detection for FGF9, and counterstaining with DAPI. At day 2.5 of pregnancy, FGF9 was primarily localized to the apical surface of epithelial cells but was also found in the glandular epithelium (GE) and some stromal cells (Fig. 2A, I-II). Next, on day 4.5 of pregnancy, when embryo attachment and implantation commences, we utilized a vascular permeability assay to identify implantation sites [18]. The implantation sites, detected as distinct blue bands, were dissected and prepared for analyses as described in the materials and methods section. The robust fluorescent staining for FGF9 was identified on the apical surface of the LE surrounding the implanting conceptus. Staining for FGF9 was also present throughout the cytoplasm of the LE, on the apical surface of the glandular epithelium (GE), and subpopulation of stromal cells (Fig. 2A, III-IV). By day 7.5 of pregnancy, FGF9 protein levels declined, but remained localized to the apical surface of the residual LE (Fig. 2A, V-VI).

Next, to analyze FGF9 distribution during early pregnancy we prepared protein extracts from uteri at day 2.5, 4.5, and 7.5 of pregnancy for western blot analyses (Fig. 2B). At day 4.5 of pregnancy, protein extracts were prepared separately from dissected uterine sections labeled with Chicago blue (implantation site) and from those that did not retain the blue color (non-implantation site). Our results revealed that of the three expected FGF9 protein species (25, 29, and 30 kDa) [20], the 25-kDa FGF9 protein species was not detected in the uterus during early pregnancy (Fig. 2B). Although we detected FGF9 protein species of 29 and 30 kDa at all analyzed time points of pregnancy, our result indicates that the 30 kDa form was

Fig. 2. Expression of FGF9 during peri-implantation. (A) Localization of FGF9 protein in mouse uteri at day 2.5 (I-II), 4.5 (III-IV), and 7.5 (V-VI) of pregnancy. Immunofluorescence staining with a mouse monoclonal antibody to FGF9 (green) is shown. Nuclei were counterstained with DAPI (blue). The arrow in panel IV shows an embryo at the implantation site. These results are representative of at least three independent samples. Scale bars = 200 μ m (I, III, V) and 20 μ m (II, IV, VI); dpc: days post coitum; (*) represents region at higher magnification. (B) FGF9 protein expression during early pregnancy. FGF9 protein was present as two separate bands of approximately 30 and 29 kDa. Actin was used as a loading control; NIS: non-implantation site; IS: implantation site. These results are representative of three independent samples.

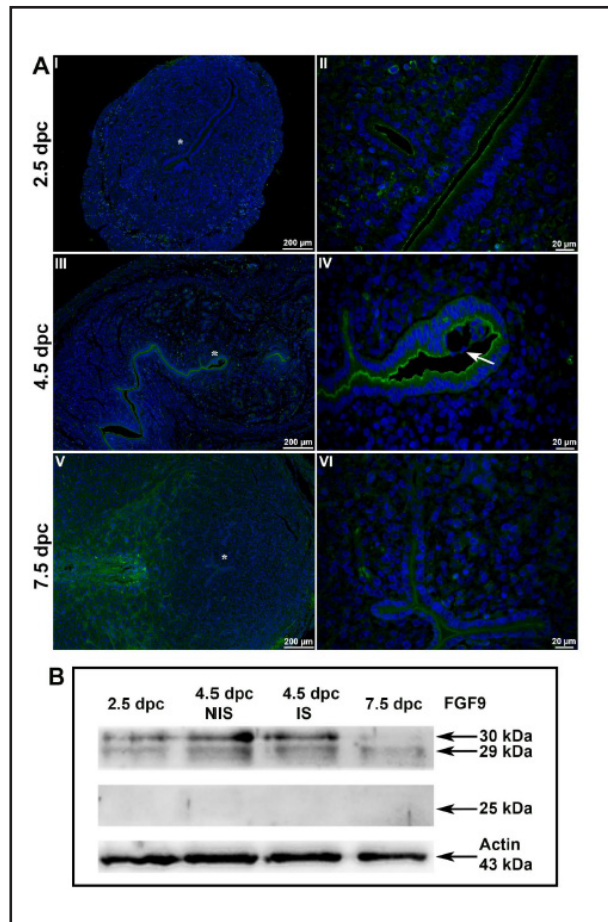
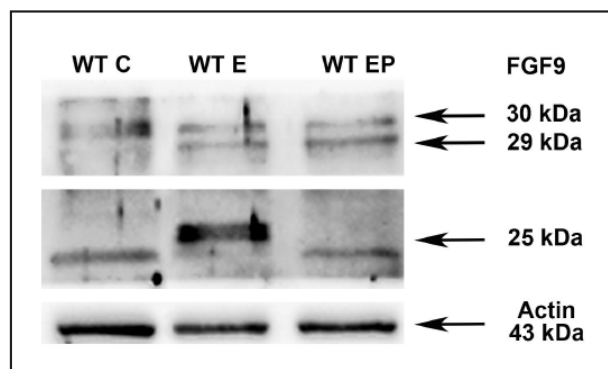


Fig. 3. Analysis of FGF9 expression in the uterus of ovariectomized mice. Animals were treated with sesame oil as a control (C), estrogen (E), or estrogen/progesterone (EP). FGF9 was present as three separate bands of approximately 30, 29, and 25 kDa. Actin was used as a loading control. These results are representative of three independent samples. No signals were observed when primary antibodies specific for FGF9 were replaced with whole rabbit IgG, confirming antibody specificity (data not shown).



dynamically expressed, whereas expression of the 29 kDa form was constant. Stronger FGF9 expression was detected at day 4.5 of pregnancy, with no appreciable difference between implantation and non-implantation sites (Fig. 2B).

We also examined FGF9 expression in ovariectomized mouse uteri after treatment with E, EP, and sesame oil as a control (Fig. 3). In addition to the 29 and 30 kDa bands, an additional 25-kDa band was present after E treatment (Fig. 3). Altogether, our results imply that FGF9 activity could be necessary for embryo implantation.

Expression of FGF9 at the implantation site

Tightly controlled spatial and temporal cellular and molecular changes induced by E and P are required for establishing the window of implantation. Consequences of ER α ablation include partial loss of PR expression and complete loss of embryo attachment and implantation [21, 22]. Previously, it was reported that along with E, PGE2 induces the expression of FGF9 [23, 24]. In addition, the rate-limiting step in PGE2 synthesis is regulated by COX2 [25]. In mice, genetic ablation of COX2 results in infertility because of defective implantation and decidualization [26]. Earlier analyses using in situ hybridization and immunohistochemistry suggested that COX2 expression was restricted to the stroma surrounding the implantation site at day 4.5 of pregnancy [27].

Here we analyzed *Cox2* and *Fgf9* mRNA and protein expression using qPCR and immunostaining with indirect immunofluorescence. We found a substantial increase in *Cox2* mRNA expression at the site of implantation, when compared to that at the non-implantation site (Fig. 4D). Immunostaining for COX2 indicated that at day 4.5 of pregnancy, very strong expression of COX2 was present in stromal cells proximal to the implanting embryo (Fig. 4, A-C). In addition, at the same point of pregnancy, our analyses indicated that lower levels of COX2 were also present throughout LE (Fig. 4B). Also, at day 4.5 of pregnancy, immunofluorescence indicative of FGF9 (Fig. 4, E-G) was found in the LE, but not in COX2-positive stromal cells (Fig. 4C). Contrary to *Cox2* mRNA, *Fgf9* mRNA expression was not significantly different between implantation and non-implantation sites (Fig. 4H). Furthermore, studies with COX2 mutant mice showed that COX2-derived PGs are important

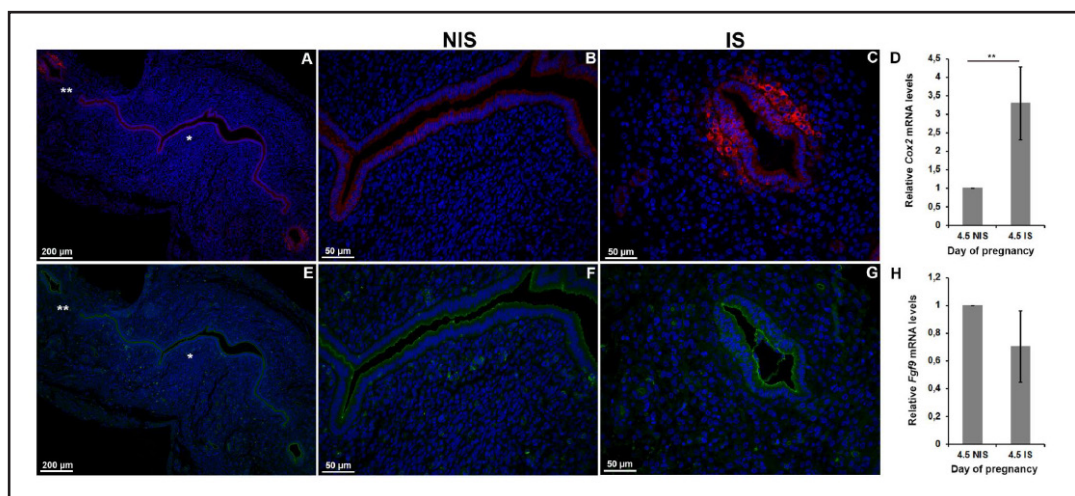


Fig. 4. Distribution of COX2 and FGF9 protein at day 4.5 of pregnancy. (A&E) Expression of COX2 and FGF9 at day 4.5 of pregnancy. (B&F) Localization of COX2 and FGF9 at non-implantation site. (C&G) Localization of COX2 and FGF9 at implantation site around blastocyst attachment. Mice were sacrificed at the indicated time point and uterine tissues were isolated. Immunofluorescence staining was performed with a rabbit monoclonal antibody to COX2 (red) and mouse monoclonal antibody to FGF9 (green). Nuclei were counterstained with DAPI (blue). These results are representative of at least three independent samples. Scale bars = 200 μ m (A, E) and 50 μ m (B, C, F, G); dpc: days post coitum; NIS: non-implantation site; IS: implantation site (*) represents region at higher magnification at non-implantation site; (**) represents region at higher magnification at implantation site. Difference in relative levels of *Cox2* and *Fgf9* mRNA expression between non-implantation and implantation site (D&H). Mice were bred and pregnancy was indicated by the appearance of a vaginal plug, after which animals were sacrificed at day 4.5 of pregnancy. Implantation sites at day 4.5 of pregnancy were visualized by intravenous injection of Chicago blue B dye. The implantation sites, detected as distinct blue bands, were dissected and prepared separately from the non-implantation site for RNA isolation. Real time reverse-transcription PCR was performed for quantification of gene expression. Bars represent mean \pm SEM and those with ** are significantly ($p < 0.01$) different based on the Student's *t* test.

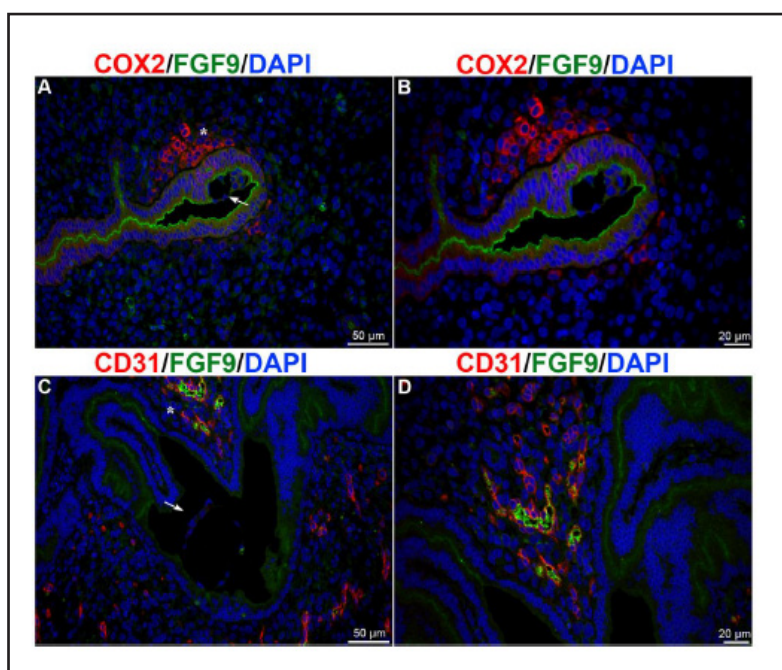
for vascular permeability and angiogenesis during implantation and decidualization [28]. Therefore, we applied triple immunofluorescence to analyze the relationship between FGF9 and COX2 at the implantation site (Fig. 5A & B). We found a strong signal for FGF9 and COX2 in cells surrounding the implantation site. In the LE, FGF9 was detected on the apical site of the cells, whereas COX2-positive cells were present in the stroma as well as in some cells of the LE. Interestingly, in the LE, COX2-positive cells were localized to the site of immediate contact between the embryo and the LE (Fig. 5A & B). This indicates that local COX2 enzymatic activity should induce adequate expression of PGs for stimulation of FGF9 activity at the site of embryo implantation.

As aforementioned, one of the first signs of implantation is increased endometrial vascular permeability at the site of embryo apposition [29]. In addition, it is known that FGFs are strong angiogenic inducers [30]. Previously, it was reported that FGF9 delivery during angiogenesis produces durable, vasoresponsive microvessels wrapped by smooth muscle cells [31]. Therefore, we hypothesized that FGF9 might be associated with angiogenesis during early pregnancy. For these analyses, we used an antibody specific for FGF9 and a marker of endothelial cells, specifically, anti-CD31 (PECAM-1). With triple-labeling immunofluorescence, we identified that FGF9- and CD31-positive cells were part of the microvasculature, but localized to a distinct population of cells (Fig. 5C & D). Therefore, FGF9-positive cells could represent a mural population of cells. Mural cells, consisting of vascular smooth muscle cells and pericytes, cover endothelial cells to regulate vascular stability and homeostasis [32]. This finding indicates that FGF9 could have an important role in microvessel maturation during angiogenesis at the implantation site.

Expression of FGF9 and its receptors at the implantation site

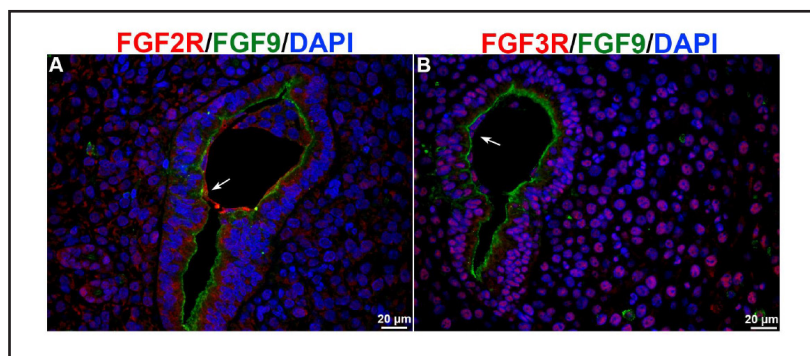
The biological activity of FGFs is mediated by four different tyrosine kinase receptors (FGFR1–4). To exert its effect, FGF9 is known to bind FGFR2, FGFR3, and FGFR4 [33]. We

Fig. 5. FGF9 expression at the implantation site at day 4.5 of pregnancy. (A&B) Co-expression of FGF9 and COX2 at the implantation site at day 4.5 of pregnancy. Mice were sacrificed at the indicated time point and uterine tissues were isolated. Triple immunofluorescence staining was performed with a mouse monoclonal antibody to FGF9 (green) and a rabbit monoclonal antibody to COX2 (red). Nuclei were counterstained with DAPI (blue). (C&D) Analysis of FGF9 co-expression with CD31 (PECAM-1) at the implantation site at day 4.5 of pregnancy. Triple immunofluorescence staining was performed with a mouse monoclonal antibody to FGF9 (green) and a rabbit monoclonal antibody to CD31 (red). Nuclei were counterstained with DAPI (blue). The arrows in panel A and C depict the embryo; dpc: days post coitum; (*) represents the region at a higher magnification. These results are representative of at least three independent samples. Scale bars represent 50 μ m (A, C) and 20 μ m (B, D).



Triple immunofluorescence staining was performed with a mouse monoclonal antibody to FGF9 (green) and a rabbit monoclonal antibody to CD31 (red). Nuclei were counterstained with DAPI (blue). The arrows in panel A and C depict the embryo; dpc: days post coitum; (*) represents the region at a higher magnification. These results are representative of at least three independent samples. Scale bars represent 50 μ m (A, C) and 20 μ m (B, D).

Fig. 6. Localization of FGF9 and its receptors, FGFR2 and FGFR3, at the implantation site at day 4.5 of pregnancy. (A) Triple immunofluorescence staining was performed using a mouse monoclonal antibody specific for FGF9 (green) and a rabbit polyclonal antibody for FGFR2 (red). Nuclei



were counterstained with DAPI (blue). (B) Triple immunofluorescence staining was performed with a mouse monoclonal antibody for FGF9 (green) and a rabbit polyclonal antibody for FGFR3 (red). Nuclei were counterstained with DAPI (blue). The arrows in panel A and B indicate the trophectoderm of the embryo; dpc: day post coitum. These results are representative of at least three independent samples. Scale bars = 20 μm.

next analyzed the spatial relationship between FGF9 and FGFR2/FGFR3, which are known to bind FGF9 with high affinity [34, 35], at the implantation site. Triple immunofluorescence staining using an antibody specific for FGF9 in combination with that recognizing FGFRs showed that FGFR2 localized to the cytoplasm (Fig. 6A), whereas FGFR3 was predominantly present in the nuclei (Fig. 6B) of stromal and epithelial cells. Strong immunofluorescence staining indicated the presence of FGF9 on the apical side of epithelial cells. In addition, intense staining for FGFR2, but not FGFR3, was present in the trophectoderm. Altogether, our results demonstrated synchronized spatial expression of FGF9 and its receptor, FGFR2. Therefore, FGF9-FGFR2 signaling could be important for communication between the uterine epithelium and trophoblast, which is necessary for embryo apposition and implantation.

Discussion

E-driven proliferation in the uterus is mediated by paracrine signaling involving FGFs [11, 15]. P suppresses E-dependent proliferation in the epithelium, which is mandatory to establish the microenvironment required for successful implantation [36]. Our data provide evidence that expression of *Fgf2* and *Fgf9* peak at two distinct time points during pregnancy: *Fgf9* at day 4.5 of pregnancy, when attachment and implantation commence, and *Fgf2* at day 7.5 of pregnancy, corresponding to decidua formation. Previously, whole mount in situ hybridization data showed that *Fgf2* expression appears in the stroma surrounding the implanting blastocyst and continues to increase with decidualization [13]. Our data presented here show for the first time that FGF9 expression is highly upregulated during implantation in the mouse uterus. Earlier, it was reported that FGF9 in human stromal cells is regulated by E [15].

Recent studies using chromatin immunoprecipitation followed by sequencing of enriched chromatin fragments showed that *Fgf9* has an ER-, but not a PR, response element [37, 38]. We assessed the effect of E and EP on FGF9 expression in ovariectomized mouse uteri, and our data showed the existence of three FGF9 isoforms detected as bands of 25, 29, and 30 kDa. Only the 25-kDa FGF9 species was strongly induced by E treatment. The existence of three protein species of FGF9, at 25, 29 and 30 kDa, was previously reported for FGF9 isolated from human glial cells [20]. Therefore, with more than 99% sequence identity between human and mouse FGF9 homologs, it is reasonable to assume that these heparin-binding proteins could be post-translationally processed in the same manner in the uteri of both species. It is known that in mice, at day 4 of pregnancy, a short surge of E is

mandatory for embryo attachment and implantation [1]. Surprisingly, the E-induced 25-kDa FGF9 isoform, detected in the uteri of ovariectomized mice, was absent from pregnant uteri during the peri-implantation period. The lack of this FGF9 protein species and the maximum expression of the other 29- and 30-kDa proteins at day 4.5 of pregnancy suggest that FGF9 isoforms have a selective role in endometrial functions.

Previously, it was shown that the COX2 product PGE2 induces FGF9, which promotes endometrial stromal cell proliferation [23]. Female mice with genetic ablation of *Cox2* exhibit pleiotropic defects including impaired implantation and decidualization [26, 39]. The spatial-localized expression of COX2 around the attachment site could be a consequence of embryonic signaling through protease-activated epithelial Na⁺ channel (ENaC), which results in Ca²⁺ influx and the induction of a COX2 and PGs surge [40, 41]. Our results suggest that at day 4.5 of pregnancy, PGs synthesized around the implantation site by COX2 might induce epithelial FGF9 expression. This upregulation of FGF9 signaling could be required to produce a highly specific, transient environment that is necessary for the functional window of implantation. As previously reported, FGF9 expression is regulated with E and PGE2 [14, 15]. PGE2 signaling is mediated by the EP3 receptor [23, 24]. In uterine stromal cells, the EP3 receptor is a product of the transcriptional activity of HOXA-10, and accordingly, the expression of EP3 is reduced in *HOXA-10*^{-/-} mice [42]. Recent reports revealed that in the endometrium of mice and humans, HOXA-10 is regulated by P [4]. Therefore, in the uterus, the activity of FGF9 could depend on a delicate balance between E and P activity.

One of the first signs of implantation is increased endometrial vascular permeability at the site of blastocyst attachment and penetration. This process is regulated with PGs [26]. In addition, it is known that FGFs have role in angiogenesis in the uterus [43]. Recent data showed that FGF9 delivery during angiogenesis produces durable, vasoresponsive microvessels wrapped by smooth muscle cells, and promotes new vascular maturation. In addition, the combination of FGF2 and FGF9 in Matrigel plugs implanted in mice resulted in a robust synergistic effect on microvessel formation [31]. Microvessels consist of a tubular endothelial-cell lining surrounded by supporting mural cells. Our data, based on triple immunofluorescence analysis of FGF9, CD31 (PECAM-1), and DAPI, showed that FGF9 is located in cells that are associated with microvessels; however, its expression does not overlap with CD31 in endothelial cells. Therefore, we speculate that FGF9-positive cells could be mural cells acting in a paracrine and autocrine fashion to stimulate microvessel maturation.

FGF9 binds with high affinity to FGFR2-IIIc and FGFR3-IIIc receptors [33]. Interestingly, FGFR3 was expressed in the nucleus of stromal cells and the LE surrounding the implantation site. Nuclear localization of FGFR3 has been reported for breast, bladder, and pancreatic cancer [44-46]. It has also been proposed that cellular mechanisms used by cells during implantation can be exploited by cancer cells to migrate through and invade tissues [36, 47]. Therefore, in addition to progressing the understanding of the intricate mechanism of embryo implantation, further study of FGFR3 could also contribute to improved understanding of cellular and molecular events leading to metastasis. A recent study showed that post-natal uterine specific ablation of FGFR2 resulted in stratified LE and pregnancy loss during peri-implantation. This demonstrates a critical role for FGFR2 in proper post-natal uterine development as well as normal uterine function [12]. Further, it was established that FGF9-FGFR2IIIc signaling in the uterine epithelium prevents epithelial stratification [12, 48]. The presence of FGFR2 was also previously detected in mouse blastocysts and was shown to play an important role in development during peri-implantation [49]. Furthermore, global deletion of *Fgfr2* results in peri-implantation death and trophoblast arrest, suggesting that the mutant dies before a visible decidual reaction. FGFR2-deficient blastocysts induce increased capillary permeability, which is an indication of the first step of implantation, but without appropriate stromal decidualization. Hence, it appears that in the absence of FGFR2, uterine trophoblast-induced decidualization is inadequate [50]. Implantation is an intricate process that requires a properly prepared endometrium and a functional embryo that is competent for implantation [41]. Our results, showing the expression of FGF9 in the

endometrium and the presence of FGFR2 in the trophoctoderm, indicate that an FGF9-FGFR2 axis could be vital for the recognition of blastocysts that are competent for implantation.

Conclusion

In summary, our data suggest that in mice, FGF9 is a crucial factor required for establishing an appropriate microenvironment for successful implantation and for pregnancy establishment. Additional analyses of pathways through which FGF9 participates in angiogenesis and endometrium-embryo communication should contribute to the understanding of molecular mechanisms that underlay process of implantation. Many pathways involved in endometrial function during implantation are conserved in mice and humans. Therefore, better knowledge of the molecular mechanisms involved in the FGF9 regulation of implantation could conceivably contribute to understanding as well as treatment of infertility.

Acknowledgements

This work was supported by a grant from the Croatian science foundation (HRZZ 3432) (BMJ) and a grant from the University of Rijeka, Croatia 13.06.1.1.08 (BMJ). The authors gratefully acknowledge Tatjana Daka and Tihana Vujnović for technical assistance and Ksenija Tulić for administrative help.

Disclosure Statement

The authors of this manuscript declare that they have no conflicts of interests.

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