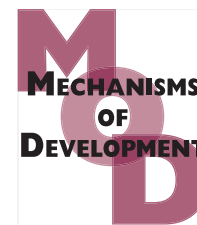


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Novel expression of EGFL7 in placental trophoblast and endothelial cells and its implication in preeclampsia

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ABSTRACT

The mammalian placenta is the site of nutrient and gas exchange between the mother and fetus, and is comprised of two principal cell types, trophoblasts and endothelial cells. Proper placental development requires invasion and differentiation of trophoblast cells, together with coordinated fetal vasculogenesis and maternal vascular remodeling. Disruption in these processes can result in placental pathologies such as preeclampsia (PE), a disease characterized by late gestational hypertension and proteinuria. Epidermal Growth Factor Like Domain 7 (EGFL7) is a largely endothelial-restricted secreted factor that is critical for embryonic vascular development, and functions by modulating the Notch signaling pathway. However, the role of EGFL7 in placental development remains unknown. In this study, we use mouse models and human placentas to begin to understand the role of EGFL7 during normal and pathological placentation. We show that *Egfl7* is expressed by the endothelium of both the maternal and fetal vasculature throughout placental development. Importantly, we uncovered a previously unknown site of EGFL7 expression in the trophoblast cell lineage, including the trophoctoderm, trophoblast stem cells, and placental trophoblasts. Our results demonstrate significantly reduced *Egfl7* expression in human PE placentas, concurrent with a downregulation of Notch target genes. Moreover, using the BPH/5 mouse model of PE, we show that the downregulation of *Egfl7* in compromised

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Abbreviations: EGFL7, Epidermal Growth Factor Like Domain 7PE, preeclampsiaTSC, trophoblast stem cellESC, embryonic stem cellMEF, mouse embryonic fibroblastCK, cytokeratinCVS, chorionic villus sample

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placentas occurs prior to the onset of characteristic maternal signs of PE. Together, our results implicate Egfl7 as a possible factor in normal placental development and in the etiology of PE.

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1. Introduction

The placenta serves as the site of contact for the maternal and embryonic circulatory systems to enable nutrient and gas exchange, and contains two primary functional cell types, trophoblasts and endothelial cells. Proper placental development requires invasion and differentiation of trophoblast cells, as well as coordinated maternal vascular remodeling and fetal vasculogenesis (Norwitz et al., 2001). Any disruption in these processes can result in placental pathologies, including preeclampsia (PE). PE is a leading cause of maternal and fetal morbidity and mortality worldwide, and the only resolute treatment is delivery of the baby and placenta. Although the pathophysiology of PE remains largely unknown, inadequate trophoblast cell invasion, endothelial cell dysfunction, dysregulated uteroplacental vascularization, and an imbalance of pro- and anti-angiogenic growth factors have been implicated in the disease (Young et al., 2010).

Recent studies have implicated Notch signaling, an evolutionarily conserved pathway that is important for cell fate specification, proliferation, and patterning (Phng and Gerhardt, 2009), in placental development and the pathogenesis of PE. Targeted mutations of Notch signaling components in the mouse result in placental defects, demonstrating their vital roles for proper placental development (Duarte et al., 2004; Fischer et al., 2004; Hunkapiller et al., 2011; Krebs et al., 2000). In addition, Notch pathway proteins are down-regulated in trophoblast cells, endothelial cells, and stromal cells of third trimester placentas from human preeclamptic patients as compared to normal patients (Cobellis et al., 2007).

Epidermal Growth Factor Like Domain 7 (EGFL7) is a secreted factor that is present in both soluble and extracellular matrix-bound forms (Fitch et al., 2004). Egfl7 expression in the developing embryo is largely restricted to endothelial progenitors and actively proliferating endothelium (Campagnolo et al., 2005; Fitch et al., 2004; Soncin et al., 2003). However, it is also found in embryonic stem cells, pre- and peri-implantation embryos, primordial germ cells, and some CNS neurons (Campagnolo et al., 2008; Fitch et al., 2004; Parker et al., 2004; Schmidt et al., 2009; Soncin et al., 2003). EGFL7 is largely down-regulated during late embryogenesis and in the quiescent endothelium of the adult mouse, and is upregulated during pathological and physiological angiogenesis, such as in the uterus (Campagnolo et al., 2005; Fitch et al., 2004; Soncin et al., 2003). Egfl7 is upregulated in response to, and has a protective effect against, hypoxia (Badiwala et al., 2010; Gustavsson et al., 2007; Xu et al., 2008). It acts as a chemoattractant for endothelial cells and plays an important role in their proliferation and migration (Campagnolo et al., 2005; Durrans and Stuhlmann, 2010; Schmidt et al., 2007). EGFL7 functions in vascular patterning, stratification, and tubulogenesis

in vitro and in the mouse and zebrafish (Campagnolo et al., 2005; Durrans and Stuhlmann, 2010; Nichol et al., 2010; Parker et al., 2004). EGFL7 has been shown to modulate the Notch signaling cascade by acting either as a Notch agonist, such as in the developing embryo, or as a Notch antagonist, such as in the postnatal retina and neural stem cells (Nichol et al., 2010; Schmidt et al., 2009).

Despite its key role in early embryogenesis, vascular development, and modulation of Notch signaling, the expression pattern and function of EGFL7 in normal and PE placentas is poorly understood. In this study, we investigated the expression pattern of EGFL7 in normal murine and human placentas. Rodents and primates both undergo hemochorial placentation (Cross et al., 2003). Despite some structural differences, the trophoblast cell types and the molecular pathways driving placental development are highly conserved between mouse and human (Cross et al., 2003; Georgiades et al., 2002; Hu and Cross, 2010; Rossant and Cross, 2001). Importantly, the labyrinth in the mouse placenta is analogous to the chorionic villi in human placentas, whereas the junctional zone in mice is analogous to the cytotrophoblast cell columns (Rossant and Cross, 2001) or the basal plate in humans (Georgiades et al., 2002).

In addition to examining the expression profile of Egfl7 during normal placental development, this study investigates a potential role for EGFL7 in preeclampsia by analyzing human PE placentas and compromised placentas from the BPH/5 murine PE model. The BPH/5 mouse strain exhibits the characteristic PE signs of late-gestational hypertension, proteinuria, and endothelial dysfunction (Davisson et al., 2002; Dokras et al., 2006). BPH/5 mice also show fetoplacental defects such as impaired endothelial cell branching, maternal spiral artery remodeling, and reduced fetal labyrinth depth (Dokras et al., 2006). Here we have described the spatiotemporal expression profile of Egfl7 in placental endothelial cells in the mouse and human. We uncovered a previously unknown site of EGFL7 localization in the non-endothelial trophoblast lineage, beginning at the blastocyst stage and becoming restricted to a subset of differentiated trophoblast cells. Furthermore, we provide evidence that a downregulation of EGFL7 is associated with human PE and the BPH/5 mouse model of PE, and this downregulation is concomitant with a decrease in Notch target gene expression.

2. Results

2.1. EGFL7 is expressed by maternal and fetal endothelial cells in the mouse placenta throughout gestation

To obtain a temporal and spatial expression profile for Egfl7 throughout pregnancy, we used the mouse as a model

system. To localize expression of *Egfl7* transcripts in the maternal and fetal placental vasculature, RNA in situ hybridization (ISH) was performed on 100 μm thick vibratome sections of C57BL/6 mouse placentas using an *Egfl7* specific riboprobe. *Egfl7* transcripts were highly expressed at E10.5, and were localized to maternal vessels, including the spiral arteries of the decidua (Fig. 1A and B), and the branching vascular structures in the fetal labyrinth (Fig. 1A and C). Following ISH, specimens were embedded in paraffin, sectioned, and counterstained to analyze the cellular morphology of *Egfl7* mRNA expression in the placenta. Results revealed *Egfl7* transcript expression in cells lining the large lumen of maternal vessels in the decidua (Fig. 1D) and in the narrow and highly branched capillaries of the fetal labyrinth (Fig. 1E). *Egfl7* transcript levels were dramatically reduced at E12.5 and nearly undetectable at E18.5 (Supplemental Fig. 1). *Egfl7* sense

controls (insets of Fig. 1A, Supplemental Fig. 1A and D) demonstrate specificity of the *Egfl7* riboprobe.

To determine the spatiotemporal expression profile of EGFL7 protein, we performed double immunofluorescent staining for EGFL7 and the pan-endothelial marker, CD31, on C57BL/6 mouse placentas (Fig. 1F–M, Supplemental Fig. 2). EGFL7 colocalized with CD31 in the maternal decidua and the fetal labyrinth at every stage examined, confirming that EGFL7 localized to maternal and fetal endothelial cells in the placenta.

Together, the results demonstrate that *Egfl7* transcript and protein expression was dynamic throughout gestation. *Egfl7* expression is high during early stages of vascular development in the placenta, with transcript expression peaking at E10.5 (Fig. 1A–E), as well as high protein expression at E10.5 and E12.5 (Fig. 1F–M, Supplemental Fig. 2A). At E18.5, a time point with fewer newly forming nascent vessels, *Egfl7*

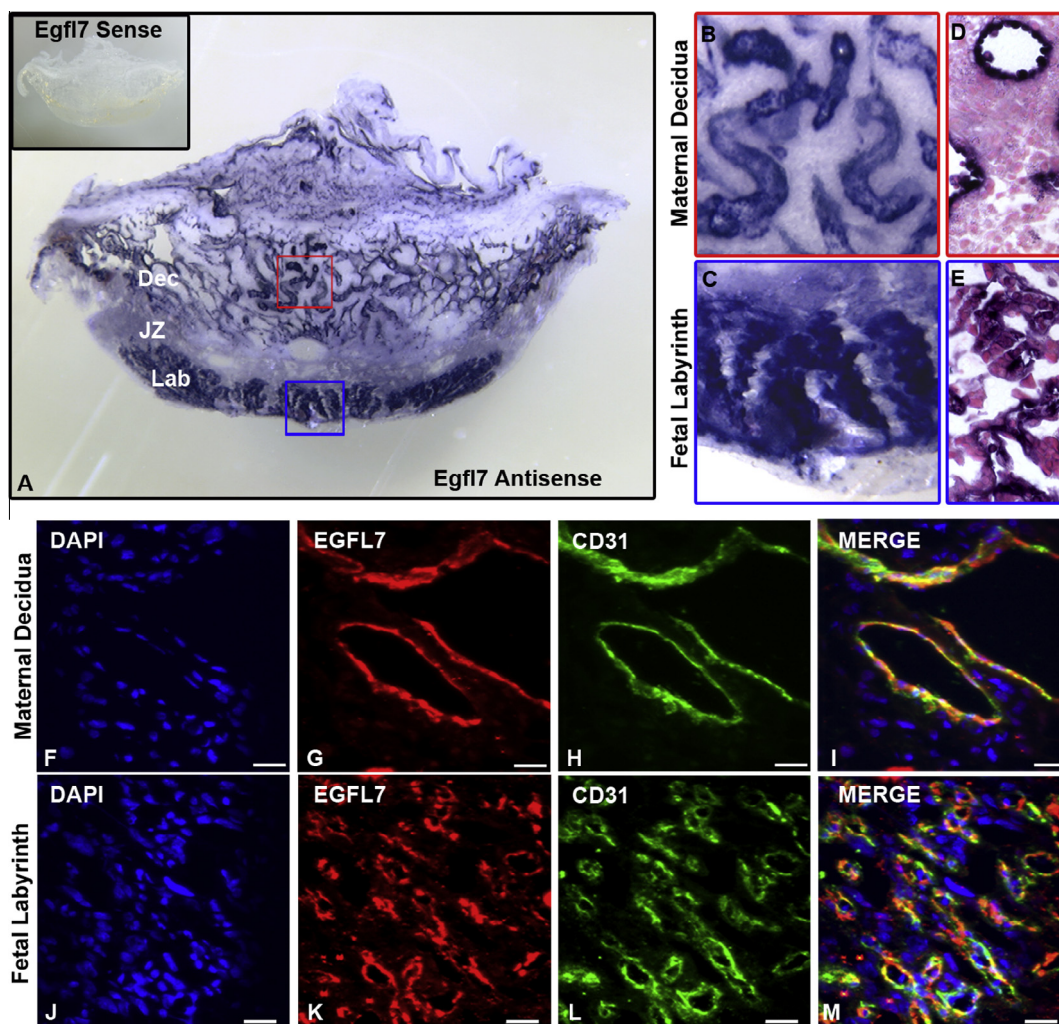


Fig. 1 – EGFL7 is expressed by maternal and fetal endothelial cells in the mouse placenta. In situ hybridization was performed to determine *Egfl7* mRNA localization, using an *Egfl7* riboprobe on 100 μm thick vibratome sections of E10.5 C57BL/6 placentas (A–E). Higher magnification images of boxes in (A) demonstrating *Egfl7* transcript is highly expressed in the maternal decidua (B) and fetal labyrinth (C). *Egfl7* sense controls (A inset) show specificity of *Egfl7* riboprobe. Paraffin sections of the vibratome sections showing cellular morphology after *Egfl7* riboprobe staining in the maternal decidua (D) and fetal labyrinth (E). To determine the cell types expressing EGFL7 protein in the placenta, double immunofluorescent staining was performed on E12.5 C57BL/6 placentas for EGFL7 (red), CD31 (green) and nuclear DAPI (blue) (F–M). EGFL7 colocalizes with the endothelial cell marker, CD31, in the maternal decidua (F–I) and the fetal labyrinth (J–M). Dec-Maternal Decidua, JZ-Junctional Zone, Lab-Fetal Labyrinth. Scale bar (F–M) = 20 μm .

transcript expression is dramatically lower (Supplemental Fig. 1D), while the protein expression persists (Supplemental Fig. 2B). Our data are consistent with previously reported Egfl7 expression data during vascular development and physiological angiogenesis, where the highest expression levels of Egfl7 are detected in actively proliferating endothelial cells, such as during formation of the initial vascular plexus during early embryogenesis (Campagnolo et al., 2005, 2008; Durrans and Stuhlmann, 2010; Fitch et al., 2004; Nichol et al., 2010; Parker et al., 2004; Schmidt et al., 2007).

2.2. Novel EGFL7 expression in placental trophoblast cells and the trophoblast cell lineage

In addition to demonstrating dynamic expression of EGFL7 in the placental endothelium, our *in situ* hybridization images at high magnification also uncovered a novel expression domain for Egfl7 in non-endothelial cells of the junctional zone of the mouse placenta, albeit at reduced levels (Fig. 2A and B). Images of Egfl7 ISH paraffin sections revealed clusters of EGFL7 positive cells located in the junctional zone, between the fetal labyrinth and trophoblast giant cells (Fig. 2B). Double immunofluorescent staining for EGFL7 and the pan-trophoblast marker CYTOKERATIN (CK), performed on mid-gestation mouse placentas confirmed EGFL7 localization to the trophoblast lineage (Fig. 2C–F). This is the first evidence that EGFL7 localizes to trophoblast cells of the placenta.

The newly uncovered expression of EGFL7 in placental trophoblast cells prompted us to examine its temporal expression pattern during trophoblast lineage development in mice. Differentiated trophoblast cells are derived from the outer cell layer of blastocysts, the trophoctoderm (Cross, 2005). We have previously shown that Egfl7 mRNA was expressed by blastocysts and embryonic stem cells, however its spatial expression in pre-implantation embryos was not defined (Campagnolo et al., 2005, 2008). Here, we isolated embryos from C57BL/6 mice at the blastocyst stage of development and performed whole-mount immunofluorescent staining for EGFL7 and the trophoctoderm marker, CDX2. Both inner cell mass cells and trophoctoderm cells expressed EGFL7 (Fig. 2G–J). Therefore, Egfl7 is expressed at the first step of trophoblast lineage development.

To further establish EGFL7 expression in the trophoblast lineage, we assayed trophoblast stem cells (TSC), a model of *in vitro* trophoblast cell differentiation. Multipotent TSC are derived from the trophoctoderm of blastocysts, and upon removal of exogenous Fibroblast Growth Factor, they recapitulate the defining characteristics of *in vivo* trophoblast cell differentiation (Tanaka et al., 1998). Semi-quantitative and Real Time RT-PCR analysis showed that TSC express Egfl7 mRNA at significantly higher levels than embryonic stem cells (ESC), whereas primary mouse embryonic fibroblasts (MEF) express very low levels of Egfl7 (Fig. 2K, $P < 0.01$). RNA *in situ* hybridization on TSC revealed specific localization of Egfl7 mRNA in TSC (Fig. 2L–O). EGFL7 protein was detected on cultured TSC, as shown by colocalization of EGFL7 and CDX2 using double immunofluorescent staining (Fig. 2P–T). Two additional antibodies that are directed against different regions of the EGFL7 protein gave comparable results (data not shown). Together, our results show that EGFL7 is

expressed by the trophoctoderm of blastocysts, in trophoblast stem cells, and a subset of differentiated trophoblast cells in the mid-gestation mouse placenta.

2.3. EGFL7 is expressed by fetal endothelial cells and trophoblast cells of human chorionic villi

To determine if EGFL7 expression in endothelial and trophoblast cells is conserved in human placentas, chorionic villi samples were obtained from normal pregnant women with an average pregnancy duration of 39 weeks. Double immunofluorescent staining analysis revealed that EGFL7 was expressed on fetal vessels, where it colocalized with the pan-endothelial marker, CD31 (Fig. 3A–D). Importantly, EGFL7 was also expressed on the syncytiotrophoblast layer of the villi, where it colocalized with the pan-trophoblast marker CYTOKERATIN (Fig. 3E–H). To confirm this observation, EGFL7 staining was performed using two antibodies directed against different regions of the EGFL7 protein, and both gave comparable results (Supplemental Fig. 3). Of note, expression of Egfl7 was not detected in the analogous mouse syncytiotrophoblasts. To further determine if EGFL7 localized to trophoblast cells of the human placenta, we isolated cytotrophoblast cells from term placentas as previously described (Hunkapiller and Fisher, 2008). Double immunofluorescent staining showed that EGFL7 and CYTOKERATIN colocalized to the isolated cytotrophoblast cell (Fig. 3I–L). Thus, our data demonstrate that EGFL7 is expressed in endothelial cells and trophoblast cells in both mouse and human placentas.

2.4. EGFL7 is downregulated in placentas of the BPH/5 mouse model of PE, prior to the onset of maternal signs of PE

To examine potential variations in the expression and spatial distribution of EGFL7 in PE, we first utilized the BPH/5 mouse model of preeclampsia. E10.5 placentas from control C57BL/6 and compromised BPH/5 fetoplacental units (see Materials and Methods) were subjected to immunofluorescent staining for EGFL7 and CD31 (Fig. 4B–I). Importantly, this time point is prior to the onset of the late-gestational signs of PE in BPH/5 mice (Davisson et al., 2002). EGFL7 protein expression was reduced in the fetal labyrinth zone of the BPH/5 placentas compared to C57BL/6 (Fig. 4C and G). Total levels of CD31 appear unchanged (Fig. 4D, E, H and I), suggesting no overall reduction in placental vascular density. The pattern of the vessels, however, appears irregular in BPH/5 placentas. In fact, a previous study reported the anatomical appearance of attenuated and irregularly branched fetal vessels in isolectin B4-stained BPH/5 placentas (Dokras et al., 2006). To provide quantitative evidence, we performed Real Time RT-PCR analysis of E10.5 placentas from C57BL/6 and affected BPH/5 fetoplacental units for Egfl7. Results demonstrated that Egfl7 mRNA levels were significantly downregulated in BPH/5 placentas compared to C57BL/6 (Fig. 4A, $P < 0.05$).

2.5. EGFL7 expression is significantly reduced in human preeclamptic placentas

Our results in the mouse model of PE prompted us to investigate potential variations in the expression of EGFL7

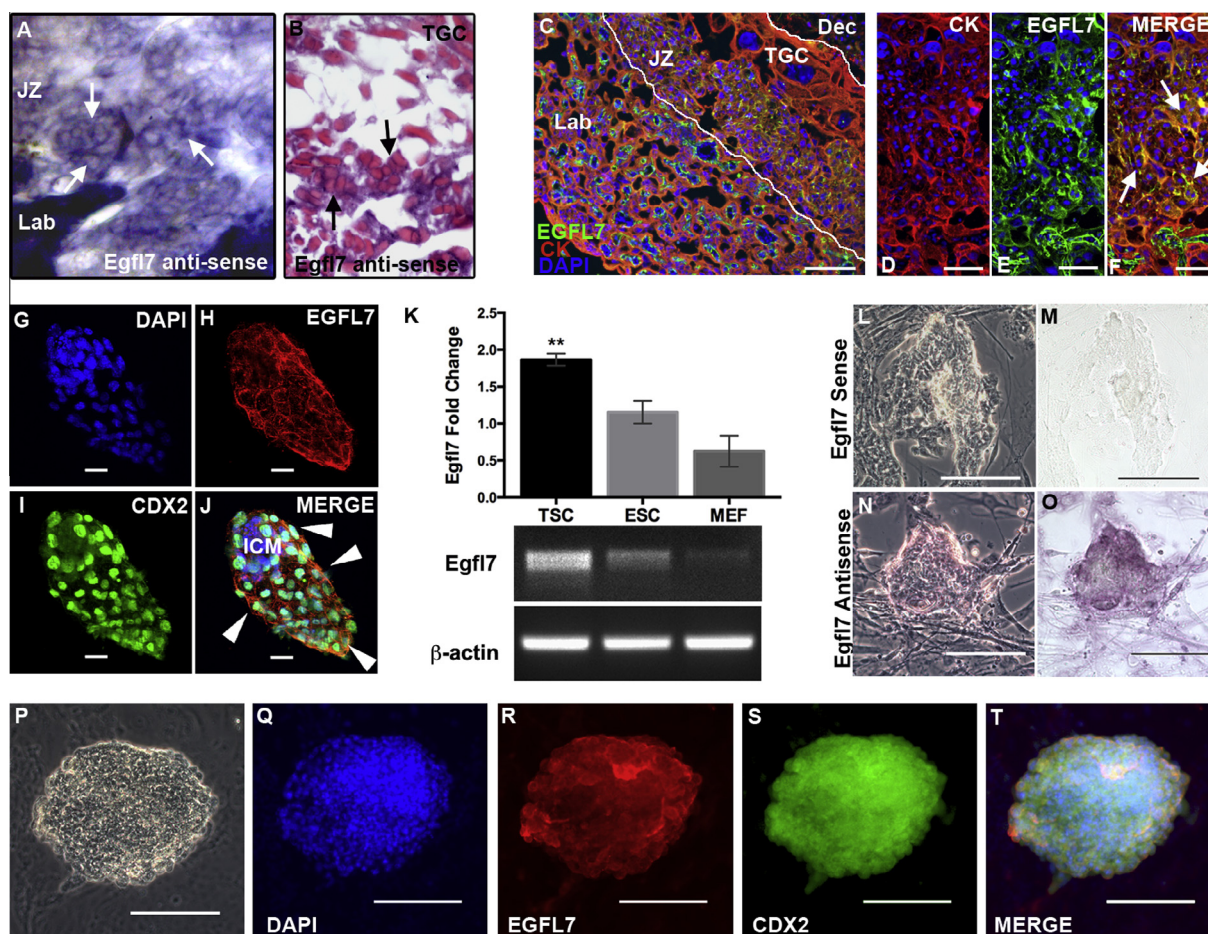


Fig. 2 – EGFL7 is expressed in the trophoblast cell lineage of the mouse. In situ hybridization was performed to determine Egfl7 mRNA localization, using an Egfl7 riboprobe on 100 μ m thick vibratome sections of E10.5 C57BL/6 placentas (A). Shown is the junctional zone of the placenta (A). Paraffin sections of the vibratome section showing cellular morphology of Egfl7 riboprobe staining in the junctional zone of the placenta (B). Arrows indicate positive Egfl7 transcript signal in the junctional zone trophoblast cells. (TGC = trophoblast giant cell). To determine the cell type expressing EGFL7 protein in the placenta, double immunofluorescent staining was performed on E10.5 C57BL/6 placentas (C–F) for the pan-trophoblast marker, CYTOKERATIN (CK, red), EGFL7 (green), and nuclear DAPI (blue). Cross-section of E10.5 placenta showing the fetal labyrinth (Lab), junctional zone (JZ, demarked by white lines) and maternal decidua (Dec). High magnification images (D–F) of the junctional zone demonstrating colocalization of EGFL7 with CK in the JZ trophoblast cells (arrows). Whole mount immunofluorescent staining of blastocyst-stage C57BL/6 embryos (G–J) for DAPI (blue), EGFL7 (red), and trophectoderm cell marker, CDX2 (green) reveals EGFL7 protein localization to both inner cell mass (ICM) cells and trophectoderm cells (arrowheads). Real Time RT-PCR data for Egfl7 transcript expression in trophoblast stem cells (TSC), embryonic stem cells (ESC), and mouse embryonic fibroblasts (MEF), shows a significantly higher level of Egfl7 expression in TSC, $P < 0.01$ (K, top). Agarose gel with products of semi-quantitative RT-PCR for TSC, ESC and MEFs are shown, using Egfl7 and β -actin specific primers (K, bottom). RNA in situ hybridization analysis of Egfl7 mRNA in TSC (L–O) defines Egfl7 transcript expression in TSC. Phase-contrast and bright-field images of TSC incubated with an Egfl7 sense control probe (L and M), and an Egfl7 antisense probe (N and O). Double immunofluorescent staining of TSC (P–T) for DAPI, EGFL7, and CDX2 demonstrating EGFL7 protein localization to TSC. Phase-contrast image of TSC colony is shown in (P). Scale bars in D–F = 25 μ m, G–J = 20 μ m, C and L–T = 100 μ m.

in human PE placentas. Biopsies from normal and PE human placentas were obtained and analyzed. Clinical characteristics of patients in this study are shown in Table 1 (Nelson and Burton, 2010). The study group consisted of ten early-onset preeclamptic patients with average pregnancy duration of 32 weeks. The control group consisted of ten healthy pregnant women with average pregnancy duration of 39 weeks. All patients underwent Cesarean sections. There were no

differences in patient age at delivery, gravidity, or parity between the study and control groups. However, the study group exhibited lower neonatal birth weight and placental weight, higher systolic and diastolic pressure, and earlier gestational age, all of which are characteristic of PE patients.

To examine the EGFL7 protein expression profile of normal and PE placentas, we performed immunofluorescent staining for EGFL7 and CD31 on human placental biopsies (Fig. 5A–F).

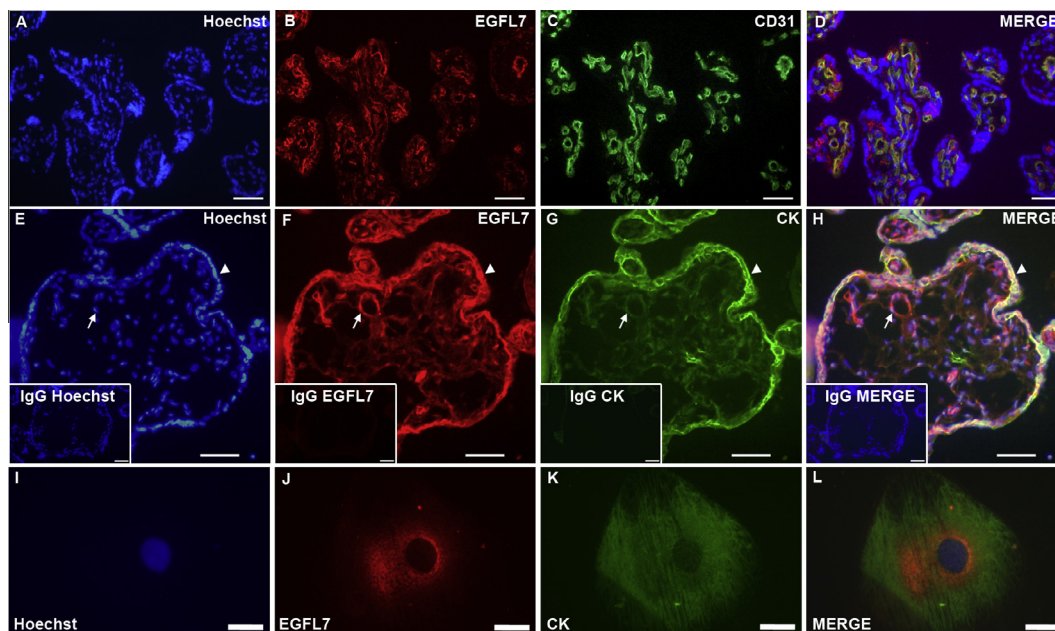


Fig. 3 – EGFL7 is expressed in endothelial cells and trophoblast cells of the human placenta. Double immunofluorescent staining of normal human placentas (A–D) for EGFL7 (red), CD31 (green), and nuclear Hoechst (blue) showing localization of EGFL7 protein on endothelial cells. Double immunofluorescent staining of human chorionic villi from placentas at 40-weeks of gestation (E–H) for nuclear Hoechst (blue), EGFL7 (red), and pan-trophoblast marker CYTOKERATIN (CK) (green) demonstrates EGFL7 protein localization to trophoblast cells (arrows-fetal vessels, arrowheads-trophoblasts). Control staining of a close-by section (insets, E–H) using IgG and secondary antibodies only and nuclear Hoechst (blue) shows specificity of the antibodies. Expression of EGFL7 protein is found in human cytotrophoblast cells isolated from term placentas (I–L). Cells were stained for Hoechst (blue), EGFL7 (red) and CYTOKERATIN (green) (I–L). Scale bars in E–H and insets = 100 μm , and I–L = 30 μm .

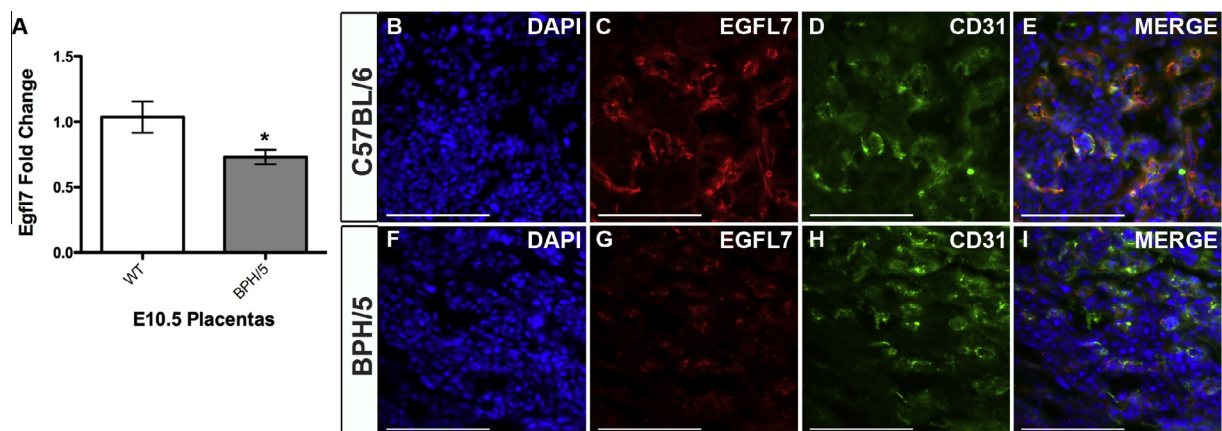


Fig. 4 – EGFL7 is downregulated in the placentas of the BPH/5 mouse model of preeclampsia. Real Time RT-PCR data for *Egfl7* transcript levels in E10.5 placentas from C57BL/6 (WT) and BPH/5 mice reveal a significant decrease in *Egfl7* in BPH/5 mice $^*P < 0.05$ (A). Immunofluorescent staining for EGFL7 protein (red), CD31 (green), and nuclear DAPI (blue) on the fetal labyrinth of E10.5 placentas from C57BL/6 (B–E) and BPH/5 mice (F–I) demonstrating a decrease in EGFL7 in BPH/5 placentas. Scale bar = 100 μm .

EGFL7 was immunodetected on the endothelium and trophoblast cells of the human placenta. When comparing immunofluorescent staining of normal and PE placentas, EGFL7 appeared lower in both endothelial and trophoblast cells of PE placentas (Fig. 5A and B), whereas no major differences were observed for CD31 (Fig. 5C and D).

To quantify this observation, we performed Real-Time RT-PCR analysis on normal and PE samples for EGFL7 (Fig. 5G) and CD31 (Fig. 5H). EGFL7 mRNA expression was significantly decreased by more than 10-fold in PE when compared to normal placentas ($P < 0.05$). In contrast, no significant difference in the level of CD31 mRNA was detected, suggesting the

Table 1 – Clinical characteristics of pregnancies for placentas studied. Data are presented as median \pm standard deviation and analyzed using a student's t-test.

Parameter	Clinical characteristics of pregnancies for placentas studied		
	Preeclampsia (n = 10)	Control (n = 10)	Significance
Parity	0 \pm 0.3	0	
Gestational age (weeks)	32 \pm 2	39 \pm 0	<0.0001
Maternal age (years)	34 \pm 4	35 \pm 6	0.54
Gravidity	2 \pm 1	2 \pm 1	0.59
Birth weight (g)	1235 \pm 359	3372 \pm 354	<0.001
Birth weight percentile	9 \pm 3	63 \pm 18	<0.001
Placental weight (g)	236 \pm 65	550 \pm 55	<0.001
Systolic blood pressure at delivery (mm Hg)	161 \pm 11	130 \pm 5	<0.001
Diastolic blood pressure at delivery (mm Hg)	109 \pm 8	82 \pm 3	<0.001

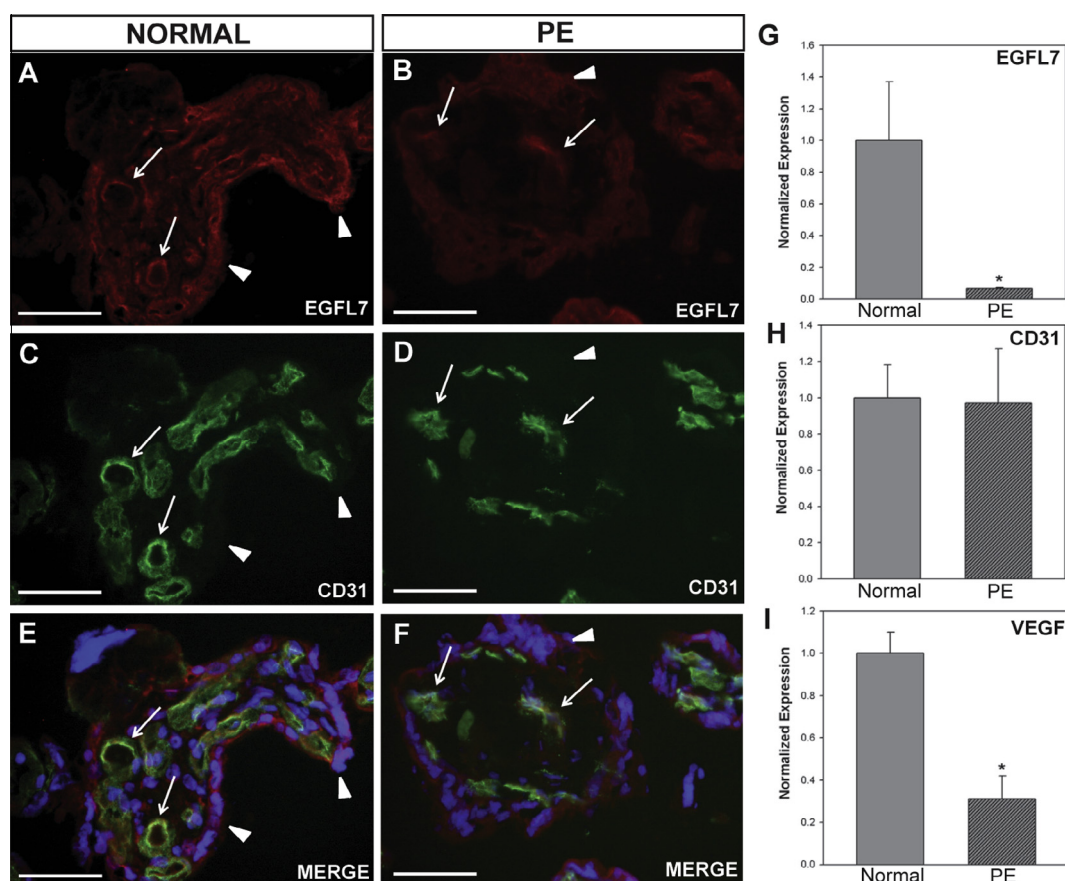


Fig. 5 – EGFL7 is significantly reduced in human preeclamptic placentas. Double immunofluorescent staining of normal and preeclamptic (PE) human placentas (A–F) for EGFL7 protein (red), CD31 (green), and nuclear Hoechst (blue) revealing a decrease in EGFL7 protein in PE placentas (arrows–fetal vessels, arrowheads–trophoblasts). Real Time RT-PCR data for EGFL7 (G), CD31 (H), and VEGF (I) on normal and preeclamptic (PE) human placentas demonstrating a significant decrease in EGFL7 transcript levels in PE placentas, even further than the known angiogenic factor, VEGF (n = 10, *P < 0.05). Scale bars = 50 μ m.

difference in EGFL7 transcript levels was not due to an overall reduced expression of endothelial-specific genes or a reduction in vascular density (Costa et al., 2013; Nadra et al., 2010; Nichol et al., 2010; van Tuyl et al., 2005). In our study, the reduction in EGFL7 levels in PE was more pronounced as compared to VEGF levels (Fig. 5I), a growth factor whose

implication in PE has been extensively studied by others, and found to be either decreased (Cooper et al., 1996; Lyall et al., 1997; Park et al., 2010) or increased (Akeran et al., 2008; Chung et al., 2004; Geva et al., 2002) in PE placentas. Thus, EGFL7 expression is significantly downregulated in both a mouse model of PE and human PE placentas.

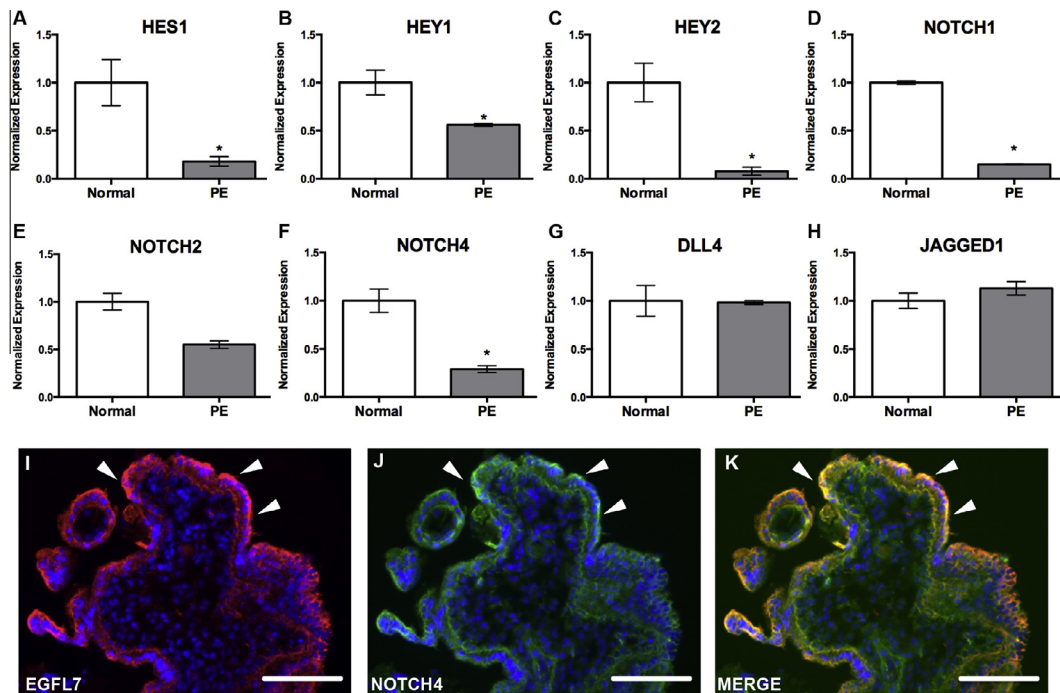


Fig. 6 – Expression of NOTCH target genes and NOTCH receptors are downregulated in PE, concomitant with EGFL7. Real Time RT-PCR data for NOTCH pathway members on normal and preeclamptic (PE) human placentas (A–H). Gene expression analysis for HES1 (A), HEY1 (B), and HEY2 (C) demonstrating a significant decrease in all NOTCH target gene transcripts in PE. Gene expression analysis for NOTCH receptors NOTCH1 (D), NOTCH2 (E), and NOTCH4 (F) demonstrating a significant downregulation in NOTCH1 and NOTCH4, and a downward trend in NOTCH2 in PE patients. Gene expression analysis indicating no change in NOTCH ligands DLL4 (G) and JAGGED1 (H). ($n = 3-7$, $P < 0.05$). Immunofluorescent staining of a cross section of human villi (I–K) for EGFL7 (red), NOTCH4 (green), and Hoechst (blue) (arrowheads-trophoblasts) shows colocalization of EGFL7 protein and NOTCH4. ($P < 0.05$) Scale bars = 50 μm .

2.6. EGFL7 expression correlates with Notch signaling in normal and PE placentas

Previous studies demonstrated that EGFL7 interacts with and modulates Notch signaling *in vitro* and *in vivo* (Nichol et al., 2010; Schmidt et al., 2009). To examine if EGFL7 and the Notch signaling pathway are concomitantly dysregulated in PE, we performed Real-Time RT-PCR analysis of normal and human PE placentas for NOTCH pathway members and NOTCH target genes. Our results demonstrated a significant decrease in HES1, HEY1, HEY2, NOTCH1, and NOTCH4 mRNA expression ($P < 0.05$), and a downward trend in NOTCH2 expression in PE samples when compared to normal placentas (Fig. 6A–F). In contrast, transcript levels for the Notch ligands DLL4 and JAGGED1 were unchanged (Fig. 6G and H). Additionally, double immunofluorescent staining of human placentas revealed that EGFL7 colocalized with NOTCH4 on the syncytiotrophoblast cells (Fig. 6I–K). Thus, downregulation of Notch signaling members and NOTCH target gene expression correlate with a downregulation of EGFL7 in PE placentas, suggesting that EGFL7 may possibly function through the Notch signaling pathway.

3. Discussion

PE is a complex placental disease that occurs in approximately 2–7.5% of pregnancies worldwide (Dolea and

AbouZahr, 2003; Wallis et al., 2008), yet the etiology of the disease remains unknown and early predictive biomarkers are lacking. Research on the human placenta has mostly been limited to studying the organ at late gestation and full term. To fully understand the early causes and progression of this placental disease, it is crucial to employ appropriate animal and *in vitro* models. Our results show that expression of EGFL7 is significantly reduced in compromised placentas from the BPH/5 mouse model of PE compared to control mice. These findings are corroborated by our results from human PE samples. Importantly, reduced EGFL7 expression in PE placentas did not correlate with reduced expression of the pan-endothelial marker CD31 or reduced density of CD31-positive vessels. This suggests that the change in EGFL7 was not a consequence of reduced vascular density. However, as a caveat, CD31 transcript expression is an indirect indicator for endothelial cell content and therefore may vary from the total vascular density in the placenta.

Of interest to this study, EGFL7 was identified in two recent microarray studies as one of the genes significantly reduced in human PE placentas compared to normal placentas, consistent with our data (Johansson et al., 2011; Junus et al., 2012). Importantly, PE is thought to begin much earlier in gestation than when the late-stage human placental biopsies used in most studies are obtained (Merviel et al., 2004; Waite et al., 2002). Our results describe changes in EGFL7 expression in phenotypically abnormal placentas of a PE

mouse model, prior to the onset of characteristic maternal signs of PE, suggesting EGFL7 may be an early biomarker for the disease. It would be of interest to determine whether EGFL7 is altered in phenotypically abnormal placentas of other placental pathologies.

Moreover, we have shown that EGFL7 is downregulated in human placentas of early-onset PE patients. Formally, we cannot rule out the possibility that the difference in EGFL7 expression is due to a difference in gestational age (PE: week-32 versus Normal: average of 39-week gestation). However, analysis of a single placenta at week-34 of gestation from a normal pregnancy indicated that EGFL7 expression levels were similar to those in normal term placentas (data not shown). Furthermore, both early onset PE and intrauterine growth restriction (IUGR) are associated with abnormal placentation and changes in plasma levels of placental angiogenic factors, but IUGR does not present with the maternal signs of hypertension and proteinuria. A recent study demonstrated that isolated IUGR with no maternal disease is associated with the same changes in placental angiogenic factors and subclinical endothelial dysfunction as in PE (Crispi et al., 2006), suggesting the only difference between these two diseases may be maternal manifestations of the disease. Early-onset PE, which manifests by maternal hypertension and proteinuria, and progresses to a systemic hypoperfusion of multiple maternal organs, is often accompanied by abnormal fetal growth (Ness and Sibai, 2006), as in the cases of our study (9/10; 90%). Therefore, reduced EGFL7 expression correlates with abnormal placentation that may be associated with either early-onset PE or isolated IUGR. Contribution of additional facilitating factors, possibly at the level of maternal predisposition, may influence the manifestation of maternal hypertension and proteinuria.

Previously, EGFL7 expression was thought to be restricted to actively proliferating vascular endothelium and to embryonic stem cells (Campagnolo et al., 2008; Fitch et al., 2004; Schmidt et al., 2009). Our study has used multiple methods to uncover a novel and dynamic expression pattern for Eglf7 during murine placental development. Eglf7 is expressed highly in the fetal and maternal vasculature of the placenta. Its highly regulated expression pattern that correlates with expression during embryonic development suggests a potential angiogenic role in placental angiogenesis. Notably, in addition to its expression in the fetal and maternal vessels of the placenta, we showed, for the first time, that Eglf7 was also expressed in the trophoblast cell lineage, beginning at the blastocyst stage and becoming restricted to a subset of differentiated trophoblast cells in the mature placenta. Trophoblast cells are the principal non-endothelial cell populations at the fetomaternal interface. Targeted mutations in genes important for trophoblast cell differentiation result in placental defects or complete loss of placenta formation (Cross, 2005). Interestingly, knockdown of Eglf7 in ESC decreases the proliferation rate of undifferentiated ESC and impairs endothelial cord formation in an embryoid body model of ESC differentiation (Durrans and Stuhlmann, 2010). Determining the role of Eglf7 in TSC and trophoblasts will be crucial for understanding the molecular mechanisms that control trophoblast lineage development and formation of the placenta.

EGFL7 has previously been shown to colocalize and functionally interact with NOTCH to modulate Notch signaling (Nichol et al., 2010; Schmidt et al., 2009). Notch signaling is a crucial pathway for placental development that affects both trophoblast and vascular cell function. For example, mouse strains with targeted deletions in the Notch receptors *Notch1/4*, their ligand *Dll4*, Notch target genes *Hey1/2*, or the Notch nuclear co-activator *RBPJK* show defects in chorioallantoic branching and fetal placental angiogenesis (Duarte et al., 2004; Fischer et al., 2004; Gale et al., 2004; Gasperowicz and Otto, 2008; Krebs et al., 2004, 2000; Limbourg et al., 2005; Oka et al., 1995). *Mash2*, a basic helix-loop-helix transcription factor that is thought to be repressed by the Notch target *Hes1* (Gasperowicz and Otto, 2008), is critical for trophoblast cell fate specification in the mouse (Guillemot et al., 1994). *Notch2* receptor function is crucial for formation of maternal blood sinuses and maternal spiral artery and arterial canal remodeling (Hamada et al., 2007; Hunkapiller et al., 2011). A recent study demonstrated that the *Notch2* receptor is prominently expressed in the junctional zone trophoblast cells at E10.5 (Gasperowicz et al., 2013), as is EGFL7. It would be of interest to determine if EGFL7 functionally interacts with NOTCH2 in the placental junctional zone. Here we showed that NOTCH4 colocalized with EGFL7 in the placenta, and that downregulation of EGFL7 coincided with a reduction in Notch target gene expression in human PE placentas. Together, our results suggest EGFL7 may act, at least in part, as a NOTCH agonist in the placenta. Interestingly, whereas Eglf7 was found to act as an antagonist of Notch postnatally and in HUVECs (Nichol et al., 2010; Schmidt et al., 2009), embryonic overexpression of Eglf7 significantly increased *Hey1* and *Hey2* transcript levels, consistent with acting as a Notch agonist during embryogenesis (Nichol et al., 2010). It will be important to further explore the functional relationship between EGFL7 and the Notch signaling pathway during placental development, in addition to other pathways through which EGFL7 may be functioning.

EGFL7 gain- and loss-of-function approaches will help to fully understand the role of EGFL7 in the placenta and in placental pathologies, and to dissect the underlying mechanism in both trophoblast cells and endothelial cells. It will be important to examine if endothelial and/or trophoblast-derived EGFL7 acts as an angiogenic factor in the placenta and if secreted EGFL7 acts in an autocrine or paracrine fashion, as has been suggested (Nichol and Stuhlmann, 2012). Of interest, partial embryonic lethality at mid-gestation was reported in the EGFL7 mutant mice that either overexpress EGFL7 or are EGFL7 deficient (Nichol et al., 2010; Schmidt et al., 2007). It is plausible that the partial lethality is, at least in part, due to a dysfunction in placental development.

In conclusion, our study has uncovered a novel expression domain for EGFL7 in non-endothelial trophoblast cell types. Its novel and dynamic expression pattern suggests a potential role in the molecular mechanisms that regulate trophoblast cell proliferation, differentiation, and invasion during placental development. Our study also links a significant downregulation of EGFL7 expression to PE placentas. We demonstrate that, in a PE mouse model, the change occurs in phenotypically abnormal placentas before the onset of the characteristic maternal signs of preeclampsia. It is tempting to speculate

that EGFL7 could play a functional role in the pathophysiology of PE, and that EGFL7 could be an early biomarker of PE.

4. Materials and methods

4.1. Mice and placental sampling

All animal protocols were approved by the Institutional Animal Care and Use Committee at Weill Cornell Medical College and Cornell University. BPH/5 mice were from in-house colonies. C57BL/6 mice were obtained from Jackson Laboratories and served as a control for the BPH/5 strain (Davisson et al., 2002). For timed pregnancies, the day of visualization of a vaginal plug was designated embryonic day 0.5 (E0.5). Fetoplacental units of BPH/5 pregnancies are of varying status as assessed by ultrasonography *in utero*, ranging from normal to compromised to resorbed (Davisson et al., 2002). A subset of placentas were dissected only from E10.5 BPH/5 fetoplacental units that were compromised, i.e., with reduced fetal size, heart rate, and blood flow as determined by ultrasound as described (Davisson et al., 2002; Dokras et al., 2006). Placentas from normal or resorbed BPH/5 fetoplacental units were not sampled.

4.2. Human placenta sampling

Biopsies of placentas from normal ($n = 10$) and early-onset PE pregnancies ($n = 10$) were obtained from the Catholic University of Rome. The study respected the principles expressed in the Declaration of Helsinki and was approved by the Bioethical Committee of the Catholic University of Rome. For the collection of samples, all patients provided written informed consent. Clinical criteria used to define preeclampsia in our study were in accordance with the definition of the International Society for the Study of Hypertension in Pregnancy (ISSHP). In short, a patient (previously normotensive) was defined as being affected by PE if she had at least two diastolic blood pressure measurements of 90 mm Hg in 24 h (≥ 4 h apart), accompanied by proteinuria of at least 300 mg in a 24-h collection of urine. Early-onset PE diagnosis was defined as the development of PE before the 34th week of gestation. All 10 PE patients displayed abnormal uterine artery Doppler velocimetry. Among PE patients, two had a normal umbilical artery Doppler velocimetry, and five patients showed altered umbilical artery Doppler data (two of which were associated with a brain sparing effect). No Doppler data were available for three PE patients. Nine out of ten (90%) of the early-onset PE pregnancies were associated with intra-uterine growth restriction, defined as an estimated fetal weight below the 10th percentile for gestational age.

Chorionic villus samples were obtained from 3 healthy patients during prenatal diagnosis on the 10th week of gestation, following standard biopsy procedures. Placental villi from the third trimester of gestation were collected from 10 normotensive, healthy patients during Cesarean section by multiple biopsies in the parasagittal plane with respect to the umbilical cord insertion. Similar procedures were followed for the collection of PE samples. Portions of all placental sample types were immediately frozen in liquid

nitrogen or embedded in OCT. Clinical characteristics of patients in this study are shown in Table 1 (Nelson and Burton, 2010).

4.3. Cell Culture

Mouse trophoblast stem cells (TSC) were gifts from Dr. Kat Hadjantonakis (Memorial Sloan Kettering Institute, New York, NY) and Dr. Janet Rossant (Hospital for Sick Children, Toronto, Canada). TSC were cultured and maintained using published methods (Tanaka et al., 1998), on mouse embryonic fibroblasts (MEF) or with 50% MEF preconditioned medium. Primary human cytotrophoblasts were isolated from term placentas of normal patients, using a published protocol (Hunkapiller and Fisher, 2008). Cytotrophoblast cells were cultured on Matrigel-coated dishes in DMEM/H-21 medium (Gibco) supplemented with 2% Nutridoma (Roche).

4.4. RT-PCR

Placentas from C57BL/6 and BPH/5 mice were dissected and flash frozen in liquid nitrogen. TSC were grown on 60-mm plates for RNA extraction. For mouse and cell culture studies, RNA was isolated using Trizol (Invitrogen) and reverse transcribed using qScript cDNA Supermix (Quanta Biosciences). Gene expression was measured quantitatively using SYBR Green (Applied Biosciences) and specific primer sets for β -actin and *Egfl7* as described (Nichol et al., 2010). Differences among target expression were quantified using the $\Delta\Delta CT$ method with normalization to β -actin. For semi-quantitative RT-PCR, cDNA was amplified using the following primers:

Egfl7 (forward) 5'-CCACAAAAGAAGAAGGCTACCC-3'
Egfl7 (reverse) 5'-TCCAAGAAGGACCCTGCTCACTC-3'
 β -actin (forward) 5'-GTGGGCCGCTCTAGGCACCAA-3'
 β -actin (reverse) 5'-CTCTTTGATGTACGCACGATTTC-3'

Products were analyzed on agarose gels.

DNase-free RNA from human placental tissue was prepared using the RNeasy Mini Kit (Qiagen). RNA quality was examined by determining the presence of ribosomal RNA bands in agarose gels. RNA was reverse transcribed using random primers and the Superscript First Strand Synthesis System (Invitrogen) following the manufacturer's protocol. Specific intron-spanning primers for EGFL7, CD31 and VEGF were designed using Primer Express software (Applied Biosystems, sequences are listed below). Gene expression was measured using Real Master Mix SYBR ROX (Eppendorf) and normalized to GAPDH.

EGFL7: 5'-TCGTGCAGCGTGTGTACCAG-3', 5'-GCGGTAGGCC
 GTCCTATAGATG-3'
 CD31: 5'-TAGCGCATGGCCTGGTTAGAG-3', 5'-GGCGGTGCT
 CCCAAGTAGTCT-3'
 VEGF: 5'-ATGACGAGGGCCTGGAGTGTG-3', 5'-CCTATGTGC
 TGGCCTTGGTGAG-3'
 GAPDH: 5'-TCGGAGTCAACGGATTTGGT-3', 5'-GAATTTGCC
 ATGGGTGGAAT-3'

NOTCH1: 5'-GCGGGATCCACTGTGAGAA-3', 5'-CCGTTGAAG CAGGAGCTCTCT-3'

NOTCH2: 5'-AAAAATGGGGCCAACCGAGAC-3', 5'-TTCATC CAGAAGCGCACAA-3'

NOTCH4: 5'-CGGGCCTCTCTGCAACCT-3', 5'-GACGTCTAT GCCTGGCTCAGT-3'

HES1: 5'-AAAGATAGCTCGCGGCATTC-3', 5'-AGGTGCTTC ACTGTCATTTCCA-3'

HEY1: 5'-CATCGAGGTGGAGAAGGAGAGT-3', 5'-GACATGG AACCTAGAGCCGA-3'

HEY2: 5'-CGACCTCCGAGAGCGACAT-3', 5'-CTTTGCCCGG AGTAATTGTTCT-3'

JAGGED1: 5'-GGAGGCGTGGGATTCCA-3', 5'-CCGAGTGAG AAGCCTTTTCAATAAT-3'

DLL-4: 5'-CCAGCCAGATGGCAACTTGT-3', 5'-CCCGAAAGAC AGATAGGCTGTT-3'

4.5. RNA in situ hybridization

Egfl7 cDNA probes were generated by RT-PCR using the following primers: *Egfl7* forward 5'-AGTTACTGGTGCCAGGG ATG-3' and *Egfl7* reverse 5'-TCCTCCAAGAAGGACACCTG-3'. Amplicons were subcloned into pCR-2.1 or pCRII-Topo-TA vectors (Invitrogen) and linearized with SpeI or EcoRV. All probes were synthesized using DIG-labeling mix (Roche) according to manufacturer's instructions.

In situ hybridization (ISH) was performed by modification of the protocol described by [Hurtado and Mikawa \(2006\)](#). C57BL/6 placentas were isolated, fixed in 4% paraformaldehyde, and methanol dehydrated. Placentas were rehydrated, incubated in 5% low melt agarose (BioRad) at 42 °C for 2 h, and embedded in 5% low melt agarose through solidification at room temperature. Blocks were cut at 100 µm thickness. Agarose was removed; sections were washed in PBT (PBS-0.01% Tween20), and treated with Proteinase K (10 µg/ml, Roche) for 20 min. After terminating Proteinase K reaction with Glycine (2 mg/ml) and postfixation with 4% paraformaldehyde–0.1% glutaraldehyde (40 min), samples were washed with PBT, preincubated with hybridization buffer (50% Formamide, 200 µg/ml Yeast RNA, 0.5% Chaps, 1.3× SSC, 5 mM EDTA, 100 µg/ml Heparin, 0.2% Tween20 in DEPC-H₂O) for 1 hour at 60 °C, and incubated overnight with DIG-labeled RNA probes (1 µg/ml) at 60 °C. Samples were washed and blocked with 2% Boehringer Blocking Reagent (BBR) (Roche) in MABT for 1 h and 2% BBR–20% NGS in MABT for 1 hour, then incubated with alkaline phosphatase conjugated anti-DIG (Roche) at 1:2000 overnight at 4 °C. Samples were washed with MABT then NTMT (100 mM NaCl, 100 mM Tris–HCl pH9.5, 50 mM MgCl₂, 1% Tween20, 20 µM Levamisole, in H₂O). Signals were detected with NBT-BCIP (Roche) chromogenic substrate in NTMT at 4 °C. Samples were post-fixed in 4% paraformaldehyde and imaged using a DKC-5000 (Sony) Digital Photo Camera.

Specimens were processed for paraffin sectioning as described by [Hurtado and Mikawa \(2006\)](#). Briefly, after ISH specimens were post-fixed with 4% paraformaldehyde in PBS, rocking for 48 h at room temperature, and then overnight at 4 °C. Specimens were then dehydrated stepwise in ethanol/

double distilled H₂O solutions as follows: 70% ethanol for 10 min, 90% ethanol for 10 min, 95% ethanol for 10 min, 3×100% ethanol for 10 min, 1:1 of ethanol/Citrisolv (Sigma) for 15 min. Specimens were then incubated in paraffin at 65 °C as follows: 1:1 of Citrisolv/paraffin for 15 min, 3×paraffin for 1 h, paraffin overnight, 2×paraffin for 1 h. Specimens were then embedded in paraffin in a sagittal orientation and sectioned at 10 µm thickness. Sections were rehydrated through a series of ethanol washes (100%, 90%, 85%, 70%), counterstained with Nuclear Fast Red (Sigma–Aldrich), dehydrated through a series of ethanol washes (70%, 85%, 90%, 100%), and mounted with Permount (Fisher Scientific).

For TSC, in situ hybridization was performed as above with the following modifications: TSC were cultured on 8-chamber slides (BD Biosciences) for 2 days prior to fixation and Proteinase K incubation was carried out for 3 min.

4.6. Antibodies

The following primary antibodies were used for all immunostaining: EGFL7 (Santa Cruz, SC-34416, 2 µg/ml; R&D Systems AF3089; 2 µg/ml, In-house EGFL7 ([Fitch et al., 2004](#)), 1:100), CD31 (BD Biosciences, 553370; 5 µg/ml (mouse studies) or 0.78 mg/ml (human studies)), CDX2 (Biogenex; CDX2-88; 25 mg/ml), Cytokeratin (DakoCytomation; 53.5 µg/ml (human studies)), Cytokeratin (DAKO Z0622; 12 µg/ml (mouse studies)), and NOTCH4 (Santa Cruz, SC-5594; 2 µg/ml). The following secondary antibodies were used: Dylight 594-donkey- α -goat (Jackson ImmunoResearch, 1.5 µg/ml), Alexa 488-donkey- α -goat (Jackson ImmunoResearch, 1.5 µg/ml) or Cy3-donkey- α -goat (Chemicon, 4 µg/ml), Alexa488 donkey- α -rabbit (Jackson ImmunoResearch, 1.5 µg/ml) or FITC-donkey- α -rabbit (Chemicon, 4 µg/ml), Alexa488 donkey- α -mouse (Jackson ImmunoResearch, 1.5 µg/ml (mouse studies) or Invitrogen, 4 µg/ml (human studies)).

4.7. Immunofluorescence

Mouse whole fetoplacental units or placentas alone were isolated and embedded in an OCT:30% sucrose (2:1) mixture. Sections were permeabilized in 0.5% Triton-X/0.1% Saponin/PBS (TSP) and blocked with 1% donkey serum in 0.1% TSP/PBS (PBS–TSP). Primary antibodies (EGFL7, CD31, CYTOKERATIN) were incubated for 3 h at 37 °C in block, then secondary antibodies in block, and mounted with Prolong Gold + DAPI (Invitrogen).

TSC were grown on 8-chamber slides (BD Biosciences) for 2–4 days and fixed with cold methanol for 5 min. TSC were washed in PBS–TSP and blocked with 1% donkey serum in PBS. Primary antibodies (EGFL7, CDX2) were incubated overnight at 4 °C or 2 h at RT in block, then secondary antibodies in block, and mounted as above.

For the human studies, sections were fixed in methanol for 5 min at –20 °C and blocked in 10% donkey serum/PBS. Sections were incubated with primary antibodies (EGFL7, CD31, CYTOKERATIN, NOTCH4) in 0.1% BSA/PBS overnight at 4 °C. Sections were incubated in secondary antibodies, stained with Hoechst33342 (Sigma–Aldrich), and mounted with Møwiol (Sigma–Aldrich).

Primary human cytotrophoblasts were cultured overnight, fixed with ice-cold methanol for 5 min at -20°C , and immunostained with EGFL7 and CYTOKERATIN antibodies.

All images were acquired using an Axioplan 2 imaging microscope (Carl Zeiss) or a LSM Live 5 line scanner confocal microscope (Carl Zeiss).

4.8. Whole mount immunostaining

Blastocyst-stage embryos were isolated by flushing uteri with M2 Medium (Sigma). Embryos were collected between E3.5 and E4.0, washed with M2 medium, fixed in 2% paraformaldehyde, and permeabilized with 0.25% Triton-X/PBS. Embryos were blocked with 2.5% donkey serum/0.5% PBS-TSP/PBS, and incubated with primary antibodies (EGFL7, CDX2) overnight at 37°C in block. Blastocysts were incubated with secondary antibodies and mounted with Prolong Gold + DAPI using Fastwell spacers (Sigma-Aldrich).

4.9. Statistics

Data are represented as mean \pm SEM. The data were analyzed using a student's *t*-test with statistical significance defined as $P < 0.05$.

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Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at <http://dx.doi.org/10.1016/j.mod.2014.04.001>.

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