Abstract. Although endogenous digitalis-like factor (EDLF) is associated with the development of various physical disorders, the role in preeclampsia remains unclear. This study investigated the effects of EDLF on vascular endothelial cell damage in patients with preeclampsia and the potential mechanisms. From July 2014 to July 2015, 120 singleton pregnancy cases underwent a prenatal examination, inpatient delivery and had normal blood pressure were included in the study, either as patients with severe preeclampsia or the control patients. Serum EDLF levels were compared in these two groups, and an \textit{in vitro} hypoxic trophocyte-induced vascular endothelial cell damage model was established to explore the changes in hypoxic trophocyte EDLF level and the subsequent effects on human umbilical vein endothelial cells (HUVECs). Nuclear factor-κB (NF-κB) p65 gene expression was silenced in hypoxic trophocytes, and EDLF levels and HUVEC damage were subsequently assessed. Serum EDLF levels were significantly higher in the severe preeclampsia cases than in the controls at the same gestational week (P<0.001). EDLF levels in hypoxic trophocytes increased with the increasing co-culture duration. Damage to the biofunctions of HUVECs co-cultured with hypoxic trophocytes also increased with co-culture duration. However, silencing of NF-κB p65 in the hypoxic trophocytes reduced the EDLF levels. Annexin A2 was highly expressed in HUVECs, and no biofunctions were significantly damaged (P<0.05) compared with the group without receiving NF-κB p65 silencing. Serum EDLF levels were significantly higher in patients with severe preeclampsia compared with the controls. The results of the current study indicate that NF-κB p65 has a role in regulating EDLF production in hypoxic trophocytes.

Introduction

Preeclampsia is a common complication during pregnancy that can occur after 20 weeks of gestation. The clinical manifestations of preeclampsia include hypertension, proteinuria and edema. This complication triggers systemic disorders of multiple organs, including the brain, eyes, liver and kidneys (1), which greatly affect the health of the mother and fetus. Preeclampsia can cause multisystem damage, is closely associated with adverse pregnancy outcomes and has a high morbidity of 5-7% (2).

According to the complex multifactorial pathogenesis of preeclampsia, symptomatic treatment remains the principle therapeutic strategy, rather than etiological treatment. Currently, the most effective treatment remains pregnancy termination (3). The pathological mechanism involves insufficient vascular remodeling of the placental bed, placental ischemia and anoxia, and secondary systemic endothelial injury and/or activation (4). The patient improves following delivery of the placenta, which may indicate that detrimental factors generated in the placenta have an important role in inducing the associated endothelial cell damage (5). These detrimental factors may directly or indirectly damage vascular epithelial cells and cause organ disorders in the mother.

Endogenous digitalis-like factor (EDLF) is a micromolecular polypeptide, most of which binds non-covalently with plasma protein. It has digitalis-like activity, and inhibits the specific binding of ATPase to its receptor, and also suppresses the cell membrane sodium pump in cardiac tissue, the vascular wall, the central nervous system, the renal tubular epithelium and others. Thus, EDLF regulates water and salt metabolism, and vascular wall tension, and thus has cardiac, natriuretic and vasoconstrictive roles (6). EDLF is associated with the development of various heart, vascular, kidney and liver diseases (7). Organ damage in preeclampsia predominantly involves the cardiovascular system, liver and kidneys, indicating the potential role of EDLF in the development of preeclampsia. EDLF biosynthesis is closely associated with cholesterol side-chain cleavage in the metabolism of progesterone and pregnenolone (8). Progesterone is a hormone produced by the
placenta during gestation. Another study reported that nuclear factor-κB (NF-κB) is activated in the placenta in preeclampsia; NF-κB participates in local inflammatory reactions in the placenta and the development of preeclampsia (9).

Activated NF-κB participates in initiating and regulating the transcription of multiple genes associated with immunity, inflammation, apoptosis and cell proliferation (10,11). Cellular proliferation and differentiation are involved in embryo implantation and development (12). Multiple inflammatory factors are involved in various physical and pathological processes, including premature delivery and the initiation of parturition (13). NF-κB p65, the most explored subunit of NF-κB, has a more crucial role than the other subunits. NF-κB p65 is involved in physical processes including embryo implantation, pregnancy maintenance and the initiation of parturition (14), and the development and progression of preeclampsia (15-18).

Nitric oxide (NO) is an effective vasodilator that regulates angiostasis and tissue blood flow; in addition, NO can regulate arterial blood pressure, maternal organ perfusion, the fetal-placental vascular response and placental blood flow (19).

The widely accepted pathogenesis of preeclampsia includes epithelial cell damage, a subsequent decrease in NO production and several further pathological changes. In the blood vessel endothelium, NO synthase (NOS) is the rate-limiting enzyme in NO synthesis, and NOS expression is dependent on NF-κB regulation (20). These data suggest that NF-κB facilitates the important role of NO in preeclampsia by regulating NOS.

Although EDLF is closely associated with vascular epithelial damage and is a natural inhibitor of sodium-potassium ATPase, it affects the blood vessel endothelium through other pathways. Various effects of EDLF on blood vessel endothelial cells were revealed by differences in protein expression, and certain potential key target proteins were discovered. Annexin A2 is an important target of EDLS (via alteration of the expression of Annexin A2 to exert influence on the functions of Annexin A2); Annexin A2 is closely associated with the blood vessel endothelium (21). A study on the association between the Annexin A family and preeclampsia demonstrated that decreased Annexin A2 production may be associated with the development of preeclampsia (22).

Based on the aforementioned data, we hypothesize that EDLF may be a detrimental factor originating from the placenta that contributes to the development of preeclampsia. Furthermore, by regulating EDLF expression in hypoxic trophocytes, NF-κB p65 may affect the production of Annexin A2 (an EDLF target protein), thus decreasing NO production and resulting in the development of preeclampsia. This is a potential mechanism underlying vascular endothelial cell damage in preeclampsia. To evaluate this hypothesis, hypoxic trophocytes and human umbilical vein endothelial cells (HUVECs) were co-cultured, and changes in EDLF production by hypoxic trophocytes and the influence of EDLF on HUVEC functions were determined. Subsequently, the potential mechanism underlying the effect of EDLF generated by hypoxic trophocytes on the biofunctions of HUVECs was explored.

Materials and methods

Patients. Among the singleton pregnancy outpatient cases in the Second Xiangya Hospital of Central South University (Changsha, China) from July 2014 to July 2015, 60 cases of severe preeclampsia (age, 33.73±4.49 years) were randomly selected and included in the severe preeclampsia group. The diagnostic criteria were based on those reported in the literature (23): i) Systolic pressure ≥160 mmHg and/or diastolic pressure ≥110 mmHg; ii) proteinuria ≥5,000 mg/24 h or random urine protein (+++); iii) persistent headache, vision disorder or other cranial nerve symptoms; iv) persistent epigastric pain, subcapsular hematoma of the liver or hepatorrhexis symptoms; v) hypertension (increased serum ALT or AST); vi) abnormal renal function (oliguria: 24 h urine volume <400 ml or urine volume per hour <17 ml; or serum creatinine >106 ng/l); vii) hypoproteinemia, hydropsortheorax or serooperationteum; viii) blood system disorders: microangiopathic hemolysis or blood platelet count <100x109/l; ix) cardiac failure and pulmonary edema; x) fetal growth restriction or oligohydramnios; or xi) early-onset preeclampsia before 34 weeks of pregnancy. The exclusion criteria included the following: i) Chronic hypertension; severe diseases of the heart, lungs, kidneys, liver, or thyroid, or other endocrine diseases experienced before pregnancy; ii) obstetric complications, including placental abruption, gestational diabetes mellitus, premature rupture of membranes, maternal-fetal blood group incompatibility, or placenta previa; iii) the use of assisted reproductive technology; and iv) administration of abortion prevention medicine.

The 60 singleton pregnancy cases with a prenatal examination or inpatient delivery and normal blood pressure during the same period were included in the control group (age, 32.73±4.36 years). Patient information, such as age, blood pressure, height, weight, gestational weeks, and parity and gravidity, was collected and recorded. Peripheral venous blood samples were centrifuged to collect sera. EDLF concentrations were determined using EDLF radioimmunooassay kit (cat. no. FM0063; Beijing North Institute of Biological Technology, Beijing, China; used in the in vitro experiments as well). As numerous factors can influence the outcome of this radioimmunoassay (such as cross reactivity with digoxin and ouabain and experimental conditions) (24-26), samples were collected strictly according to the inclusion criteria, the blood collection time and conditions were highly standardized, and the experimental procedures (such as incubation and centrifugation conditions) were performed in strict accordance with the instructions provided with the kit.

This study was approved by the Ethics Committee of the Second Xiangya Hospital of Central South University. Informed consent was obtained from all participants.

Experimental cells. Human BeWo placental choriocarcinomata trophocyte cells (Shanghai Sixin Biotechnology, Ltd., Shanghai, China) were used in this study (27). HUVECs were purchased from the Physical Laboratory of Xiangya School of Medicine, South Central University.

Trophocyte and HUVEC cultures. HUVECs were cultured in Dulbecco's modified Eagle's medium (Hyclone, Logan, UT, USA), and BeWo cells were cultured in F12K medium (Life Technologies; Thermo Fisher Scientific, Inc., Waltham, MA, USA); both cultures were supplemented with fetal bovine serum (FBS; Gibco, Invitrogen, Grand Island, NY, USA) and
incubated in saturated humidity and 5% CO₂ at 37°C. Culture media were changed every 2 days. Cells were passaged with 0.25% trypsin upon reaching 80% confluence.

**HUVEC confluence.** HUVECs in the logarithmic growth phase were diluted to a concentration of 1x10⁵/l, seeded into 5 wells of a 12-well plate at 1 ml/well and incubated in saturated humidity and 5% CO₂ at 37°C. Cell growth and confluence were observed regularly and frequently by light microscopy (BX53; Olympus Corporation, Tokyo, Japan). The basic features of healthy cells included the following: Proper transparency, low intracytoplasmic granular substance level, small cytoplasmic volume and no hypertrophy. In addition, the healthy cells adhered well and grew evenly, with no or few cells floating in the culture medium. Healthy cells were co-cultured with hypoxic trophocytes.

**Treatment of BeWo trophocytes with hypoxia.** Following trypsinization (0.25%), centrifugation (at 350 x g at room temperature for 5 min) and resuspension (in complete medium containing 10% FBS), BeWo cells in the logarithmic growth phase were diluted to a concentration of 4x10⁵/l, seeded into 5 wells of a 12-well plate at 1 ml per well and cultured overnight for adherence in saturated humidity and 5% CO₂ at 37°C. A sterile stock solution of CoCl₂ (1 mol/l) was prepared and diluted in complete culture medium to a concentration of 10 mmol/l to form the working solution. The medium was replaced with 1 ml of fresh medium. Then, 30 µl of the working CoCl₂ solution was added for a final concentration of 300 µmol/l. For the hypoxia treatment, these cells were incubated in saturated humidity and 5% CO₂ at 37°C for 24 h.

**Transfection of hypoxic trophocytes with the NF-κB plasmid.** Hypoxia-treated BeWo cells were divided into 3 groups, and culture media were replaced with antibiotic-free F12K medium (2 ml/bottle). The NF-xBsi-2 plasmid (Shanghai GeneChem Co., Ltd., Shanghai, China) was transfected into these cells as follows. NF-xBsi-2 (20 µl) was added to a sterile Eppendorf tube, and Lipofectamine® 2000 transfection reagent (5 µl; GenStar, Beijing, China) was added and diluted in complete culture medium to a concentration of 10 nmol/l to form the working solution. The medium was replaced with 1 ml of fresh medium. Then, 30 µl of the working CoCl₂ solution was added for a final concentration of 300 µmol/l. For the hypoxia treatment, these cells were incubated in saturated humidity and 5% CO₂ at 37°C for 24 h.

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**Evaluation of HUVEC barrier function.** The culture supernatant in the lower chamber was harvested (500 µl) at the designated time-points, and a fluorospectrophotometer (excitation wavelength, 488 nm; sensitivity, 50) was used to determine the fluorescence intensity of FITC. All the operations were performed in the dark.

**MTT assay to evaluate HUVEC proliferation.** HUVEC proliferation was evaluated using MTT assays, which were performed in accordance with the manufacturer's instructions (Sigma-Aldrich; Merck KGaA). The endothelial cells in the upper chamber were treated with trypsin (0.25%), resuspended, seeded into a 96-well plate at a concentration of 1x10³ cells/well and incubated for 48 h. Then, 20 µl MTT solution (5 mg/ml) was added to each well, and the medium was discarded after 4 h of incubation. Subsequently, dimethyl sulfoxide (150 µl) was added, and after 10 min of incubation at room temperature, 100 µl from each well was transferred into the coated wells of a 96-well plate. Absorbance values were determined at a wavelength of 490 nm measured using a 960PRO fluorospectrophotometer (INESA; Shanghai, China; http://en.inesa-instrument.com/index.html). The other groups were analyzed via the procedures described above.

**Nitrate reductase activity assay to determine the NO concentration.** The NO level in the HUVEC culture supernatant was detected using nitrate reductase activity assays (NO detection kit; cat. no. A012; Nanjing Jiancheng Bioengineering Institute, Nanjing, China). Endothelial cell culture supernatants from the upper chambers were harvested at the designated time-points (500 µl/group), and the reagents were mixed well with the specimens according to the manufacturer's instructions (Nanjing Jiancheng Bioengineering Institute) and incubated at room temperature for 10 min. Then, the absorbance values were determined (550 nm wavelength and 0.5 cm optical path) and normalized to that of double-distilled water as the blank.

**Western blot analysis of Annexin A2 expression in HUVECs.** Cells were suspended (20 µl) and treated with radioimmuno-precipitation lysis buffer (100 µl; Beyotime, Shanghai, China) containing 1 µl each of phenylmethylsulfonyl fluoride and protease and protein tyrosine phosphatase inhibitors. Then, the cells were incubated on ice for 30 min and centrifuged at 350 x g at 4°C for 5 min. The supernatant was harvested and stored at -20°C; the Laemmli system was used. The concentrations of the spacer and separation gels were 5 and 12%, respectively. The prepared separation gel was gently infused into the space between the fixed double-paned glass, and isopropanol was used to seal the top; the gels were allowed to set for 30 min at room temperature. Protein levels were determined using a bicinchoninic acid protein determination kit (Wellbio, Changsha, China). A total of 25 µg protein was loaded per lane, with 15% gel. NC membranes were used for protein transfer. TBST containing 5% skim milk was used for blocking. After immersing, the membranes were placed
Table I. Comparison of serum EDLS levels between the two groups.

<table>
<thead>
<tr>
<th>Group</th>
<th>Serum EDLS (ng/l)</th>
<th>P-valuea</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>66.49±10.36</td>
<td>P&lt;0.01</td>
</tr>
<tr>
<td>Severe preeclampsia</td>
<td>125.09±14.33</td>
<td></td>
</tr>
</tbody>
</table>

aHealthy control vs. severe preeclampsia. EDLS, endogenous digitalis-like factor.

at room temperature for 1 h. The following procedures included primary antibody incubation, secondary antibody incubation, development (or staining) and exposure (28). The primary antibodies were as follows: NF-κB (cat. no. 6956S; source, mouse; dilution, 1,000; molecular weight, 65 kDa; Cell Signaling Technology, Beverly, MA, USA), Annexin A2 (cat. no. 11256-1-AP; source, rabbit; dilution, 1,000; molecular weight, 36 kDa; Proteintech, Rosemont, IL, USA), Actin (cat. no. 60008-1-Ig; source, mouse; dilution, 4,000; molecular weight, 42 kDa; Proteintech). The membranes were incubated with the primarily antibodies at 4°C overnight. After incubation, they were washed thrice with TBS-T, for 15 min per wash. HRP-conjugated goat anti-rabbit IgG was used as the secondary antibody (dilution: 1: 3,000; Proteintech) (incubation for 45-60 min). The samples were washed thrice, for 15 min per wash. The exposure was performed using superECL Plus (Thermo Pierce, Rockford, IL, USA). Western blot analysis was performed using Quantity One software (4.6.2; Bio-Rad Laboratories, Inc., Hercules, CA, USA).

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR). In addition, Annexin a2mRNA was detected using a reverse transcription kit [Fermentas (Thermo Fisher Scientific, Inc., Pittsburgh, PA, USA)]. Annexin A2 mRNA levels in HUVECs were determined by RT-qPCR. Total RNA was extracted with TRIzol reagent (Life Technologies; Thermo Fisher Scientific, Inc.) according to the manufacturer's instructions. The purity and concentration of total RNA were determined by ultraviolet spectrophotometry. Actin was used as a normalization control. Reverse transcription was performed with a kit from Fermentas. Primers were designed with Primer 5.0 software (PREMIER Biosoft, Vancouver, Canada) and synthesized by Invitrogen (Thermo Fisher Scientific, Inc.). The extension length of the Annexin A2 primers (forward, 5'-GCCCTGGCAAAGGTAGA-3' and reverse, 5'-ACGCTCCGCTGTTCA-3') generated a product of 149 bp. The actin primers (forward, 5'-CATCCCTGCCGCT GGACCTG-3' and reverse 5'-TAAATGTACACGCACGA TTTCC-3') generated a product of 107 bp. One sample was analyzed in three parallel experimental wells. The total reaction system of 30 µl contained RT product as a template (1 µl), forward primer (10 µM, 0.5 µl), reverse primer (10 µM, 0.5 µl), PCR-grade H₂O (13 µl) and 2X SYBR-Green PCR Master Mix (15 µl; Applied Biosystems, Foster City, CA, USA). The qPCR amplification conditions were as follows: 95°C for 10 min, followed by 40 cycles of 95°C for 10 sec and 60°C for 50 sec. The melting curves were collected at 60-95°C. The qPCR (5 µl) products were analyzed via 2% agarose gel electrophoresis. Annexin A2 mRNA was detected with a Gel Doc XR gel-imaging system (Bio-Rad Laboratories, Inc.). Data were analyzed using the qPCR analytical tool, and quantification was performed using the 2⁻ΔΔCt method.

Statistical analysis. The data were analyzed using SPSS 19.0, and data are presented as the mean ± standard deviation. Normality tests were performed with the Kolmogorov-Smirnov and Shapiro-Wilk tests. The homogeneity of the variance between groups was assessed via Levene's method. The inter-group differences of paired data were analyzed using the t-test. Pairwise comparisons of differences among multiple groups were performed using the Student-Newman-Keuls-q test. P<0.05 was considered to indicate a statistically significant difference.

Results

EDLF levels. Serum EDLF levels in the severe preeclampsia patients and the normal pregnancy controls are presented in Table I. No significant differences were identified in gestational weeks, age, weight, height, or parity and gravidity between the two groups (P>0.05). Serum EDLF levels were significantly higher in severe preeclampsia cases compared with normal pregnancy cases (P<0.05).

HUVEC morphology. HUVEC morphology was observed by light microscopy (Fig. 1). The cells grew well to confluency and exhibited a cobblestone-like arrangement, sharp edges, abundant cytoplasm, elliptical shape, centralized nuclei and clear nucleoli.

Morphological changes in trophocytes following transfection. BeWo cells exhibiting the typical characteristics were passaged after 3 days of culture and observed by light microscopy after an additional 2 days (Fig. 2). The cells showed healthy and compact growth, and had formed multiple layers.

Silencing efficiency of NF-κB siRNA. The gene silencing efficiency of NF-κB siRNA was evaluated by western blot analysis (Fig. 3 and Table II). NF-κBsi-2 exerted the best gene-silencing effect and was therefore used in the subsequent experiments.

Morphology of trophocytes following transfection. Fluorescently-labeled siRNA-transfected trophocytes were observed by fluorescence microscopy. Green fluorescence was successfully expressed in the trophocytes (Fig. 4).

EDLF levels in hypoxic trophocytes at different time points following NF-κB p65 gene silencing. The culture supernatant of hypoxic trophocytes was harvested at different time points, and EDLF concentration was determined by radioimmunoassay. EDLF expression levels were increased over time in untransfected hypoxic trophocytes co-cultured with HUVECs. However, in the cells transfected with NF-κBsi-2 the cells exhibited no significant changes in EDLF expression over time under the same conditions (Table III).

Effect of NF-κB p65 silencing in hypoxic trophocytes on HUVEC functions
Proliferative activity. The proliferation of HUVECs co-cultured with hypoxic trophocytes changed according to the co-culture time (Table IV). Based on the MTT assay results, HUVEC proliferation gradually decreased with increasing time of co-culture with untransfected trophocytes. After transfection, the proliferation of HUVECs co-cultured

![Figure 1. HUVEC morphology under light microscopy. (A) Confluent HUVECs fused into a monolayer (x100). (B) Confluent HUVECs (x200). (C) HUVECs growing well (x100). (D) HUVECs growing well (x400). HUVEC, human umbilical vein endothelial cell.](image1)

![Figure 2. BeWo morphology as viewed by light microscopy. (A) x100. (B) x200.](image2)

![Figure 3. Silencing efficiency of NF-κBsi. NF-κB, nuclear factor-κB; si, small interfering RNA.](image3)

![Figure 4. BeWo cells following transfection (x100). (A) Prior to and (B) following transfection with nuclear factor-κB small interfering RNA plasmid. Control, normally cultured cells (cells and corresponding medium only); Con, cultured cells transfected with anti-sense strand vectors.](image4)

*Proliferative activity.*
with transfected trophocytes gradually increased, exhibiting significant differences after 24 h (P<0.05) and 48 h (P<0.01).

**Barrier function.** The barrier function of HUVECs co-cultured with hypoxic trophocytes changed over time (Table V). FITC-conjugated glucan was applied to the co-culture with untransfected hypoxic trophocytes and gradually moved through the HUVEC monolayer from the upper chamber into the lower chamber, which increased the concentration of FITC-glucan in the lower chamber over time, and provided an indicator of the HUVEC barrier function. However, in the culture using NF-κBsi-2 trophocytes, the amount of FITC-glucan passing through the HUVEC monolayer decreased gradually, and the barrier function damage was relatively improved by 24 h of culture (P<0.05) compared with the untransfected cells at 24 h; significant differences were also observed after 48 h of co-culture (P<0.01).

**NO level.** Changes were observed in NO production by HUVECs co-cultured with hypoxic trophocytes (Table VI). In HUVECs cultured with untransfected trophocytes, the relative NO level gradually decreased with increasing co-culture time. However, in HUVECs cultured with NF-κBsi-2 trophocytes, the relative NO production increased gradually, and was increased compared with the untransfected group by 24 h (P<0.05), and significant differences were also observed after 48 h (P<0.01).

Annexin A2 expression in co-cultured HUVECs. Annexin A2 expression in HUVECs was determined by western blot analysis (Fig. 5; Table VII). In the untransfected group, Annexin A2 protein expression in co-cultured HUVECs decreased over time. Following NF-κBsi-2 transfection of the trophocytes, Annexin A2 levels in HUVECs were increased, beginning at 24 h, and the differences between the untransfected and NF-κBsi-2 groups became significant at 72 h (P<0.01).

Relative Annexin A2 mRNA expression levels in HUVECs. HUVECs were harvested prior to trophocyte transfection and following transfection with NF-κBsi-2, and total RNA was extracted and transcribed into cDNA. Relative Annexin A2 mRNA levels were detected via qPCR, and actin was used for data normalization (Table VIII). Annexin A2 mRNA expression in HUVECs decreased gradually over time when co-cultured with untransfected trophocytes. After 24 h of co-culture with NF-κBsi-2-transfected trophocytes, Annexin A2 mRNA levels in HUVECs increased over time compared with the untransfected group (P<0.05).

**Discussion**

Preeclampsia is one of the major causes of mortality in pregnant women and perinatal infants. The fundamental pathological changes include systemic damage to vascular endothelial cells, inflammation-associated systemic small vessel spasms and a hypercoagulable state. These changes can cause various clinical manifestations, including hypertension, edema and proteinuria, and subsequent severe ischemia in vital organs.
including the heart, brain, liver and kidneys. Furthermore, in severe cases, this condition can result in sequelae or mortality in the mother or infant. Although numerous research efforts have been directed toward investigating this condition, its pathogenesis remains unclear (29,30). Therefore, pregnancy termination is often considered the only feasible strategy because more efficient treatments have yet to be developed. However, terminating the pregnancy is not always an acceptable solution. Due to the high mortality and disability rates in severe preeclampsia cases, early screening, prevention and diagnosis have more important roles than active treatment.

The findings of the current study indicated that serum EDLF levels were significantly higher in patients with severe preeclampsia patients compared with normal pregnant women, which indicated a potential association between EDLF and preeclampsia development. Subsequently, a Transwell system was used to co-culture hypoxic trophocytes and HUVECs for 12, 24, 48, 72, 96 or 120 h to detect changes in EDLF protein levels. The results demonstrated that EDLF increased with increasing duration of co-culture with hypoxic cells, accompanied by a decrease in free NO. In addition, HUVEC proliferation and cellular monolayer barrier function, as well as NO production by HUVECs, changed markedly during co-culture with hypoxic trophocytes, indicating the potential for cytokine-mediated signaling between trophocytes and HUVECs.

### Table V. Changes in human umbilical vein endothelial cell barrier function post-transfection.

<table>
<thead>
<tr>
<th>Co-culture time</th>
<th>Untransfected</th>
<th>NF-κBsi-2</th>
</tr>
</thead>
<tbody>
<tr>
<td>12 h</td>
<td>12.64±0.06</td>
<td>12.08±0.06</td>
</tr>
<tr>
<td>24 h</td>
<td>13.79±0.56</td>
<td>11.53±0.43</td>
</tr>
<tr>
<td>48 h</td>
<td>15.55±0.16</td>
<td>8.66±0.39</td>
</tr>
<tr>
<td>72 h</td>
<td>16.36±0.48</td>
<td>7.50±0.63</td>
</tr>
<tr>
<td>96 h</td>
<td>17.64±0.51</td>
<td>5.85±0.64</td>
</tr>
<tr>
<td>120 h</td>
<td>18.68±0.63</td>
<td>4.69±0.55</td>
</tr>
</tbody>
</table>

*P<0.05 vs. untransfected; *P<0.01 vs. untransfected. n=3. NF-κB, nuclear factor-κB; si, small interfering RNA.

### Table VI. Relative nitric oxide levels in human umbilical vein endothelial cells.

<table>
<thead>
<tr>
<th>Co-culture time</th>
<th>Untransfected</th>
<th>NF-κBsi-2</th>
</tr>
</thead>
<tbody>
<tr>
<td>12 h</td>
<td>0.77±0.01</td>
<td>0.79±0.00</td>
</tr>
<tr>
<td>24 h</td>
<td>0.73±0.02</td>
<td>0.85±0.01</td>
</tr>
<tr>
<td>48 h</td>
<td>0.64±0.05</td>
<td>0.83±0.02</td>
</tr>
<tr>
<td>72 h</td>
<td>0.62±0.02</td>
<td>0.85±0.02</td>
</tr>
<tr>
<td>96 h</td>
<td>0.52±0.02</td>
<td>0.87±0.02</td>
</tr>
<tr>
<td>120 h</td>
<td>0.45±0.02</td>
<td>0.90±0.01</td>
</tr>
</tbody>
</table>

*P<0.05 vs. untransfected; *P<0.001 vs. untransfected. n=3. NF-κB, nuclear factor-κB; si, small interfering RNA.

### Table VII. Relative Annexin A2 protein expression levels in human umbilical vein endothelial cells.

<table>
<thead>
<tr>
<th>Co-culture time</th>
<th>Untransfected</th>
<th>NF-κBsi-2</th>
</tr>
</thead>
<tbody>
<tr>
<td>12 h</td>
<td>0.51±0.00</td>
<td>0.52±0.01</td>
</tr>
<tr>
<td>24 h</td>
<td>0.47±0.00</td>
<td>0.56±0.01</td>
</tr>
<tr>
<td>48 h</td>
<td>0.41±0.00</td>
<td>0.58±0.01</td>
</tr>
<tr>
<td>72 h</td>
<td>0.39±0.01</td>
<td>0.59±0.01</td>
</tr>
<tr>
<td>96 h</td>
<td>0.36±0.01</td>
<td>0.60±0.01</td>
</tr>
<tr>
<td>120 h</td>
<td>0.31±0.01</td>
<td>0.63±0.01</td>
</tr>
</tbody>
</table>

*P<0.05 vs. untransfected; *P<0.01 vs. untransfected. Annexin A2 was normalized to the actin gray level. NF-κB, nuclear factor-κB; si, small interfering RNA.

### Table VIII. Relative Annexin A2 mRNA expression levels in human umbilical vein endothelial cells.

<table>
<thead>
<tr>
<th>Co-culture time</th>
<th>Untransfected</th>
<th>NF-κBsi-2</th>
</tr>
</thead>
<tbody>
<tr>
<td>12 h</td>
<td>0.49±0.01</td>
<td>0.51±0.01</td>
</tr>
<tr>
<td>24 h</td>
<td>0.44±0.01</td>
<td>0.54±0.00</td>
</tr>
<tr>
<td>48 h</td>
<td>0.42±0.01</td>
<td>0.55±0.01</td>
</tr>
<tr>
<td>72 h</td>
<td>0.38±0.01</td>
<td>0.57±0.01</td>
</tr>
<tr>
<td>96 h</td>
<td>0.38±0.01</td>
<td>0.62±0.02</td>
</tr>
<tr>
<td>120 h</td>
<td>0.29±0.01</td>
<td>0.66±0.00</td>
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*P<0.05 vs. untransfected; *P<0.01 vs. untransfected. Values are presented as the mean ±standard deviation. n=3. NF-κB, nuclear factor-κB; si, small interfering RNA.
endothelial cells and corresponding effects on endothelial cell biofunctions. Using RNAi technology, NF-κB p65 gene expression was silenced in hypoxic trophocytes, which downregulated the expression of EDLF. This led to upregulation of Annexin A2, an important EDLF target protein (38), and a decrease in HUVEC damage. Therefore, it is inferred that EDLF is a toxic factor originating from the placenta in patients with severe preeclampsia. NF-κB p65 may regulate EDLF expression in hypoxic trophocytes, thus influencing Annexin A2 expression and subsequent NO production. These events cause endothelial cell damage and the development of preeclampsia.

In patients with severe preeclampsia, hypoxic trophocytes produce high levels of EDLF, which is regulated by NF-κB p65, to elicit cytokine secretion by endothelial cells, which affects vasomotion and cell proliferation (53). Various endothelial cell functions are realized through the balance of NO and endothelin (ET) (20). A decrease in NO production and release may evoke an imbalance in vascular factors, with subsequent arteriole spasms and increased peripheral resistance, particularly in the kidneys, uterus and placenta, which promotes the development of hypertension and preeclampsia (31). A previous study reported that EDLF levels are significantly higher in newborn umbilical cord blood than in the peripheral blood of pregnant women (32).

In addition, the previous study reported that EDLF induces the expression and release of ET-1 to initiate vasocostriction; this upregulated the sensitivity of the vasculature to endogenous vasopressin and downregulated the relaxant effects of NO, prostacyclin and acetylcholine, which may result in disordered vasodilatation and vasoconstriction during the development of preeclampsia (32). As a physiological stress, pregnancy creates tension in sympathetic nerves, which promotes the development of preeclampsia through exciting sympathetic nerves and eliciting a series of physiological responses, such as vasoconstriction and water and sodium retention. A previous study reported that EDLF evokes a group of sympathetic nerve excitation reactions and water and sodium retention. A previous study reported that EDLF expression in hypoxic trophocytes, thus influencing Annexin A2 expression and subsequent NO production. These events cause endothelial cell damage and the development of preeclampsia.

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expression of various genes, including those for epidermal growth factor and platelet-derived growth factor. These events are often followed by vascular smooth muscle cell proliferation, intramembrane signaling for vascular remodeling, and subsequent increases in the level of collagenous fibers, the thickness of vessel walls and the mean pressure in arterioles. These changes subsequently elicit the clinical manifestations of hypertensive disorders complicating pregnancy, including hypertension, proteinuria and edema (18,55).

In the current study, hypoxic trophocytes were co-cultured with HUVECs. RNAi technology was used to silence the NF-κB p65 gene in hypoxic trophocytes, which resulted in reduced EDLF levels. By contrast, the expression of Annexin A2, an EDLF target protein, was upregulated in HUVECs, and NO production increased. Furthermore, the monolayer barrier function and HUVEC proliferation were increased by NF-κB silencing in hypoxic trophocytes. These results suggested that NF-κB p65 may affect Annexin A2 expression in HUVECs by regulating EDLF production in hypoxic trophocytes, which impairs vascular endothelial cell function and NO secretion.

The results of the current study demonstrated the effects of EDLF only in terms of vascular endothelial cell damage. However, EDLF also plays a maintenance role in normal pregnancy. The levels of EDLF that lead to positive and negative effects remain to be determined via analysis of a large number of samples. The results of such an analysis may lead to the development of a sensitive and specific marker for convenient, rapid and reproducible early diagnosis and monitoring, and for assessing therapeutic effects. EDLF causes vascular endothelial cell damage through multiple pathways; this study focused on only the NF-κB p65 pathway. Further comprehensive studies of multiple pathways may contribute more valuable information regarding preeclampsia pathogenesis.

In conclusion, high levels of EDLF may have an important role in the multifactorial pathogenesis of preeclampsia, and NF-κB p65 is a potential key factor in the regulation of EDLF concentration in preeclampsia. The results of this study may provide novel strategies for intervening in the development of preeclampsia in clinical practice.

References


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