

CLDN3 expression and function in pregnancy-induced hypertension

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Abstract. This aim of the present study was to investigate the expression and function of claudin 3 (CLDN3) in pregnancy-induced hypertension. The mRNA expression levels of CLDN3 in the placental tissue and peripheral blood of patients with pregnancy-induced hypertension were measured using reverse transcription-quantitative PCR. Human trophoblast HTR8/SVneo cells overexpressing CLDN3 were generated using a lentiviral vector. Cell Counting kit-8 (CCK-8) assay, flow cytometry, Transwell chamber assays, confocal laser scanning microscopy and western blot analysis were performed to detect cell proliferation, invasion, migration and apoptosis, in addition to matrix metalloproteinase (MMP) expression and ERK1/2 phosphorylation. The mRNA expression levels of CLDN3 were significantly reduced in the placental tissues and peripheral blood samples of patients with pregnancy-induced hypertension compared with healthy pregnant controls. CLDN3 overexpression significantly increased HTR8/SVneo cell proliferation, invasion and migration whilst reducing apoptosis. HTR8/SVneo cells overexpressing CLDN3 also exhibited increased myofiber levels, increased MMP-2 and MMP-9 expression and increased ERK1/2 signaling activity. CLDN3 downregulation may be associated with the pathogenesis of pregnancy-induced hypertension. In conclusion, CLDN3 promotes the proliferative and invasive capabilities of human trophoblast cells, with the underlying mechanisms possibly involving upregulation of MMP expression via the ERK1/2 signaling pathway.

Introduction

Pregnancy-induced hypertension is a common disease observed during the gestational period. If not treated in time,

pregnancy-induced hypertension may cause significant harm to the mother and fetus (1,2). Recent clinical studies have shown that maternal mortality as a result of pregnancy-induced hypertension to be 4.2-10 million worldwide as of 2017, accounting for ~9% of all maternal deaths and 2.2% of perinatal child mortality (3). Thus, this disease poses a serious threat to maternal and child health, and represents one of the main causes for the death of pregnant women and neonates (4). The clinical symptoms of pregnancy-induced hypertension mainly include transient hypertension and proteinuria in pregnant women, which normally disappear following delivery (5). At present, pregnancy-induced hypertension is generally divided into five types: Gestational hypertension, pre-eclampsia (mild and severe), eclampsia, chronic hypertension complicated by pre-eclampsia and chronic hypertension combined with pregnancy (6). Pregnant women with severe pregnancy-induced hypertension may suffer from hemolysis, thrombocytopenia, liver and kidney dysfunction, pulmonary edema and visual disturbances (5). Risk factors of the disease include obesity, pre-pregnancy hypertension, diabetes and old age (7). The causes for pregnancy-induced hypertension remain to be elucidated; however, they may be associated with changes in the immune system such as histocompatibility antigen-associated immunological abnormalities (8). In addition, genetic susceptibility may also be involved in the pathogenesis of this disease (9,10).

Trophoblast cells are one of the components of the maternal placental architecture, involved in the regulation of placental microenvironment remodeling, implantation of embryos and normal fetal development (11). Previous studies have shown that trophoblast cells can differentiate into two types of trophoblast cells at the early stages of blastocyst implantation, namely cytotrophoblast cells and syncytiotrophoblast cells (12,13). Cytotrophoblast cells can fuse with syncytiotrophoblast cells, which differentiate into extravillous trophoblasts (EVTs). Some EVT cells infiltrate the deeper layers of the endometrium, and are known as interstitial trophoblast cells (14), whilst others invade the maternal uterine spiral artery, and are called endovascular trophoblast cells (15). Trophoblast cells exhibit similar migratory capacities to tumor cells, which are closely related in biological function (16). Studies have shown that EVT cells migrate from the placental villi and invade the endometrium and maternal spiral arteries, where they participate in uterine artery revascularization and regulate placental blood flow and immune responses (17).

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Downregulation of the invasive ability of trophoblast cells can result in defects in uterine spiral artery remodeling and placental insufficiency, leading to pregnancy-induced hypertension, eclampsia and miscarriage (18). Therefore, it is of great clinical significance to study changes in trophoblast cell invasion and the associated mechanism in the pathogenesis of pregnancy-induced hypertension.

Tight junction proteins are important in the maintenance of cell-to-cell connections, which serve important roles in cell polarity and the formation of cellular barriers such as the intestinal epithelial barrier (19,20). In particular, claudins (CLDNs) are members of the tight junction protein family that serve important roles in the formation of tight junctions. In total, 24 CLDNs have been identified (21). In recent years, studies have demonstrated that CLDN3 is abnormally expressed in a number of tumor tissues, including hepatocellular carcinoma and breast cancer (22,23) and closely associated with the invasion and metastasis of tumor cells (24,25). Notably, when tumor cells metastasize, the tight junctions between cells must be destroyed, and CLDN3 is an important component of these junctions (26). During trophoblast cell invasion, the breaking of tight junctions is also an important prerequisite for detachment (27). However, the function of CLDN3 in this process remain unclear. Therefore, in the present study, the regulatory role of CLDN3 in the invasive abilities of trophoblast cells was investigated, on tissue and cellular levels.

Materials and methods

Study subjects and sample collection. A total of 51 pregnant women with hypertension, including 25 patients diagnosed with pregnancy-induced hypertension, 11 patients with mild pre-eclampsia and 15 patients with severe pre-eclampsia, and 30 normal pregnant women were included in this study, all of whom were admitted to Laiwu Maternal and Child Health Hospital (Laiwu, China) for delivery from December 2016 to December 2017. In these patients, the pregnancy hypertension was defined as: i) BP $\geq 140/90$ mmHg during pregnancy, which returned to normal within 12 weeks after delivery; ii) urine protein (-); and iii) cases that may be associated with upper abdominal discomfort or thrombocytopenia. Mild pre-eclampsia was defined as: i) BP $\geq 140/90$ mmHg appeared during pregnancy, which returned to normal within 12 weeks after delivery; ii) urine protein (-); and iii) cases that may be associated with upper abdominal discomfort or thrombocytopenia. Severe pre-eclampsia was defined as maternal convulsions that cannot be explained by other reasons. None of the pregnant women had previous pregnancies or suffered from hypertension, diabetes or underlying diseases, including liver, kidney and autoimmune diseases, prior to the present pregnancy. All these patients received cesarean section due to pregnancy-induced hypertension, abnormal fetal position or other social factors. There were no significant differences in age, gestational period or neonatal weight between the pregnancy-induced hypertension and control group (mean age of patients with pregnancy-induced hypertension, 29.5 ± 1.08 years; mean gestational period, 38 ± 2.08 weeks; mean neonatal weight, 7.6 ± 2.3 kg. Mean age of patients with mild-pre-eclampsia, 33.1 ± 1.08 years; gestational period, 39 ± 3.08 weeks; neonatal weight, 7.1 ± 1.3 kg. Mean age of

patients with severe pre-eclampsia, 31.45 ± 1.08 years; gestational period, 40 ± 2.60 weeks; neonatal weight, 7.7 ± 1.7 kg). Prior written informed consent was obtained from every patient and the study was approved by the ethics review board of Laiwu Maternal and Child Health Hospital. In total 10 ml peripheral blood sample was obtained from each subject and the collection of placental tissues was performed 10 min after the placenta was delivered, with a $2 \times 2 \times 2$ cm sample of tissue dissected from the central area (avoiding infarction and calcification) of the maternal placenta. The tissue was washed with saline immediately after collection, and then stored at -80°C .

Reverse transcription-quantitative PCR (RT-qPCR). Total RNA was extracted from the placental tissue and peripheral blood samples using TRIzol® (Thermo Fisher Scientific, Inc.) according to the manufacturer's protocol. RNA concentrations were quantified using the NanoDrop method. Reverse transcription was performed with $0.5 \mu\text{g}$ RNA to obtain cDNA using BeyoRT™ cDNA First Chain Synthesis kit (cat. no. D7166; Beyotime Institute of Biotechnology). Subsequent qPCR was performed using BeyoFast™ SYBR Green qPCR Mix (cat. no. 7260; Beyotime Institute of Biotechnology) in a StepOnePlus™ Real-Time PCR instrument. The following primer sequences were used: CLDN3 forward, 5'-GCCACCAAGGTCGTCTACTC-3' and reverse, 5'-CCTGCGTCTGTCCCTTAGAC-3' and GAPDH forward, 5'-CGGAGTCAACGGATTTGGTTCGTAT-3' and reverse, 5'-AGCCTTCTCCATGGTGGTGAAGAC-3'. The total $20 \mu\text{l}$ PCR mixture consisted of $10 \mu\text{l}$ RT-qPCR-Mix, $0.5 \mu\text{l}$ each primer, $2 \mu\text{l}$ cDNA and $7 \mu\text{l}$ ddH₂O. The thermocycling conditions were as follows: Initial denaturation at 95°C for 10 min; followed by 40 cycles of 95°C for 1 min and 60°C for 1 min. Target gene expression levels were calculated using the $2^{-\Delta\Delta\text{C}_q}$ method (28). GAPDH was used as internal reference.

Human trophoblast cell culture. Normal human trophoblast HTR8/SVneo cells were purchased from American Type Culture Collection. Cells were cultured using RPMI-1640 medium (Gibco; Thermo Fisher Scientific, Inc.) containing 10% fetal bovine serum (FBS; Gibco; Thermo Fisher Scientific, Inc.) and supplemented with 100 U/ml penicillin and 100 U/ml streptomycin, in a humidified atmosphere at 37°C containing 5% CO₂. When cell confluence reached 90%, the cells were passaged using 0.5% trypsin.

Construction and transfection of lentivirus vector for CLDN3. The lentiviral Lv-GFP-CLDN3 vector was constructed by Hanbio Biotechnology Co., Ltd., with the titer of 1×10^8 pfu. HTR8/SVneo cells were first seeded into 24-well plates at a density of 1×10^5 cells/well and cultured with RPMI-1640 medium containing 10% FBS. When 70% confluence was achieved, the HTR8/SVneo cells were transfected with either empty vector or Lv-GFP-CLDN3 at a multiplicity of infection of 20. The medium was changed to fresh RPMI-1640 medium containing 10% FBS after 6 h, and the cells were cultured at 37°C under 5% CO₂ for a further 48 h.

Cell counting kit (CCK)-8 assay. CCK-8 assay (Beyotime Institute of Biotechnology) was performed to measure the

viability of the HTR8/SVneo cells. Briefly, at 48 h after transfection, the cells were seeded into 96-well plates at 2×10^3 cells/well. A total of 200 μ l RPMI-1640 complete medium containing 10% FBS and 100 U/ml penicillin and streptomycin was added and the cells were then incubated at 37°C under 5% CO₂. Following cell adhesion, at 0, 24, 48 and 72 h, the cells were incubated with 20 μ l CCK-8 reaction solution for 30 min. The optical density values at 490 nm were then obtained for each well using a microplate reader, which were used to produce a cell viability curve.

Flow cytometry. Flow cytometry was performed to measure HTR8/SVneo cell proliferation and apoptosis. For cell proliferation, the cells were collected and washed with PBS. Following fixation with 4% formaldehyde for 10 min at room temperature, the cells were centrifuged at 1,000 \times g for 5 min at room temperature. The cells were treated with 0.5% Triton X-100 at room temperature for 5 min and then incubated with BD Cytotfix/Cytoperm™ Plus reagent (cat. no. 555028; BD Biosciences) at room temperature in the dark for 15 min. Fluorescence was detected and the percentage of Ki-67 calculated from it using the BD FACSVerse™ flow cytometer (BD Biosciences) and the FlowJo™ VX10 software (FlowJo LLC).

For apoptosis detection, 48 h after transfection, 1×10^6 HTR8/SVneo cells were collected and cultured with a shaker for 2 h at a speed of 200 rpm. After washing twice with ice-cold PBS, the cells were labeled using the BD Pharmingen™ FITC Annexin V Apoptosis Detection Kit I (cat. no. 556547; BD Biosciences), according to the manufacturer's protocol. Flow cytometry was conducted to detect the fluorescence, from which apoptosis rate was calculated.

For Ki-67 detection, the cells were collected and treated with BD Cytotfix/Cytoperm™ Plus reagent (cat. no. 555028; BD Biosciences) according to the manufacturer's protocols. The cells were then incubated with PE-Cy™ 7-conjugated mouse anti-Ki-67 antibody (cat. no. 561283; BD Biosciences) in the dark at room temperature for 15 min, followed by detection of the fluorescence with flow cytometry.

Transwell chamber assay. The infiltration ability of the cells was detected using Transwell chamber assays. Matrigel® solution was diluted in serum-free RPMI-1640 medium (v:v, 1:3), evenly smeared onto the upper chamber and incubated at 37°C for 60 min. The HTR8/SVneo cells were then seeded onto the upper chamber at a density of 1×10^5 cells/well and cultured in 200 μ l serum-free RPMI-1640 medium while 500 μ l RPMI-1640 medium containing 20% FBS was present in the lower chamber. After 24 h, the cells were fixed with 4% formaldehyde at room temperature for 10 min and subjected to Giemsa staining at room temperature for 2 min. Following rinsing for 2 min, the cells were air-dried. HTR8/SVneo cells that infiltrated to the lower chamber were observed and counted under a light microscope, from a total of five random fields of view under high magnification (x200). For the measurements of cell migration, identical procedures were performed as described aforementioned using Transwell chambers that were not coated with Matrigel.

Confocal laser scanning microscopy (CLSM). The cytoskeleton was imaged using CLSM. After transfection,

when 80% confluence was reached, HTR8/SVneo cells were fixed with 4% formaldehyde at room temperature for 10 min. After washing with PBS, the cells were permeabilized using 0.5% Triton X-100 at room temperature for 5 min and incubated with 200 μ l 100 nmol/l Rhodamine Phalloidin at room temperature in the dark for 30 min. The cytoskeleton was then observed using a confocal microscope (no. of fields taken, 5; magnification, x400; model SP8; Leica Microsystems GmbH).

Western blot analysis. At 48 h after transfection, the cells were lysed using RIPA buffer (Beyotime Institute of Biotechnology) supplemented with PMSF (100 mM) on ice. The protein concentration was determined using Bicinchoninic Acid assay (Beyotime Institute of Biotechnology). Then, 20 μ g sample was separated by 10% SDS-PAGE, and then transferred onto PVDF membranes. After blocking with 50 g/l fat-free milk diluted in 0.1% TBST at room temperature for 1 h, the membranes were incubated with rabbit anti-human anti-CLDN3 (1:1,000 dilution; cat. no. 83609), rabbit anti-human anti-matrix metalloproteinase (MMP)-2 (1:1,000 dilution; cat. no. 40994), rabbit anti-human anti-MMP-9 (1:1,000 dilution; cat. no. 13667), rabbit anti-human anti-ERK1/2 (1:1,000 dilution; cat. no. 5013), mouse anti-human anti-p-ERK1/2 (1:1,000 dilution; cat. no. 9106) or rabbit anti-human anti-GAPDH (1:5,000 dilution; cat. no. 5174) primary antibodies (all from Cell Signaling Technology, Inc.) at 4°C overnight. After washing with TBST, the membranes were incubated with horseradish peroxidase-conjugated goat anti-rabbit (cat. no. A0208) or goat anti-mouse (cat. no. A0216) secondary antibody (both 1:4,000 dilution; Beyotime Institute of Biotechnology) at room temperature for 1 h. Protein bands were visualized using the ECL method (Pierce; Thermo Fisher Scientific, Inc.) and analyzed using the Quantity one V4.6.7 software (Bio-Rad Laboratories, Inc.).

Statistical analysis. Data are presented as the mean \pm SD. SPSS 20.0 software (IBM Corp.) was used for statistical analysis. Pairwise comparisons were performed using the Student's t-test, whereas one-way ANOVA followed by Dunnett's test was used for multiple group comparisons. $P < 0.05$ was considered to indicate a statistically significant difference.

Results

Expression of CLDN3 in placental tissues from patients with pregnancy-induced hypertension. To investigate the expression of CLDN3 in the placenta of the pregnant patients with and without hypertension, RT-qPCR was performed. Compared with the healthy control group, the expression levels of CLDN3 mRNA in the placental tissue were significantly reduced in the pregnant patients with hypertension ($P < 0.01$; Fig. 1A). In addition, relative CLDN3 mRNA expression levels in the placental tissues from the severe pre-eclampsia, mild pre-eclampsia and pregnancy-induced hypertension groups were significantly reduced compared with the healthy control group ($P < 0.05$; Fig. 1A). In the peripheral blood, compared with the control group, the relative CLDN3 mRNA expression levels in the pregnancy-induced hypertension, mild pre-eclampsia and severe pre-eclampsia groups were significantly lower ($P < 0.05$; Fig. 1B). These results suggest that at the onset of pregnancy-induced hypertension, CLDN3 expression

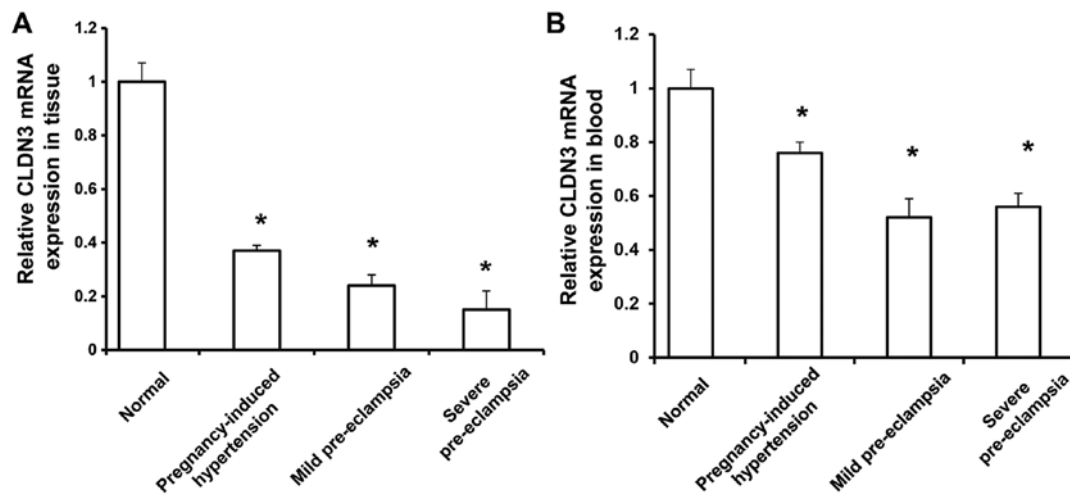


Figure 1. Expression levels of CLDN3 mRNA in the placental tissues and peripheral blood of patients. The mRNA expression levels of CLDN3 in (A) placental tissues and (B) peripheral blood of normal control, pregnancy-induced hypertension, mild pre-eclampsia and severe mild pre-eclampsia groups were measured using reverse transcription quantitative PCR. Experiments were performed in triplicates. * $P < 0.05$ vs. normal. CLDN3, claudin 3.

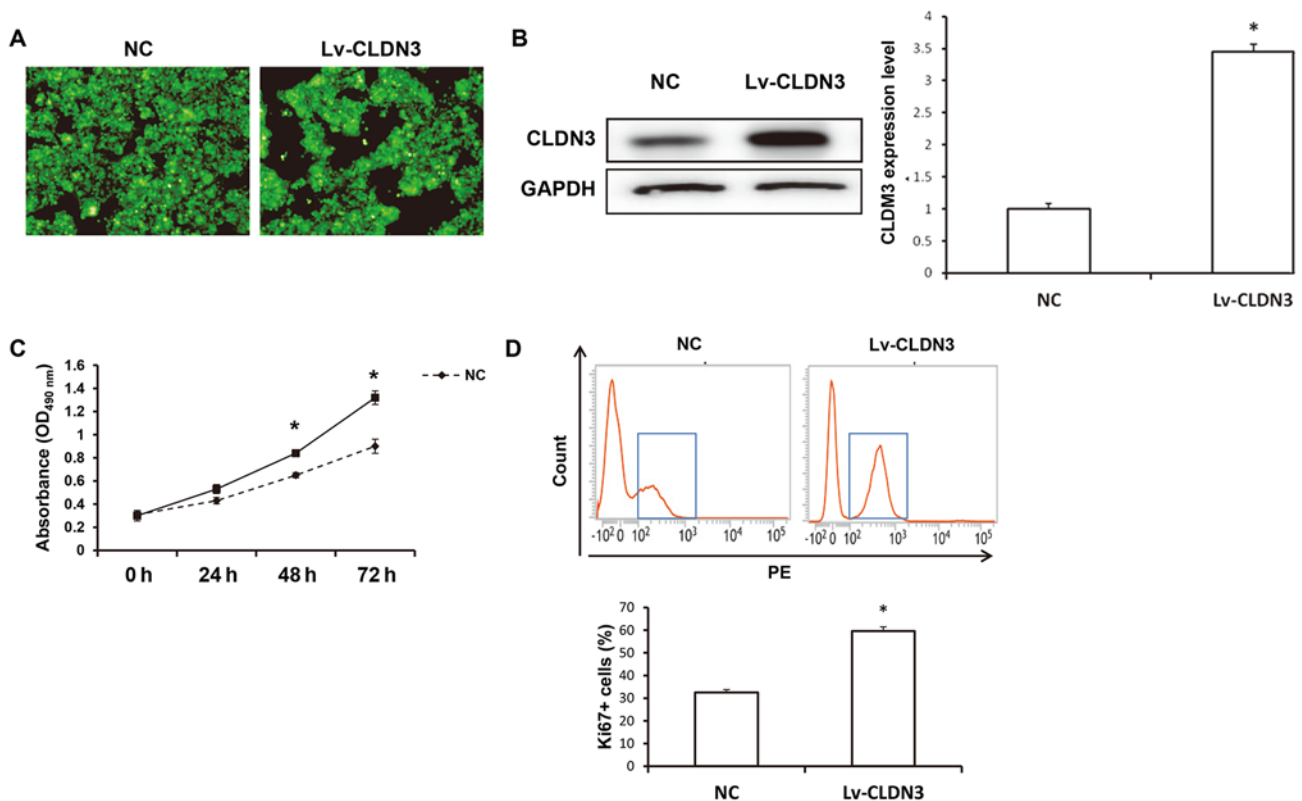


Figure 2. Effects of CLDN3 overexpression on the proliferation of HTR8/SVneo cells. (A) After transfection with Lv-GFP-CLDN3, GFP fluorescence was detected using confocal laser scanning microscopy (magnification, x200). (B) CLDN3 overexpression in HTR8/SVneo cells was verified using western blot analysis. (C and D) Following transfection, cell viability was determined using (C) Cell Counting kit-8 assay and (D) proliferation was assessed via the quantification of Ki-67 using flow cytometry. Experiments were performed in triplicates. * $P < 0.05$ vs. NC. CLDN3, claudin 3; GFP, green fluorescent protein; NC, cells transfected with empty vector; OD, optical density; SSC, side scatter; PE, phycoerythrin.

was reduced in placental tissues, and may be associated with disease pathogenesis.

CLDN3 overexpression enhances of HTR8/SVneo cell viability. The effects of CLDN3 on the viability of HTR8/SVneo cells were next investigated. At 48 h after transfection with Lv-GFP-CLDN3, GFP fluorescence was distributed evenly in

the cytoplasm of the HTR8/SVneo cells (Fig. 2A). Western blot analysis confirmed that CLDN3 expression was elevated in HTR8/SVneo cells after transfection with Lv-GFP-CLDN3 compared with those in the empty vector-transfected control ($P < 0.05$; Fig. 2B). The viability of the transfected cells was measured using a CCK-8 assay. Compared with the control group, CLDN3 overexpression significantly increased

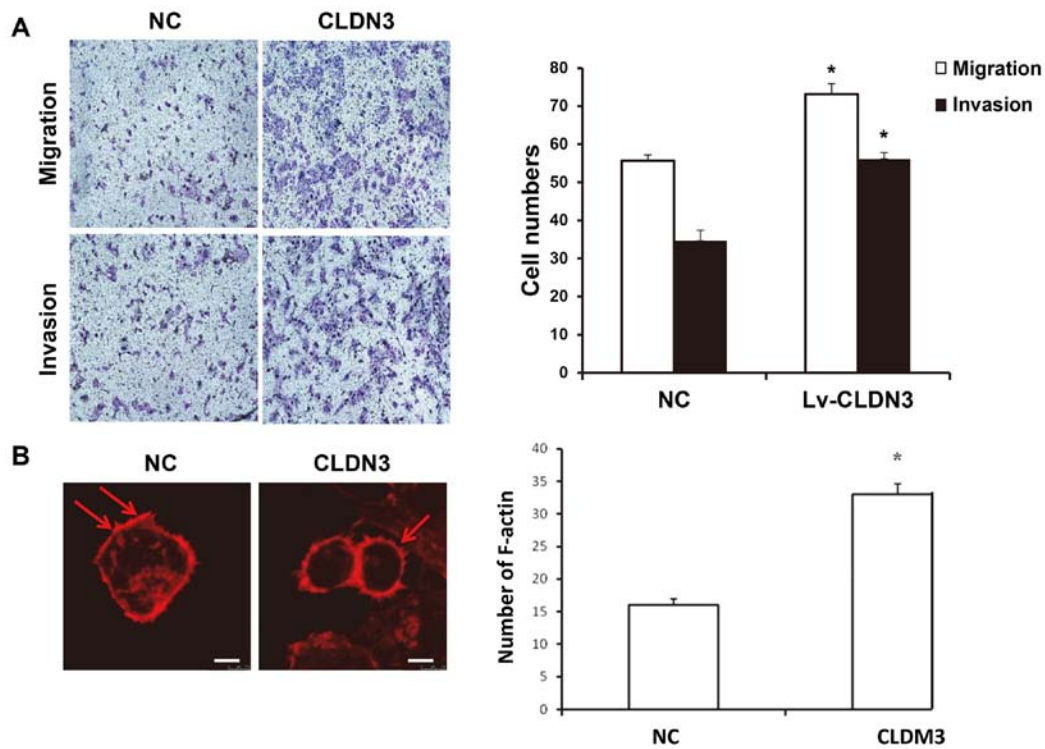


Figure 3. Effects of CLDN3 overexpression on HTR8/SVneo cell invasion and migration. (A) The invasive and migratory capabilities of transfected HTR8/SVneo cells were measured using Transwell chamber assays. (B) The F-actin cytoskeleton (red fluorescence) was imaged using confocal laser scanning microscopy. Scale bar, 2 μ m. * P <0.05 vs. NC. CLDN3, claudin 3; NC, cells transfected with empty vector.

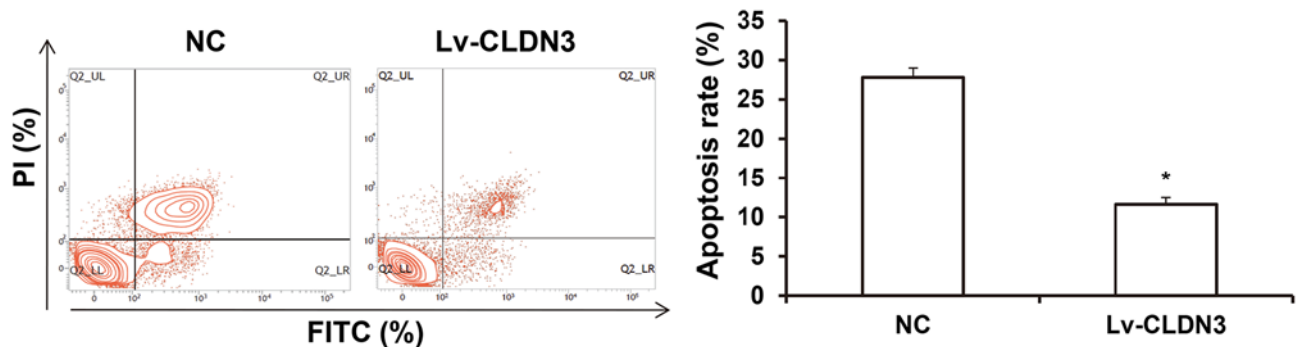


Figure 4. Effects of CLDN3 overexpression on the apoptosis of HTR/SVneo cells. The apoptosis of transfected HTR/SVneo cells was detected using flow cytometry and the apoptotic rates were calculated accordingly. Experiments were performed in triplicates. * P <0.05 vs. NC. CLDN3, claudin 3; FITC, fluorescein isothiocyanate; PI, propidium iodide. NC, cells transfected with empty vector.

HTR8/SVneo cell viability at 24, 48 and 72 h after transfection (P <0.05; Fig. 2C). Consistent results were observed using flow cytometry, which indicated that the overexpression of CLDN3 significantly upregulated Ki-67 expression in HTR8/SVneo cells (P <0.05; Fig. 2D). These results indicate that CLDN3 overexpression significantly promoted HTR8/SVneo cell proliferation.

CLDN3 overexpression promotes the invasive and migratory abilities of HTR8/SVneo cells. To investigate the effect of CLDN3 on the invasive and migratory abilities of HTR8/SVneo cells, Transwell chamber assays were performed. Compared with the control group, CLDN3 overexpression significantly promoted the migratory and invasive capabilities of HTR8/SVneo cells (P <0.05; Fig. 3A). In addition, CLSM

results showed that in HTR8/SVneo cells overexpressing CLDN3, the number of F-actin myofibers was significantly increased, indicating increased migratory ability (P <0.05; Fig. 3B). These results suggest that CLDN3 overexpression promotes HTR8/SVneo cell migration and invasion.

Effects of CLDN3 overexpression on HTR8/SVneo cell apoptosis. Apoptosis is also an important factor affecting the migratory and invasive abilities of HTR8/SVneo cells (29). To investigate the effects of CLDN3 on apoptosis in HTR8/SVneo cells, flow cytometry was performed. Compared with the control group, the overexpression of CLDN3 significantly reduced HTR8/SVneo cell apoptosis (P <0.05; Fig. 4). These results suggest that CLDN3 overexpression can suppress apoptosis in HTR8/SVneo cells.

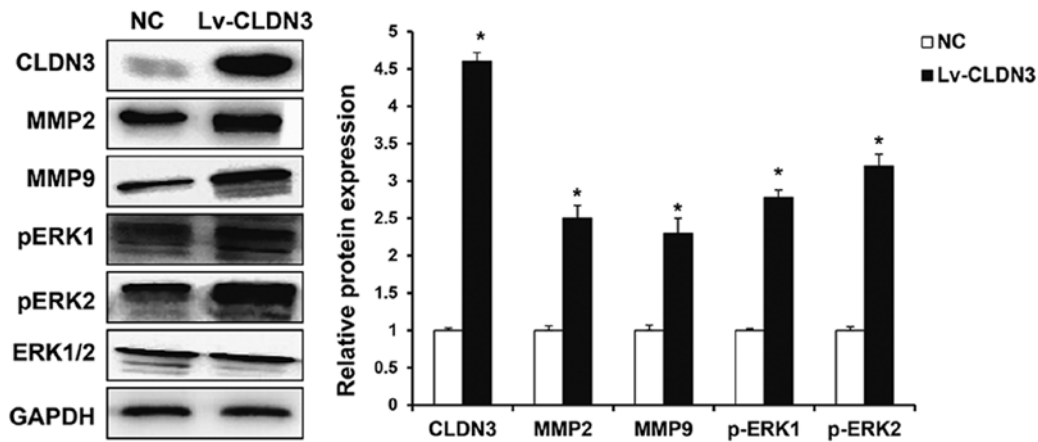


Figure 5. Effects of CLDN3 on MMP expression and ERK phosphorylation in HTR8/SVneo cells. Following transfection, the expression levels of MMP-2 and MMP-9 proteins, in addition to ERK1/2 phosphorylation were detected using western blot analysis. Experiments were performed in triplicates. * $P < 0.05$ vs. NC. CLDN3, claudin 3; MMP, matrix metalloproteinase; NC, cells transfected with empty vector.

Effects of CLDN3 on the ERK1/2 signaling pathway and MMP expression in HTR8/SVneo cells. The ERK1/2/MMP pathway serves an important role in the cell migration and is regulated by tight junction proteins (6). To investigate the effects of CLDN3 on the ERK1/2 signaling pathway (30) in addition to MMP-2 and MMP-9 expression in HTR8/SVneo cells, western blot analysis was performed following CLDN3 overexpression. The overexpression of CLDN3 significantly elevated the expression levels of MMP-2 and MMP-9 in HTR8/SVneo cells. ERK1/2 phosphorylation was also significantly increased following CLDN3 overexpression (Fig. 5). These results suggest that CLDN3 can significantly affect the expression of MMPs and ERK activation in HTR8/SVneo cells.

Discussion

Pregnancy-induced hypertension is a common complication in obstetrics (31). Clinical manifestations of pregnancy-induced hypertension include proteinuria and fetal growth restriction after 20 weeks of gestation (32). At present, the pathogenesis of this disease remains poorly characterized (33). A previous study has demonstrated that in most women with pregnancy-induced hypertension, the symptoms of hypertension typically disappear following the delivery of placenta, suggesting that the placenta may serve an important role in the development of pregnancy-induced hypertension (34). Trophoblast cells are important for the structure and function of placenta, where the tight junction proteins serve important roles in maintaining the cellular polarity and integrity of cell barriers. However, whether or not the tight junction proteins regulate the biological function of trophoblast cells remains unclear. The present study demonstrated that the expression levels of the tight junction protein CLDN3 were significantly downregulated in the placental tissues of patients with pregnancy-induced hypertension compared with healthy pregnant controls. Supporting this, *in vitro* experiments using human trophoblast cell line HTR8/SVneo cells showed that CLDN3 overexpression promoted cell proliferation and invasion, possibly by activating the ERK1/2/MMP-2/MMP-9 signaling pathway.

The placenta is the location where gas and nutrient exchange occurs and where metabolic products are eliminated between the mother and the fetus. As part of the main placental structure, invasion of trophoblast cells into the endometrium is an important physiological process in the formation of the placenta (35,36). Studies have shown that extravillous trophoblast cells invade into the uterine decidua and myometrial spiral artery, where they gradually replace the endothelial cell layer and part of the smooth muscle tissue in the maternal spiral arteries, resulting in thickened blood vessels to provide sufficient blood supply to the fetus (37,38). Trophoblast cell invasion is a key factor to this process, and factors hindering this physiological process may lead to placental dysplasia, pregnancy-induced hypertension and eclampsia (39). In particular, one of the pathological changes in the placenta during pre-eclampsia is the shallow and limited invasion of trophoblast cells into the uterine spiral artery (40).

Tight junction proteins, of which the CLDN family of proteins is an important example, serve important roles in the maintenance of cell barriers and polarity. In recent years, a large number of studies have shown that CLDNs are closely associated with tumor cell invasion and metastasis (41). Indeed, CLDN2 has been reported to promote the self-renewal of colon cancer stem cells (42), whereas the downregulation of CLDN12 expression by interleukin (IL)-18 has been demonstrated to activate the p38 pathway to promote the invasion and metastasis of breast cancer in another recent study (43). CLDN3 is an important member of the CLDN family. Studies have confirmed the involvement of CLDN3 in the regulation of tumor cell invasion and metastasis through multiple signaling pathways. Ahmad *et al* (44) showed that the loss of CLDN3 expression activated the IL-6/glycoprotein 130/STAT3 pathway to promote the malignant transformation of colon cancer. In lung squamous cell carcinoma, CLDN3 could inhibit the invasion and metastasis of lung cancer cells (45). However, it remains to be elucidated whether CLDN3 is involved in the regulation of human trophoblast cell invasion. Results from the present study showed that the mRNA expression levels of CLDN3 in the placental tissues and peripheral blood of patients with pregnancy-induced hypertension were significantly reduced compared with those in healthy controls,

indicating a negative association with disease pathogenesis. The *in vitro* experiments of the present study indicated that the overexpression of CLDN3 promoted HTR8/SVneo cell proliferation, migration and invasion. During invasion, human trophoblast cells would detach from adjacent cells or the extracellular matrix. Normal cells undergo apoptosis to maintain the stability of the tissue environment, whereas tumor cells or other migratory cells may suppress apoptosis to promote their migration. In the present study, CLDN3 overexpression significantly inhibited the apoptosis of HTR8/SVneo cells. These observations suggest that CLDN3 promotes the proliferation, in addition to the invasive and migratory abilities of HTR8/SVneo cells *in vitro*. The present study demonstrated changes in CLDN3 expression in human tissues and cells, and also detected similar changes in the levels of CLDN3 in the peripheral blood. From these observations it could be hypothesized that the placental tissue released CLDN3 into the circulation; however, there is no direct evidence that the placenta is the sole source of CLDN3.

It has previously been found that human trophoblast cells secrete MMPs to degrade the extracellular matrix during invasion (46). MMP-2 and MMP-9 are the main enzymes involved in the degradation of the basal membrane of the uterus (47). It has been reported that MMP-2 and MMP-9 could assist in the invasion of human trophoblast cells, and MMP-2 and MMP-9 can degrade extracellular matrix, making cells more susceptible to metastasis and infiltration (48,49). In the present study, results from western blot analysis showed that CLDN3 overexpression significantly elevated the expression levels of MMP-2 and MMP-9 in human trophoblast cells. Regulation of MMPs involves a number of different factors, including cytokine profiles, signaling pathway activation and MMP-inhibiting factors. A previous study found that both the expression and activity of MMP-9 are regulated by the ERK1/2 signaling pathway (50). Indeed, the present study showed that the overexpression of CLDN3 significantly elevated the phosphorylation levels of ERK1 and ERK2 in HTR8/SVneo cells, suggesting that CLDN3 activates the ERK1/2 signaling pathway.

However, in the present study, although the expression levels of CLDN3 in the placental tissue and the effects on the biological function of trophoblast cells were investigated, only a single trophoblast cell line was used, and further in-depth studies on primary trophoblast cells are required to confirm the regulatory effects of CLDN3.

In conclusion, the present study showed that CLDN3 can promote human trophoblast cell proliferation, migration and invasion, with the underlying mechanism possibly involving the upregulation of MMP-2 and MMP-9 expression levels via the ERK1/2 signaling pathway. These findings suggest that downregulation of CLDN3 may be associated with the pathogenesis of pregnancy-induced hypertension, which may serve to assist in the design of therapeutic interventions for the treatment of this disease in the clinic.

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Availability of data and materials

All data generated or analyzed during this study are included in this published article.

Authors' contributions

AZ, YQ and KL contributed to the study design, experiment performance, data collection and analysis, and manuscript preparation. All authors read and approved the final manuscript.

Ethics approval and consent to participate

The study was approved by the ethics review board of Laiwu Maternal and Child Health Hospital (Shandong, China). Prior written informed consent was obtained from every patient.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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