miRNA-34a-5p downregulation of VEGFA in endometrial stem cells contributes to the pathogenesis of endometriosis

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Abstract. Endometrial-derived stem cells (EnSCs) serve an important role in the development of endometriosis via retrograde menstruation. Abnormal expression of miRNAs in EnSCs is involved in the etiology of endometriosis, however, the mechanisms remain unclear. The aim of the present study was to investigate the expression of miR-34a-5p and VEGFA in endometrial samples from patients with or without endometriosis, and then examine the underlying mechanism of microRNA-34a-5p regulation of VEGFA in EnSCs. Endometrial samples from patients with or without endometriosis were collected, and miR-34a-5p expression in the two groups was measured using RT-PCR. Human endometrial-derived stem cells (hEnSCs) were isolated from these endometrial samples, and hEnSCs were transfected with the miR-34a-5p mimics or control miRNAs. qPCR and western blotting were performed to assess the effects of miR-34a-5p on the expression of VEGFA in hEnSCs, and cell growth was assessed by an MTT assay. miR-34a-5p was significantly downregulated in patients with endometriosis when compared with that of those without endometriosis. VEGFA expression levels in hEnSCs with an overexpression of miR-34a-5p were significantly reduced when compared with those in the negative control (P<0.01). In addition, the upregulation of miR-34a-5p suppressed EnSCs proliferation by targeting the 3' untranslated region of VEGFA. miR-34a-5p provides a novel avenue for the understanding of the development of endometriosis, and may facilitate the development of potential therapeutics against endometriosis.

Introduction

Endometriosis is a common, and multifactorial gynaecological disorder that affects 10-15% of reproductive-age women (1). The development of this condition is endometrial tissue grows outside of the uterine cavity, which can induce varying degrees of painful symptoms and infertility in infected individuals (2). Endometriosis also has a propensity to recur and may be associated with ovarian cancer (3). The impact of endometriosis is great, likely exceeding our expectations, as it can dramatically impair the quality of life.

The pathogenesis of endometriosis was initially defined as the presence of endometrial tissues in extra-uterine cavity, including pelvic peritoneum, bladder and ureters (4). There are several hypotheses have been proposed, including lymphatic and vascular metastasis, iatrogenic direct implantation, coelomic metaplasia, embryonic rest and mesenchymal cell induction (5). Moreover, the development of endometriosis was also involved in stem cells (6). This theory postulates that the endometrial stem cells are abnormally shed during menses, where they gain access to the peritoneal cavity by retrograde menstruation and ectopic implantation. The presence of mesenchymal stem cells (MSCs) in stromal cells of ectopic endometrial tissues was reported (7). These results indicated that ectopic endometrial stem cells (EnSCs) are regarded as responsible for the pathogenesis of endometriosis.

Endometriosis has also been defined as an angiogenic disease, since the ectopic survival of endometrium requires the formation of new blood vessels. Angiogenesis, therefore, is a critical step in developing endometriotic lesions (8). Vascular endothelial growth factor (VEGF) is a member of VEGF family, coding by a 28 kb-long gene which is located on chromosome 6p21.3. It is known as a crucial regulator of angiogenesis, endothelial cells growth and migration (9). Several studies have reported that VEGF play an important role in the angiogenesis of endometriosis (10). miRNAs such as miR-126, Let7-f, miR-27b, miR-17-92 cluster and miR-130a were identified as proangiogenic miRNAs that regulate the translation of angiogenic factors (VEGF-A) (11). miRNAs have recently emerged as an important factor in endometriosis (12). Toloubeydokhti et al (13) assessed the expression of miR-17-5p, miR-23a, miR-23b and miR-542-3p, and found that miR-23b and miR-542-3p are downregulated, whereas miR-17-5p is upregulated in ectopic endometrium, affecting the stability of their target genes’ expression, and playing an important role in the pathogenesis of endometriosis. Moreover, miRNAs are aberrantly expressed in endometriotic stromal cells play an important role in the pathogenesis of.
endometriosis (14). miR-34a-5p is highly expressed in multiple types of cancer, which inhibited tumor angiogenesis by blocking VEGF production by directly inhibiting endothelial cell functions (15).

In this study, the aim was to investigate the expression of miR-34a-5p and VEGFA in endometriosis tissues, and analyzed the function and mechanism of miR-34a on the endometriosis stem cells (EnSCs).

**Materials and methods**

**Endometriosis tissues.** Ten endometriosis patients were diagnosed by laparoscopic surgical examination, and undergone surgical excision of endometriosis tissues. The control tissue samples were collected from 10 premenopausal patients without endometriosis. All these patients had not received any preoperative hormonal therapy or taken any medicine for at least three months. The resected tissues were minced and part of samples was stored at -80°C before the total RNA and protein extraction. Written consent was obtained from each patient before the study. This study was approved by the Research Ethics Committee of the Zhujiang Hospital of Southern Medical University (Guangzhou, China).

**Endometrial stem cell isolation, culture and transfection.** Human endometriosis tissue was washed in PBS (Gibco, USA), minced, and digested in collagenase (1 mg/ml, Gibco, USA) for 30-45 min at 37°C with agitation. Then, resultant cell solutions were filtered and centrifuged, and mononuclear cells were separated by Ficoll (Sigma, St. Louis, MO, USA) and washed in PBS. The isolated cells were cultured in Dulbecco's modified Eagle's medium (DMEM)/F-12 medium containing 10% fetal bovine serum (FBS) and 1% penicillin/streptomycin antibiotic (all Gibco, Grand Island, NY, USA) and then incubated at 37°C in 5% CO₂. When the cells were approximately 80% confluent, they were trypsinized (Gibco) and halved for characterization by flow cytometry and expansion. EnSCs were seeded into a 24-well plate and transfected with the miR-34a-5p mimics by Lipofectamine™ 2000 Transfection Reagent (Invitrogen, Carlsbad, CA, USA).

**Flow cytometry.** The EnSCs were characterized by flow cytometry for cell surface markers. Cells were incubated with the specific antibody at concentrations recommended by the respective manufacturers for 1 h and analyzed by flow cytometry. The antibodies used were: FITC-conjugated anti-CD146 (eBioScience Inc., San Diego, CA, USA; endometrial stem cell markers), FITC-conjugated anti-CD140b (Abcam, Cambridge, MA, USA; endometrial stem cell markers), FITC-conjugated anti-CD34 (Abcam; hematopoietic marker), FITC-conjugated anti-CD31 (Novus Biologicals, Littleton, CO, USA; endothelial marker).

**RNA extraction and real-time PCR.** To investigate the expression level of VEGFA mRNA and miR-34a-5p, we extracted the total RNA from endometriosis tissues or EnSCs. Tissues or cells were lysed with the Trizol reagent (Life Technologies, Grand Island, NY, USA) according to the manufacturer's specification. mRNAs were dissolved in RNase-free water and stored at -80°C before utilize. The quantitative analysis of RNA was performed by RT-PCR using One Step SYBR PrimeScript PLUS RT-PCT kit (Takara, Tokyo, Japan) for each sample according to the manufacturer's manual. After the reaction, the VEGFA mRNA level and the miR-34a-5p level was calculated and presented as the relative level of VEGFA to β-actin (as control) by ∆∆Ct method, each sample was measured for three independent experiments.

**Western blot analysis.** Western blot analysis was performed to determine the expression of VEGFA on the protein level. Harvested the EnSCs post transfecting, and lysed the cells into cell lysis buffer (Bio-Rad, Hercules, CA, USA). Centrifuged the samples at 12,000 g for 15 min at 4°C and collected the supernatant. Protein extracts were boiled in SDS/β-mercaptoethanol and separated in a 10% SDS-PAGE (sodium dodecyl sulphate-polyacrylamide gel electrophoresis). Then, transferred the protein sample to a nitrocellulose membrane (Millipore, Bedford, MA, USA), blocked with 5% skimmed milk powder overnight at 4°C. The membrane was incubated with VEGFA-spe-cific antibody in TBST (mouse monoclonal antibody, Abcam, Cambridge, UK, 1:500) at 37°C for 1 h, washed with TBST, next incubated with the HRP-linked secondary anti-mouse antibody (New England Biolabs, Ipswich, UK) for 30 min at 37°C. ECL kit (Life Science, Woodland Hills, CA, USA) was used to carry out the chemiluminescence reaction. The membrane was scanned by a Smart Chemi-TM larm Analysis System (Thermo Scientific, Rockford, IL, USA) and quantified according to the band density by Quantity One software with β-actin as loading control.

**Dual luciferase assay.** The sequence of 3'UTR of VEGFA from hs-miR-34a-5p were download from Genebank (NC-BI), aligned by Megalign (DNASTAR; GATC Biotech, Konstanz, Germany). The sequence of mutant 3'UTR of VEGFA was synthesized by Sangon Biotech (Shanghai, China). All the sequences in this study were amplified by PCR (polymerase chain reaction) with Phusion polymerase (New England Biolabs). 3'UTR of VEGFA and the mutant 3'UTR of VEGFA were cloned into the upstream of firefly luciferase (Fluc) gene behind the Cytomegalovirus promoter in pmirGLO (Promega, Madison, WI, USA) with restriction endonuclease and DNA ligase (New England Biolabs), received VEGFA-Fluc and mutant-VEGFA-Fluc recombinant plasmid. The ESCs seeded in a 24-well plate were co-transfected with the miR-34a-5p mimics/miR-Ctrl and VEGFA-Fluc/mutant-VEGFA-Fluc by Lipofectamine™ 2000 Transfection Reagent (Invitrogen). 48 h after transfecting, collected the cells and assayed with the Dual-Luciferase Assay kit (Promega) used GLOMAX (Promega) and recorded data, the relative luciferase activity (%) represented the expression level of VEGFA.

**MTT assay.** Transfected EnSCs were diluted to a certain concentration and then were placed over a 96-well plate, at a density of 5x10⁴ cells/well. Each well was inoculated with 100 µl cell suspension, apart from one blank well (instead with 100 µl medium containing 10% fetal bovine serum). Then they were cultured in a 5% CO₂ incubator at 37°C. After 0, 12, 24 and 48 h of culture, the proliferation of
ESCs was detected by MTT assay. With an addition of 20 \( \mu l \) 3-(4,5-Dimethylthiazol-2-yl)-5-(3-Carboxymethoxyphenyl)-2-(4-Sulfophenyl)-2H-Tetrazolium, Inner Salt (MTS) reagent (Promega) to each well, cells were then cultured in an incubator for 1-4 h. After that, the optical density (OD) at 580 nm of each well was obtained with a microplate reader (Biotek Instruments, Inc., Vermont, USA). The cell proliferation was evaluated by OD value. All of these experiments were repeated three times.

**Statistical analysis.** All the analyses were performed using SPSS software, version 17.0 (SPSS Inc., Chicago, IL, USA). SPSS was used to compare miR-34a-5p expression levels between samples taken from patients with or without endometriosis. Data are represented as mean ± standard deviation (SD) unless indicated otherwise. Statistical significance was tested using Student’s t-test, and \( P<0.05 \) was considered to indicate a statistically significant difference.

**Results**

**Characterization of isolated human EnSCs.** EnSCs isolated from endometriosis tissue were positive for CD140b, CD146 and negative for CD31 and CD34, as demonstrated by flow cytometry analysis (Fig. 1).

**Expression of miR-34a-5p and VEGFA in endometriosis.** To investigate the expression of VEGFA in endometriosis, we examined the mRNA level of VEGFA and miR-34a-5p level by quantitative RT-PCR in 10 eutopic endometrial tissues and 10 ectopic endometrial tissues. We focused on miR-34a-5p because it was significantly downregulated miRNA in ectopic endometrial tissues by microarray analysis. It had shown that the relative mRNA level of VEGFA was significantly upregulated in ectopic endometrial tissues than in eutopic endometrial tissues (\( P<0.01 \)) (Fig. 2A). The relative level of miR-34a-5p was significantly lower in endometriosis tissues than that in eutopic endometrial tissues (\( P<0.01 \)) (Fig. 2B). By analyzing the correlation of relative VEGFA mRNA level with the miR-34a-5p level in ectopic endometrial tissues, we found an inverse correlation between them.

The high mRNA level of VEGFA accompanied a low expression level of miR-34a-5p (Pearson correlation, \( R^2=0.4648, P=0.0299 \) (Fig. 2C).

**Expression of VEGFA in EnSCs after transfection with miR-34a-5p mimics.** To further determined whether the expression of VEGFA was downregulated by miR-34a-5p in EnSCs. We examined the relative level of miR-34a-5p and the expression of VEGFA in both mRNA and protein levels after 48 h transfection. The level of miR-34a-5p was significantly increased after the transfection with miR-34a-5p mimics (\( P<0.01 \)), compared with the miR-control group. However, the VEGFA mRNA was reduced post the transfection with miR-34a-5p mimics (\( P<0.01 \)). The expression level of VEGFA protein was decreased in EnSCs transected with miR-34a-5p mimics rather than miR-control group. The band intensity of VEGFA was downregulated in EnSCs transected with miR-34a-5p mimics (\( P<0.01 \)). Therefore, miR-34a-5p suppressed the expression of VEGFA in both mRNA and protein levels in ESCs (Fig. 3).

**miR-34a-5p targets the 3’UTR of VEGFA gene.** To verify miR-34a-5p inhibited the expression of VEGFA by targeting the 3’UTR of VEGFA, miR-34a-5p mimics and miR-control were transfected into EnSCs which already had been transfectected with the 3’UTR of VEGFA linked a luciferase reporter. The results showed that the relative luciferase activity was significantly decreased post transfecting miR-34a-5p mimics (\( P<0.001 \)). However, there was no significant difference in the relative luciferase activity between miR-34a-5p mimics and miR-control in the EnSCs, which were transfected with the mutant 3’UTR of VEGFA. All these data demonstrates that miR-34a-5p targets the 3’UTR of VEGFA gene and inhibits the VEGFA expression effectively (Fig. 4).

**Effect of miR-34a-5p mimics on the proliferation of EnSCs.** miR-34a-5p could modulate the level of VEGFA, which is a regulator of cell growth, so miR-34a-5p may also influence proliferation of EnSCs. Then, we curved the ESCs number at 0, 12, 24 and 48 h after transfecting with miR-control and miR-34a-5p mimics. Fig. 5 showed that the proliferation ability...
of EnSCs was significantly reduced post the transfection with miR-34a-5p mimics (P<0.01).

**Discussion**

The present study is the first to investigate the expression of miR-34a-5p and VEGFA in endometriosis. The results showed that miR-34a-5p expression levels significantly decreased in endometriosis compared with that in without endometriosis patients, but the VEGFA expression levels significantly increased in endometriosis compared with without endometriosis patients. Negative correlation was observed between relative VEGFA mRNA level and miR-34a-5p level in ectopic endometrial tissues. Moreover, over-expression of miR-34a-5p...
miR-34a belongs to a miRNA family, and its as a strong antitumor regulator of cell growth in multiple myeloma was observed (16, 17). miR-34a is, to date, one of the most characterized tumor suppressor miRNAs in a variety of tumors (18). miR-34a-5p expression has been shown to be directly regulated by p53 and antagonizes processes, including cervical, ovarian and testicular cancer (19, 20). Ma et al conducted a series of studies on miR-34a-5p, and were the first to confirm its pro-apoptotic function (21). Moreover, miR-34a-5p has been found to inhibit cell proliferation and invasion in vitro, which suggested that miR-34a-5p might play a role in inhibiting tumor recurrence (22). miR-34a-5p has been reported to be a direct transcriptional target of p53 and is downregulated in various types of tumors (23). We measured miR-34a-5p expression levels in endometrial tissue and ESCs. Regardless of the sample types, miR-34a-5p expression levels were consistently downregulated in patients with endometriosis. The results support previous reports that miR-34a-5p suppresses disease progression, including diseases associated with the endometrium.

Angiogenesis is a crucial determinant in tumor initiation, progression, and metastasis. Angiogenesis plays an important role in endometriosis progression (30). VEGF is thought to be the primary stimulator of angiogenesis, during the course of development and in a variety of pathological conditions. A large number of researches have been done showing that miRNAs play important roles in vascular development and angiogenesis (31). It is reported that miRNAs (eg miR-17-5p...
and miR-199a-5p) regulate angiogenesis (32). In addition, miR-203 suppresses tumor growth and angiogenesis by targeting VEGFA in cervical cancer (33). We found an inverse correlation between levels of VEGFA and miR-34a-5p. To our knowledge, this is the first demonstration that VEGFA is a functional target of miR-34a-5p. As such, miR-34a-5p regulates cell proliferation and angiogenesis through its effect on VEGFA.

In summary, VEGFA was overexpressed in endometriosis tissues, and we identified miR-34a-5p is able to target the VEGF gene, preventing the latter to function as an inhibitor of angiogenesis. These results imply that miR-34a-5p might regulate VEGFA in ESCs and might contribute to the pathogenesis of endometriosis. This study may provide a potential biomarker for endometriosis therapeutics.

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References