miR-494-3p promotes the progression of endometrial cancer by regulating the PTEN/PI3K/AKT pathway

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Abstract. MicroRNAs (miRs) are essential regulators in the development and progression of cancer. The role of miR-494-3p in endometrial cancer (EC) has not yet been investigated. In the present study, the expression levels of miR-494-3p were significantly upregulated in EC tissues compared with adjacent normal tissues. Furthermore, upregulation of miR-494-3p in patients with EC indicated poorer prognosis; miR-494-3p overexpression significantly promoted the proliferation, migration and invasion of HHUA and JEC cells in vitro. Consistently, inhibition of miR-494-3p in HHUA cells significantly suppressed tumor growth in vivo in a xenograft model. Additionally, phosphatase and tensin homolog (PTEN) was revealed to be a direct target of miR-494-3p in EC cells. Furthermore, overexpression of miR-494-3p inhibited PTEN expression and consequently activated the downstream phosphoinositide 3-kinase/protein kinase B (PI3K/AKT) signaling pathway. Restoration of PTEN or inhibition of PI3K/AKT pathway also abolished miR-494-3p-mediated proliferation, migration and invasion of HHUA and JEC cells. In summary, the results of the present study revealed the importance of the miR-494-3p/PTEN/PI3K/AKT axis in the progression of EC, which may provide novel insight into potential therapeutic targets for the treatment of EC.

Introduction

Endometrial cancer (EC) is the fourth most prevalent and aggressive type of gynecological cancer, which has become a great threat against women's health (1,2). Annually, >280,000 women are diagnosed with EC worldwide; EC results in numerous cancer-associated mortalities (3). Unremitting efforts have been made to develop effective methods for treating EC in the past decades. Radical surgery may be the only effective therapeutic method for early stage EC. At present, the incidence of EC is still rising due to tumor recurrence and metastasis (4,5). Therefore, investigating the molecular mechanisms of endometrial carcinogenesis, screening out effective diagnostic biomarkers and developing promising therapeutic targets are particularly important for the treatment of EC.

MicroRNAs (miRNAs/miRs) are a member of the family of noncoding RNAs and have a length of 21-23 nucleotides (6). Numerous reports have demonstrated that miRNAs could regulate gene expression by directly associating with the 3'-untranslated region of target mRNAs (7,8). By modulating the protein expression levels of target genes, miRNAs are widely involved in various biological processes and human diseases (9). In addition, accumulating evidence has indicated that miRNAs are essential regulators in a variety of cancers by serving as oncogenes or tumor suppressors (10). For instance, miR-199a/b-3p suppresses the proliferation of gastric cancer cells by regulating P21 activated kinase 4/mitogen activated protein kinase/extracellular signal-regulated kinase signaling pathway (11). Li et al (12) indicated that miR-34a directly targeted high-mobility group box 1 and inhibited the proliferation, migration and invasion of cancer cells in cutaneous squamous cell carcinoma. In EC, Zhao et al (13) revealed that miR-126 inhibited the migration and invasion of EC cells by targeting insulin receptor substrate 1. In summary, the aforementioned findings highlighted the importance of miRNAs in the progression of cancer, which suggests that miRNAs may be promising biomarkers and therapeutic targets in the treatment of cancers.

Recent studies have indicated that miR-494-3p could induce lung carcinogenesis and regulate the proliferation, invasion, migration and apoptosis of human glioblastoma cells (14,15). On the contrary, it has been suggested that miR-494-3p acts as a tumor suppressor in some cancers, such as prostate (16) and lung cancer (14). For example, Shen et al (16) have reported that miR-494-3p suppresses the proliferation, invasion, and migration of prostate cancer; however, the role of
miR-494-3p remains unknown in EC. In the present study, it was revealed that miR-494-3p was significantly upregulated in EC tissues and was associated with the prognosis of patients with EC. In addition, overexpression of miR-494-3p markedly promoted the proliferation, migration and invasion of EC cells. Furthermore, miR-494-3p directly targeted phosphatase and tensin homolog (PTEN) and consequently regulated phosphoinositide 3-kinase (PI3K)/protein kinase B (AKT) pathway in EC cells. Additionally, inhibition of the PI3K/AKT pathway abolished miR-494-3p-mediated effects on HHUA and JEC cells. In summary, the results of the present study revealed the pivotal role of the miR-494-3p/PTEN/PI3K/AKT pathway in EC progression, which may provide novel insight in the identification of therapeutic targets for the treatment of EC.

Materials and methods

Patient samples. The present study was approved by the Human Studies Committee at The Second Affiliated Hospital of Zhengzhou University (Zhengzhou, China). Informed consent was obtained from each patient prior to surgery. A total of 43 pairs of EC samples (all females; age, 54±16 years) were obtained from patients; Patients that were treated with chemotherapy or radiotherapy before surgery were excluded. Samples were histologically validated for type and grade. These samples were collected from May 2012 to July 2016.

Cell lines and cell culture. Human EC cell lines HHUA and JEC were obtained from the Cell Bank of Type Culture Collection of Chinese Academy of Science (Shanghai, China) and cultured in DMEM (Sigma-Aldrich; Merck KGaA, Darmstadt, Germany) supplemented with 15% of fetal bovine serum (FBS), 100 U/ml of penicillin and 100 µg/ml of streptomycin (all from Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA). Cells were incubated at 37°C in a humidified atmosphere of 5% CO2. To inhibit the PI3K/AKT signaling pathway, 50 µM LY294002 (cat. no. 9901; Cell Signaling Technology, Inc., Danvers, MA, USA) was added to the lower chamber at 37°C. Non-migratory cells were removed using a cotton swab following overnight incubation, while the migratory cells in the lower chamber were fixed with paraformaldehyde for 30 min at 25°C and stained with 0.1% crystal violet for 30 min at 25°C. Colony numbers were manually counted.

Cell transfection. miR-494-3p mimics (5'-UGAAACAUAACACGGGAAACCUC-3'), inhibitors (5'-GAGGGUUCCCCGUGUAUGUUUCAC-3') and negative controls (NCs; 5'-ACAUCUCAAGAUAUGAGUCAUA-3') were synthesized by Invitrogen (Thermo Fisher Scientific, Inc.). For PTEN overexpression, the coding sequence of PTEN was amplified by PCR and constructed into the pcDNA3 vector (Invitrogen; Thermo Fisher Scientific, Inc.) between EcoRI and XhoI. All transfections were performed using Lipofectamine® 2000 (Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer's protocol, at a concentration of 50 nM for miRNAs and 1 µg for pcDNA3-PTEN. After 48 h at 37°C, the transfection efficiency was validated using qRT-PCR as described below.

In vivo nude mouse xenograft assay. All animal experiments were approved by the Ethics Committee of The Second Affiliated Hospital of Zhengzhou University. Female BALB/c nude mice (4 mice per group; weight, 20.00±1.92 g), 4–6-weeks-old were obtained from Beijing Vital River Laboratories Animal Technology Co., Ltd. (Beijing, China) and were routinely housed in light-(12 h dark/12 h light) and temperature-controlled rooms (23°C). These mice were given free access to sterile food and water during the experiment process. HHUA cells (1x10^7) were transplanted with miR-494-3p inhibitors or controls resuspended in 200 µl FBS-free culture medium and subcutaneously injected into the right flanks of mice. miR-494-3p inhibitors or controls were injected into the formed tumor tissues every three days for 42 days. The tumor volume and weight were determined routinely following inoculation using direct measurement and calculated using the formula (length x width^2)/2. The mice were then sacrificed; tumor tissues were obtained and tumor weights were determined.

Cell proliferation. Cell proliferation was examined using a Cell Counting Kit-8 (CCK8) assay (R&D Systems, Inc., Minneapolis, MN, USA) according to the manufacturer's protocols. Proliferation was determined through measuring absorbance at 450 nm using an ELx808 absorbance reader (BioTek Instruments, Inc., Winooski, VT, USA). And a colony formation assay was also conducted. In brief, 1,000 HHUA or JEC cells were seeded into 6-well plates and cultured for 14 days in DMEM medium at 37°C. Then the colonies were fixed with paraformaldehyde for 30 min at 25°C and stained with 0.1% crystal violet for 30 min at 25°C. Colony numbers were manually counted.

Transwell assay. To examine cell migration, cells were plated onto a 24-well Transwell chamber (Corning Incorporated, Corning, NY, USA). A total of 2x10^4 cells were diluted in serum-free medium at 24 h post-transfection and subsequently inoculated onto the upper chamber. A total of 600 µl Dulbecco's modified Eagle's medium (Sigma-Aldrich; Merck KGaA) with 10% FBS (Gibco; Thermo Fisher Scientific, Inc.) was added to the lower chamber at 37°C. Non-migratory cells were removed using a cotton swab following overnight incubation, while the migratory cells in the lower chamber were fixed with paraformaldehyde for 30 min at 25°C and stained with 0.1% crystal violet for 30 min at 25°C. A cell invasion assay was performed in a similar manner, but Matrigel (Collaborative Research, Bedford, MA, USA) was added to the upper chamber. Finally, the number of migrated and invasive cells were observed and counted using an optical microscope (magnification x200, Nikon Corporation, Tokyo, Japan). Three random fields were counted per group.

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR). Total RNA from tissues or cells was extracted using TRIzol® reagent (Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer's protocols. RNA (0.5 µg) was reverse transcribed using a PrimeScript RT Kit (Takara Bio, Inc., Otsu, Japan) according to the manufacturer's protocols. Then, the transcripts were analyzed on an ABI 7300 qPCR system (Applied Biosystem; Thermo Fisher Scientific, Inc.) using specific primers synthesized by Beijing Sunbiotech Co., Ltd. (Beijing, China) using the TaqMan™ MicroRNA Assay kit (Applied Biosystems; Thermo Fisher Scientific, Inc.) for miRNAs and the Fast SYBR™-Green Master Mix (Applied Biosystems; Thermo Fisher Scientific Inc.) for mRNAs. The thermocycling conditions were as follows: Denaturation at
95°C for 10 min; followed by 40 cycles of denaturation at 95°C for 15 sec and elongation at 60°C for 1 min. Three repeats were performed. Relative expression was calculated and normalized to endogenous β-actin (ACTB) or U6. Relative expression levels were evaluated via the 2^−ΔΔCq method (17). Primer sequences were as follows: miR-494-3p, forward, 5′-AACGAGACGACGACAGAC-3′ and reverse, 5′-TGAAATACACGGGGAAACCTC-3′; U6 forward, 5′-AACGAGACGACGACAGAC-3′ and reverse, 5′-GCAATTTCTGAAGCGTCCCTA-3′; PTEN forward, 5′-TCCCGAGACATGACGACCATC-3′ and reverse, 5′-TGCTTGAATCCAAAACTCTTACT-3′; and ACTB forward, 5′-CGGCGGCTCTAATAAAAATGA-3′ and reverse, 5′-GAGGGCTACGGGATAGCC-3′.

Western blotting. Total protein was isolated from cultured HHUA and JEC cells using radioimmunoprecipitation assay buffer (Thermo Fisher Scientific, Inc), and the supernatant was collected via centrifugation at 13,282 x g for 10 min at 4°C. Protein concentration was evaluated using a Pierce BCA Protein Assay kit (Pierce; Thermo Fisher Scientific, Inc.). Subsequently, the extracted protein was mixed with loading buffer and boiled at 100°C for 5 min. Total protein (30 µg) was separated using a 10% SDS-PAGE gel and transferred onto polyvinylidene fluoride membranes (EMD Milipore, Billerica, MA, USA). Subsequently, the membranes were blocked with 5% (w/v) non-fat milk for 2 h at 25°C and incubated with antibodies against PTEN (1:2,000; cat. no. 9188), AKT (1:2,000; cat. no. 4691), phosphorylated-AKT (1:2,000; cat. no. 4060), P13K (1:2,000; cat. no. 4249), p-P13K (1:2,000; cat. no. 4228), BCL2 (1:2,000; cat. no. 2872), caspase-3 (1:2,000; cat. no. 9664) or GAPDH (1:2,000; cat. no. 5174) (all from Cell Signaling Technology, Inc.) for 2 h at 25°C. The membranes were then probed with a horseradish peroxidase-conjugated secondary antibody (1:5,000; cat. no. ab7090; Abcam, Cambridge, UK) at 25°C for 1 h, and signals were visualized using an enhanced chemiluminescence kit (Beiyetime Institute of Biotechnology, Beijing, China) according to manufacturer's protocol.

Luciferase reporter assay. The potential binding site for miR-494-3p in PTEN 3′-UTR region was predicted using the TargetScan7 tool (www.targetscan.org/vert_71/). Cells were seeded on a 6-well-plate at a density of 1x10^4 cells/well and transfected with 50 nM miR-494-3p mimics or negative controls, 100 ng pGL3-PTEN 3′-UTR-Wild type or Mutant vector and 1 ng pRL-TK luciferase plasmid (Promega Corporation, Madison, WI, USA) according to the manufacturer's protocols (Promega Corporation). After 48 h, luciferase assays were performed using the dual-luciferase reporter assay system (Promega Corporation) according to the manufacturer's protocols. Luminescent signals were quantified with a luminometer (Glomax, Promega Corporation), and the value of firefly luciferase was normalized to that of Renilla luciferase.

Statistical analysis. Statistical analysis was performed using SPSS 17.0 (SPSS Inc., Chicago, IL, USA). The differences among groups, in at least three separate experiments, were analyzed using a Student's t-test or one-way analysis of variance followed by Dunnett's multiple comparison test, as appropriate. The samples were divided into miR-494-3p low and high expression groups according to the median value of miR-494-3p. Then Kaplan-Meier analysis and log-rank test were used for survival evaluation. Spearman's rank correlation analysis was performed to analyze correlation between miR-494-3p and PTEN expression levels. P<0.05 was considered to indicate a statistically significant difference.

Results

miR-494-3p is upregulated in EC tissues. To investigate the function of miR-494-3p in EC, 43 EC samples were employed and the expression of miR-494-3p was determined by RT-qPCR in the present study. The results indicated that miR-494-3p was significantly upregulated in EC tissues compared with adjacent normal tissues (Fig. 1A). Then, these tissues were divided into low- or high-expression subgroups according to miR-494-3p expression, followed by Kaplan-Meier survival analysis. As presented in Fig. 1B, increased expression of miR-494-3p in patients with EC indicated poorer prognosis.

Overexpression of miR-494-3p promotes EC cell proliferation, migration and invasion. To further determine the effects of miR-494-3p on EC cells, miR-494-3p was overexpressed in EC cell lines (HHUA and JEC; Fig. 2A). Then, CCK8 and colony formation assays were performed using HHUA and JEC cells transfected with miR-494-3p mimics or controls. The results revealed that overexpression of miR-494-3p significantly promoted cellular proliferation and increased the number of the colonies compared with the control group (Fig. 2B and C). In addition, the effects of miR-494-3p on cellular apoptosis were evaluated by analysing the protein expression levels of caspase-3 and B-cell lymphoma-2 (Bcl-2). The results revealed that overexpression of miR-494-3p did not affect caspase-3 and Bcl-2 expression (data not shown), indicating that apoptosis was not affected by miR-494-3p. Tumor metastasis is one of the main causes of tumor malignancy (18). The effects of miR-494-3p on tumor cell metastasis were investigated using a Transwell assay. As presented, overexpression of miR-494-3p significantly promoted the migration and invasion of HHUA and JEC cells compared with the control group (Fig. 2D and E). Furthermore, the effects of miR-494-3p inhibition were evaluated by CCK8 and Transwell assays.
expression levels were significantly inhibited following transfection with miR-494-3p inhibitors (Fig. 2F). In addition, the results demonstrated that miR-494-3p inhibitor significantly suppressed the proliferation and invasion of HHUA and JEC cells compared with the control (Fig. 2G and H).

**Inhibition of miR-494-3p suppresses tumor growth in vivo.** To further investigate the physiological function of miR-494-3p in vivo, a xenograft experiment was performed using HHUA cells. miR-494-3p-silenced or control HHUA cells were injected into nude mice. At indicative time points following injection, the tumor volumes were measured; miR-494-3p knockdown significantly delayed tumor growth in vivo compared with the control (Fig. 3A). In addition, the tumor weights were determined at the endpoints of experiments. As presented in Fig. 3B, miR-494-3p knockdown resulted in significantly reduced tumor weight compared with the control.

miR-494-3p activates the PI3K/AKT signaling pathway by targeting PTEN. The miR-494-3p-regulated molecular mechanisms associated with EC were investigated. According to bioinformatic prediction, PTEN was identified to be a potential target gene of miR-494-3p. There were two conserved potential binding sites in the 3'-UTR of PTEN mRNA (Fig. 4A); then, luciferase reporter assays were conducted. As presented in Fig. 4B, overexpression of miR-494-3p significantly inhibited luciferase activity in HHUA and JEC cells compared with the control. Furthermore, the present study reported that overexpression of miR-494-3p significantly downregulated the mRNA expression levels of PTEN in HHUA and JEC cells compared with the control. Then, the expression levels of miR-494-3p and PTEN in EC tissues were evaluated. The results of RT-qPCR indicated that there was an inverse correlation between the expression levels of miR-494-3p and PTEN in EC tissues (Fig. 4D). A recent study revealed that the downstream signaling of PTEN comprised the PI3K/AKT pathway (8), which was widely involved in various human cancers, such as EC (19,20). Furthermore, the effects of miR-494-3p on PTEN protein expression and PI3K/AKT activation were evaluated using western blot analysis. Overexpression of miR-494-3p notably downregulated the protein expression levels of PTEN, and upregulated the phosphorylation of PI3K and AKT (Fig. 4E). In summary,
the aforementioned data indicated that miR-494-3p activates the PI3K/AKT pathway by targeting PTEN.

**Restoration of PTEN or inhibition of the PI3K/AKT pathway abolishes miR-494-3p-mediated effects on EC cells.** To determine whether miR-494-3p affects EC cell proliferation, migration and invasion via the PTEN/PI3K/AKT pathway, the protein expression of PTEN was restored or the PI3K/AKT pathway was inhibited using a specific inhibitor (LY294002) in miR-494-3p-overexpressed HHUA and JEC cells (Fig. 5A). Then, CCK8 and colony formation assays were performed to evaluate cellular proliferation. As presented in Fig. 5B and C, overexpression of miR-494-3p significantly promoted cell proliferation compared with the control; however, restoration of PTEN or inhibition of the PI3K/AKT pathway abrogated the effects of miR-494-3p overexpression on EC cells. Similarly, overexpression of miR-494-3p significantly promoted cellular migration and invasion compared with the control; however restoration of PTEN or inhibition of the PI3K/AKT pathway in miR-494-3p-overexpressed HHUA and JEC cells reversed these effects (Fig. 5D and E). In summary, the results in the present study demonstrated that miR-494-3p promoted the progression of EC via the PTEN/PI3K/AKT signaling pathway.

**Discussion**

MiRNAs can regulate the development and progression of EC (21); however, the functions of numerous miRNAs in EC have not been determined. In the present study, the expression levels of miR-494-3p were significantly upregulated in EC tissues compared with adjacent normal tissues. Additionally, the expression of miR-494-3p was positively associated with...
the poor prognosis of patients with EC. Therefore, miR-494-3p may serve as an oncogene in EC, consequently promoting tumor progression.

Numerous studies demonstrated that miR-494-3p was an essential regulator in numerous types of cancer. For example, a recent study indicated that miR-494-3p promoted the development of lung cancer (14). Liu et al (22) reported that miR-494 promoted cell proliferation, migration and invasion, and increased sorafenib resistance in hepatocellular carcinoma by targeting PTEN. Li et al (15) revealed that miR-494-3p regulated cellular proliferation, invasion, migration, and apoptosis via PTEN/AKT signaling in human glioblastoma cells. Additionally, some studies have indicated that miR-494 served as a tumor suppressor in certain types of cancer. For instance, miR-494-3p could induce cellular senescence and enhance radiosensitivity of oral squamous carcinoma cells (23). Shen et al (16) demonstrated that miRNA-494-3p targeted C-X-C chemokine receptor type 4 (CXCR4) to suppress the proliferation, invasion and migration of prostate cancer cells. These reported contrary functions of miR-494-3p that may be due to different target genes in various types of cancer. In the present study, overexpression of miR-494-3p significantly promoted cellular proliferation in EC in vitro and in vivo. In addition, overexpressed miR-494-3p induced cellular migration and invasion in vitro. The findings of the present study indicated an oncogenic role of miR-494-3p in EC.

Furthermore, previous studies revealed that PTEN could regulate a variety of biological processes, including the cell cycle, apoptosis, migration and invasion by inhibiting the PI3K/AKT signaling pathway (24, 25). PTEN mutations, deletions or silencing by promoter hypermethylation often result in the tumorigenesis of several cancers, such as EC (26). A previous study indicated that PTEN was mutated in 83% of EC tissues (27). Downregulation of PTEN was associated with activation of the PI3K/AKT pathway, consequently promoting tumor development and progression (28). For example, E3 ubiquitin-protein ligase regulated PTEN/PI3K/AKT signaling to promote the cell growth and migration of hepatocellular carcinoma cells (29). Liu et al (30) reported that Sal-like protein 4 suppressed PTEN expression to promote glioma cell proliferation via the PI3K/AKT pathway. Another study revealed that miRNA-1297 contributed to tumor growth
of human breast cancer by targeting the PTEN/P13K/AKT signaling pathway (28). In addition, miRNA-92a promoted epithelial-mesenchymal transition via the PTEN/P13K/AKT pathway in metastatic non-small cell lung cancer (18). In the present study, miR-494-3p directly targeted PTEN and down-regulated its expression. Furthermore, restoration of PTEN and inhibition of the P13K/AKT pathway inhibited the proliferation, migration and invasion of EC cells overexpressing miR-494-3p, which indicated that miR-494-3p regulated the progression of EC in a PTEN/P13K/AKT-dependent manner.

In summary, the findings of the present study revealed the essential role of miR-494-3p and its functional mechanisms in EC. The results suggested that the miR-494-3p/PTEN/P13K/AKT axis may be a promising therapeutic target for the treatment of EC.

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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

LZ initiated, designed this work, analyzed and interpreted the results. LZ wrote this manuscript. XW, TW, WZ and XZ performed the experiments. All authors read and approved the final manuscript.

Ethics approval and consent to participate

For the use of human samples, the protocol for the present study was approved by the Institutional Ethics Committee of the Second Affiliated Hospital of Zhengzhou University (Zhengzhou, China) and all enrolled patients signed a written informed consent document. All animal experiments were approved by the Ethics Committee of the Second Affiliated Hospital of Zhengzhou University.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

References


