

# miR-26a inhibits ovarian cancer cell proliferation, migration and invasion by targeting TCF12

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**Abstract.** Epithelial ovarian cancer (OC) is a common cause of death from gynecological tumors. MicroRNAs (miRNAs) may function as either oncogenes or tumor suppressors, playing crucial role not only in tumorigenesis, but also in tumor invasion and metastasis. miR-26a and transcription factor 12 (TCF12) have been reported to be involved in carcinogenesis, but the regulatory role of miR-26a/TCF12 in OC remains unknown. The aim of the present study was to investigate the expression profiles of TCF12 and miR-26a in OC patients and the correlation between TCF12 and miR-26a expression, and to demonstrate the effects of miR-26a binding on TCF12, to further reveal the miR-26a/TCF12 regulatory effects on the proliferation, migration, invasion and apoptosis in OC cells. In the present study, the expression of miR-26a was found to be low, while TCF12 was highly expressed in OC patient tissues and cell lines, and low miR-26a expression was statistically significantly correlated with high TCF12 expression. To the best of our knowledge, the present study was the first to demonstrate that TCF12 is a direct target of miR-26a, and upregulation of miR-26a resulted in TCF12 inhibition in OC cells. Furthermore, the proliferation, migration and invasion were inhibited and apoptosis was induced by miR-26a upregulation in OC cells. These results indicated that miR-26a may act as a tumor suppressor in OC, and TCF12 targeting by miR-26a may be a new therapeutic strategy for OC.

## Introduction

Epithelial ovarian cancer (OC) is the most common gynecological tumor with a high mortality rate worldwide (1).

Epithelial OC is associated with a high rate of metastasis at an early stage, and metastasis is the major cause of the dismal prognosis (2); however, the detailed mechanisms underlying the propensity of OC for metastasis remain unclear. There is an urgent need to identify methods for early diagnosis and effective treatment for OC.

MicroRNAs (miRNAs) are an endogenous class of single-stranded small RNAs ~20-24 nucleotides in length, which play a variety of important regulatory roles in cells (3). The first miRNA was identified in *Caenorhabditis elegans* (4), whereas hundreds of miRNAs have been identified to date and found to be highly conserved in all types of organisms (5). In recent years, research on miRNAs has markedly progressed; in particular, the mechanism of target gene regulation, and numerous miRNAs and their target genes have been identified and investigated. miRNAs are promising targets for the diagnosis and treatment of diseases caused by multiple factors, particularly cancer (6), and they have also been studied in the context of OC (7). miR-26a has been reported to act as a tumor suppressor in numerous human cancers, such as prostate cancer (8-10), lung cancer (11-13), thyroid cancer (14), breast cancer (15-17), bladder cancer (18), hepatocellular carcinoma (19-21), colorectal cancer (22,23), glioblastoma (24), melanoma (25) and esophageal squamous cell carcinoma (26). However, there are few reports of the role of transcription factor 12 (TCF12) in OC.

As a transcription factor, TCF12 belongs to the basic helix-loop-helix (bHLH) E protein family, which can recognize the E-box sequence CANNTG (27). TCF12 has been proven to be a regulator of cell development and differentiation (28), and has been demonstrated to be highly expressed in several human cancers (29-31). However, the role of TCF12 and the association of miR-26a and TCF12 expression in OC have not been extensively investigated to date.

The aim of the present study was to investigate the expressions of miR-26a and TCF12 and their correlation in OC patients. In addition, the regulatory effect of miR-26a on TCF12 expression was determined, and the proliferation, migration, invasion and apoptosis of OC cells were also investigated.

## Materials and methods

**Clinical patient samples and cell lines.** A total of 27 cases of human fresh OC tissues and paired normal ovarian tissues

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were collected from patients (mean age 55.10±11.10 years, range 35-78 years) who were diagnosed with primary OC and underwent initial surgery at the Department of Obstetrics and Gynecology of the Affiliated Hospital of Nantong University. Written informed consent was obtained from all the patients and the study protocol was approved by the Ethics Committee of the Affiliated Hospital of Nantong University.

The human OC cell lines SK-OV-3 and A2780, the normal ovarian epithelial cell line IOSE80 and 293 cells were cultured in Dulbecco's modified Eagle's medium (DMEM; Thermo Fisher Scientific, Inc.) and supplemented with 10% fetal bovine serum (FBS; Thermo Fisher Scientific, Inc.) at 37°C in a humidified incubator with 5% CO<sub>2</sub>.

**Oligos, plasmids and transfection.** The human miR-26a mimic (sense: 5'-UUCAAGUAAUCCAGGAUAGGCU-3', antisense: 5'-AGCCUAUCCUGGAUUACUUGAA-3') was used to regulate the expression of TCF12 in OC cells and a scrambled miR-26a mimic sequence used as negative control (NC; sense: 5'-GGUCGUCUGAUUAUCGAUACA-3', antisense: 5'-UUGUAUCGUUAUCAGACGACC-3'). The miR-26a binding site of TCF12 was predicted using TargetScan online software (<http://www.targetscan.org/>). The 50 base pairs (bp) cDNA of the 3' untranslated region (UTR) sequence containing miR-26a-binding sites from TCF12 mRNA was cloned into a pGL3-control vector (Promega Corporation) downstream of the luciferase gene, which was used as the wild-type TCF12 (wtTCF12) plasmid. A point mutant miR-26a-binding site of TCF12 was constructed and was used as the mutant TCF12 (muTCF12) plasmid. Cell transfection of 50 pmol miRNA or 80 ng plasmid was performed by Lipofectamine® 2000 transfection reagent (Thermo Fisher Scientific, Inc.) according to the manufacturer's instructions. miRNA, NC and DNA oligos were obtained from Biomics Biotechnologies Co., Ltd.

**Reverse transcription-quantitative PCR (RT-qPCR) analysis.** Small RNAs were isolated from tissues and cell lines using the mirVana miRNA Isolation Kit (Thermo Fisher Scientific, Inc.) according to the manufacturer's instructions, and the miR-26a expression levels were detected by the stem-loop RT-qPCR method (32). U6 small RNA was used as an internal control. Total RNA was extracted from cells using TRIzol® reagent (Thermo Fisher Scientific, Inc.) and RT-qPCR reactions were carried out using the One-Step RT-qPCR kit (Thermo Fisher Scientific, Inc.) according to the manufacturers' instructions. The thermocycling conditions were as follows: Pre-denaturation at 95°C for 10 min, followed by 40 cycles of denaturation at 95°C for 20 sec, annealing at 58°C for 20 sec and extension at 72°C for 30 sec.  $\beta$ -actin was used as an internal control. miRNA and mRNA level were analyzed by the 2<sup>- $\Delta\Delta C_q$</sup>  method (33). The primer sequences were as follows: TCF12 forward: 5'-CTAATGAAGATGAGGATT-3' and reverse: 5'-GATGAAGAATAAGGAGTT-3';  $\beta$ -actin forward: 5'-TTGCCGACAGGATGCAGAGGA-3' and reverse: 5'-AGGTGGACAGCGAGGCCAGGAT-3'; miR-26a forward: 5'-GCGTTGTCTGGAATGTAAGG-3' and reverse: 5'-TGACGAGTTTAGAGCCGATAG-3'; and U6 forward: 5'-AACGCTTCACGAATTTGCGT-3' and reverse: 5'-CTCGCTCGGCAGCACA-3'.

**Dual luciferase reporter assay.** The 293 cells were used to measure luciferase activity. Briefly, 293 cells were plated into 24-well plates at 1×10<sup>5</sup> per well and cultured for 24 h. Subsequently, 80 ng wtTCF12 or muTCF12 plasmid and 50 nmol/l miR-26a mimic or NC were co-transfected into 293 cells using Lipofectamine® 2000 transfection reagent (Thermo Fisher Scientific, Inc.) along with pRL-TK plasmid (Promega Corporation). Cells were collected 48 h after transfection and firefly and Renilla luciferase activities were measured using a dual luciferase reporter assay system according to the manufacturer's instructions (Promega Corporation). The results of firefly luciferase activity were normalized to the Renilla luciferase activity.

**Western blot analysis.** Briefly, following treatment for 48 h as described above, SK-OV-3 and A2780 cells were collected, and total protein was extracted with RIPA lysis and extraction buffer (Thermo Fisher Scientific, Inc.). Total proteins were quantified using the BCA method, then separated by 10% SDS-PAGE with 50  $\mu$ g per lane and transferred onto a PVDF membrane (GE Healthcare). Subsequently, after blocking with 5% non-fat dry milk for 2 h at room temperature, the membrane was incubated with rabbit anti-human TCF12 antibody (1:500 dilution, cat no. ab245540) or mouse anti-human  $\beta$ -actin antibody (1:1,000 dilution, cat no. ab179467) at 4°C overnight. After washing with TBST with 0.1% Tween-20, the PVDF membrane was incubated with a horseradish peroxidase-conjugated IgG (1:2,000 dilution, cat no. ab205718) for 2 h at room temperature; all antibodies were from Abcam, Inc. Finally, the specific proteins were detected using Pierce ECL Western Blotting Substrate (Thermo Fisher Scientific, Inc.), and ImageJ software 1.51 (National Institutes of Health) was used for densitometry analysis.

**Cell proliferation assay.** Cell proliferation was detected using the MTT method. In brief, 1.5×10<sup>3</sup> cells per well were grown in 96-well plates overnight, and then the cells were treated as indicated above. At 24, 48 and 72 h after treatment, 10  $\mu$ l MTT solution (Promega Corporation) were added to each well. At 4 h after incubation at 37°C protected from light, 150  $\mu$ l DMSO were added to each well and incubated at 37°C for 10 min. Finally, the OD value of each well was measured using a microplate reader at 490 nm.

**Transwell assay.** Transwell assay was used to evaluate cell migration and Matrigel-based Transwell assay was used to evaluate cell invasion. The cells were plated onto 6-well plates and cultured for 24 h. After treatment for 48 h, as indicated above, cells were collected and suspended in DMEM at a density of 1×10<sup>6</sup> cells/ml. Prior to treatment, Transwell chambers (8- $\mu$ m pore size; Corning, Inc.) were incubated with DMEM for 1 h at 37°C. For cell invasion detection, Matrigel (BD Biosciences) was thawed at 4°C overnight, and 100  $\mu$ l (1:8 diluted in DMEM) were added into the Transwell upper chamber and incubated at 37°C for 1 h. Cell suspension (100  $\mu$ l) was added into the upper chamber and 600  $\mu$ l DMEM supplemented with 10% FBS was added into the lower chamber. Following incubation for 24 h at 37°C, the cells on upper surface of the membrane were carefully removed using a cotton swab, and the cells on the lower surface were fixed

in 10% formaldehyde for 20 min at room temperature. After washing with PBS, the cells were stained in 0.5% crystal violet solution for 10 min at room temperature and washed with PBS. Finally, the stained cells on the lower surface of upper chamber were observed under a light microscope (magnification, x100).

**Cell apoptosis assay.** Annexin V-FITC and propidium iodide (PI) double staining followed by flow cytometry was used to evaluate cell apoptosis. Briefly, after treatment for 48 h as described above,  $1 \times 10^5$  SK-OV-3 or A2780 cells were collected and washed with PBS, then transferred into centrifuge tubes and centrifuged at  $800 \times g$  for 5 min at  $4^\circ\text{C}$ . After washing with PBS, the cells were re-suspended in  $195 \mu\text{l}$  1X Annexin V-FITC binding buffer and  $5 \mu\text{l}$  Annexin V-FITC (Sigma-Aldrich; Merck KGaA) and incubated for 10 min at room temperature in the dark. Following centrifugation for 5 min at  $800 \times g$  at  $4^\circ\text{C}$ , the cells were re-suspended in  $190 \mu\text{l}$  1X Annexin V-FITC binding buffer followed by  $10 \mu\text{l}$  PI (Sigma-Aldrich; Merck KGaA). Finally, cell apoptosis was detected using a flow cytometer (FACSCalibur; BD Biosciences) and the results were analyzed by BD CellQuest software, version 5.1 (BD Biosciences).

**Statistical analysis.** All data are presented as mean  $\pm$  standard deviation. Statistical analyses were performed using SPSS 20.0 software (IBM Corp.). Differences between two groups were tested by paired Student's two-tailed t-test. One-way ANOVA followed by Dunnett's post hoc test was used to compare multiple groups. The correlation of miR-26a with TCF12 expression was analyzed by Spearman's correlation analysis.  $P < 0.05$  was considered to indicate statistically significant differences.

## Results

**Low miR-26a and high TCF12 expression is observed in OC patients.** To investigate the expression of miR-26a and TCF12 in OC patients, 27 cases of fresh OC samples and paired normal ovarian tissues were used for RT-qPCR analysis. The results demonstrated that the miR-26a expression levels were low in OC tissues, while the TCF12 expression levels were high, compared with those in normal ovarian tissues ( $P < 0.05$ ; Fig. 1A and B). Spearman's correlation analysis revealed that miR-26a was negatively correlated with TCF12 expression ( $r = -0.695$ ,  $P < 0.05$ ; Fig. 1C).

**miR-26a directly targets the 3'UTR region of TCF12 mRNA.** As predicted by software, TCF12 mRNA contains a miR-26a-binding site at position 124 in the 3'UTR region. The luciferase reporter plasmids wtTCF12 and muTCF12 were constructed (Fig. 2A) and co-transfection of wtTCF12 or muTCF12 plasmid and miR-26a mimic into 293 cells was performed. Following transfection for 48 h, the luciferase activity of wtTCF12 and miR-26a co-transfected cells was obviously decreased compared with that in cells co-transfected with NC and wtTCF12 or muTCF12 ( $P < 0.05$ ; Fig. 2B).

**miR-26a inhibits the expression of TCF12 in OC cells.** Furthermore, the miR-26a expression levels were found to be low in the OC cell lines SK-OV-3 and A2780, while the

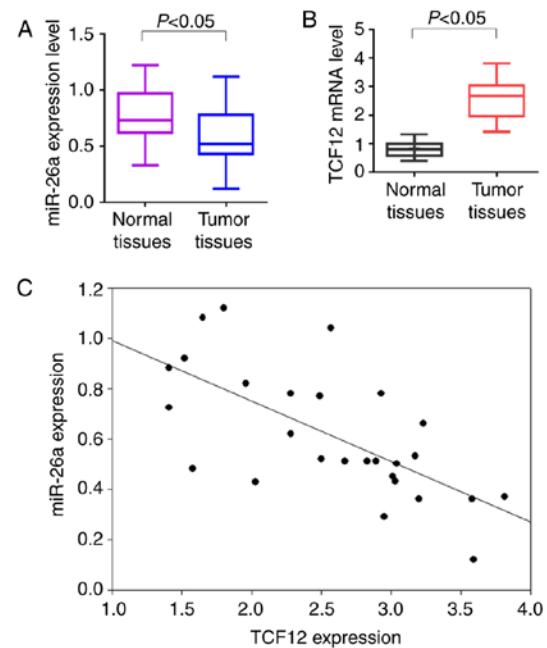


Figure 1. miR-26a and TCF12 expression in OC and paired normal ovarian tissues. (A) miR-26a levels in 27 cases of OC tissues and paired normal ovarian tissues were detected by stem-loop RT-qPCR; (B) The expression levels of TCF12 in 27 cases of OC tissues and paired normal ovarian tissues were detected by RT-qPCR; (C) Correlation of miR-26a and TCF12 levels in OC patients. OC, ovarian cancer; TCF12, transcription factor 12; RT-qPCR, reverse transcription-quantitative PCR.

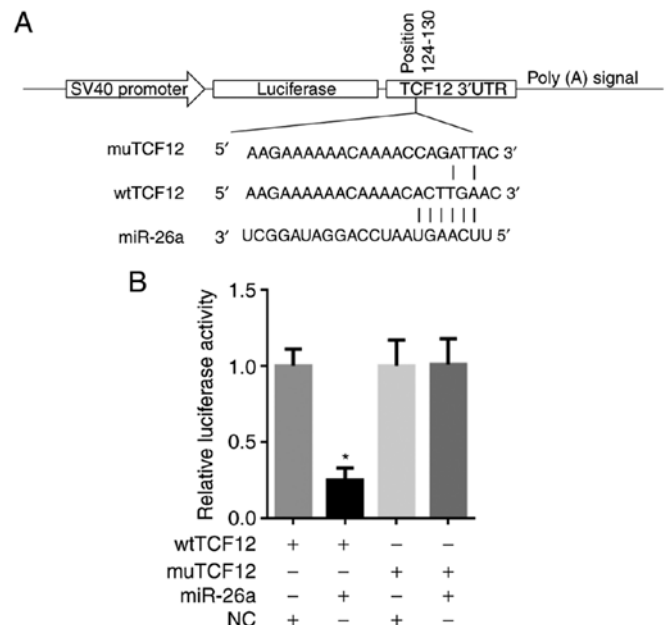


Figure 2. The regulatory effect of miR-26a on the expression of TCF12 was detected by dual luciferase reporter assay. (A) The miR-26a-binding site of TCF12 and the constructed reporter plasmids. (B) The relative luciferase activity was detected by the dual luciferase reporter assay. \* $P < 0.05$  compared with muTCF12 and pGL3 control-treated cells. TCF12, transcription factor 12; NC, negative control; UTR, untranslated region; wt, wild-type; mu, mutant type.

TCF12 expression levels were high, compared with those in IOSE80 cells ( $P < 0.05$ ; Fig. 3A and B). To further observe regulation of TCF12 expression by miR-26a in OC cells, human

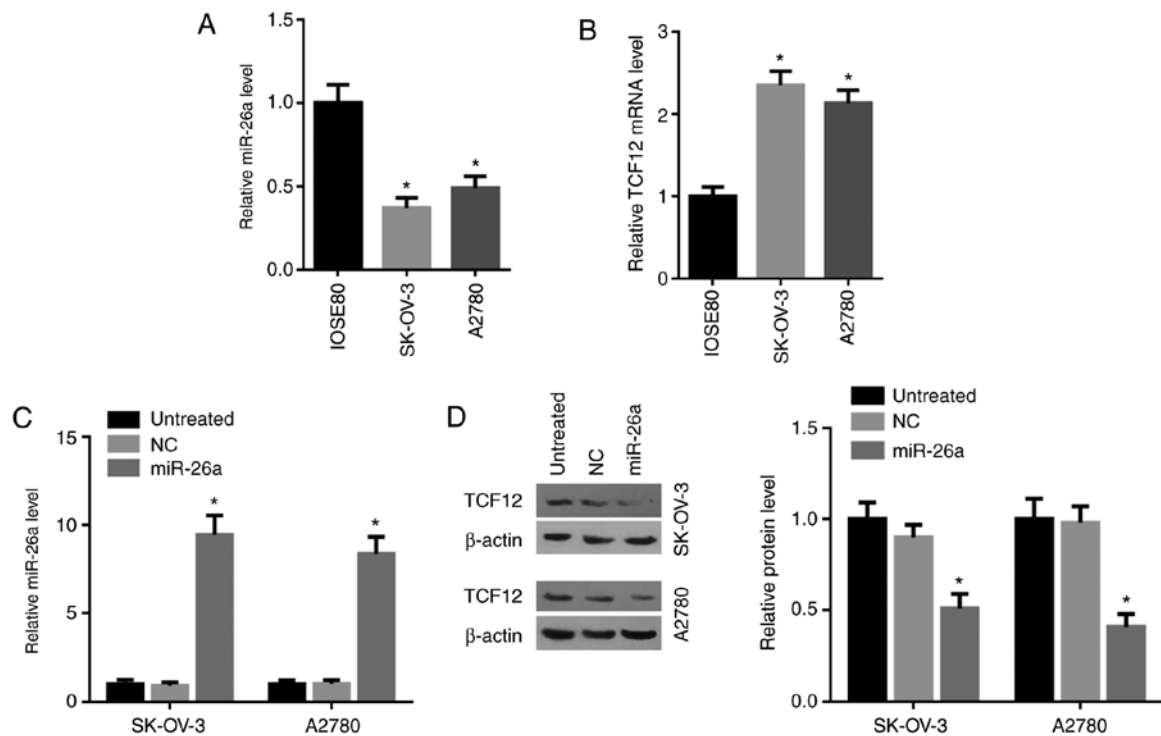


Figure 3. miR-26a and TCF12 expression levels and their interaction in OC cells. (A) miR-26a levels in the SK-OV-3 and A2780 OC cells and in the normal ovarian cell line IOSE80. (B) TCF12 mRNA levels in SK-OV-3, A2780 and IOSE80 cells. \* $P < 0.05$  compared with IOSE80 cells. (C) miR-26a levels in SK-OV-3 and A2780 OC cells with miR-26a mimic transfection, \* $P < 0.05$  compared with NC-treated or untreated cells. (D) The protein expression of TCF12 was inhibited by miR-26a. \* $P < 0.05$  compared with NC-treated or untreated cells. OC, ovarian cancer; TCF12, transcription factor 12; NC, negative control.

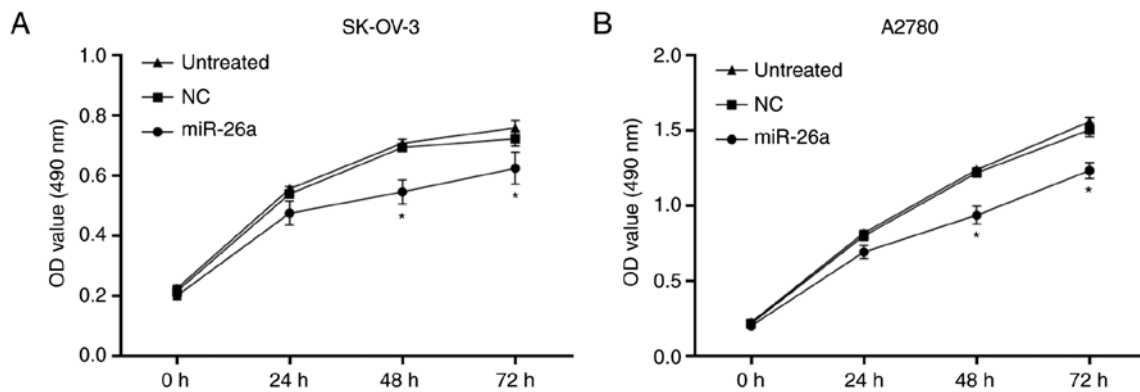


Figure 4. Effects of TCF12 inhibited by miR-26a on OC cell proliferation. (A) SK-OV-3 cell proliferation was inhibited by miR-26a; (B) A2780 cell proliferation was inhibited by miR-26a. \* $P < 0.05$  compared with NC or untreated cells. TCF12, transcription factor 12; NC, negative control; OD, optical density.

miR-26a mimic was used to increase endogenous miR-26a in OC cells, and the miR-26a levels in SK-OV-3 and A2780 cells were significantly upregulated following miR-26a mimic transfection (Fig. 3C). The protein expression of TCF12 was significantly inhibited by miR-26a in both SK-OV-3 and A2780 cells, compared with NC-treated or untreated cells ( $P < 0.05$ ; Fig. 3D).

**miR-26a inhibits the proliferation of OC cells.** MTT assay was used to investigate the inhibitory effects of miR-26a on OC cell growth. TCF12 inhibition by miR-26a clearly inhibited the growth of SK-OV-3 (Fig. 4A) and A2780 (Fig. 4B) cells at 48 and 72 h, compared with NC-treated or untreated cells ( $P < 0.05$ ).

**miR-26a inhibits the migration and invasion of OC cells.** Transwell and Matrigel-based Transwell assay was used to evaluate the inhibitory effects of miR-26a on OC cell migration and invasion, respectively. The results demonstrated that miR-26a treatment significantly decreased SK-OV-3 and A2780 cell migration and invasion, compared with NC-treated or untreated cells ( $P < 0.05$ ; Fig. 5A and B).

**miR-26a induces the apoptosis of OC cells.** Annexin-V-FITC/PI flow cytometry was used to evaluate the effects of miR-26a on OC cell apoptosis. Treatment with miR-26a for 48 h significantly increased apoptosis of both SK-OV-3 and A2780 cells compared with cells treated with NC ( $P < 0.05$ ; Fig. 5C).



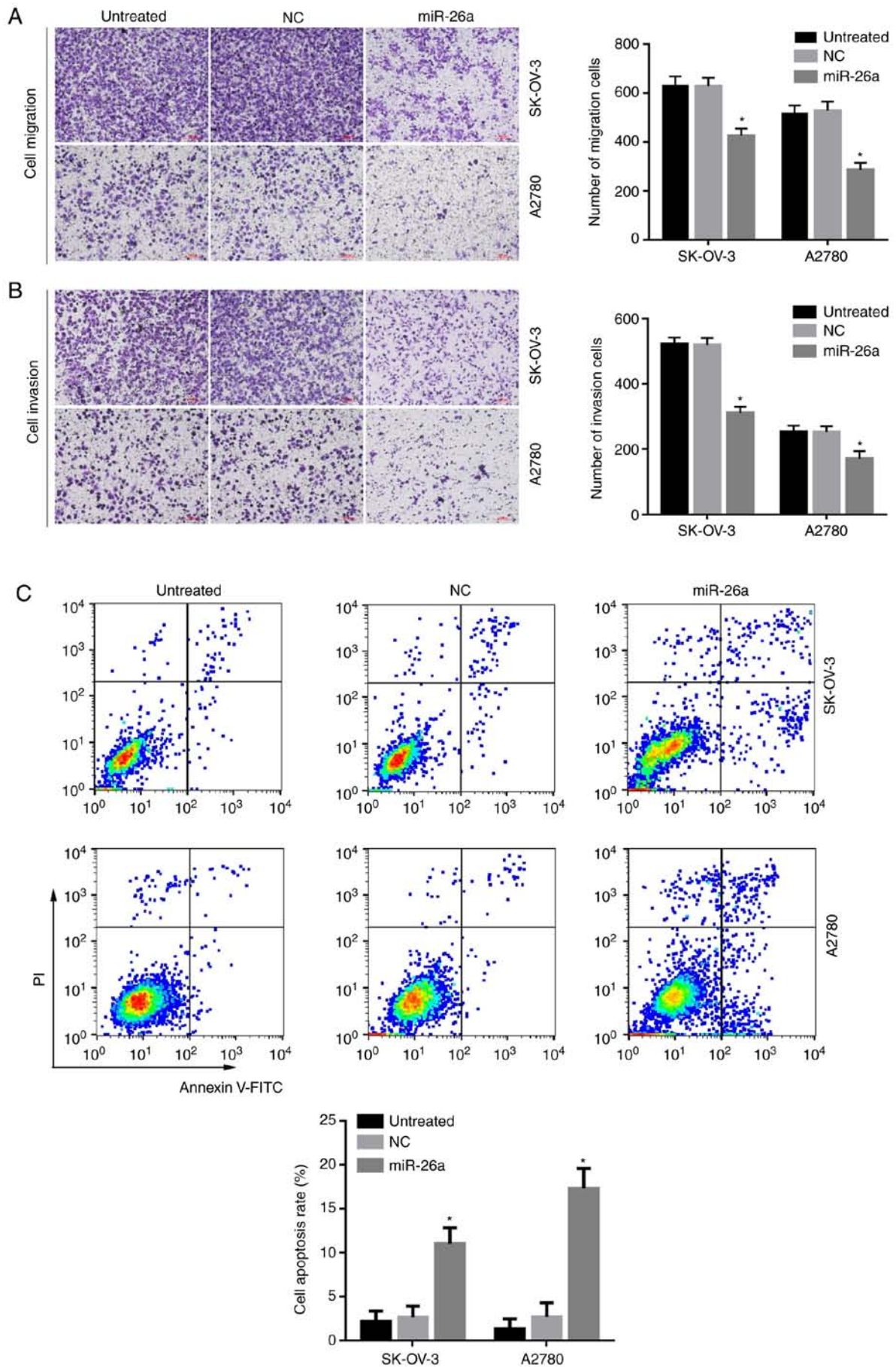


Figure 5. Effects of TCF12 inhibited by miR-26a on OC cell migration, invasion and apoptosis. (A) The migration of SK-OV-3 and A2780 cells was inhibited by miR-26a; (B) the invasion of SK-OV-3 and A2780 cells was inhibited by miR-26a; (C) apoptosis of SK-OV-3 and A2780 cells was induced by miR-26a. \* $P < 0.05$  compared with NC or untreated cells. TCF12, transcription factor 12; FITC, fluorescein isothiocyanate; PI, propidium iodide; NC, negative control. Scale bar, 100  $\mu$ m.

## Discussion

Lack of early diagnosis and effective treatment is a major cause of death from OC among women with malignant gynecological tumors (2). Individualized cancer treatment based on molecular targeted therapy is crucial for improving therapeutic efficacy, and molecules that are heavily involved in the uncontrolled cell growth and proliferation, inhibition of apoptosis and tumor metastasis may represent promising targets for cancer therapy (34).

miRNAs are small endogenously expressed RNAs in mammalian cells, which have been used in regulation of gene expression at the post-transcriptional level in a sequence-specific manner (35). miRNAs have been demonstrated to play multiple functional roles in biological processes, such as cell developmental control, regulation, and the occurrence of various diseases, particularly cancer, due to their high abundance, sequence conservation and tissue specificity (5). Due to their different expression patterns and functions, miRNAs involved in tumorigenesis may be classified as oncogenes or tumor suppressors in different human cancers (36); thus, miRNAs may be used as novel targets or markers for detection, prognosis, diagnosis and therapy (37). In recent years, a number of studies have identified aberrant expression of miRNAs in ovarian tumorigenesis leading to OC (38). miR-26a was identified as a tumor suppressor in a number of human cancers. miR-26a was shown to exert an antiproliferative effect on prostate cancer cells by decreasing survival and migration and inducing cell cycle arrest and apoptosis (9). In gastric cancer, miR-26a promoted cell proliferation, migration and invasion, and phosphatase and tensin homolog was identified as a direct target of miR-26a (10). miR-26a was demonstrated to enhance the metastatic potential of lung cancer cells by targeting glycogen synthase kinase (GSK)3 $\beta$  of the  $\beta$ -catenin pathway; miR-26a expression was found to be negatively correlated with GSK3 $\beta$  expression in patients with lung cancer, and overexpression of GSK3 $\beta$  reversed the enhancing effect of miR-26a on lung cancer cell migration and invasion. In addition, integrin  $\beta$ 8 (13), lin-28 homolog B (39) and matrix metalloproteinase-2 (11) have also been identified as targets in lung cancer. miR-26a has been proven to be associated with metastasis, survival and apoptosis in breast cancer, acts as a tumor suppressor and serves as a prognostic marker (15,16). The proliferation, migration and invasion of hepatocellular carcinoma (HCC) cells can also be inhibited by miR-26a by targeting diverse genes and pathways, such as the epithelial-to-mesenchymal transition (40), interleukin-6-Stat3 (13) and PI3K/Akt (23) pathways, and low miR-26a levels were found to be closely associated with metastasis and progression of HCC (19-21). In bladder cancer (18), colorectal cancer (23) and melanoma (25), miR-26a was found to play the same functional roles. In recent years, the expression of miR-26a has been reported to be low in OC tissues and cells, further investigation demonstrated that the proliferation of OC cells was inhibited while their apoptosis was induced by miR-26a, suggesting that miR-26a downregulation may be predictive of poor prognosis in epithelial OC (41,42). Moreover, TCF12 was found to be highly expressed in OC patients in our recent study (43), and its overexpression was correlated

with histological grade and metastasis of OC, whereas TCF12 downregulation inhibited the growth, migration and invasion of OC cells and promoted apoptosis. In the present study, the expression of miR-26a was also found to be low in OC patients and cell lines, whereas the expression of TCF12 was high, and low miR-26a expression was statistically significantly correlated with high TCF12 expression. To the best of our knowledge, the present study was the first to demonstrate that TCF12 is a direct target of miR-26a by the dual luciferase reporter assay. The expression of TCF12 in OC cells was inhibited by miR-26a at both the mRNA and protein levels. Furthermore, the proliferation, migration and invasion of SK-OV-3 and A2780 cells were inhibited by miR-26a through suppression of TCF12, whereas apoptosis was induced.

Therefore, the findings of the present study suggest that upregulation of miR-26a may inhibit metastasis and promote apoptosis in OC cells by targeting TCF12, which may be a potential strategy for OC treatment.

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## Availability of data and materials

The datasets generated and analyzed during the present study are available from the corresponding author upon reasonable request.

## Authors' contributions

SG, TB and YZ contributed to conception and design of this study; YL and TB collected the clinical samples and patient information; SG, TB, MS and YL performed the experiments; SG, TB, MS, YL and YZ acquired, analyzed and interpreted the data; SG was a major contributor to drafting and writing the manuscript; TB, MS, YL and YZ revised the manuscript. All authors have read and approved the final version of the manuscript for publication.

## Ethics approval and consent to participate

Written informed consent was obtained from all the patients and the study protocol was approved by the Ethics Committee of the Affiliated Hospital of Nantong University.

## Patient consent for publication

Not applicable.

## Competing interests

The authors declare that they have no competing interests.

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