miR-28-5p suppresses cell proliferation and weakens the progression of polycystic ovary syndrome by targeting prokineticin-1

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Abstract. Prokineticin-1 (PROK1) serves important roles in the pathogenesis of polycystic ovary syndrome (PCOS); however, the association between microRNA (miR)-28-5p and PROK1 remains unclear. In the present study, the roles of miR-28-5p and PROK1, and their interaction in PCOS were investigated. Rat ovary granule cells were transfected with miR-28-5p mimics, and PROK1 expression levels were measured by reverse transcription-quantitative PCR and western blotting. A dual-luciferase reporter assay was performed to determine the association between miR-28-5p and PROK1. Additionally, pcDNA-PROK1 was co-transfected into rat ovary granule cells with miR-28-5p mimics. Cell proliferation, apoptosis, cell cycle and the expression of signaling proteins were investigated using cell counting Kit-8 assays, 5-ethynyl-2'-deoxyuridine staining, flow cytometry and western blotting, respectively. PROK1 expression was suppressed in rat ovary granule cells by miR-28-5p mimics, but upregulated following transfection with miR-28-5p inhibitors. The dual-luciferase reporter assay revealed that miR-28-5p binds to the 3'-untranslated region of PROK1. Proliferation activity was increased in PROK1-overexpressing cells; this effect was eliminated by co-transfection with miR-28-5p mimics. PROK1-overexpressing rat ovary granule cells exhibited significantly suppressed cell apoptosis and a decreased number of cells in G1; miR-28-5p mimics reversed these effects. Western blotting revealed that the PI3K/AKT/mTOR signaling pathway was activated by PROK1. The present results suggested that miR-28-5p attenuated the progression of PCOS by targeting PROK1, which may promote the pathogenesis of PCOS via the PI3K/AKT/mTOR pathway, indicating that the miR-28-5p/PROK1 axis may be a potential therapeutic target for patients with PCOS.

Introduction

Polycystic ovary syndrome (PCOS) exhibits high morbidity and affects ~4% of women of reproductive age worldwide (1,2). PCOS is frequently accompanied by a number of alterations in the reproductive, endocrine and metabolic pathways. Patients with PCOS typically present with polycystic ovaries, insulin resistance, chronic anovulation and infertility (3,4). Previous studies revealed that PCOS pathogenesis is due to environmental and genetic factors (5); however, efforts to understand the underlying pathogenic mechanisms are complicated by the heterogeneity of PCOS.

Prokineticin 1 (PROK1), also known as endocrine gland-derived vascular endothelial growth factor (EG-VEGF), is involved in ovarian physiology, endometrial receptivity, embryo implantation and successful pregnancies, together with its receptors (PROK1 receptors (PROKRs)) (6,7). Previous research revealed that PROK1 is an angiogenic and proliferative factor predominantly expressed in the organs of the female reproductive system (8). PROK1 is also involved in female reproductive pathophysiological processes (8). In addition, PROK1 was reported to be differentially expressed during the cycle of vascular morphogenesis in the primate ovary and polycystic human ovaries, compared with VEGF (9-11). There is also evidence that PROK1 mediates placental angiogenesis directly or indirectly (11); however, the precise mechanisms of PROK1 in PCOS require further investigation.

MicroRNAs (miRNAs/miRs) are widely expressed small RNAs that negatively regulate the post-transcriptional expression of functional genes (12). Aberrant expression of miRNAs has been broadly identified as an important factor in the development of metabolic disorders, such as insulin resistance and obesity (13,14). A recent study reported that miRNAs serve important roles in PCOS, and that the concentration of miR-155 in serum may be a biomarker for monitoring estroprogestinic
treatment (15). Dysregulated expression of another miRNA, miR-28-5p, has been frequently reported in the development of various types of cancers, including hepatocellular carcinoma, colorectal and prostate cancer (16-18), and most recently ovarian cancer (19). Shi and Teng (18) demonstrated that miR-28-5p expression was downregulated in hepatocellular carcinoma, and that cell proliferation and tube formation capacity were attenuated in cells expressing miR-28-5p. Our previous research indicated that hypermethylation of the miR-28-5p promoter reduces miR-28-5p expression and promotes cell proliferation in rat ovary granulosa cells (Meng et al, unpublished data); however, the downstream mechanisms underlying the effects of miR-28-5p remain unclear.

A previous study indicated that cell proliferation and follicular growth are activated during all stages of ovarian development in PCOS, indicating that cell death in the ovary may mediate follicular atresia (20). There is also evidence that granulosa cell death occurs following oocyte apoptosis (21). Therefore, cell proliferation and survival may be responsible for the development of PCOS.

In the present study, decreased expression of PROK1 was reported in cells transfected with miR-28-5p mimics. A dual-luciferase reporter assay indicated that miR-28-5p targeted the 3'-untranslated region (3'-UTR) of PROK1, attenuating PROK1 expression. Finally, cell proliferation, apoptosis and cell cycle distribution were evaluated following transfection with a PROK1-overexpressing plasmid, in the presence or absence of miR-28-5p mimics. The results of the present study indicated that miR-28-5p inhibited the progression of PCOS by targeting PROK1, which acts via the PI3K/AKT/mTOR signaling pathway, suggesting that PROK1 may be a potentially useful target in the treatment of PCOS.

Materials and methods

Cell culture and transfection. The rat ovary granulosa cells used in the present study were primary cells prepared by Procell Life Science & Technology Co., Ltd. (cat. no. CP-R050) via mechanical separation, collagenase digestion and differential adhesion. The cells were cultured in DMEM/F12 (Hyclone; GE Healthcare Life Sciences) with 15% fetal bovine serum (FBS; Fisher Scientific, Inc.) according to the manufacturer's protocols.

Animals. Female Sprague Dawley rats (age, 21 days; weight, 50±10 g; n=12) had free access to food and water under a controlled temperature of 23±2°C and humidity (55-65%) with a 12-h light/dark cycle. A PCOS model was established as previously described (22).

Plasmid construction. Total RNA was extracted using TRIzol reagent (Takara Bio, Inc.) from ovary granulosa cells (cat. no. CP-R050; Procell Life Science & Technology Co., Ltd.) according to the manufacturer's protocols. Reverse transcription reaction was performed using a Bester™ qPCR RT Kit (cat. no. DBI-2220; DBI Bioscience). The reverse transcription protocol was as follows: RNA denaturation at 65°C for 5 min; genomic DNA removal at 37°C for 5 min; reverse transcription at 37°C for 15 min; reverse transcriptase inactivation at 98°C for 5 min. The vector pCDNA3.0 (Guangzhou Vipotion Biotechnology, Co., Ltd.) was used for the overexpression of PROK1. The coding sequence of PROK1 was cloned using the following primers: Forward 5'-CGGGATCCATGAGGAGTGCTGTGCGAAGCT-3' and reverse, 5'-CCGCTGGACTGAAGCTTGGTACAAGTTGACATTCTCTAGGGCC-3'. The PCR mixture contained: 0.25 µl PrimeSTARHS DNA Polymerase (Clontech Laboratories, Inc.), 5 µl 5X PrimeSTAR buffer (Clontech Laboratories, Inc.), 2 µl dNTP Mix (Clontech Laboratories, Inc.); 0.5 µl primers (Sangon Biotech Co., Ltd.), 1 µl cDNA and 15.75 µl ddH2O, (Takara Bio, Inc.). The thermocycling conditions as follows: Initial denaturation at 94°C for 5 min, followed by 30 cycles of 94°C for 30 sec; 58°C for 30 sec; 72°C for 30 sec; 72°C for 5 min. Then, amplification products were ligated into the vector using BamHI and Xhol restriction sites and identified via double enzyme digestion. Empty vector was used as a NC. Animal experiments were performed in compliance with the ARRIVE guidelines (23), once ethical approval from the Nanfang Hospital Animal Ethics Committee had been obtained.

RNA extraction and reverse transcription-quantitative PCR (RT-qPCR). Total RNAs were extracted using TRIzol® reagent (Takara Bio, Inc.) according to the manufacturer's protocols. Total RNA (~400 ng) was reverse transcribed into cDNA using a Bestar qPCR RT kit (cat. no. 2200; Kay Biological Technology Co., Ltd.). Samples were incubated at 37°C for 15 min and at 85°C for 5 min. U6 small nuclear RNA was used as an internal reference for miR-28-5p expression; primers were obtained from Takara Bio, Inc. GAPDH was used as internal reference; primers were synthesized by Sangon Biotech Co., Ltd. A SYBR Green qPCR kit (Kay Biological Technology Co., Ltd.) was used to quantify miRNA or mRNA expression using a Stratagene Real time PCR system (Mx3000P; Agilent Technologies, Inc.). The amplification reactions were performed under the following conditions: Initial denaturation at 95°C for 5 min, followed by 40 cycles of 95°C for 15 sec, 58°C for 20 sec, and 72°C for 30 sec. All qPCR reactions were repeated at least three times. Relative miRNA or mRNA expressions were calculated by 2^−ΔΔcq method (24). The primers are listed in Table I.
Western blotting. Cells (~10^4) were lysed using protein lysis buffer (cat. no. P0013B; Beyotime Institute of Biotechnology) and total protein was then extracted. Subsequently, a Pierce BCA Protein Assay kit (cat. no. 23227; Pierce; Thermo Fisher Scientific, Inc.) was used to determine the total protein concentration. Equal amounts of total protein (10 µg) were separated by 12% SDS-PAGE. The PVDF membrane was placed in a 5% BSA blocking solution for 1.5 h at room temperature. The membranes were incubated at 4°C for 12 h with primary antibodies against PROK1 (1:500; cat. no. ab72807; Abcam), Bcl-2 (1:1,000; cat. no. ab196495; Abcam), Bax (1:1,000; cat. no. ab32503; Abcam), caspase-3 (1:500; cat. no. ab13847; Abcam), p62 (1:1,000; cat. no. ab1677; Cell Signaling Technology), cyclin D1 (1:200; cat. no. ab16663; Abcam), PI3K (1:1,000; cat. no. 4255S; Cell Signaling Technology), phosphorylated (p-)PI3K (1:1,000; cat. no. 4228T; Cell Signaling Technology), AKT (1:1,000; cat. no. 4691S; Cell Signaling Technology), p-AKT (1:2000; cat. no. 4060S; Cell Signaling Technology), mTOR (1:1,000; cat. no. 2983S; Cell Signaling Technology), p-mTOR (1:1,000; cat. no. 5536S; Cell Signaling Technology) and GAPDH (1:1,000; cat. no. ab8245; Abcam). The PVDF membrane was placed in a diluted horseradish peroxidase-conjugated rabbit anti-mouse immunoglobulin G secondary antibody solution and incubated for 1 h at room temperature. The intensities of labeled proteins were visualized using X-ray films (cat. no. 4741023951; Yestar Healthcare Holdings Co., Ltd.; http://www.yestarcorp.com/) with an ECL chemiluminescence detection kit (BeyoECL Plus; cat. no. P0018M; Beyotime Institute of Biotechnology) and ImageJ (version 1.8.0; National Institutes of Health) was used to quantify expression by densitometry.

Cell Counting Kit-8 (CCK-8). Cells (2x10^5) transfected with exogenous sequences were suspended in 200 µl culture medium and seeded into 96-well plates. The CCK-8 assay was conducted following culture under standard conditions (37°C) for a further 48 h. Then, 10 µl CCK-8 reagent (Beijing Solarbio Science & Technology Co., Ltd.) was added to each well, followed by incubation for a further 1 h at 37°C. Finally, the absorbance at 450 nm was detected for quantification of cell proliferation rates using a plate reader (BioTek Instruments, Inc.).

5-Ethynyl-2'-deoxyuridine (EdU) staining and flow cytometry. Cells (10^4) transfected with exogenous sequences were plated into 96-well plates and cultured for 48 h. Then, 100 µl EdU reagent (cat. no. B23151; Invitrogen; Thermo Fisher Scientific, Inc.) was added to each well after washing with PBS. The cells were incubated at 37°C for 2 h. 100 µl glycerol was added to each well for neutralization after cells were fixed with 4% paraformaldehyde for 15 min at room temperature. Subsequently, 0.5% Triton X-100 was added into each well, and cells were incubated for 10 min at room temperature, followed by washing with PBS. Apollo reagents and Hoechst 33342 staining reagent were used in an EdU kit (cat. no. CA1170; Beijing Solarbio Science & Technology Co., Ltd.) were used according to the manufacturer protocols. Then, the cells were incubated for 30 min at room temperature. Cells were analyzed via flow cytometry and the software used for the analysis was Cytomics FC500 Flow Cytometry CXP Software v2.0 (Beckman Coulter, Inc.).

Cell apoptosis and cell cycle distribution. Following transfection, 10^4 cells were cultured for 48 h, and the apoptosis of cells was then analyzed using the Annexin V-FITC/propidium iodide (PI) cell apoptosis detection kit (BD Biosciences) containing according to the manufacturer's protocols, and cells were incubated at 4°C for 30 min. Cells with positive signals for Annexin V-FITC and PI were in the late stage of apoptosis, whereas cells with positive signals for Annexin V-FITC only were in the early apoptosis stage. The total apoptosis ratio was defined by the summation of late-apoptotic and early-apoptotic cells.

Following transfection of cells with exogenous sequences, cells were stained with PI dye was incubated at 4°C for 30 min to determine the cell cycle distribution. A flow cytometer was used to analyze the cells and the software used for the analysis was Cytomics FC500 Flow Cytometry CXP Software v2.0 (Beckman Coulter, Inc.).

Dual-luciferase reporter assay. TargetScan 7.2 (http://www.targetscan.org) was used to identify the putative targets of miR-28-5p, and PROK1 was found to be one of the targets of miR-28-5p. The software was used according to previous report (25). The 3‘-UTR sequences of the PROK1 gene were obtained using primers (forward, 5’-CCGTCCTGAGTCCTCAACGGTGTATCGGTTGTTCACTTATCTGAG-3’; Sangon Biotech co., ltd.). RT-qPCR was performed using primers (reverse, 5’-GATGACACCAAACAGGGAATAG-3’ and forward, 5’-GACAGAAGATGGTTTCTATTCGCTTATTAG-3’; Sangon Biotech Co., Ltd.). Amplification products including the wild-type (WT) and Mut 3’-UTRs of PROK1 were then separately ligated into Psi-cHECK2 plasmids (cat. no. C8021; Promega Corporation). Psi-CHECK2 plasmids were co-transfected with miR-28-5p mimics or NC into cells using Lipofectamine 2000™ and incubated for 48 h at 37°C. The luciferase activities of firefly and Renilla were measured with the Dual-Luciferase Reporter Assay system (cat. no. C8021; Promega Corporation) according to the manufacturer’s protocol. Firefly luciferase activity was normalized.
to Renilla luciferase activity. Cell lysates were collected using a GloMax machine (Promega Corporation).

Statistical analysis. Data from >3 biological replicates for each assay were presented as the mean ± standard deviation, and data were analyzed using SPSS 16.0 software (SPSS, Inc.). Student’s t-test was used to analyze differences between two groups, and one-way analysis of variance followed by Tukey’s test was performed to compare differences between three or more groups.

Results

miR-28-5p suppresses PROK1 expression by targeting its 3’-UTR. To investigate the relationship between miR-28-5p and PROK1, the mRNA expression levels of PROK1 were analyzed by qPCR, and the protein expression was evaluated via western blotting following transfection of ovary granule cells with miR-28-5p mimics. It was demonstrated that mRNA expression of PROK1 was significantly suppressed in miR-28-5p overexpressing cells compared with the NC (Fig. 1A). Similarly, the protein expression of PROK1 was significantly downregulated following the transfection of ovary granule cells with miR-28-5p mimics compared with the NC (Fig. 1B). Conversely, transfection with miR-28-5p inhibitor induced the opposite results in PROK1 expression in ovary granule cells (Fig. 1A and B).

To determine whether miR-28-5p binds to PROK1, the 3’-UTR of PROK1 was cloned and inserted into psi-CHECK2 plasmids. For comparison, the binding site predicted by the online database TargetScan was mutated, and then also inserted into psi-CHECK2 plasmids (Fig. 1C). Cells co-transfected with WT 3’-UTR and miR-28-5p exhibited significantly reduced luciferase activity compared with cells transfected with NC or MUT 3’-UTR (Fig. 1D). The results indicated that miR-28-5p negatively regulated PROK1 expression via direct binding to its 3’-UTR.

PCOS development promoted by PROK1 overexpression is inhibited by miR-28-5p via the PI3K/AKT/mTOR signaling pathway. PCOS is a common endocrine-reproductive disease in females and it has been demonstrated that ovary granular cells in PCOS patients showed increased apoptotic rates (17,26). To verify whether PROK1 promotes PCOS in rat ovary granule cells, a PROK1 overexpression plasmid was constructed. Additionally, a rescue experiment series was designed to further demonstrate the influence of the interaction between miR-28-5p and PROK1 on rat ovary granule cell survival. As presented in Fig. 2A, a western blot assay verified the overexpression of the PROK1 plasmid, revealing that PROK1 was significantly upregulated in cells transfected with the PROK1 overexpression plasmid compared with empty vector. The overexpression of PROK1 protein was accompanied by significantly increased cell proliferation rates in rat ovary granule cells (Fig. 2B and C); however, the effects of PROK1 upregulation on proliferation were attenuated by co-expression of PROK1 and miR-28-5p mimics, compared with cells transfected with pcDNA-ProK1 alone.

Additionally, apoptosis was significantly suppressed in rat ovary granule cells overexpressing PROK1 compared with

Figure 1. miR-28-5p suppresses PROK1 expression by targeting the 3’-UTR of PROK1. (A) mRNA expression of miR-28-5p and PROK1 in rat ovary granule cells. (B) Protein expression of PROK1. (C) WT and MUT sequences in the 3’-UTR of PROK1 involved in binding with miR-28-5p. **P<0.01, as indicated. (D) Dual-luciferase reporter assay confirming an interaction between miR-28-5p and the PROK1 3’-UTR. All data are presented as the mean ± SD. **P<0.01 vs. NC (n=3). miR-28-5p, microRNA-28-5p; PROK1, prokineticin 1; 3’-UTR, 3’-untranslated region; NC, negative control; WT, wild type; MUT, mutant.
empty vector (Fig. 3A and B), and a significantly increased percentage of these cells were in S phase (Fig. 3C and D). These tendencies were attenuated by co-transfection with miR-28-5p mimics, which elevated cell apoptosis, and redistributed cells to the G0/G1 phase compared with pcdna-ProK1 alone. The expression of apoptosis- and cell cycle-associated proteins was investigated via western blotting (Fig. 3E). Compared with empty pcdna3.0 vector, transfection with pcDNA3-ProK1 significantly upregulated Bcl-2 and cyclin d1 expression, whereas caspase-3, Bax and p62 expression levels were downregulated; however, co-transfection with miR-28-5p mimics attenuated these effects. Additionally, it was revealed via western blotting that PI3K/AKT/mTor activation was also potentiated in ProK1-overexpressing cells compared with the control (Fig. 3F). Conversely, this activation was reversed by co-transfection with miR-28-5p mimics. These results indicated that ProK1 promoted the proliferation and suppressed the apoptosis of rat ovary granule cells via the PI3K/AKT/mTor signaling pathway.

Discussion

Abnormal expression of miRNAs has been reported in a variety of cancers, resulting in increased attention regarding the potential pathogenic effects of miRNAs on cancer initiation and progression (27,28). MiR-28-5p has been reported to be closely associated with tumor carcinogenesis. For example, hepatocellular carcinoma progression was suppressed by miR-28-5p via the targeting of insulin-like growth factor-1 (IGF-1) (18); miR-28-5p also exerts a number of antitumor effects in renal cell carcinoma via the regulation of Ras-related protein Rap-1B (29). Additionally, miR-28-5p was reported to be downregulated in colorectal cancer tissue compared with in normal colon tissue, and to suppress colorectal cancer cell proliferation, migration and invasion (30). Conversely, certain studies have suggested that miR-28-5p may also serve as a potential marker for the development and progression of cancers (19,31). For example, it was reported that miR-28-5p may promote the development and progression of renal cell carcinoma by increasing the risk of chromosomal instability and promoting checkpoint weakness (31). Additionally, miR-28-5p acted as a carcinogenic biomarker to enhance ovarian cancer cell proliferation, inhibit cell apoptosis, and contribute to migration and invasion (19). In the present study, it was demonstrated that miR-28-5p suppressed cell proliferation, and induced cell cycle arrest and apoptosis in PCOS. Therefore, miR-28-5p regulate PCOS and inhibit PCOS progression.

It has been previously reported that miRNAs contribute to the regulation of various physiological processes by targeting mRNAs (32,33). And the activation or suppression of the mRNA by these miRNAs could induce positive or negative effects on the progression of various diseases, including PCOS (34-36). For example, miR-324-3p expression was observed to be reduced in PCOS; miR-324-3p inhibited the proliferation, and promoted the apoptosis of granulosa cells by targeting Wnt-2B (37). In addition, miR-93 promoted ovarian granulosa cell proliferation by targeting cyclin-dependent kinase inhibitor 1A in PCOS (38), miR-483 suppressed cell proliferation in PCOS by targeting IGF-1 (39). In the present study, it was demonstrated that miR-28-5p negatively regulated ProK1. The present findings suggested that miR-28-5p was involved in PCOS by regulating ProK1. ProK1 serves an important role in angiogenesis, and it was observed that the concentration of VEGF in the ovaries of females with PCOS
was increased compared with healthy females (40). In addition, previous studies also demonstrated that PROK1 promoted human umbilical vein endothelial cell proliferation, differentiation and survival (7,41). Furthermore, PROK1 also promoted the morphogenesis of vascular endothelial cells into tube-like structures, as observed in 2D-model experiments (42). In the present study, it was demonstrated that the overexpression of PROK1 promoted cell proliferation, and inhibited cell cycle arrest and apoptosis in PCOS. The present findings further indicated that PROK1 may accelerate PCOS progression.

The cell cycle is a series of important cellular events that occurs during various physiological and pathological processes, and the initiation of G1 phase is critical during cell cycle progression (43,44). Cyclins are important regulators of cell cycle progression. Cyclin D1, as a key regulatory protein of G1 phase progression, has been established as a major indicator of cell cycle progression, with increased sensitivity when compared with other cyclins (45). Specifically, cyclin D1 promotes the transition from G1 to S phase by binding and activating cyclin-dependent kinase 4 (CDK4), which accelerates cell proliferation (46). Previous studies have reported that p62 serves important roles in cell cycle progression by reducing the ubiquitinylated-exacerbated degradation of CDK1 and accelerating G2/M phase progression (47,48). In the present study, it was demonstrated that PROK1 upregulated cyclin D1 expression and downregulated p62 expression in ovary granule cells, whereas miR-28-5p mimics reversed these effects, which indicated that miR-28-5p promoted cell cycle arrest by targeting PROK1 in PCOS.

Apoptosis is an important biological death process that is genetically controlled, which serves as an important mechanism for maintaining a stable internal environment (49). Caspase-3, Bcl-2 and Bax are central functional proteins in apoptosis pathways (50). Caspases are a class of cysteine aspartic acid-specific proteases in the cytoplasm; caspase-3, also known as 32 kDa cysteine protease, is a member of the interleukin-lp-converting enzyme family (51). Caspase-3 is a common downstream factor in various apoptotic pathways. Bcl-2 exhibits anti-apoptotic effects, whereas Bax protein is structurally similar to Bcl-2 but induces opposing effects on cell apoptosis (52). In the present study, decreased apoptosis in cells co-transfected with miR-28-5p and PROK1 was observed when compared with cells transfected with PROK1 alone, as determined by flow cytometry. Cell cycle distribution in cells co-transfected with mimics and PROK1 compared with PROK-transfected cells alone, as determined by flow cytometry. (E) Altered expression of Bax, Bcl-2, Caspase 3, p21 and cyclin D protein induced by PROK1 overexpression were reversed by miR-28-5p. (F) Activation of the PI3K/AKT/mTOR signaling pathway following transfection with pcDNA-PROK1, or co-transfection with pcDNA-PROK1 and mimics. All data are presented as the mean ± SD. *P<0.05, **P<0.01 and ***P<0.001, as indicated (n=3). miR-28-5p, microRNA-28-5p; PROK1, prokineticin 1; mimics, miR-28-5p mimics; p-, phosphorylated; PI, propidium iodide.
study, PROK1 decreased caspase-3 and Bax expression, and increased Bcl-2 expression, whereas treatment with miR-28-5p mimics attenuated these alterations in protein expression. Therefore, it was further indicated that miR-28-5p promoted cell apoptosis by targeting PROK1 in PCOS.

The PI3K/AKT pathway serves an important role in mediating ovarian granular cell function (53-56). PI3K activation induces the phosphorylation of AKT proteins and mTOR gene expression, leading to the activation of downstream signaling associated with cell proliferation, survival and angiogenesis (57). Patients with PCOS show decreased ovary granular cells caused by decreasing viability of cell proliferation or survival (20). Previous studies showed that the PI3K/AKT signaling pathway is activated in PCOS (53,54). Upregulation of Wnt-5a, which acts as a proinflammatory factor, is a signal to PI3K/AKT, activating inflammation and oxidative stress in the granulosa cells of patients with PCOS via NF-κB (53). Additionally, Xin et al (55) reported that patients with insulin resistance underwent significant alterations to the PI3K/AKT/GSK3 signaling pathway. The present study demonstrated that PROK1 affect the progression of PCOS involved in the PI3K/AKT/mTOR signaling pathway.

In conclusion, these findings suggested that miR-28-5p could attenuate the progression of PCOS by targeting the 3'-UTR of PROK1 which may involved in the PI3K/AKT/mTOR pathway, indicating that the miR-28-5p/PROK1 axis may be a potential therapeutic target for patients with PCOS.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

LM made substantial contributions to the conception and design of the study. HY contributed to data acquisition, data analysis and data interpretation. CJ performed some experiments, analyzed the data and drafted the article. SQ was involved in the design of the experiments and critically revised the manuscript for important intellectual content in the submission process. All authors agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of the work are appropriately investigated. All authors have read and approved the final manuscript.

Ethics approval and consent to participate

Animal experiments were approved by the Nanfang Hospital Animal Ethic Committee, and conducted in accordance with the ARRIVE guidelines.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

References


