miR-139 Controls Viability Of Ovarian Cancer Cells Through Apoptosis Induction And Exosome Shedding Inhibition By Targeting ATP7A

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Fang Xiao
Songshu Xiao
Min Xue

Department of Obstetrics and Gynecology, Third Xiangya Hospital of Central South University, Changsha, Hunan 410013, People’s Republic of China

Background: Emerging proof suggests that microRNA (miRNA) malfunction is correlated to the generation and development of multiple malignancies. It has been proven that miRNA (miR)-139 represses a variety of malignancies. However, the understanding of its impact on ovarian cancer (OC) is insufficient. We aimed to investigate the expression and function of miR-139 in OC.

Methods: The expression of mir-139 was detected and recorded, and the relationship of mir-139 remaining OC cells was explored. At the same time, we studied the correlation between ATP7A and mir-139 by the luciferase reporter test, Western blot, and quantitative real-time reverse transcription PCR.

Results: The expression of miR-139 was remarkably downregulated in OC specimens. Furthermore, excessive miR-139 expression noticeably inhibited the migration, colony generation, proliferation, and invasion of OC cells. In addition, excessive miR-139 expression remarkably repressed the death and the expression of proteins related to cell death in OC cells, as well as inhibited the shedding of exosomes. According to the luciferase reporter test, Western blot, and quantitative real-time reverse transcription PCR, miR-139 directly targeted ATP7A. Furthermore, the expression of ATP7A was found to be negatively related to miR-139 levels in OC specimens. It was revealed via a rescue experiment that excessive ATP7A expression counteracted the repressive effect of miR-139 in OC cells.

Conclusion: It was revealed via an in vivo study that miR-139 remarkably inhibited the growth of malignancies by downregulating ATP7A in nude mice. miR-139 represses the development of malignancies in OC by directly targeting ATP7A, offering an innovative approach for molecular therapy of OC.

Keywords: apoptosis, ATP7A, exosome shedding, miR-139, ovarian cancer

Introduction
Ovarian cancer (OC) has a high mortality rate worldwide. Despite huge progress in OC therapy, the 5-year overall survival for OC patients is 35% to 38%, which can be mainly attributed to diagnosis at terminal stage, frequent metastasis, and the fast generation of chemoresistance. Consequently, elucidating the etiology of the illness and discovering crucial agents that can assist in the diagnosis and innovative therapy of OC are necessary.

MicroRNAs (miRNAs) are a group of small endogenous RNAs without coding functions that consist of 18 to 22 nucleotides and inhibit the expression of target genes via direct binding with the 3’ untranslated regions (3’-UTRs) of their mRNA, thereby...
downregulating gene expression both at the mRNA and protein levels. Previous research has proven that miRNAs contribute to the modulation of multiple malignancies and thus offer promising treatment options. It has been demonstrated that some miRNAs can malfunction, such as miR-98-5p, miR-92, miR-141, and miR-93-5p, and result in the generation and development of OC. It was revealed that miR-139, which modulates various cellular processes including invasion, cell cycle, proliferation, and vessel generation, malignancies in some kinds of malignancies. Emerging proofs about colorectal cancer, hepatocellular carcinoma, bladder cancer, oral tongue squamous cell carcinoma, and esophageal squamous cell carcinoma have indicated that miR-139 suppresses malignancies and is thus commonly downregulated in malignancies.

Through mediating Wnt/β-catenin signaling pathway, miR-139 is involved in the cell growth and metastasis of bladder cancer cells. By direct targeting NOB1, miR-139 can induce the cell apoptosis and inhibit the metastasis of cervical cancer cells. In glioblastoma multiforme, miR-139 can hinder the cell migration and invasion of cancer cells through targeting ZEB1/2.

Copper-transporting ATPases are intracellular transporters that maintain cellular copper homeostasis. Dysregulated expression of ATP7A in ovarian cancer cells can alter the amount of cellular copper, which could then impact the activities of other transporters that import or export platinum derivatives. Increased expression of ATP7A in ovarian cancer cells predicts short survival for patients treated with platinum derivatives and suggests a role for ATP7A in OC. Whether miR-139 regulated ATP7A in ovarian cancer cells, thus to affect the cell proliferation and apoptosis? This remains to be studied. Consequently, our research aimed to explore miR-139 expression in clinical OC specimens and OC cell lines to elucidate its molecular mechanism and impact on OC development and metastasis.

Materials And Methods

Clinical Specimens

Biopsy specimens of OC and surrounding non-malignant tissues were obtained from 31 patients treated at our hospital. The surrounding tissues were 3 cm away from the malignancy fringe with no obvious malignant cells, as assessed by an expert pathologist. All specimens were immediately frozen at −80°C after surgery using liquid nitrogen. No patients had undergone radiation therapy or chemotherapy prior to surgery. Fully informed consent was acquired from every participant, which won approval from Research Ethics Committee of our hospital.

Cell Culture And Transfection

Human OC cell lines HO-8910 and SKOV3 were purchased from Tumor Cell Bank of the Chinese Academy of Medical Science (Beijing, P.R. China). RPMI-1640 medium with 10% FBS was used and incubated at 37°C with 5% CO2. MiR-139 mimics, inhibitor, and negative control (NC) mimics were purchased from GenePharma (Shanghai, P.R. China). ATP7A overexpression plasmids (pCDNA3.1-ATP7A) were purchased from the same company. SKOV3 cells were seeded in 24-well plates and were transfected with vectors (5 µg) or miRNA mimics/NC (100 mM) using Lipofectamine 2000 in serum-free media.

RNA Isolation And Quantitative Real-Time Reverse Transcription PCR (qRT-PCR)

TRIZol reagent was used to isolate total RNA from the cultivated cells or tissues as instructed. SmartSpec Plus spectrophotometer was used for RNA quantification and quality check by purity assessment based on A260/A280 ratio. SYBR PrimeScript miRNA RT PCR Kit was used to measure the expression of miR-139-5p. The corresponding primers of miR-139-5p and U6 were acquired from Applied Biosystems. Two micrograms of total RNA underwent reverse transcription (RT) to generate cDNA with the help of PrimeScript RT Reagent Kit using oligo (dT) primers (Takara). SYBR Green Real-Time Master Mix was used for qRT-PCR. GAPDH served as an internal control. The expression levels of ATP7A mRNA and miR-139 were normalized against GAPDH or U6 levels based on the 2−ΔΔCt approach.

Luciferase Reporter Assay

Wild type (WT) ATP7B or ATP7A luciferase reporter gene vectors (called ATP7B-WT-3′UTR or ATP7A-WT-3′ UTR), as well as mutant ones were constructed. HEK293T cells were co-transfected with the corresponding vectors and miR-139 inhibitors or mimics, and then cultured overnight in 24-well plates. Dual Luciferase Reporter Assay (Promega, USA) was performed 48 h after transfection.
Separation Of Exosomes
Exosomes were separated using differential centrifugation approaches. Cultures were centrifuged and the supernatant was then centrifuged at 2000 ×g for 20 min to acquire apoptotic bodies, and then at 12,200 ×g for 60 min to harvest the microvesicles. Another supernatant was passed through 0.22-μm filters and was centrifuged at 120,000 ×g for 2 h to obtain the exosomes. One milliliter of TRizol reagent was added to every tube of acquired pellet to isolate total RNA as per the manufacturer’s instructions.

Proliferation Test
Cell Counting Kit-8 was used to assess cell proliferation. In brief, cells were cultured in 96-well plates. CCK-8 reagent was added to the corresponding wells, and allowed to incubate for 2 h. Media with CCK-8 were subsequently transferred to new 96-well plates and absorbance was measured.

Cell Cycle Test
Cells went through 12-h starvation to synchronize the cultures prior to 24-h re-activation with 10% FBS. Cells were fixed and FACS Caliber flow cytometer was subsequently used to categorize the cells. Flowjo software (Treestar Inc., USA) was used to evaluate the cell phase distribution.

Colony Generation Test
Cells were incubated with 0.25% trypsin. Nearly five hundred cells were seeded in 6-well plates (250 cells/mL). Cells were fixed for 10 min using anhydrous ethanol and stained for 30 min with 0.1% crystal violet. Colonies comprising 50 cells were counted and the relative colony number was acquired. Colony-generating capability was assessed by obtaining the ratio of the number of colonies generated in the transfection group to that in the control group multiplied by 100.

Flow Cytometry (FC)
PBS was used to wash the acquired cells. A million cells were isolated from each specimen, and stained with an Annexin V-FITC/propidium iodide (PI) kit. FC (BD FACS Aria; BD Biosciences, Franklin Lakes, NJ) was then performed to detect positive cells 48 h after transfection.

Cell Migration Test
Transwell test was performed to examine cell migration. In brief, 5×10⁴ cells were suspended in DMEM without serum and were seeded on the top well of 24-well poly-carbonate transwell filters. DMEM with 10% FBS was supplemented to the bottom well. Cells at the top surface were scraped off after a 24-h incubation and those at the bottom surface were fixed, stained, and quantified.

Cell Invasion Test
Twenty-four-well transwell chambers with 8-µm pore size polycarbonate membranes were used for the invasion test. Transwell chambers initially coated with Matrigel were seeded with cells. Cells suspended in 200 µL of DMEM without serum were seeded on the top chamber, while those in 8 µL of DMEM with 10% FBS were seeded on the bottom chamber. Cells that did not undergo invasion were eliminated from top chamber using a cotton swab after 24 h of incubation, and those that invaded the bottom were fixed and stained with 0.1% crystal violet. Cells were quantified in 6 random fields for every well. Relative folds of cells with invasion were displayed.

Xenograft Malignancy Model
BALB/c mice were purchased from Peking Union Medical College (Beijing, China) and reared in sterile conditions. Mice received subcutaneous injection of SKOV3 cells (5×10⁶ per 0.1 mL) through their back. The volume and weight of the mice were examined. The volume of the malignancies of the NC or miR-139 groups (n=6 each) were measured every 3 days based on the following formula: volume = 0.5 × length × width × width. Mice were sacrificed 21 days subsequent to injection, and the malignancies were excised and prepared. All mice were treated in accordance with protocols approved by Institutional Animal Care and Use Committee of Third Xiangya Hospital of Central South University, and the committee approved the experiment.

Western Blot (WB)
Lysates were homogenized using a lysis buffer (Beyotime, China), while Bradford assay was used for protein quantification. Proteins were assessed via standard SDS-PAGE. Various concentrations (15%–80%) of Tris–HCl polyacrylamide gels (Bio-Rad) were used to isolate proteins, which were then transferred onto PVDF membranes. After blocking, blots were incubated with primary antibodies, including anti-ATP7A, anti-Bax, anti-β-actin, and anti-Bcl2 antibodies. Another incubation was carried out with secondary antibodies. Enhanced chemiluminescence (ECL) plus detection reagent (Pierce, Rockford, IL, USA) was used to examine immunoreactive bands.
**Statistical Analysis**
Results were displayed as mean ± SEM. Differences between various groups were evaluated using two-tailed, unequal-variance Student’s t-test or ANOVA prior to Tukey’s post hoc analysis. A p-value<0.05 was regarded to be statistically significant.

**Results**
**miR-139 Was Downregulated In OC Specimens**
Our research explored miR-139 expression in 31 OC specimens matched to normal specimens to reveal the impact of miR-139 on the generation and development of OC. MiR-139 was found to be downregulated in OC specimens compared to control specimens (Figure 1). Moreover, both transcription and translation of ATP7A were remarkably increased in OC specimens in comparison with those in the control specimens. These findings indicate that the expression of miR-139 was inhibited in OC and that it potentially modulated ATP7A.

**miR-139 Directly Targeted ATP7A To Impair Exosome Secretion-Modulated miR-139 Trafficking In SKOV3 Cells**
Bioinformatics analysis indicated that miR-139 targeted ATP7A. Excessive miR-139 expression impaired ATP7A 3′-UTR function in SKOV3 cells, while miR-139 inhibition improved it (Figure 2A and B). Consequently, miR-139 regulated ATP7A expression probably via direct 3′-UTR binding. Moreover, excessive miR-139 expression inhibited ATP7A translation. In contrast, miR-139 inhibition increased ATP7A translation.

![Figure 1](image1.png)

**Figure 1** MiR-139 was downregulated in OC specimens. (A,B) Expression levels of miR-139 (A) and ATP7A (B) in OC specimens and in matched surrounding non-malignant tissues were measured using qRT-PCR. C-D, Representative immunoblots (C) and quantitative assessment of ATP7A (D) in OC specimens and in matched surrounding non-malignant tissues. Results are displayed as means ± SEM, n=31. **p<0.01 vs tumor group.
Figure 2. MiR-139 directly targeted ATP7A to impair exosome secretion-modulated miR-139 trafficking of SKOV3 cells. (A,B) Link between miR-139 and 3′-UTR of ATP7A mRNA was assessed by luciferase reporter assay. (C-D) Representative immunoblots and quantitative assessment of ATP7A in SKOV3 cells that underwent 48-h transfection with miR-139 (C) or miR-139 inhibitor (D). (E-F) Expression of miR-139 in exosomes from SKOV3 cells that underwent 48-h transfection with miR-139 (E) or miR-139 inhibitor (F) was assessed using qRT-PCR. Results are displayed as means ± SEM. **p<0.01 vs NC group.
translation in SKOV3 cells (Figure 2C and D). We isolated exosomes and examined miR-139 expression to explore the impact of the miR-139-ATP7A interaction on exosome secretion-modulated miR-139 trafficking. Excessive miR-139 expression decreased miR-139 levels in exosomes (Figure 2E and F), but miR-139 inhibition increased it. These findings indicated that miR-139 directly targeted ATP7A to impair exosome secretion-modulated miR-139 trafficking in SKOV3 cells.

**miR-139 Repressed Proliferation Of SKOV3 Cells**

SKOV3 cells were transfected with miR-139 mimics, then CCK8 and FC were performed to assess and explore the impact of miR-139 on OC proliferation. It was revealed via CCK8 that the proliferation of SKOV3 cells was remarkably repressed with excessive miR-139 expression compared to that of NC (Figure 3A). It was revealed via FC that the proportion of cells in S phase was lower in specimens transfected with miR-124 mimics than in those transfected with NC (Figure 3B). Furthermore, excessive miR-139 expression remarkably repressed colony generation of SKOV3 cells (Figure 3C and D). These findings suggested that miR-139 repressed proliferation of SKOV3 cells.

**miR-139 Induced Death Of SKOV3 Cells**

FC was carried out with Annexin V/PI staining to evaluate the impact of miR-139 on cell death. Cells that were transfected with a miR-139 mimic displayed a remarkably elevated death rate in comparison to those of NC (Figure 4A and B). Proteins involved in cell death are crucial factors; thus, our research examined alterations in the expression of Bax and Bcl2 in SKOV3 cells. Bax was noticeably upregulated while Bcl2 was remarkably downregulated in SKOV3 cells that were transfected with a miR-139 mimic compared to those in the NC (Figure 4C–E). These findings proved that miR-139 induced death of SKOV3 cells.

**miR-139 Repressed Invasion And Migration Of SKOV3 Cells**

Our research subsequently explored the contribution of miR-139 to the invasion and migration of SKOV3 cells. Cell migration was assessed via migration test using transwell chambers without coating. It was revealed that
excessive miR-139 expression remarkably inhibited migration (Figure 5A and B). Cell invasion was assessed via invasion test using chambers with preliminary Matrigel coating. Invasion by cells transfected with miR-139 was less than those transfected with NC (Figure 5C and D). These findings suggested that miR-139 repressed the invasion and migration capabilities of SKOV3 cells.

Excessive ATP7A Expression Counteracted Cell Growth Arrest And Migration Inhibition Induced By miR-139 Mimics In SKOV3 Cells

In SKOV3 cells transfected with miR-139 mimics, migration, proliferation, and cell death tests were carried out with or without pcDNA3-ATP7A to explore whether ATP7A was crucial for the impact of miR-139 on repressing malignancy in OC. Excessive ATP7A expression counteracted cell growth arrest and cell death induced by miR-139 mimics (Figure 6A and B). Furthermore, excessive ATP7A expression also counteracted migration inhibition induced by miR-139 mimics (Figure 6C and 6D). These findings proved that ATP7A modulation was crucial to miR-139 in regulating cell death, migration, and proliferation of SKOV3 cells.

miR-139 Repressed OC Malignancy Growth In Vivo

Male BALB/c nude mice were injected with SKOV3 cells in their flanks to verify whether miR-139 repressed OC growth. The malignancy volume and weight was remarkably lower and the expression of ATP7A in mice with excessive miR-139 expression than in those with the NC (Figure 7A–D). This in vivo finding indicated that miR-139 repressed OC generation.

Discussion

Our research indicated that miR-139 expression was repressed in OC, bringing about excessive ATP7A expression and increasing cell death, proliferation as well as migration of OC cells. We revealed that miR-139 could modulate the generation and development of OC via the
regulation of ATP7A. Our research sheds light on the impact of miR-139 on OC generation and indicates that miR-139/ATP7A could serve as a promising target for OC therapy.

Whether a certain miRNA is oncogenic or malignancy-repressing is mainly dictated by its target genes in various cellular reactions.\(^{23}\) MiR-139 has been previously found to repress some kinds of malignancies.\(^{24}–^{26}\) MiR-139 targeted multiple crucial genes that contribute to motility.\(^{27,28}\) MiR-139-5p bound to NOTCH1 and AMFR mRNA, thus inhibiting the migration of cancer cells.\(^{29}\) Expression of miR-139 was repressed in glioblastoma, resulting in the inhibition of the migration of glioblastoma cells by targeting ZEB1.\(^{30}\) Furthermore, miR-139 expression was repressed in prostate cancer, resulting in the inhibition of migration and proliferation by targeting SOX5.\(^{31}\) Our study suggested that the expression of miR-139 was repressed in OC. Excessive miR-139 expression induced the death and inhibited the migration and proliferation of OC cells. Furthermore, in vivo findings indicated that excessive miR-139 expression remarkably impaired malignancy growth. These findings indicated that miR-139 exerted an essential malignancy-repressing impact on OC.

It has been demonstrated that miR-139 counteracts malignancy and represses proliferation and invasion by targeting various crucial genes.\(^{32}\) It was suggested via TargetScan that miR-139 targeted ATP7A. Some studies have indicated that ATP7A was upregulated and involved in the development of various malignancies, such as esophageal squamous cell cancer and breast cancer.\(^{33}–^{35}\) A recent study revealed that ATP7A expression was upregulated and contributed to the invasion and proliferation of OC cells.\(^{33,36}\)

Figure 5 miR-139 inhibited OC cell invasion and migration. (A) Images revealing migrated SKOV3 cells transfected with NC or miR-139 mimics (miR-139) on the bottom surface of transwell membranes. (B) Number of migrated SKOV3 cells of various groups in 5 random fields under a microscope. (C) Images revealing invaded SKOV3 cells transfected with NC or miR-139 mimics (miR-139) on the bottom surface of transwell membranes. (D) Number of invaded SKOV3 cells of various groups in 5 random fields under a microscope. Results are displayed as means ± SEM. \(^{**}\)p<0.01 vs NC group.
which counteracted the effect of miR-139. Our research proved through a luciferase reporter assay that miR-139 directly targeted ATP7A. Additionally, excessive miR-139 expression downregulated the transcription and translation of ATP7A, while excessive ATP7A expression impaired miR-139-mediated inhibition of proliferation and migration. Clinical samples showed that the expression of miR-139 was inversely related to that of ATP7A.

Figure 6 Excessive ATP7A expression counteracted the effects of miR-139 mimics. (A) Transfection of ATP7A counteracted cell growth arrest mediated by miR-139 mimics in SKOV3 cells. (B) Transfection of ATP7A counteracted cell death triggered by miR-139 mimics in SKOV3 cells. (C–D) Transfection of ATP7A counteracted migration prohibition triggered via miR-139 mimics in SKOV3 cells. Results are displayed in the form of means ± SEM. **p<0.01 vs NC group, *p<0.05 vs miR-139 group.

Figure 7 miR-139 inhibited malignancy growth in vivo. (A) Photographs revealed malignancies in xenografts of SKOV3 cells with and without excessive miR-139 expression. (B) Volume of the xenograft malignancy. (C) Weight of the xenograft malignancy. (D) Representative immunoblots of ATP7A in the xenograft. Results are displayed in the form of means ± SEM. n=6, *p<0.05 vs NC group.
Conclusions
In summary, this research indicated that miR-139 represses OC invasion and proliferation by targeting the expression of ATP7A. These findings could provide a potential innovative target for the treatment of OC.

Ethics Approval And Informed Consent
Fully informed consent was acquired from every participant, which won approval from Research Ethics Committee of Third Xiangya Hospital of Central South University (Changsha, China).

Author Contributions
All authors contributed to data analysis, drafting or revising the article, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

Disclosure
The authors declare that there are no conflicts of interest.

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