Original Article
MiR-302a inhibits the tumorigenicity of ovarian cancer cells by suppression of SDC1

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Abstract: MicroRNA plays an important role in tumor proliferation and cell cycle. In this study, we suggested the level of miR-302a was increasing in the human ovarian cancer cells compared to the normal cells. We aimed to explore the role of miR-302a downregulation in human ovarian cancer cells. Functional studies demonstrate overexpression of miR-302a could significantly suppress ovarian cancer cells proliferation and promote the cell cycle progress. In vitro reporter assay suggested SDC1 is a direct target gene of miR-302a. Furthermore, the expressions of miR-302a in ovarian cancer cells were inversely correlated with that of SDC1. Upregulation of SDC1 could rescue the effect of overexpressed miR-302a in the ovarian cancer cells. These findings provide evidence that miR-302a plays a key role in inhibition of the ovarian cancer cells proliferation, and enhancing the cells’ apoptosis through targeting SDC1, and strongly suggest that exogenous miR-302a may have therapeutic value in treating ovarian cancer.

Keywords: miRNA, miR-302a, SDC1, ovarian cancer

Introduction

Ovarian cancer (OC) is the most leading cause of gynecologic cancer-related deaths in Europe and North America, and the burden of this devastating cancer is expected to increase further in the coming years. Due to the difficulty of effectively diagnosing OC at its early stage, and optimal debulking with platinum-based chemotherapy remains the cornerstone of management. Unfortunately, despite frequent initial responses to chemotherapy, these tumors almost invariably relapse. Therefore, elucidating the molecular mechanisms involved in OC is essential for developing cancer prevention strategies and possible guiding disease management in the clinic [1].

In this several years, a class of novel non-coding RNAs called MicroRNAs (MiRNAs) had been discovered in plants and animals. MicroRNAs (miRNAs) include 18-26 nucleotides, which post-transcriptionally regulate gene expression in multicellular organisms by affecting both the stability and translation of mRNAs. In the process of tumor formation, the abnormal expression or the loss of the dynamic balance between oncogenes and tumor suppressor genes, leads tumorigenesis and development of cancer. MiRNA, as an important regulation factor of gene expression, also involved in tumor formation and progression. Considerable evidence showed critical functions for miRNAs in diverse biological processes, such as proliferation [2-10], apoptosis [11-18], angiogenesis [19-25], cell differentiation [26-28], adhesion and metastasis [29] of tumor cells. Therefore, downregulation of certain miRNA expression may cause a variety of imbalance, including many kinds of cancers. Previous studies have confirmed cancer-specific miRNAs in many types of cancers, involved breast cancer [30], lung cancer [31], hepatocellular carcinoma [32] and so on.

The miR-302-367 cluster consists of four highly-homologous miRNA members, which are transcribed together as a noncoding RNA cluster containing mir-302b, mir-302c, mir-302a, mir-302d, and mir-367 in a 5’-to-3’ direction [33]. Most studies of this cluster have focused on the maintenance of stemness [34-38] and the ability of the cluster to reprogram somatic cells into induced pluripotent stem cells (iPSCs)
To date, miR-302s have been proven to post-transcriptionally regulate CCND1 and CDK4, therefore affecting cell cycle progression. Other studies have demonstrated the tumor suppressive activity of miR-302 in human pluripotent stem cell by both the CCNE-CDK2 and CCND-CDK4/6 pathways in G1-S cell cycle transition [43]. In contrast, a recent study demonstrated that the miR-302-367 cluster compromised the maintenance of glioma-initiating cells (GiCs), strongly inhibited the clonogenicity of GiCs and promoted the loss of stem-like proteins, including OCT4 and NANOG, as well as the down-regulation of SOX1 and SHH [44, 45]. Consistent with these findings, another report demonstrated the importance of the miR-302-367 cluster in cell differentiation by showing that the cluster controls mes-endodermal fate specification [46]. The subclass comprised of miR-302a-d (miR-302s) has also been shown to inhibit tumorigenicity and induce apoptosis in various tumors and cancer cells, including MCF7 breast cancer cells, HepG2 hepatocellular carcinoma cells, and Tera-2 embryonal teratocarcinoma cells [43]. So, the relationship between miR-320a and cancers needs to be investigated further.

In this research, we found that miR-302a was frequently down-regulated ovarian cancer cells. Next, we analyzed the miR-302a targets by bioinformatics software, and found that miR-302a can target SDC1. Further, in vitro experiments proved that the re-expression of miR-302a inhibited OC cells proliferation dramatically, and arrested the OC cell cycle at the G1/S phase. The dual-luciferase reporter assays further demonstrated that SDC1 was a novel target of miR-302a. The over expression of SDC1 led to rescue the effect of apoptosis related to miR-302a, such, strongly suggesting that miR-302a suppresses the growth of OC cells by targeting SDC1.

Materials and methods

Bioinformatics analysis

MicroRNA target prediction was performed using TargetScan v6.2 25, Microcosm v5.0 26, and miRanda 27, followed by expression correlation between miRNA-mRNA pairs. MatInspector (http://www.genomatix.de) and GeneGo (http://www.genego.com/metacore.php) were used for searching the transcription factor binding sites as well as analyzing the miRNA-mRNA network, respectively. MatInspector is a software tool that utilizes a large library of matrix descriptions for transcription factor binding sites (TFBS) to locate matches in DNA sequences. By introducing a matrix family concept, optimized thresholds, and comparative analysis, the program produces concise results that avoid redundant and false-positive matches. It assigns a quality rating to matches and thus allows quality-based filtering and selection of matches. GeneGo’s MetaCore™ is an integrated “knowledge-based” platform for pathway analysis of OMICs data and gene lists. MetaCore™ is based on a proprietary manually curated database of human protein-protein, protein-DNA, and protein compound interactions, as well as metabolic and signaling pathways for human, mouse, and rat, supported by proprietary ontologies and controlled vocabulary.

Cell culture, transfection

OVCAR3 and SKOV3 cells were cultured in RPMI 1640 (GIBCO) and McCoy’s 5A Media, with 10% heat-inactivated fetal bovine serum, 100 IU penicillin/ml, 0.1 mg streptomycin/ml in a humidified 5% (v/v) atmosphere of CO2 at 37°C. Transfected with Lipofectamine 2000 Reagent (Invitrogen) followed the manufacturer’s protocol.

Fluorescent reporter assay

The Luciferase expression vector pGL3/Luciferase was constructed. The fragment of SDC1 3’UTR wild-type or mutant was cloned into pGL3/Luciferase at the same sites. Cells were transfected with miR-302a or control vector in 48-well plates, and with the reporter vector SDC1-WT or SDC1-MUT. The intensities of luciferase fluorescence were detected with Fluorescence Spectrophotometer F-4500 (HITACHI).

Quantitative RT-PCR

To detect the relative level of transcript, real-time RT-PCR was performed. Briefly, a cDNA library was generated through reverse transcription using M-MLV reverse transcriptase (Promega) with 2 μg of the large RNA extracted from the cells. The cDNA was used for the amplification of SDC1 gene and the β-actin gene was used as an endogenous control for
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the PCR reaction. PCR was performed under the following conditions: 94°C for 4 min followed by 40 cycles of 94°C for 1 min, 56°C for 1 min, 72°C for 1 min. SYBR Premix Ex Taq™ Kit (TaKaRa) was used following the manufacturer’s instructions, and the real-time PCR was performed and analyzed by 7300 Real-Time PCR system (ABI). All primers were purchased from AuGCT Inc.

Western blotting

Cultured cells were lysed by RIPA (0.1% SDS, 1% Triton X-100, 1 mM MgCl₂, 10 mM Tris-HCl (pH 7.4) in 4°C for 25 min. Collected the lysates were centrifuged, and protein concentration was determined. Total cell lysates (50 μg) were fractionated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis. Proteins were electroblotted onto nitrocellulose membranes. Nonspecific binding sites of membranes were saturated with 5% skim milk in TBST solution (100 mmol/L Tris-Cl, pH 7.5, 150 mmol/L NaCl, 0.1% Tween 20) and incubated for 2 hours with antibodies at room temperature. The following antibody was used: anti-SDC1 and anti-GAPDH. After 4 washes with TBST, the filters were incubated with goat anti-mouse peroxidase-conjugated secondary antibody (Sigma) in 5% skim milk in TBST solution for 1 hour at room temperature; reactions were developed using enhanced chemoluminescence (Perkin Elmer, USA).

Cell proliferation assay and colony formation assay

SKOV3 and OVCAR3 cells were seeded in 96-well plate at 6,000, 7,000 and 8,000 cells per well the day before transfection. The cells

Figure 1. Identification of differential expression of miR-302a in human ovarian cancer. A. We use quantitative Real-time PCR to detect miR-302a differential expression level in human ovarian cancer cells (C13K, 3AO, SKOV3 and OVCAR3) and normal cells. U6 snRNA was regarded as an endogenous normalizer and the relative miR-302a expression level of the 4 kinds of human ovarian cancer cells (means ± SD) is shown (*P < 0.05). B. We use Northern blot and Western blot to detect miR-302a and SDC1 differential expression level in 7 human ovarian cancer tissue samples (*P < 0.05). C. We use quantitative Real-time PCR to detect miR-302a differential expression level in human ovarian cancer stage III/IV tissue samples and the stage I/II ones (*P < 0.05). D. We use Western blot to detect SDC1 differential expression level in human ovarian cancer stage III/IV tissue samples and the stage I/II ones (*P < 0.05).
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were transfected with miR-302a or control vector 0.2 µg per well. MTT assay was used to measure the viable, proliferating cells at 12 h, 24 h, 48 h after transfection. The absorbance at 570 nm was measured using a μQuant Universal Microplate Spectrophotometer (Biotek Instruments). After transfection, SKOV3 and OVCAR3 cells were counted and seeded in 6-well plates (in triplicate) at 50, 60 and 75 cells per well. Fresh culture medium was

Figure 2. Overexpression of miR-302a enhances ovarian cancer cells proliferation and promotes the cell cycle. A. The relative level of miR-302a expressed in OC cells after the transfection with miR-302a or control vector. B. The cell independent growth activity in vitro was assessed by colony formation assay. OC cells were transfected with miR-302a or control vector. The colony formation assay was shown. C. OC cells were transfected with miR-302a or control vector, and then seeded in 6-well plates. Colonies were counted only if they contained more than 50 cells, and the number of colonies was counted from the 6th day after seeding. The number of colonies was counted from the 6th day after seeding. The colony formation rate was calculated and was shown (*P < 0.05). D. OC cells were transfected with miR-302a or control vector. Cell growth activity was determined at 12 h, 24 h and 48 h post-transfection by MTT assay. Values are means ± SD of three duplications and the relative cell growth activity is shown (*P < 0.05). E. The effect of miR-302a on apoptosis was examined by FCM analysis. OC cells were transfected with miR-302a mimics or control, and then the medium was replaced with serum-free DMEM for 48h. F. OC cells after transfected were analyzed for apoptotic rate after staining with Annexin V-FITC and PI. Data represent means ± S.D. from four independent experiments (*P < 0.05). G. The histogram showed the percentages of OC cells after miR-302a transfection in G1/S, S and G2/M phases (n=3, mean ± SD).
replaced every 3 days. Colonies were counted only if they contained more than 50 cells, and the number of colonies was counted from the 6th day after seeding and then the cells were stained using crystal violet. The rate of colony formation was calculated with the equation: colony formation rate = (number of colonies/number of seeded cells) ×100%.

Flow cytometry analysis

After 48 hours transfection as earlier described, the cells were harvested and washed twice with PBS. Washed cells were resuspended in 0.6 mL PBS, and fixed by the addition of 1.4 mL 100% ethanol at 4°C overnight. The fixed cells were rinsed twice with PBS, and resuspended in propidium iodine (PI) solution, including 50 mg/mL PI and 50 mg/mL RNaseA (Sigma) in PBS without calcium and magnesium, and incubated at 37°C for 30 minutes in the dark. Stained cells were passed through a nylon-mesh sieve to remove cell clumps and analyzed by a FACScan flow cytometer and Cell Quest analysis software (Becton Dickinson, San Jose, CA, USA). Flow cytometry analysis was repeated 3 times.

Statistical analysis

Data are expressed as means ± standard deviation (SD), and P < 0.05 is considered as statistically significant by Students-Newman-Keuls test.

Results

MiR-302a expression level in human ovarian cancer cells and their correlation analysis with clinicopathological characteristics

We use quantitative Real-time PCR to detect miR-302a differential expression level in 4 kinds of human ovarian cancer cells, C13K, 3AO, SKOV3 and OVCAR3. The results showed that, the expression level of miR-302a in human ovarian cancer cells was significantly lower than the normal cells (Figure 1A). Staging of ovarian cancer is based on clinical and radiologic examination. Ovarian cancer at its early stages (I/II) is difficult to diagnose until it spreads and advances to later stages (III/IV) because most symptoms are nonspecific and thus of little use in diagnosis. Most patients present with Stage III or IV disease. Stage I is a small tumor completely confined to ovary.

Stage II is a tumor has pelvic extension (must be confined to the pelvis) or primary peritoneal tumor, involves one or both ovaries. Stage III is cancer found outside the pelvis or in the retroperitoneal lymph nodes, involves one or both ovaries. Stage IV is any distant metastasis (i.e. outside of the peritoneum). The expression level of miR-302a was associated with TNM stage. MiR-302a has lower expression level in the stage III/IV than stage I/II (Figure 1C). These data suggested that alterations of miR-302a could be involved in ovarian cancer progression.

Overexpression of miR-302a suppresses ovarian cancer cells proliferation in vitro

In order to study the effects of miR-302a on ovarian cancer cells proliferation, we constructed a overexpression vector: miR-302a. After transfection of ovarian cancer cells, we tested the validity of miR-302a ectopic expression by quantitative Real-time PCR in SKOV3 and OVCAR3, respectively. The results revealed that miR-302a expression level was significantly higher than the control group (Figure 2A). To test the effects of miR-302a on ovarian cancer cells proliferation, we investigated cell growth by colony formation assay and MTT assay. The colony formation rate of SKOV3 and OVCAR3 cells transfected with miR-302a were significantly lower than the control group (Figure 2B, 2C). We performed MTT assay to further confirm the effects of miR-302a on cell proliferation. We found that miR-302a could obviously suppressed SKOV3 and OVCAR3 cells growth (Figure 2D). These two experiments showed that miR-302a played a role in suppressing cell growth and proliferation in ovarian cancer cells. Up-regulating the miR-302a, cell viability and proliferation were significantly inhibited.

MiR-302a mediates cell cycle arrest and potentiates apoptosis in human ovarian cancer cells

To validate whether miR-302a is able to influence apoptosis, Flow cytometry assay was performed (Figure 2E). The results indicated that the significant increase in the apoptosis was observed in the SKOV3 and OVCAR3 cells transfected with miR-302a (Figure 2F). These results strongly suggested that introduction of miR-302a could inhibit human ovarian cancer cells growth by promoting apoptosis of cancer cells.
To confirm that the expression of miR-302a can cause G1/S arrest, SKOV3 and OVCAR3 cells transfected with miR-302a mimics were synchronized at the G1/S transition by serum starvation and hydroxyurea (HU). DNA content was examined from the time of HU release. The results showed that all cells transfected with miR-302a mimics began to arrest at G1 phase and inhibited the transfection from G1 phase to S phase (Figure 2G).

**MiR-302a directly inhibits expression of SDC1 via its 3' UTR**

We used bioinformatics methods to predict miR-302a potential target genes. The 3'UTR region of SDC1 mRNA contains miR-302a complementary binding sites (Figure 3A). Luciferase reporter assay has been widely used in verification of miRNA target genes [47, 48]. To investigate whether SDC1 can be directly targeted by miR-302a, we performed luciferase reporter assay, engineering luciferase reporters, that have either the wild-type 3' UTR of SDC1, or the mutant UTR with a 4 base pair for site-directed mutagenesis in the complementary seed sequence (Figure 3A). First, OVCAR3 cells were transfected with SDC1-wt, miR-302a or control vector. MiR-302a-suppressed the EGFP fluorescence intensity of SDC1-wt (*P < 0.05), the group transfected with SDC1-mut was not significant different to the group. C. OC cells were transfected with the mutant miR-302a or miR-302a-mut could not able to significantly suppress the EGFP fluorescence intensity of SDC1-Luc (*P < 0.05). D. OC cells were transfected with miR-302a and control vector, the expression of SDC1 mRNA and protein expression level were measured by quantitative RT-PCR and Western blot. β-actin mRNA was regarded as an endogenous normalizer and the relative SDC1 mRNA expression level is shown (*P < 0.05). GAPDH protein was regarded as endogenous normalizer and the relative SDC1 protein quantity is shown (*P < 0.05).
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Figure 4. Overexpression of SDC1 could rescue the effects which miR-302a upregulated and led to cell survival in human ovarian cancer. A. We use Western blot to detect the SDC1 differential expression level after co-transfection with miR-302a and SDC1 in human ovarian cancer cells. B. OC cells were co-transfected with miR-302a and SDC1. Cell growth activity was determined at 12 h, 24 h and 48 h post-transfection by MTT assay. Values are means ± SD of three duplications and the relative cell growth activity is shown (* P < 0.05). C. OC cells were co-transfected with miR-302a and SDC1, and then seeded in 6-well plates. Colonies were counted only if they contained more than 50 cells, and the number of colonies was counted from the 6th day after seeding. The number of colonies was counted from the 6th day after seeding. The colony formation rate was calculated and was shown (* P < 0.05). D. The effect of co-transfected with miR-302a and SDC1 on apoptosis was examined by FCM analysis. OC cells after transfected were analyzed for apoptotic rate after staining with Annexin V-FITC and PI. Data represent means ± S.D. from four independent experiments (* P < 0.05). E. The histogram showed the percentages of OC cells after co-transfected...
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3'UTR, so that luciferase activities were decreased. In contrast, mutant reporters were not repressed by miR-302a (Figure 3B). After a 4-base pair mutant of miR-302a, the luciferase activities were not decreased as well (Figure 3C). These all results suggested that, miR-302a could combine with the specific SDC1 mRNA 3'UTR binding sites and play a role in inhibiting the expression of SDC1 gene.

MiR-302a plays a negative regulatory role at SDC1 post-transcriptional level

MiRNAs regulate the target genes at the post-transcriptional level by binding their target genes 3’UTR to silence the gene function. We transfected OVCAR3 cells with miR-302a, in order to examine whether miR-302a depress endogenous SDC1 through translational repression, the expression of SDC1 protein was examined by Western blot. The results showed that overexpression of miR-302a made the expression level of SDC1 protein decreased (Figure 3D), suggesting that miR-302a negatively regulates endogenous SDC1 protein expression through translational repression mechanism. Meanwhile, high expression level of miR-302a in OVCAR3 cells could also decrease the endogenous SDC1 mRNA level (Figure 4C). Furthermore, in the 7 pairs of human ovarian cancer tissues, we found the expression level of SDC1 had a negative correlation with miR-302a expression (Figure 1B). All these data suggest that miR-302a negatively regulates the expression of SDC1 through mRNA cleavage mechanism at the post-transcriptional level.

MiR-302a repressed OC cells proliferation and promoted apoptosis by negatively regulating SDC1 expression

Previous studies show that miR-302a plays an important role in suppressing tumor cell proliferation and promoting the apoptosis. Accordingly, we detected whether SDC1 affects ovarian cancer cell growth. We constructed the SDC1 plasmid to up regulate the expression level of SDC1 and then identified the expression of SDC1 protein by Western blot. While we enhanced the expression level of miR-302a in OC cells, and then transfected with SDC1, Western blot displayed an increase protein level of SDC1 (Figure 4A). The SDC1 plasmid was confirmed effectively increase the SDC1 protein expression level. We further performed a “rescue” study in OC cells, namely, co-transfection of miR-302a mimics and SDC1 partially reverse the growth inhibition effects of OC cells from miR-302a mimics (Figure 4B, 4C). Additionally, the co-transfection with miR-302a and SDC1 showed a similar result in the “rescue” apoptosis assay (Figure 4D, 4E). These data suggests that miR-302a repressed OC cells proliferation and promoted apoptosis in OC cells through targeting SDC1.

Discussion

Transformation process of malignant tumors is regulated by the synergy of multiple genes, including overexpression of oncogenes and low expression or even loss of function of tumor suppressor genes. Recent research uncovered that the regulation of oncogenes and tumor suppressor genes was not only in the transcriptional level, but also in the post-transcriptional level, which was more important and accurate. MicroRNAs (miRNA), as important regulation factors, participate gene expression in human carcinogenesis. In recent years, the research about miRNAs has been more in-depth. MiRNA-mediated post-transcriptional gene silencing (PTGS) and the relevance with tumor formation have become the focus of attention. Tumor cells and normal cells have a significant difference in miRNA expression profiling. Most miRNA genes locate in chromosomal regions frequently display amplification, deletion or translocation in human cancers. Detection of differentially expression of miRNAs in human ovarian cancer, to determine the role of miRNAs in cancer mechanism and function of their target genes, provide a new direction for the diagnosis and treatment of in human ovarian cancer.

In the present study, we tried to identify a novel miRNA which regulates the expression of SDC1, and evaluate its effects on cell phenotype using OC cells. Initially, we used real-time PCR to find that miR-302a was significantly down-regulated in human OC cells, compared with the normal cells. The results suggested that altera-
tions of miR-302a could be involved in ovarian cancer progression. Therefore, we hypothesized that miR-302a was a negative factor of carcinogenesis in human ovarian cancer cells due to the low expression levels in human ovarian cancer. We calculated the cell growth viability through the MTT and colony formation assay to detect the relationship between miR-302a and the growth capacity of ovarian cancer cell line SKOV3 and OVCAR3. The cell growth viability of OC cells transfected with the miR-302a was significantly decreased when compared to control group (Figure 2C, 2D). We further showed that overexpressed miR-302a in OC cells induced G1 arrest, suppressed cell proliferation, and induced apoptosis. These data indicate that miR-302a may act as a tumor suppressor to inhibit cell proliferation by blocking the G1/S transition of OC cells. In other words, reduced miR-302a expression in OC cells and tissues may promote cell proliferation by activating the cell cycle (Figure 2F, 2G).

Secondly, bioinformatics analyses predicted a miR-302a binding site on the SDC1 transcript. Experimental evidence indicated that SDC1 was indeed a target of miR-302a. On one hand, the ability of miR-302a to regulate SDC1 expression was likely direct because it bound the 3′UTR of SDC1 mRNA complementarily to the miR-302a seed region. The EGFP fluorescence intensity of SDC1-wt was specifically responsive to miR-302a overexpression (Figure 3B). Furthermore, mutation of the miR-302a binding site abolished the effect of miR-302a on the regulation of EGFP fluorescence intensity. On the other hand, the endogenous SDC1 protein expression was decreased in OC cells transfected with miR-302a (Figure 3D). These results suggested that miR-302a regulated SDC1 protein expression at the post transcription level.

Syndecan 1 (SDC1) is a member of a transmembrane heparan sulfate proteoglycan family, which expresses in epithelia, and plays a critical role in cellular processes including differentiation, cell adhesion, migration and invasion, and angiogenesis [49-51]. Functions have been ascribed to the extracellular domain that carries glycosaminoglycan (GAG) side chains, to the transmembrane domain and to the cytoplasmic domain that transduces signals from extracellular ligand binding. Initial reports demonstrated that expression of syndecan-1 in vitro was associated with maintenance of epithelial morphology, anchorage-dependent growth, and inhibition of invasiveness [52]. This “tumor suppressor” function was subsequently supported by immunohistochemical studies that correlated loss of syndecan-1 with reduced survival in patients with malignant mesothelioma [53], squamous cell carcinoma of the head and neck [54], and laryngeal cancer [55]. Altered SDC1 expression has been reported gradually in a number of malignant tumor types and has been associated with differentiation stage and grade [56-58]. Marzioni D et al [59], have reported a positive correlation of SDC-1 with fibroblast growth factors (FGFRs) in bladder tumors, these factors are thought to be key molecules in low-grade BCa. Shimada et al [60] have investigated the biologic role of SDC-1 in human BCa cells. In their study, the BCa cell lines, UMUC2 and UMUC3 had SDC-1 expression silenced by siRNA transfection, which led to an induction of apoptosis in vitro and a reduction in mouse orthotopic bladder tumor growth. In addition, studies in breast and gastric cancer have demonstrated an association between increased stromal SDC1 expression, loss of cancer cell SDC1 expression, and an adverse clinical outcome [61-63]. But the association between SDC1 status and human ovarian cancer has not been extensively studied. To our knowledge, our study is the latest study to date to evaluate the effect between miRNA and SDC1 in human ovarian cancer cells. A high expression level of SDC1 could repress OC cells proliferation and promoted apoptosis in OC cells.

In summary, we demonstrated that miR-302a played an important role in the regulation of SDC1 gene expression. The effect of miRNAs on OC cells expression occurred at both mRNA and transcription levels, and at least in part through targeting SDC1. However, we emphasize that miR-302a may be capable of controlling tumor-specific gene(s), consequently favoring cell apoptosis. Therefore, our study suggests that unregulation of miR-302a may provide a better strategy to block tumor proliferation and cell cycle.

Disclosure of conflict of interest

None.
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