

Long non-coding RNA miR155HG silencing restrains ovarian cancer progression by targeting the microRNA-155-5p/tyrosinase-related protein 1 axis

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Abstract. Ovarian cancer (OC) is the third commonest gynecological malignancy worldwide. The long non-coding (lnc)RNA microRNA (miR)155HG functions as an oncogene in different human cancers. However, the function and molecular mechanism of miR155HG in OC remain elusive. The present study indicated that the expression levels of miR155HG and tyrosinase-related protein 1 (TYRP1) were significantly increased, whereas that of miR155-5p was decreased in OC tissues and cells, as detected by real-time quantitative polymerase chain reaction. It was demonstrated that knockdown of miR155HG markedly inhibited OC cell viability, migration and invasion while promoting apoptosis, as indicated by 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide, wound healing, Transwell and western blot assays. Mechanistically, it was revealed that miR155HG and TYRP1 were both targeted by miR-155-5p with complementary binding sites in the 3' untranslated region. A dual-luciferase reporter assay was used to confirm the targeting relationship between miR155HG, miR-155-5p and TYRP1. In addition, the interaction between miR155HG and miR-155-5p was further demonstrated by radioimmunoprecipitation and pull-down assays. In addition, feedback approaches determined that miR-155-5p inhibition or TYRP1 overexpression markedly reversed the inhibitory effects of miR155HG knockdown on OC cell viability, migration and invasion as well as

weakened the promotive effect of miR155HG knockdown on OC cell apoptosis. Thus, miR155HG silencing inhibited the malignant biological behavior of OC cells by targeting the miR-155-5p/TYRP1 axis. The present study provides novel insights into the underlying mechanism of OC progression.

Introduction

Ovarian cancer (OC) is the third commonest gynecological malignancy worldwide (1). Patients are frequently diagnosed with OC at an advanced stage and the 5-year survival rate is only ~35% (2). Despite advances in chemotherapy and surgery, the prognosis of patients with OC remains unsatisfactory (3). High recurrence rates and poor outcomes due to OC metastasis pose serious challenges (4). Therefore, it is necessary to explore the molecular mechanisms of OC progression to discover new therapeutic strategies.

Long non-coding (lnc) RNAs serve multiple roles in the occurrence and development of OC. LncRNA PVT1 regulates EZH2 to downregulate microRNA (miRNA/miR)-214, resulting in the suppression of OC progression (5). LncRNA LINC00319 contributes to OC progression by inhibiting miR-423-5p and enhancing nucleus accumbens-associated 1 expression (6). LncRNA PCAT6 restrains PTEN expression to accelerate the initiation and progression of OC (7). Numerous studies have shown that lncRNA miR155HG (miR155HG) serves critical roles in diverse tumors (8-10). miR155HG promotes pancreatic cancer cell growth and inhibits apoptosis by suppressing miR-802 expression (8). miR155HG deficiency attenuates glioblastoma tumorigenesis by downregulating Annexin A2 expression via an increase in miR-185 expression (9). Notably, the expression of miR155HG is enhanced in OC tissues and cells (10).

miRNAs have been shown to participate in different gynecological malignancies, including OC. miR-126-3p modulates PLXNB2 expression to attenuate OC progression (11). miR-603 suppresses the malignancy of OC cells by targeting hexokinase-2 (12). miR-122 inhibits OC cell growth by repressing prolyl 4-hydroxylase subunit α 1 (13). miR-155-5p has been regarded as a pivotal regulator of multiple

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cancers. miR-155-5p elevation impedes gastric cancer cell proliferation and triggers apoptosis (14). miR-155-5p regulates IGF2 through the PI3K pathway to exert tumor-repressing roles in Wilms tumors (15). Notably, miR-155-5p reduces OC cell viability by inhibiting HIF1 α (16).

Tyrosinase-related protein (TYRP) 1 is a melanogenic enzyme and protein (17). Previous research has reported on the relationship between melanogenic proteins and cancers, such as TYRP1 in breast cancer (18) and TYRP2 in retinoblastoma (19). TYRP1 exerts tumor-promoting functions in different types of cancer. Increased TYRP1 expression in lymph node metastases from melanoma patients is related to unfavorable prognosis (20). TYRP1 has emerged as an oncogene in colon cancer and high levels of TYRP1 are associated with decreased overall survival rates (21). Notably, El Hajj *et al.* (22) reported that TYRP1 is a target of miR-155.

Despite the aforementioned studies, the molecular mechanisms of miR155HG, miR-155-5p and TYRP1 in OC progression remain unclear. The present study assessed the expression and roles of miR155HG in OC. In addition, the relationships between miR155HG, miR-155-5p and TYRP1 in OC were determined. The present study aimed to reveal the molecular mechanism of miR155HG in OC cell viability, migration, invasion and apoptosis.

Materials and methods

Ethics statement. The study was approved by the ethics committee of the Affiliated Hospital of North Sichuan Medical College [approval no. 2020ER (A) 066]. Written informed consent was obtained from all participants. The study was conducted in accordance with the principles of the Declaration of Helsinki.

Clinical samples. Patients with OC (n=55) who underwent ovariectomy between September 2017 and October 2019 at the Affiliated Hospital of North Sichuan Medical College (Nanchong, China) were enrolled in this study. Among them, 41 cases were the serous subtype, 8 cases were the endometrioid subtype and 6 cases were other subtypes. The histopathological diagnosis of the cases was in accordance with the diagnostic categories of the World Health Organization 2020 (23). OC tissues (n=55; tumor group) and paired adjacent non-tumor tissues (tissue which were 1-2 cm away from the tumor tissues; adjacent group) were obtained from patients with OC who underwent ovariectomy. Prior to ovariectomy, radiotherapy or chemotherapy was not administered to the patients.

Cell culture. The OVCAR3 and SK-OV-3 OC cell lines and normal IOSE80 ovarian cell line (Chinese Academy of Sciences) were cultured in Roswell Park Memorial Institute (RPMI)-1640 medium (Invitrogen; Thermo Fisher Scientific, Inc.) supplemented with 10% exosome-free fetal bovine serum (FBS; Invitrogen; Thermo Fisher Scientific, Inc.) at 37°C with 5% CO₂.

Cell transfection. For cell transfection, OVCAR3 and SK-OV-3 cells grown to 85% confluence were transfected with 100 nM of negative control (NC) small interfering (si)

RNA (si-NC; 5'-UUCUCCGAACGUGUCACGU-3'), 100 nM of miR155HG siRNA (si-miR155HG-1; 5'-CUGGGAUGU UCAACCUUAA-3'; si-miR155HG-2; 5'-UCUAAAAGGGAA ACUGAAA-3'), 100 nM of mimics NC (5'-UCACAACCU CCUAGAAAGAGUAGA-3'), 100 nM of miR-155-5p mimics (5'-UUA AUGCUAAUCGUCAUAGGGGU-3'), 100 nM of inhibitor NC (5'-CAGUACUUUUGUGUAGUACAA-3'), 100 nM of miR-155-5p inhibitor (5'-ACCCCUAUCACGAU AGCAUUA-3'), 1 μ g of empty plasmid (pcDNA3.1-NC) and 1 μ g of TYRP1 overexpression plasmid (pcDNA3.1-TYRP1), or co-transfected with 100 nM of si-miR155HG-1 and miR-155-5p inhibitor, 100 nM of si-miR155HG-1 and 1 μ g of pcDNA3.1-TYRP1 using Lipofectamine[®] 3000 reagent (Invitrogen; Thermo Fisher Scientific, Inc.) for 6 h at 37°C. All oligonucleotides or plasmids were purchased from Shanghai GenePharma Co., Ltd. At 48 h post-transfection, the cells were harvested and reverse transcription-quantitative (RT-q) PCR was conducted to determine the transfection efficiency.

RT-qPCR. RT-qPCR procedures were performed according to the corresponding manufacturer's protocols. In brief, total RNA was extracted from tissues and cells using TRIzol[®] reagent (Thermo Fisher Scientific, Inc.). RNA was reverse transcribed into complementary DNA using the Prime Script RT reagent kit (Takara Biotechnology Co., Ltd.). The SYBR-Green PCR kit (Takara Biotechnology Co., Ltd.) and TaqMan MicroRNA Assay Kit (Applied Biosystems; Thermo Fisher Scientific, Inc.) were used for qPCR analysis. The following thermocycling conditions were used for the qPCR: Initial denaturation at 95°C for 3 min; followed by 40 cycles at 95°C for 15 sec, annealing at 60°C for 30 sec, elongation at 72°C for 1 min; and a final extension at 72°C for 5 min. GAPDH, U6 and β -actin were used for the normalization of miR155HG, miR-155-5p and TYRP1, respectively (8,24,25). Relative expression was calculated using the 2^{- $\Delta\Delta$ C_q} method (26). The primers used are presented in Table I.

Western blot analysis. The transfected OVCAR3 and/or SK-OV-3 cells were lysed with RIPA buffer (Beyotime Institute of Biotechnology) to extract total protein. The protein concentration was detected via the BCA Protein Assay kit. Subsequently, a total of 50 μ g protein/lane was separated by 10% SDS-PAGE and then transferred onto PVDF membranes. Following blocking with 5% skimmed milk for 2 h at 25°C, the membranes were incubated overnight at 4°C with primary antibodies, including anti-TYRP1 (1:1,000; ab235447; Abcam), anti-Bax (1:1,000; ab32503; Abcam), anti-Bcl-2 (1:2,000; ab182858; Abcam) and anti- β -actin (1:1,000; ab265588; Abcam). Thereafter, the membranes were incubated with horseradish peroxidase-conjugated secondary antibodies (1:5,000; ab97080; Abcam) at 25°C for 1 h. The proteins bound by their respective antibodies on the immunoblots were measured using an enhanced ECL kit (Thermo Fisher Scientific, Inc.) and quantified using ImageLab software (version 2.3; Bio-Rad Laboratories, Inc.).

3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay. OVCAR3 and SK-OV-3 cells (2x10³/well) were seeded into 96-well plates and incubated at 37°C with 5% CO₂. At each time point (0, 24, 48 and 72 h post-transfection), cell

Table I. Primers sequences.

Name of primer	Sequences (5'-3')
miR155HG-F	CCCAAATCTAGGTTCAAGTTC
miR155HG-R	CATCTAAGCCTCACAAACAAC
GAPDH-F	AGGTGAAGGTCGGAGTCAACG
GAPDH-R	AGGGGTCATTGATGGCAACA
miR-155-5p-F	GTGCAGGGTCCGAGGTATT
miR-155-5p-R	GCCGCTTAATGCTAATCGTGATAG
U6-F	GCTTCGGCAGCACATATACTAAAAT
U6-R	CGCTTACGAATTTGCGTGTCAT
TYRP1-F	GCTCAGTGCTTGGAAAGTTGGT
TYRP1-R	AGTTTGTCTCCAGTTCCGTTTAG
β-actin-F	GACCCTGCCATCTGTGC
β-actin-R	CGGGTGGAGGAGTTTCA

miR, microRNA; F, forward; R, reverse.

viability was determined using the MTT cell viability assay kit (Sigma-Aldrich; Merck KGaA) under an inverted light microscope (magnification x400; Olympus Corporation).

Wound healing assay. OVCAR3 and SK-OV-3 cells (1×10^6 /well) were incubated in 6-well plates. The cell monolayer was then wounded with a 10- μ l pipette tip and cultured in serum-free medium. Images of the different stages of wound healing were captured by a light microscopy (magnification, x400; Olympus Corporation) at 0 and 48 h.

Invasion assay. Transwell chambers (24-well; 8 μ M pore size; BD Biosciences) coated with Matrigel at 37°C for 30 min (BD Biosciences) were used to evaluate cell invasion. OVCAR3 and SK-OV-3 cells (1×10^5) were seeded into the upper chamber of Transwell plates (Corning, Inc.) in serum-free RPMI-1640 medium. Exosome-free FBS (10%) RPMI-1640 medium was added to the lower chamber of the Transwell plates. After 24 h, cells that had invaded the pores were fixed with methanol and stained with 0.5% crystal purple at 37°C for 30 min. Stained cells were imaged using an inverted light microscope (magnification, x400; Olympus Corporation).

Dual-luciferase reporter assay. The putative binding sites of miR-155-5p on miR155HG and the TYRP1 3' untranslated region (UTR) were predicted using StarBase (version 2.0; <http://starbase.sysu.edu.cn>) and TargetScan (release 7.2; http://www.targetscan.org/vert_72/), respectively. miR155HG and TYRP1 sequences were generated with wild-type (WT) or mutant miR-155-5p binding sites and cloned them into pmirGLO vectors (Shaanxi Youbio Technology Co., Ltd.). OVCAR3 and SK-OV-3 cells were co-transfected with the luciferase vectors and NC mimics or miR-155-5p mimics using Lipofectamine® 3000 (Invitrogen; Thermo Fisher Scientific, Inc.) for 48 h at 37°C. Relative luciferase activity was examined using a dual-luciferase reporter assay system (Promega Corporation). The activity of firefly luciferase was normalized to that of *Renilla* luciferase.

Statistical analysis. All statistical analyses were performed using GraphPad Prism 8.0 software (GraphPad Software, Inc.). Data are expressed as means \pm standard deviations. The miR155HG, miR-155-5p and TYRP1 expression of OC tissues and paired adjacent non-tumor tissues were assessed using paired Student's t-test. In addition, differences between two groups were analyzed using unpaired Student's t-test. Differences among multiple groups were assessed by a one-way analysis of variance followed by Tukey's post-hoc test. In the analysis of clinicopathological features, the age, diameter, lymph node metastasis, International Federation of Gynecology and Obstetrics (FIGO) stage (27) and histological grade were analyzed using the χ^2 test. The pathological subtype was analyzed using the Fisher's exact test. The significance of the correlations was determined using Pearson's correlation analysis. All experiments were performed in triplicate, and each experiment was repeated three times. $P < 0.05$ was considered to indicate a statistically significant difference.

Results

miR155HG is upregulated in OC. To confirm whether miR155HG is differentially expressed in OC tissues, miR155HG expression was analyzed by RT-qPCR in 55 patients with OC. The results showed that miR155HG expression was considerably upregulated in OC tissues compared with that in adjacent non-tumor tissues ($P < 0.001$; Fig. 1A). Additionally, miR155HG expression was notably elevated in tumors at FIGO stage III/IV ($P < 0.01$; Fig. 1B). Furthermore, miR155HG expression was clearly enhanced in OVCAR3 and SK-OV-3 cells compared with that in IOSE80 cells ($P < 0.001$; Fig. 1C). According to the analysis of clinicopathological features shown in Table II, the expression of miR155HG was not correlated with age, pathological subtype, diameter and lymph node metastasis ($P > 0.05$) but was closely associated with the FIGO stage ($P < 0.05$) and histological grade ($P < 0.01$) of patients with OC. Notably, the number of patients with OC with high miR155HG expression was greater than the number of patients with OC with low miR155HG expression in the 'low-grade' (grade G3) type of OC ($P < 0.01$).

miR155HG silencing restrains OC cell viability, migration and invasion while promoting apoptosis. Loss-of-function experiments were performed to investigate whether miR155HG knockdown affects OC progression *in vitro*. Fig. 2A shows that miR155HG was effectively silenced following si-miR155HG-1 ($P < 0.001$) and si-miR155HG-2 ($P < 0.01$) transfection in OVCAR3 and SK-OV-3 cells. Si-miR155HG-1 was used for subsequent assays because of its high silencing efficiency. The MTT assay revealed that the viability of OVCAR3 and SK-OV-3 cells was markedly reduced after si-miR155HG-1 transfection ($P < 0.01$; Fig. 2B). The wound healing and invasion assays revealed that the migration and invasion of OVCAR3 and SK-OV-3 cells were visibly suppressed by miR155HG deficiency ($P < 0.001$; Fig. 2C and D). Bax and Bcl-2 are biomarkers of apoptosis. miR155HG silencing markedly increased the Bax protein expression level and decreased the Bcl-2 protein expression level in OVCAR3 and SK-OV-3 cells ($P < 0.05$; Fig. 2E).

Table II. Correlation between miR155HG expression and clinicopathological features in ovarian cancer patients.

Characteristics	n	miR155HG (Low) 27	miR155HG (High) 28	P-value
Age				0.698 ^a
<55 years	23	12	11	
≥55 years	32	15	17	
Pathological subtype				0.749 ^a
Serous	41	19	22	
Endometrioid	8	5	3	
Others	6	3	3	
Diameter				0.891 ^a
<5 cm	27	13	14	
≥5 cm	28	14	14	
Lymph node metastasis				0.341 ^a
No	29	16	13	
Yes	26	11	15	
FIGO stage				0.022 ^b
I + II	24	16	8	
III + IV	31	11	20	
Histological grade				0.004 ^c
G1-G2	25	15	10	
G3	30	12	18	

^aNot significant; ^bP<0.05; ^cP<0.01. FIGO, Federation International of Gynecology and Obstetrics.

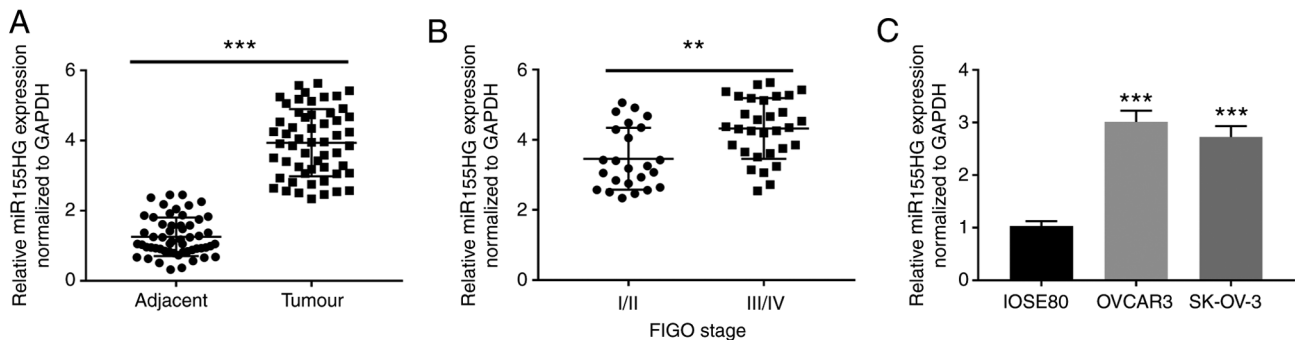


Figure 1. miR155HG is upregulated in OC. (A) The expression of miR155HG in OC tissues and adjacent non-tumor tissues was measured by RT-qPCR. ***P<0.001 vs. Adjacent; (B) Relative expression of miR155HG in patients with OC at the FIGO stage I/II and III/IV was detected by RT-qPCR. **P<0.01 vs. I/II. (C) RT-qPCR was performed to measure the expression of miR155HG in IOSE80, OVCAR3 and SK-OV-3 cells. ***P<0.001 vs. IOSE80. miR, microRNA; OC, ovarian cancer; RT-qPCR, reverse transcription-quantitative PCR; FIGO, International Federation of Gynecology and Obstetrics.

miR155HG directly targets miR-155-5p. To confirm the miR155HG mechanism of action in OC development, miR-155-5p was identified as a potential miR155HG target via StarBase (Fig. 3A). The dual-luciferase reporter assay showed that miR-155-5p upregulation evidently attenuated the activity of the WT-miR155HG reporter in OVCAR3 and SK-OV-3 cells (P<0.001; Fig. 3B). In addition, si-miR155HG-1 transfection markedly enhanced miR-155-5p expression in OVCAR3 and SK-OV-3 cells (P<0.001; Fig. 3C). In addition, miR-155-5p expression was dramatically inhibited in OC tissues compared with that in adjacent non-tumor tissues (P<0.001; Fig. 3D). An inverse correlation between miR155HG and miR-155-5p expression was observed in OC tissues (Fig. 3E). miR-155-5p

expression was considerably downregulated in OVCAR3 and SK-OV-3 cells compared with that in IOSE80 cells (P<0.001, Fig. 3F).

miR-155-5p elevation impedes OC cell viability, migration and invasion while facilitating apoptosis. To determine the biological function of miR-155-5p in OC, miR-155-5p was enhanced or blocked after miR-155-5p mimic (P<0.001) or miR-155-5p inhibitor (P<0.01) transfection in OVCAR3 and SK-OV-3 cells (Fig. 4A). miR-155-5p elevation significantly attenuated the viability of OVCAR3 and SK-OV-3 cells (P<0.01; Fig. 4B). Furthermore, miR-155-5p elevation markedly retarded the migration and invasion of OVCAR3 and

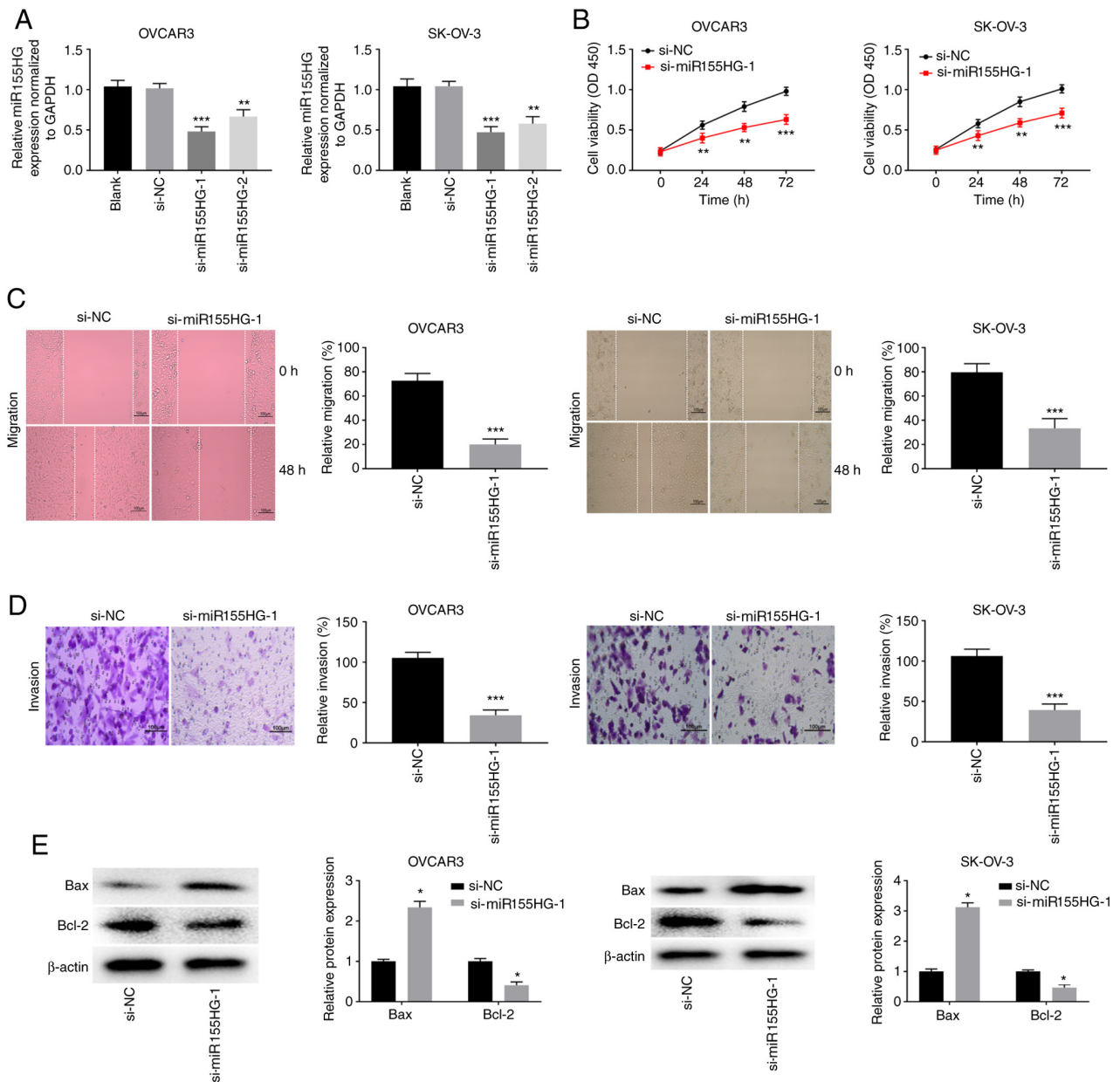


Figure 2. miR155HG silencing restrains the viability, migration and invasion, while promoting apoptosis of OC cells. (A) The transfection efficiency of si-NC, si-miR155HG-1 and si-miR155HG-2 in OVCAR3 and SK-OV-3 cells was measured by reverse transcription-quantitative PCR. ** $P < 0.01$, *** $P < 0.001$ vs. si-NC. (B) The viability of OVCAR3 and SK-OV-3 cells was assessed by MTT assay. ** $P < 0.01$, *** $P < 0.001$ vs. si-NC. The (C) migration and (D) invasion of OVCAR3 and SK-OV-3 cells were analyzed by wound-healing assay and invasion assay. *** $P < 0.001$ vs. si-NC. Scale bar=100 μm ; (E) The protein expression of Bax and Bcl-2 in OVCAR3 and SK-OV-3 cells were measured by western blotting. * $P < 0.05$ vs. si-NC. miR, microRNA; OC, ovarian cancer; si, short interfering; NC, negative control; OD, optical density.

SK-OV-3 cells ($P < 0.001$; Fig. 4C and D). Overexpression of miR-155-5p not only significantly elevated the Bax protein expression level but also reduced the Bcl-2 protein expression level in OVCAR3 and SK-OV-3 cells ($P < 0.05$; Fig. 4E).

TYRPI is a target of miR-155-5p. To demonstrate whether TYRPI is a direct target of miR-155-5p in OC, TargetScan was used to predict the binding site for miR-155-5p on the 3'-UTR of TYRPI (Fig. 5A). The dual-luciferase reporter assay showed that miR-155-5p elevation clearly hindered the activity of the WT-TYRPI reporter in OVCAR3 and SK-OV-3 cells ($P < 0.01$; Fig. 5B). Additionally, miR-155-5p deficiency visibly enhanced TYRPI expression in OVCAR3 and SK-OV-3 cells ($P < 0.001$;

Fig. 5C). Furthermore, TYRPI expression was upregulated in OC tissues compared with that in adjacent non-tumor tissues ($P < 0.001$; Fig. 5D). A negative correlation between TYRPI and miR-155-5p expression (Fig. 5E) and a positive correlation between TYRPI and miR155HG expression (Fig. 5F) were observed in OC tissues. TYRPI expression was upregulated in OVCAR3 and SK-OV-3 cells compared with that in IOSE80 cells ($P < 0.001$; Fig. 5G).

miR155HG silencing hampers the malignant biological behavior of OC cells by targeting the miR-155-5p/TYRPI axis. To ascertain whether miR155HG modulates TYRPI expression by affecting miR-155-5p repression activity,

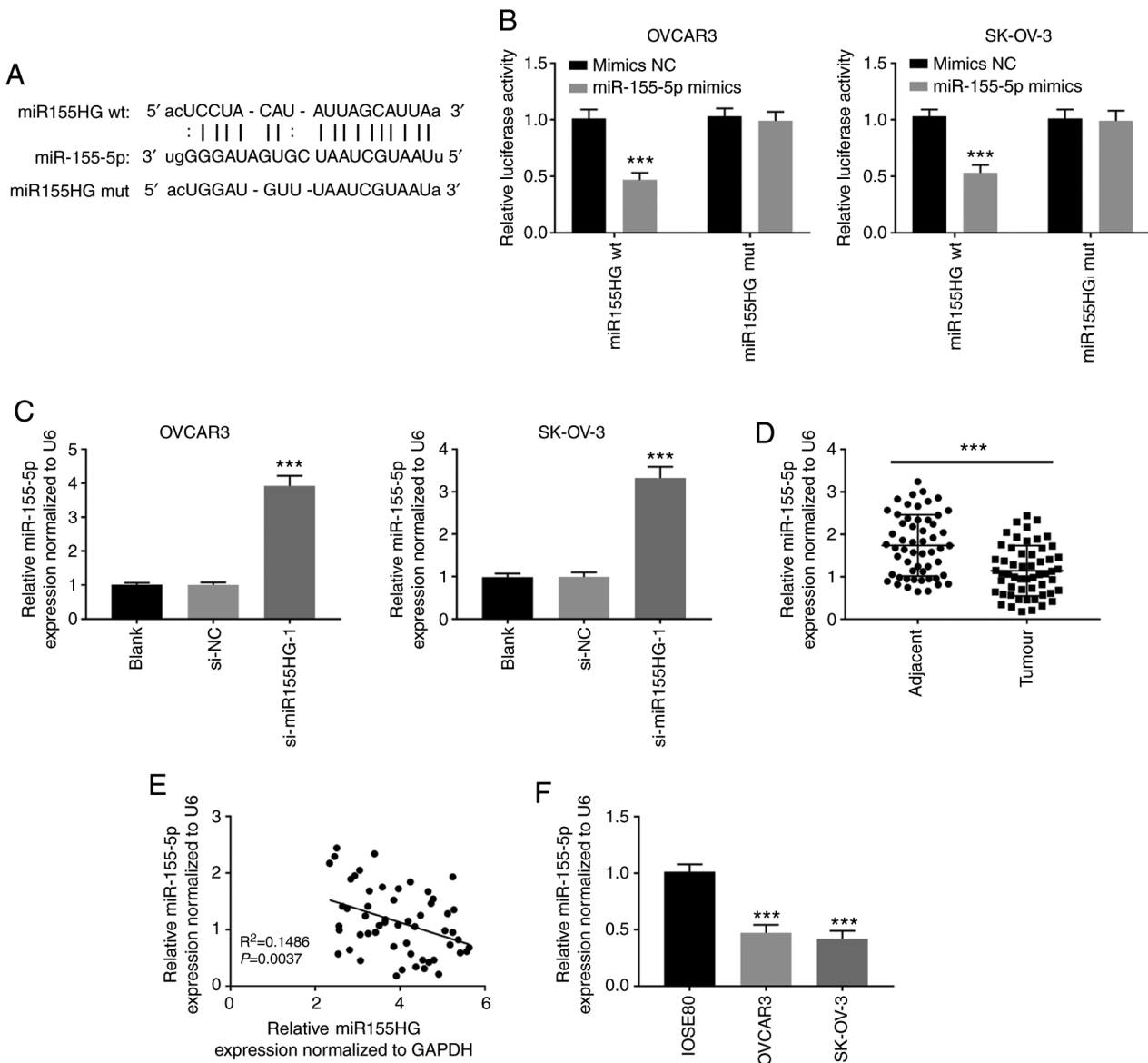


Figure 3. miR155HG directly targets miR-155-5p. (A) StarBase showed the predicted binding site between miR155HG and miR-155-5p. (B) Relative luciferase activity in OVCAR3 and SK-OV-3 cells was evaluated by dual-luciferase reporter assay. $^{***}P<0.001$ vs. mimics NC. (C) The expression of miR-155-5p was increased by the transfection of si-miR155HG-1 in OVCAR3 and SK-OV-3 cells. $^{****}P<0.0001$ vs. si-NC; (D) RT-qPCR was performed to detect the expression of miR-155-5p in OC tissues and adjacent non-tumor tissues. $^{***}P<0.001$ vs. Adjacent; (E) The expression of miR155HG was negatively correlated with miR-155-5p in OC tissues; (F) The expression of miR-155-5p in IOSE80, OVCAR3 and SK-OV-3 cells was detected by RT-qPCR. $^{***}P<0.001$ vs. IOSE80. miR, microRNA; NC, negative control; si, short interfering; RT-qPCR, reverse transcription-quantitative PCR; OC, ovarian cancer; wt, wild-type; mut, mutant.

feedback approaches were used, in which the effects of miR155HG silencing were reversed by TYRP1 overexpression or miR-155-5p inhibition. OVCAR3 cells were transiently transfected with pcDNA3.1-NC or pcDNA3.1-TYRP1. The overexpression efficiency of pcDNA3.1-TYRP1 was high in OVCAR3 cells. TYRP1 expression was effectively enhanced following transfection with pcDNA3.1-TYRP1 in OVCAR3 cells ($P<0.001$; Fig. 6A). In addition, the protein expression level of TYRP1 was markedly increased in OVCAR3 cells following transfection with pcDNA3.1-TYRP1 ($P<0.01$; Fig. 6B). As shown in Fig. 6C, miR155HG silencing visibly suppressed TYRP1 expression in OVCAR3 cells ($P<0.001$), whereas miR-155-5p deficiency weakened the inhibitory effect of miR155HG silencing on TYRP1 expression ($P<0.05$). The feedback approaches showed that TYRP1 elevation or

miR-155-5p deficiency considerably mitigated the inhibitory effects of miR155HG silencing on OVCAR3 cell viability, migration and invasion ($P<0.01$; Fig. 6D-G). In addition, TYRP1 overexpression or miR-155-5p inhibition markedly weakened the promotive effect of miR155HG silencing on the Bax protein expression level and reversed the reduction effect of miR155HG silencing on the Bcl-2 protein expression level in OVCAR3 cells ($P<0.05$; Fig. 6H).

Discussion

It has been documented that the expression of lncRNAs, such as PTAR (28), TP73-AS1 (29) and CCAT1 (30), is increased in OC. In the present study, miR155HG expression was upregulated in OC, suggesting that miR155HG may be

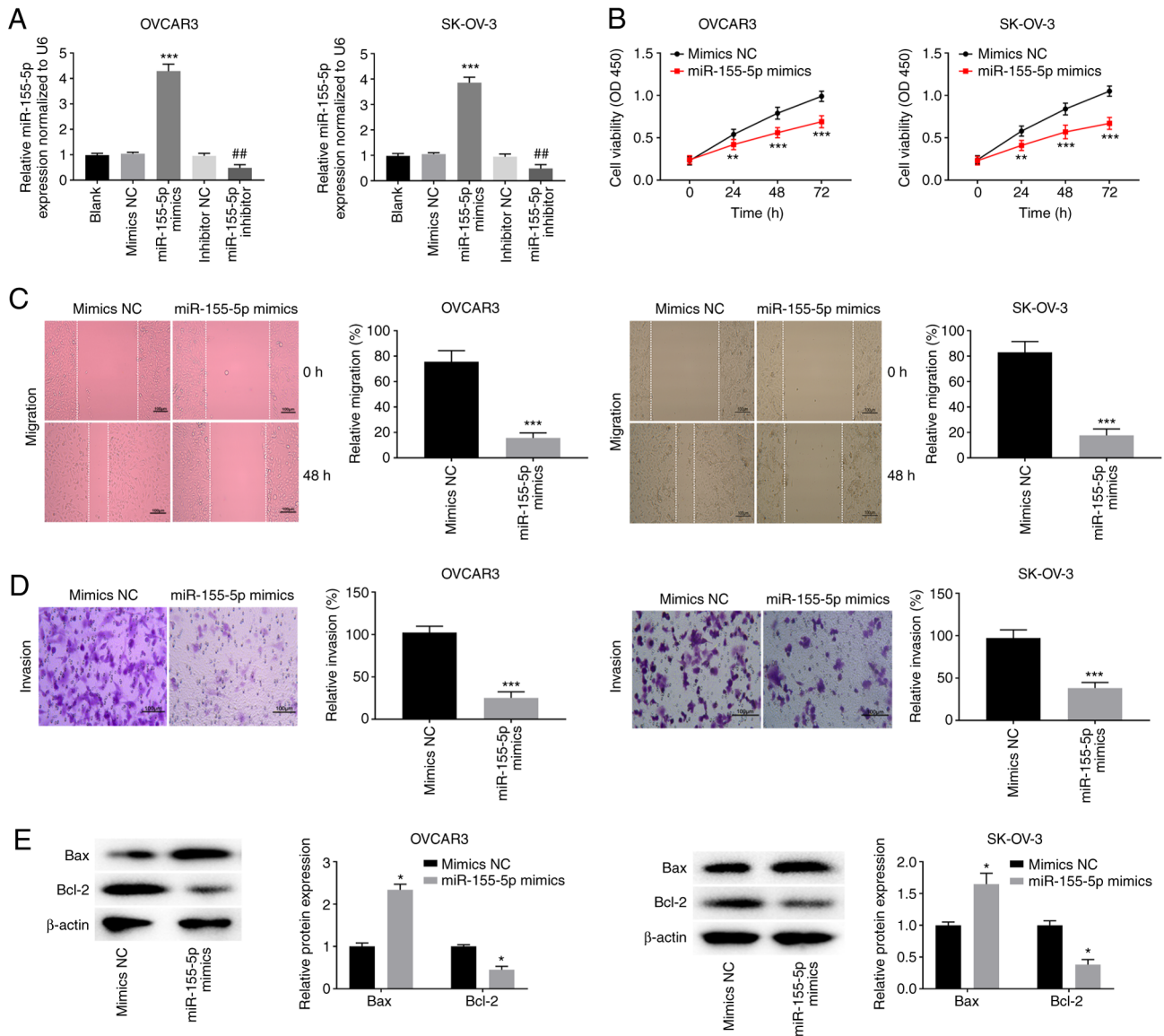


Figure 4. miR-155-5p elevation impedes the viability, migration and invasion while facilitating apoptosis of OC cells. (A) The transfection efficiency of mimics NC, miR-155-5p mimics, inhibitor NC and miR-155-5p inhibitor was evaluated by reverse transcription-quantitative PCR in OVCAR3 and SK-OV-3 cells. *** $P < 0.001$ vs. mimics NC. ** $P < 0.01$ vs. inhibitor NC; (B) MTT assay were performed after transfected with mimics NC or miR-155-5p mimics in OVCAR3 and SK-OV-3 cells. ** $P < 0.01$, *** $P < 0.001$ vs. mimics NC. The effects of miR-155-5p overexpression on the (C) migration and (D) invasion of OVCAR3 and SK-OV-3 cells were assessed. *** $P < 0.001$ vs. mimics NC. Scale bar=100 μm ; (E) Western blotting was performed to detect the protein expression of Bax and Bcl-2 in OVCAR3 and SK-OV-3 cells. * $P < 0.05$ vs. mimics NC. miR, microRNA; OC, ovarian cancer; NC, negative control; si, short interfering; OD, optical density.

an oncogene. In addition, a high level of miR155HG was correlated with FIGO stage in patients with OC. Certain lncRNAs are similar to miR155HG. For instance, high lncRNA PVT1 expression is related to poor prognosis and advanced FIGO stage in patients with OC (5). Overexpression of lncRNA HOTTIP is markedly correlated with advanced FIGO stage in patients with OC (31). Above all, the present study suggested that miR155HG expression may be associated with OC development. Previous studies have demonstrated that miR155HG participates in the malignant biological behavior of diverse cancers (24,32,33). miR155HG silencing diminishes cell viability while facilitating cell apoptosis by targeting PTBP1 to restrain glioma development (32). miR155HG knockdown upregulates miR-155-3p and downregulates TP53INP1, thus retarding the biological behavior of non-small cell lung cancer (33). miR155HG downregulation hinders

laryngeal squamous cell carcinoma progression by increasing miR-155-5p and reducing SOX10 expression (24). In the present study, miR155HG knockdown attenuated OC cell viability, invasion and migration while promoting OC cell apoptosis, indicating that miR155HG silencing may inhibit the development of OC.

Some lncRNAs serve as competing endogenous RNAs or molecular sponges to regulate miRNAs in OC. For instance, lncRNA EWSAT1 decreases miR-330-5p expression to accelerate OC progression (34). lncRNA CCAT1 exerts a tumor-promoting role in OC progression by sponging miR-490-3p (35). Notably, lncRNA NORAD induces OC cell proliferation and cell cycle transition by downregulating miR-155-5p (36). In the present study, miR-155-5p was a target of miR155HG and inversely correlated with miR155HG expression, suggesting that miR155HG may influence OC by

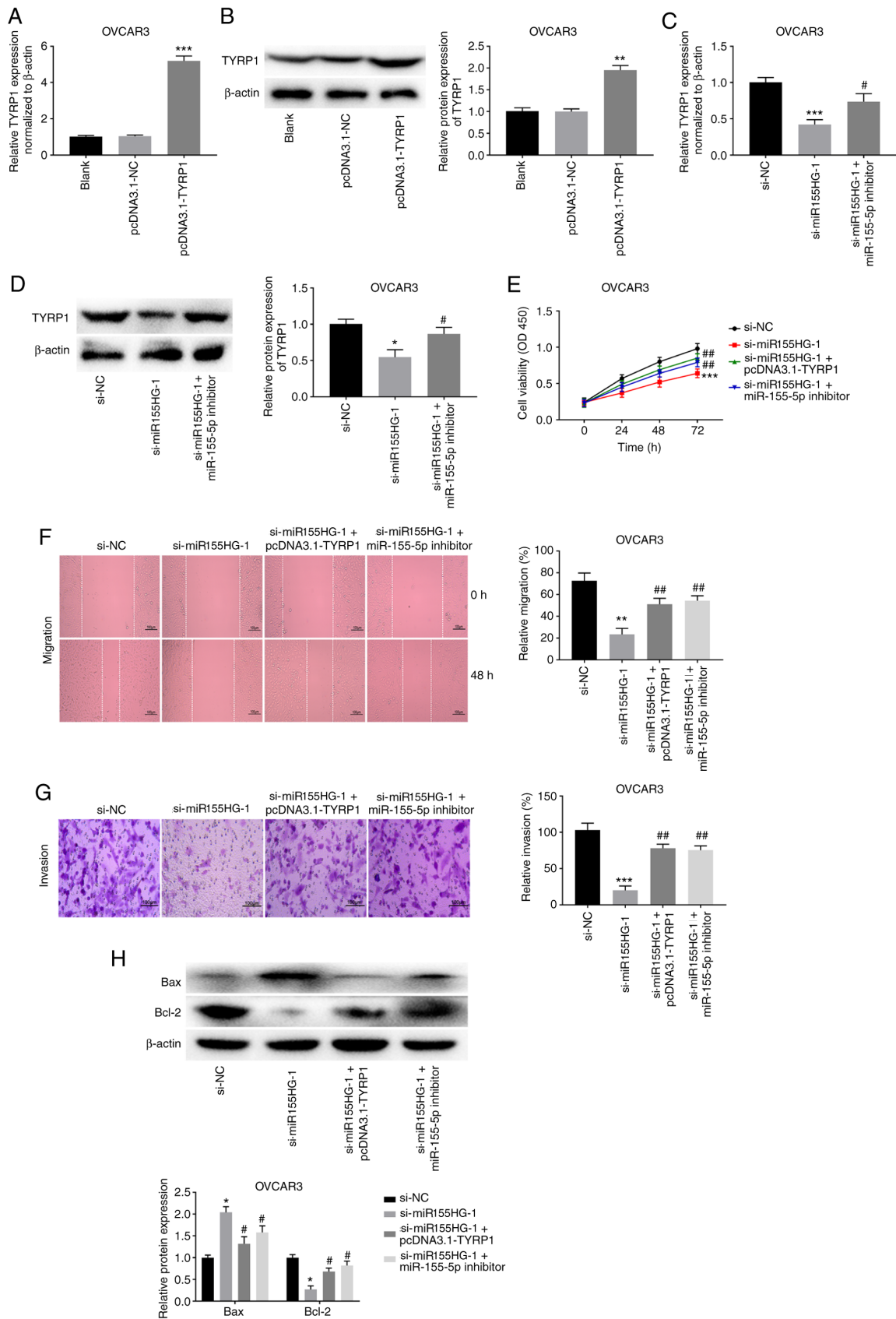


Figure 6. miR155HG silencing hampers the malignant biological behavior of OC cells by targeting the miR-155-5p/TYRP1 axis. (A) The transfection efficiency of pcDNA3.1-NC and pcDNA3.1-TYRP1 was measured by RT-qPCR in OVCAR3 cells. ***P<0.001 vs. pcDNA3.1-NC. (B) The protein expression of TYRP1 in OVCAR3 cells was detected by western blotting. **P<0.01 vs. pcDNA3.1-NC. (C) RT-qPCR was used to detect the expression of TYRP1 in OVCAR3 cells. ***P<0.001 vs. si-NC, #P<0.05 vs. si-miR155HG-1. (D) Downregulation of miR-155-5p or upregulation of TYRP1 reversed the inhibitory effects of miR155HG silencing on (E) viability, (F) migration and (G) invasion of OVCAR3 cells. *P<0.05, **P<0.01, ***P<0.001 vs. si-NC. #P<0.05, ##P<0.01 vs. si-miR155HG-1. Scale bar=100 μ m. (H) Overexpression of TYRP1 or inhibition of miR-155-5p reversed the effects of miR155HG silencing on the protein expression of Bax and Bcl-2 in OVCAR3 cells. *P<0.05 vs. si-NC; #P<0.05 vs. si-miR155HG-1. miR, microRNA; OC, ovarian cancer; NC, negative control; RT-qPCR, reverse transcription-quantitative PCR; si, short interfering; OD, optical density.

to compare with the disease group. Third, the present study did not evaluate the effect of radiotherapy or chemotherapy on miR155HG-silenced cells. Fourth, there are a number of other downstream targets of miR155HG that have not yet been evaluated in OC. Fifth, the present study was limited to the cellular level and further *in vivo* experiments are needed to confirm the role of miR155HG in OC.

In summary, miR155HG expression was enhanced in OC. In addition, miR155HG knockdown appeared to exert tumor-repressing roles in OC progression by regulating the miR-155-5p/TYRP1 axis. Thus, the current study may enhance our understanding of the mechanism underlying miR155HG in OC progression.

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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

AW and LL conceived and designed the present study, analyzed data, performed the experiments and data analyses and wrote the manuscript. CD and XL contributed significantly to analysis and manuscript preparation, performed the experiments and data analyses and revised the manuscript. AW and LL confirm the authenticity of all the raw data. All authors reviewed and approved the final manuscript.

Ethics approval and consent to participate

The present study was conducted after obtaining ethical approval from the Affiliated Hospital of North Sichuan Medical College [Nanchong, China; approval no. 2020ER (A)066]. Written informed consent was obtained from all participants.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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