

FOXK1, Regulated by miR-365-3p, Promotes Cell Growth and EMT Indicates Unfavorable Prognosis in Breast Cancer

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Background: Forkhead box K1 (FOXK1) is members of the FOX transcription factor family. Previous work has found out that FOXK1 promotes cell proliferation, migration and invasion in several cancers, such as gastric cancer, glioma cancer and lung cancer; however, the exact role of FOXK1 in breast cancer is still poorly known.

Methods: Here, the association between FOXK1 expression and the clinicopathological characteristics of patients with breast cancer was identified. To further decipher the functional roles of FOXK1, it was overexpressed or knocked down in MCF-7, MDA-MB-231 and MCF-10A cells. Cell Counting Kit-8, colony formation and cell cycle assays were performed to examine the proliferation of breast cancer cells. Moreover, wound-healing and Transwell invasion analyses were carried out to explore the effect of FOXK1 on breast cancer cell migration and invasion.

Results: Our findings discovered that FOXK1 promotes cell proliferation, migration and invasion in breast cancer. In addition, consistent with the previous report, FOXK1 also facilitates EMT in breast cancer. TargetScan was used to predict up-stream of FOXK1, indicating that miR-365-3p could regulate FOXK1 expression in breast cancer.

Conclusion: The findings of the present study demonstrated that miR-365-3p-FOXK1 axis plays a key role in breast cancer.

Keywords: FOXK1, EMT, miR-365-3p, migration, breast cancer

Introduction

Breast cancer has become one of the most common diagnosed cancer among women in the world.¹ Metastasis accounts for the majority of morbidity and mortality in breast cancer.² Although progress has been made in the diagnosis and treatment of breast cancer, the survival rate is still low.³ Therefore, it is an emergency to investigate the underlying molecular mechanism of breast cancer and develop novel therapeutic targets for the treatment.

Forkhead box K (FOXK) family is a subgroup of the Forkhead transcription factors, known to regulate multiple biological processes,⁴ including cell cycle progression, apoptosis, differentiation, metabolism, survival and senescence.⁵ FOXK1, also known as MNF, has been found to regulate expression, such as p21, c-myc and cdc2, through recognizing the DNA sequence motif, WRTAAAAYA.⁶ Previous studies have revealed that FOXK1 induces the epithelial-mesenchymal transition (EMT) in tumour cells, maintains the invasive potential of GC and promotes cell proliferation in ovarian cancer.^{7,8} These data support a role for human FOXK1 in regulating the developmental process as well as the

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potential involvement of FOXX1 in tumorigenesis. However, the function and mechanism of FOXX1 in breast cancer remain little known.

EMT is a complicated process necessary during wound healing, embryonic development and tumor progression.^{9,10} Epithelial cells lose their epithelial features and gain mesenchymal characteristics and become invasive. At the molecular level, the expression of the epithelial marker, such as E-cadherin, is decreased and the expression of mesenchymal markers, such as N-cadherin and vimentin, is increased.^{11–13} EMT alters the polarity, adhesion ability and improves their migratory and invasive ability.^{9,10,14} Several transcription factors have been found to be involved in the progress of EMT, such as Slug, Snail and ZEB1.^{15–18}

MicroRNAs (miRNAs) are a class of 22-nucleotide noncoding RNAs encoded by endogenous genes.¹⁹ They are known to induce mRNA degradation by binding to the 3'-untranslated region (UTR) of the target.²⁰ Aberrant microRNA expression influences tumor development and progression in breast cancers.²¹ miR-365 has been found to inhibit breast cancer cell growth and chemo-resistance through GALNT4.²² And miR-365 is considered as a potential diagnostic biomarker for breast cancer.²³

In the present study, we examine the function of FOXX1 in the development and progression of breast cancer. We discovered that FOXX1 is up-regulated in breast cancer tissues and cell lines. FOXX1 promotes cell migration and invasion by regulating EMT process in breast cancer. Interestingly, our work found that miR-365-3p is up-stream of FOXX1, so we provided that miR-365-3p-FOXX1 axis plays key roles in the progression of breast cancer.

Materials and Methods

Cell Lines

Human breast cancer MCF-7 cells, MDA-MB-231 and human normal breast epithelial cell line MCF-10A were available from American Type Culture Collection (Manassas, VA, USA). MCF-10A cells were cultured in DMEM/F12 medium supplemented with insulin (10 µg/mL), EGF (10 ng/mL), heat-inactivated horse serum (5%), cholera toxin (1 µg/mL) and hydrocortisone (1 µg/mL). MCF-7 and MDA-MB-231 cells were cultured in DMEM (Fisher Scientific International Inc., Hampton, NH, USA) supplemented with 100 µg/mL penicillin/streptomycin (Fisher Scientific International Inc.) and 10% fetal bovine serum (FBS, Fisher Scientific International Inc.). All cells were cultured at 37°C in a humidified atmosphere with 5% CO₂.

Tissue Samples

A total of 93 tissue samples were collected from patients with breast cancer which was diagnosed at the Linyi Central Hospital from March 2005 to April 2013. None of the patients underwent therapy prior to specimen collection, such as chemotherapy and radiotherapy. We had obtained the approval of the ethics committee of the Linyi Central Hospital before this study. Informed consent has been signed by all patients participated in this study.

Western Blotting

Tissue samples and cells were lysed in ice-cold RIPA lysis buffer (Beyotime Institute of Biotechnology) to extract whole protein according to the manufacturer's protocol. The protein concentration was measured using Pierce™ BCA Protein Assay Kit (Pierce, Waltham, MA, USA). Fifty-microgram protein samples were loaded in each lane, which was further separated by 8% sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and transferred to nitrocellulose filter membranes (Millipore, Billerica, MA, USA). After blocking in 5% skimmed milk at room temperature for 1 h, the membranes were incubated with indicated antibodies at 4°C overnight and then incubated with horseradish peroxidase (HRP)-conjugated secondary antibodies for 1 h at room temperature. Finally, the blots were detected by enhanced chemiluminescence (Cell Signaling Technology) according to the manufacturer's protocol. β-Actin was used as an internal control. Antibodies as followed: FOXX1 (no. HPA018864; 1:2000; Sigma-Aldrich, St. Louis, MO, USA), E-cadherin (no. HPA004812; 1:1000; Sigma-Aldrich, St. Louis, MO, USA), N-cadherin (no. HPA058574; 1:1000; Sigma-Aldrich, St. Louis, MO, USA), vimentin (no. HPA001762; 1:2000; Sigma-Aldrich, St. Louis, MO, USA), Slug (no. SAB1305973; 1:2000; Sigma-Aldrich, St. Louis, MO, USA), Snail (no. SAB1306281; 1:2000; Sigma-Aldrich, St. Louis, MO, USA) β-actin (no. A1978; 1:5000; Sigma-Aldrich, St. Louis, MO, USA), Goat Anti-Mouse IgG H&L (HRP) (no. ab205719; 1:3000; Abcam, Cambridge, UK) and Goat Anti-Rabbit IgG H&L (HRP) (no. ab6721; 1:5000; Abcam, Cambridge, UK).

RNA Extraction, Reverse Transcription, and Real-Time PCR

Total RNA was extracted from tissues or cells using TRIzol (Life Technologies) according to the manufacturer's protocol. Two micrograms RNA was used to synthesize complementary DNA (cDNA) using a poly-A

polymerase-based First-Strand Synthesis kit (Takara, Dalian, China). Subsequently, SYBR Green I (Applied Biosystems) was used to perform RT-qPCR. Primers for U6 (no. miRAN0002-1-100) and miR-365-3p (no. MQPS0000357-1-100) were purchased from RiboBio (Guangzhou, China). GAPDH or U6 was used as internal controls for mRNA or miRNA, respectively. Relative fold expressions were calculated using $2^{-\Delta\Delta Ct}$ method.²⁴

Cell Counting Kit (CCK-8) Assay

To determine the effect of FOXK1 on cell proliferation, CCK-8 assay was performed according to the manufacturer's protocol. MCF-7 and MDA-MB-231 cells were transfected with FOXK1 plasmid or FOXK1 siRNA, respectively. After transfection for 24 h, cells were placed into a 96-well plate at a density of 3×10^3 cells in 200 μ L DMEM supplement with 10% FBS. Twenty-microliter CCK-8 was added into each well and cells were incubated for 60 min at 37°C. At 0 h, 24 h, 48h and 72 h, absorbance was measured at 450 nm using an ELx800 microplate reader (BioTek Instruments, Inc., Winooski, VT, USA). Each experiment was carried out three times.

Cell Cycle Analysis

Approximately 4×10^5 transfected cells were harvested by 0.25% trypsinization and washed with ice-cold phosphate-buffered saline (PBS) three times. After fixation in 75% ice-cold ethanol in PBS at 4°C overnight, cells were washed with ice-cold and incubated with RNase at 37°C at least for 30 min, followed by incubation with propidium iodide (PI) for 20 min at room temperature. Finally, the cell cycle was determined using FACSCanto II flow cytometer (Becton-Dickinson & Co., Franklin Lakes, NJ, USA). Each experiment was carried out three times.

Transwell Invasion Assay

Transwell chamber (BD Biosciences) was used to measure the ability of cells to invade. First, 80 μ L Matrigel (Matrigel: DMEM=1:8) was added into Transwell chambers and incubated at 37°C for 30 min to solidify. The upper chamber contained almost 2×10^4 cells in 200 μ L serum-free DMEM. The lower chamber contained 500 μ L DMEM with 10% FBS. After incubation at 37°C in an atmosphere containing 5% CO₂ for 48 h, cells on the surface of membranes were removed by swab cotton softly and stained with 0.1% crystal violet for 15 min at room temperature. Finally, the invading cells were imaged and counted under a light microscope at

$\times 200$ magnification (Olympus Corporation). Each experiment was carried out three times.

Wound-Healing Assay

MDA-MB-231 cells were transfected with FOXK1 plasmid or FOXK1 siRNA for 48 h and placed into 6-well plates at the density of 6×10^6 cells/well with serum-free DMEM. After scratching using a pipette tip, cells were incubated at 37°C in an atmosphere containing 5% CO₂ for 48 h. Images of wound healing were captured under an inverted microscope at $\times 100$ magnification (Olympus Corporation, Tokyo, Japan) at 0 h and 48 h after scratching. Results were analysed with ImageJ software (version 1.2; National Institutes of Health). Each experiment was carried out three times.

Dual-Luciferase Reporter Assay

MCF-7 cells were placed in 12-well plates and co-transfected with miR-365 mimics, renilla and WT-FOXK1 3'-UTR or MT-FOXK1 3'-UTR. After transcription for 24 h, the cells were harvested. The luciferase activity was evaluated using a dual-luciferase reporter assay system (Promega, Madison, WI) according to the manufacturer's instruction. Firefly luciferase activity was normalized to Renilla luciferase activity. Each experiment was carried out three times.

Statistical Analysis

The results are expressed as the mean \pm standard deviation from at least three separate experiments. All data were analyzed using GraphPad Prism 8.0 software (GraphPad Software, Inc., La Jolla, CA, USA). Student's *t*-test and one-way analysis of variance followed by the Turkey post hoc test was performed to determine the statistical significance among two groups or multiple groups, respectively. Kaplan–Meier method followed by log-rank test was used to perform survival curves (gepia.cancer-pku.cn) $P < 0.05$ was considered as a statistically significant difference.

Results

FOXK1 Is Up-Regulated in Breast Cancer Tissues and Cells

To explore the potential role of FOXK1 in breast cancer, we first detected the FOXK1 mRNA and protein levels in 13 paired primary breast cancer tissues and the corresponding adjacent normal tissues using RT-qPCR and Western blotting analyses. The result revealed that the expression of FOXK1 in primary breast cancer tissues was remarkably higher than those in the adjacent normal breast tissues (Figure 1A).

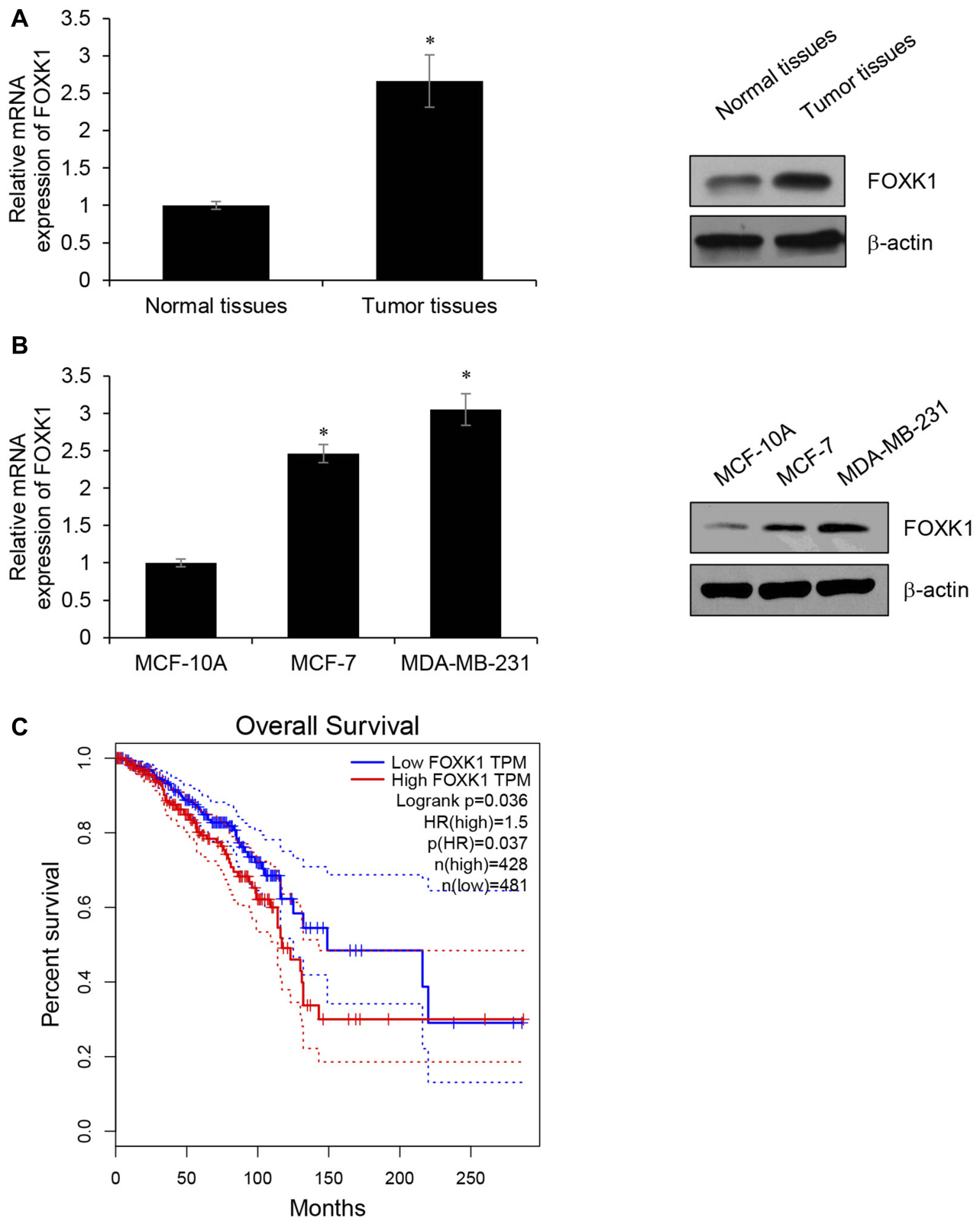


Figure 1 FOXK1 is up-regulated in breast cancer tissues and cells. **(A)** The expression of FOXK1 in breast cancer tissues and adjacent normal tissue samples as detected by RT-qPCR and Western blotting. * $p<0.05$ vs Adjacent normal tissues. **(B)** The expression of FOXK1 in breast cancer cells as detected by RT-qPCR and Western blotting. * $p<0.05$ vs MCF-10A. **(C)** The survival curve was analyzed on GEPIA website (gepia.cancer-pku.cn).

Moreover, we also determine the expression of FOXK1 in breast cancer cell lines, including MCF-7 and MDA-MB-231. Consistent with the expression of FOXK1 in breast cancer tissues, both mRNA and protein levels of FOXK1 were up-regulated in MDA-MB-231 and MCF-7 cells compared with MCF-10A cells (Figure 1B). In addition, as shown in Table 1, we found that the expression of FOXK1 is positively associated with tumor size, TNM stage and lymph node metastasis (Table 1). We also assessed the correlation of FOXK1 and the survival rate using GEPIA website (gepia.cancer-pku.cn), suggesting that high expression of FOXK1 predicted a poor prognosis (Figure 1C). These results suggest that FOXK1 is up-regulated and plays critical roles in breast cancer.

FOXK1 Promotes the Proliferation of Breast Cancer Cells

Because of the correlation of FOXK1 expression and tumor size, we assumed that FOXK1 might promote cell proliferation in breast cancer. To verify our hypothesis, we overexpressed or knocked down FOXK1 in MCF-7 and MDA-MB-231 cells, respectively. The expression of FOXK1 was established using RT-qPCR and Western blotting analyses (Figure 2A). To determine the effect of FOXK1 on cell proliferation, we performed CCK-8 assay and colony formation assay. As shown in Figure 2B, ectopic expression of FOXK1 obviously increased the proliferation rate of MCF-7 cells and inhibition of FOXK1 decreased the proliferation

rate (Figure 2B). The similar results were found in MDA-MB-231 cells (Figure 2B). The result of colony formation assay revealed that overexpression of FOXK1 resulted in an elevated number of colonies and inhibition of FOXK1 led to a reduced number of colonies (Figure 2C). Next, we further decipher whether FOXK1 promotes cell proliferation through regulation of cell cycle, so we then detected the effect of FOXK1 on cell cycle. We found that FOXK1 could facilitate G₁/S phase transition (Figure 2D). Together, our finding indicates that FOXK1 promotes the cell proliferation in breast cancer through facilitating G₁/S phase transition.

FOXK1 Improves the Invasive Ability of Breast Cancer Cells

We next determined the potential impact of FOXK1 on breast cancer cell invasion capacity using wound-healing assay and Transwell invasion assay. The number of invaded MDA-MB-231 cells was significantly increased after transfection with ectopic expression-FOXK1 when compared with the mock group (Figure 3A). And inhibition of FOXK1 also promotes the invasion of MDA-MB-231 cells (Figure 3A). Moreover, as shown in Figure 3B, the wound-healing assay indicated that the relative migration distance was increased upon FOXK1 overexpression in MDA-MB-231 cells (Figure 3B). Matrix metalloproteinases (MMPs) are key enzymes for invasion and metastasis, which are the major causes of mortality in breast cancer patients.²⁵ Among them, MMP-2/9 expression is linked with increased metastasis in various tumors, including the brain, prostate, and breast.^{25,26} So, we then determined the effects of FOXK1 on the expression of MMP-2 and MMP-9 in MDA-MB-231 cells. As shown in Figure 3C, FOXK1 overexpression significantly increased the secretion of MMP-2 and MMP-9, in contrast, FOXK1 inhibition decreased the secretion of MMP-2 and MMP-9 in MDA-MB-231 cells (Figure 3C).

FOXK1 Regulates EMT in Breast Cancer

As previously described, FOXK1 increases breast cancer cell migration and invasion capacity. We next explored whether FOXK1 may also be involved in the process of EMT in breast cancer. As shown in Figure 4A, our results demonstrated that ectopic expression of FOXK1 led to increase the mRNA expression of N-cadherin and vimentin; whereas decreased expression of E-cadherin in MCF-10A cells (Figure 4A and B). In contrast, inhibition of FOXK1 in

Table 1 Clinic Pathologic Variables in 93 BC Patients

| Variables | No. (n=93) | FOXK1 Protein Expression | | P value |
|-----------------------|------------|--------------------------|-------------|---------|
| | | Low (n=30) | High (n=63) | |
| Age | | | | |
| ≥ 40 | 51 | 18 | 33 | 0.49 |
| < 40 | 42 | 12 | 30 | |
| Tumor size | | | | |
| Large (≥ 5 cm) | 48 | 10 | 38 | 0.015 |
| Small (< 5 cm) | 45 | 20 | 25 | |
| Pathological grade | | | | |
| I-II | 41 | 18 | 23 | 0.033 |
| III-IV | 52 | 12 | 40 | |
| Lymph node metastasis | | | | |
| Yes | 50 | 11 | 39 | 0.022 |
| No | 43 | 19 | 24 | |

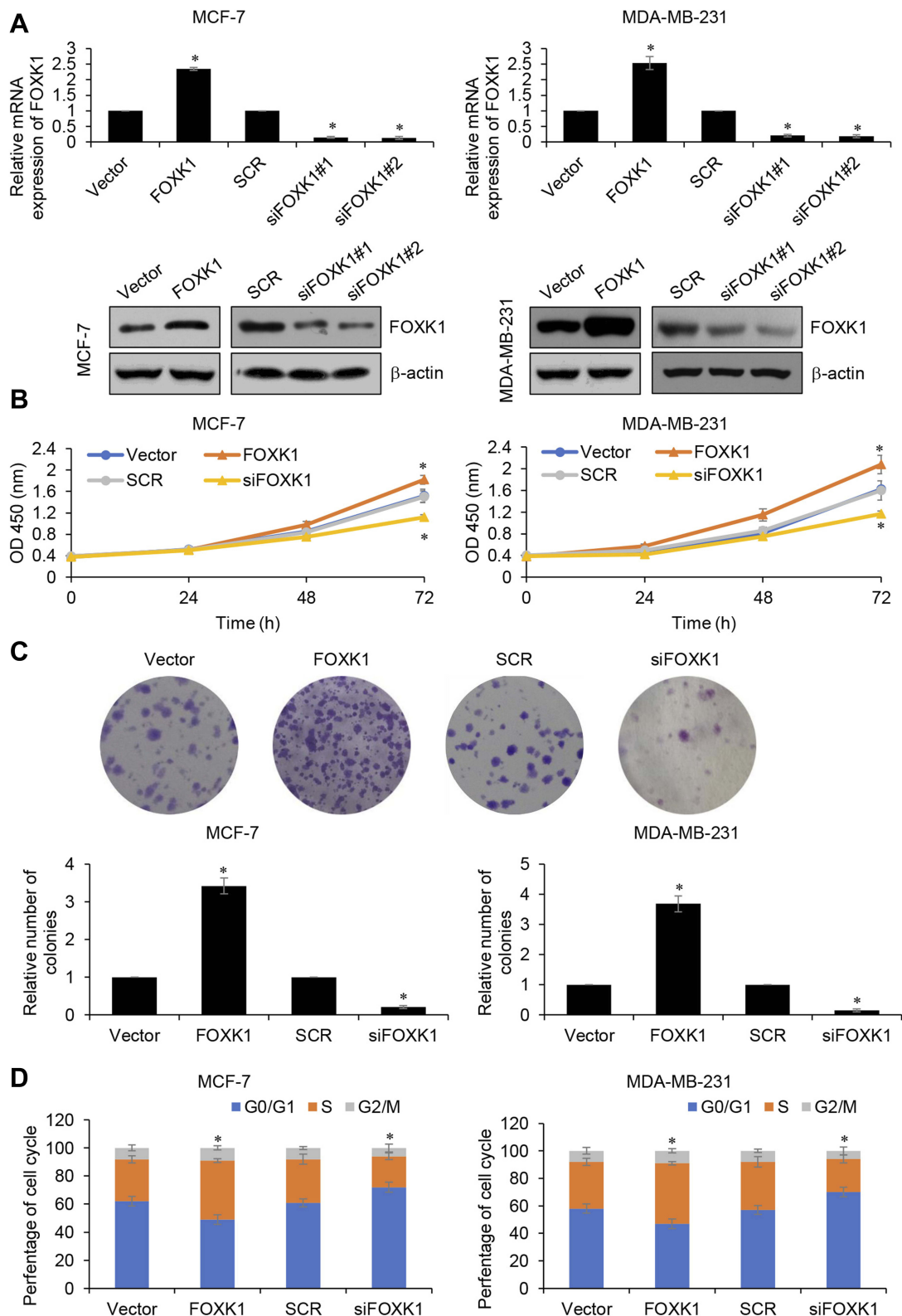


Figure 2 FOXK1 promotes the proliferation of breast cancer cells. (A) FOXK1 was overexpressed or knocked down in MCF-7 and MDA-MB-231 cells, the expression of FOXK1 was established using RT-qPCR and Western blotting. * $p < 0.05$ vs vector or scramble siRNA (SCR). (B) CCK-8 analysis of FOXK1-transfected MCF-7 and MDA-MB-231 and control cells. * $p < 0.05$ vs vector or SCR. (C) Colony formation analysis of FOXK1-transfected MCF-7 and MDA-MB-231 and control cells. * $p < 0.05$ vs vector or SCR. (D) After transfection, cell cycle assay was performed. * $p < 0.05$ vs vector or SCR.

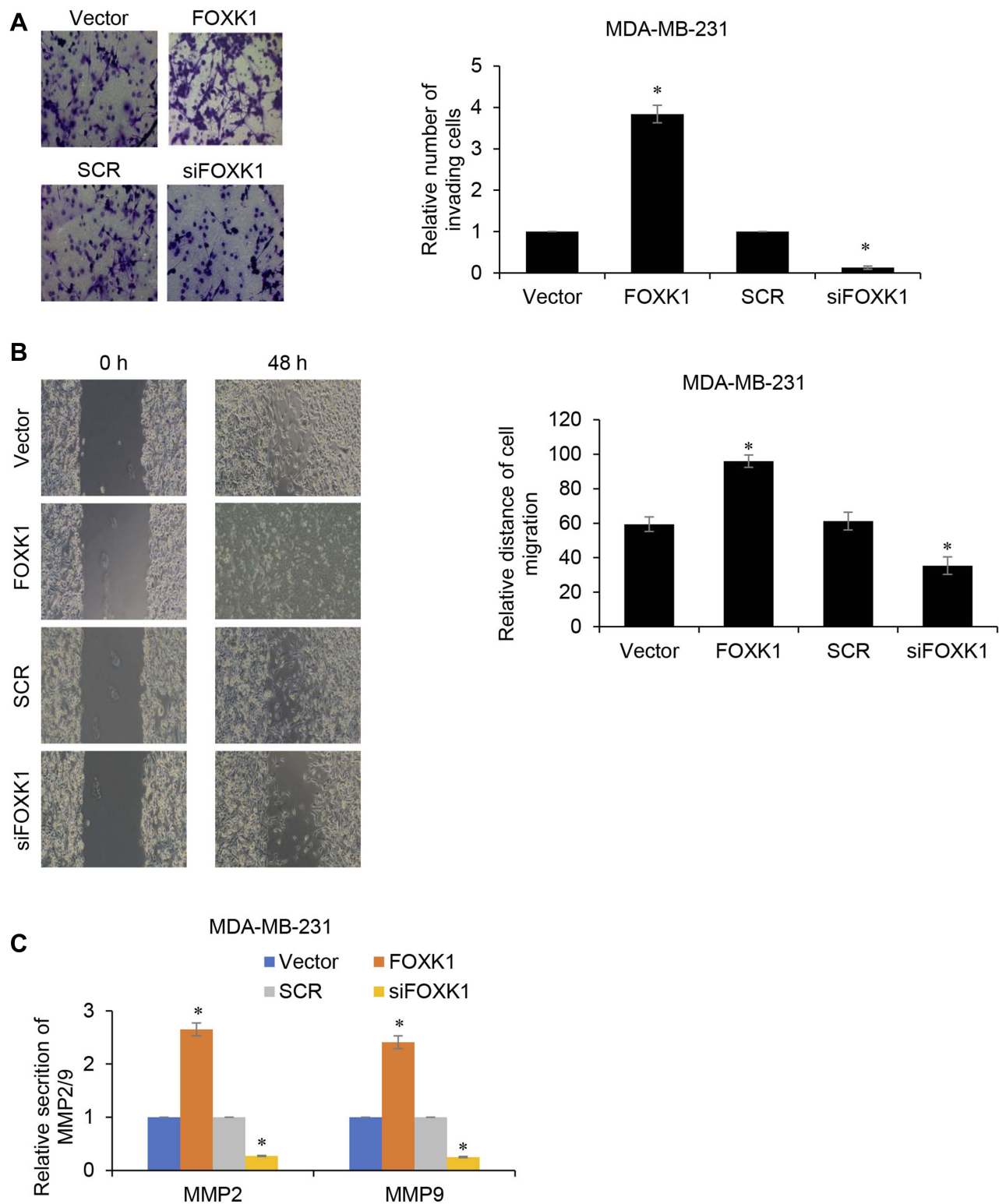


Figure 3 FOXK1 improves the invasive ability of breast cancer cells. **(A)** Transwell invasion assay in FOXK1-transfected MDA-MB-231 and control cells. * $p < 0.05$ vs vector or SCR. **(B)** Wound-healing assay in FOXK1-transfected MDA-MB-231 and control cells. * $p < 0.05$ vs vector or SCR. **(C)** The effect of FOXK1 on MMP2/9 secretion was identified by MMP2/9 ELISA kit. * $p < 0.05$ vs vector or SCR.

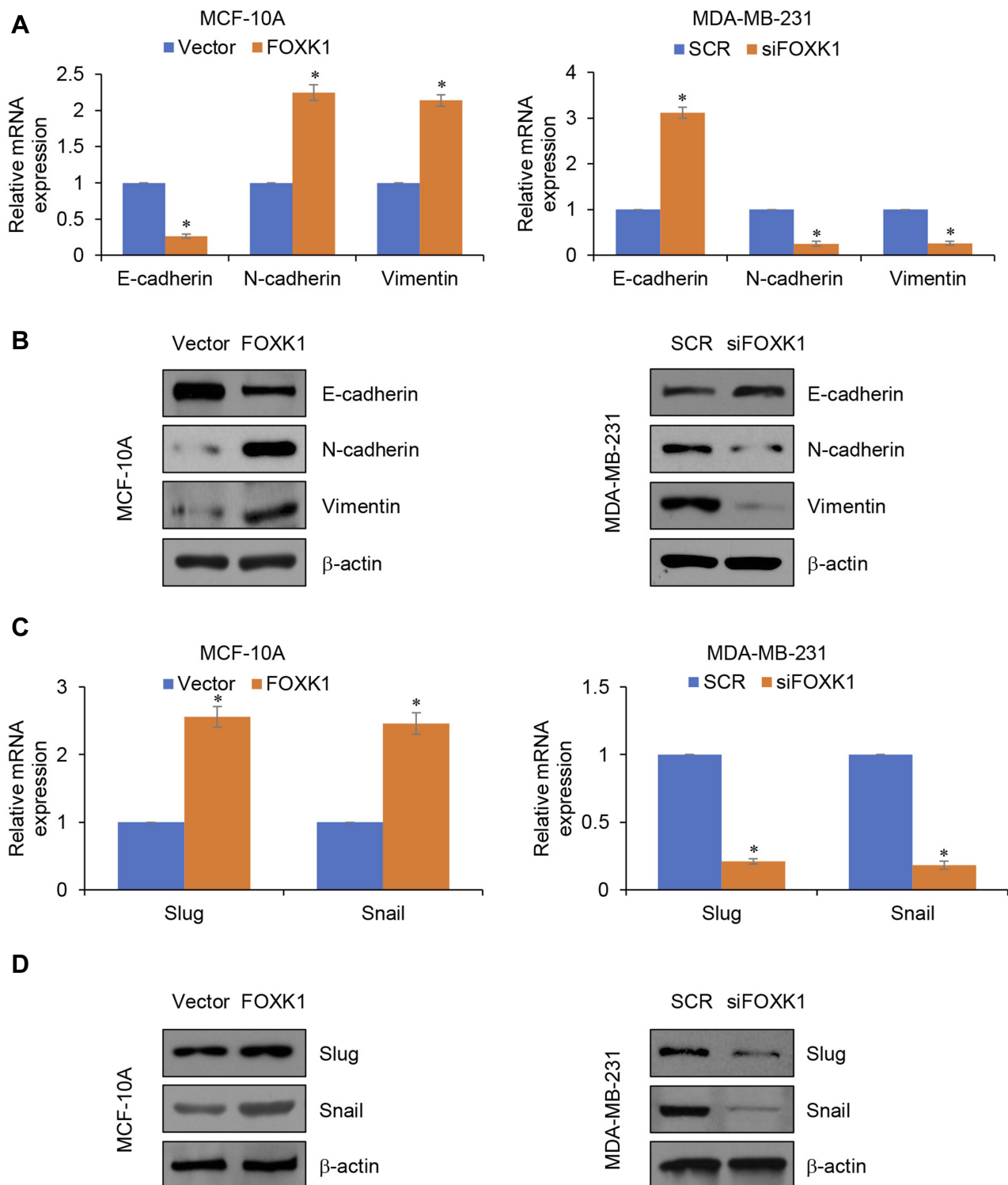


Figure 4 FOXC1 regulates EMT in breast cancer: **(A and B)**, FOXC1 was overexpressed or knocked down in MCF-10A or MDA-MB-231 cells, respectively. The expression of EMT markers was established using RT-qPCR and Western blotting. * $p < 0.05$ vs vector or SCR. **(C and D)**, The effect of FOXC1 on slug and snail expression was identified by RT-qPCR and Western blotting. * $p < 0.05$ vs vector or SCR.

MDA-MB-231 cells significantly decreased the expression of mesenchymal markers N-cadherin and vimentin and improved the expression of epithelial marker E-cadherin

(Figure 4A and B). In addition, the expression of EMT-related transcription factors, such as Slug and Snail, was obviously elevated in the FOXC1-overexpressed MCF-10A

cells when compared with that of the control cells (Figure 4C and D). In contrast, inhibition of FOXK1 in MDA-MB-231 reduced the expression of Slug and Snail in mRNA and protein levels (Figure 4C and D). Together, our work indicates that FOXK1 regulates EMT in breast cancer.

FOXK1 Is a Target of miR-365-3p in Breast Cancer

Increasing evidence supports that micro-RNA (miR) plays a key role in human cancer through targeting transcription factors. To further explore the underlying mechanism of FOXK1 in breast cancer, we predicted the up-stream of FOXK1 using TargetScan. The result indicated that FOXK1 might be a target of miR-365-3p (Figure 5A). To

further confirm this target association, the dual-luciferase reporter gene plasmid containing the wild-type (WT) or mutant-type MT of FOXK1 3'-UTR (Figure 5B) was constructed. The result of dual-luciferase reporter assays in MCF-7 cells suggested that inhibition of FOXK1 only obviously increased relative luciferase activity of reporter gene with wild-type, but not MT of FOXK1 3'-UTR, suggesting the essential role of miR-365-3p binding site within FOXK1 3'-UTR (Figure 5C). Meanwhile, we also detected the effects of miR-365-3p on the expression of FOXK1. As shown in Figure 5D, we found FOXK1 expression was negatively regulated by miR-365-3p (Figure 5D). To sum up, our finding reveals that FOXK1 is a downstream target of miR-365-3p in breast cancer.

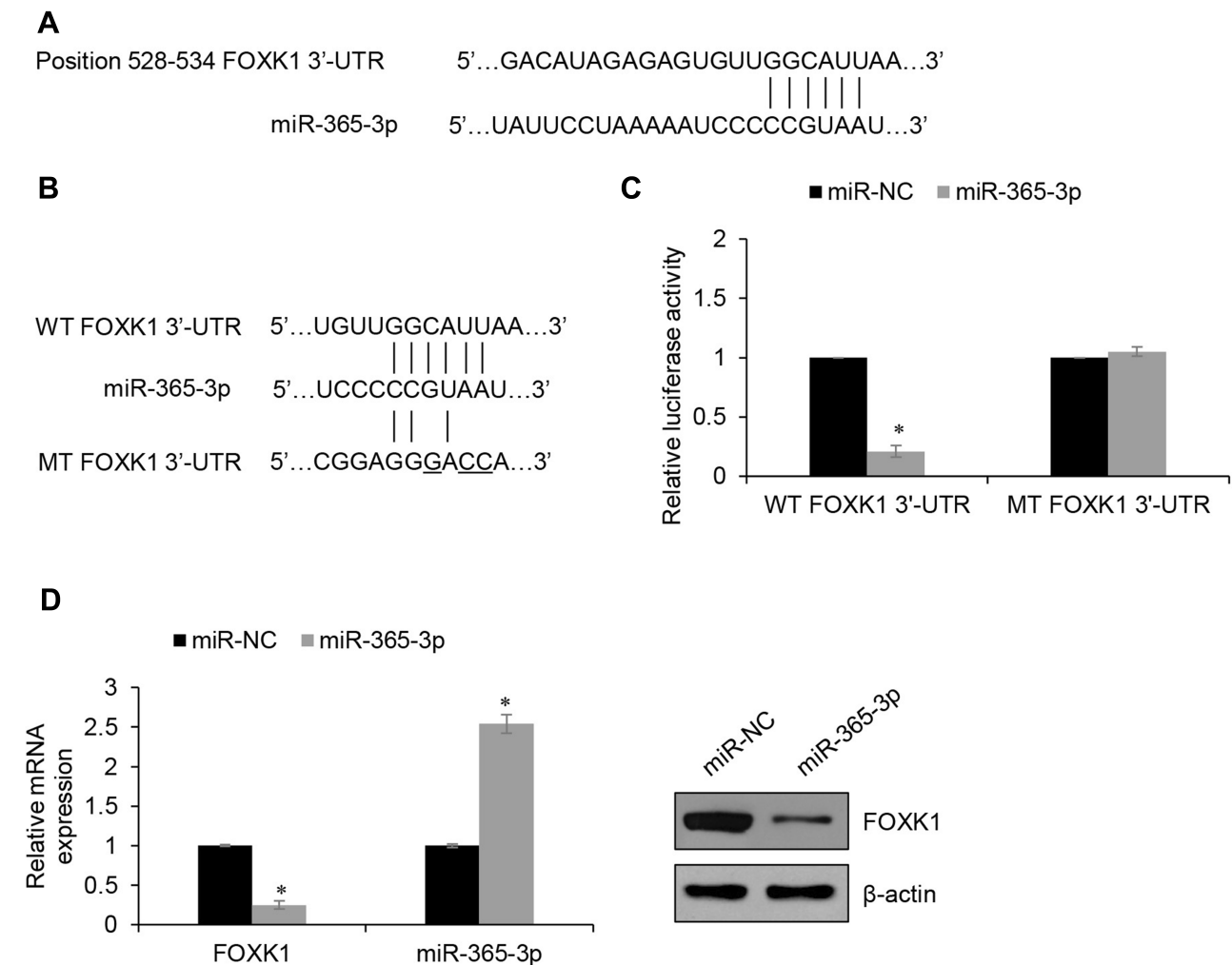


Figure 5 FOXK1 is a target of miR-365-3p in breast cancer. (A) FOXK1 is predicted to be a potential target of miR-365-3p. (B) The luciferase reporter gene plasmids containing the wild type (WT) and mutant type (MT) of FOXK1 3'-UTR were generated. (C) Dual-luciferase reporter assay revealed that the luciferase activity was decreased in MCF-7 cells co-transfected with miR-365-3p mimics and WT 3'-UTR of FOXK1 reporter plasmid, but unaltered in cells co-transfected with miR-365-3p mimics and MT FOXK1 3'-UTR plasmid. * $p < 0.05$ vs miR-NC. (D) RT-qPCR and Western blotting analyses were conducted to examine the effect of miR-365-3p on the expression of FOXK1. * $p < 0.05$ vs miR-NC.

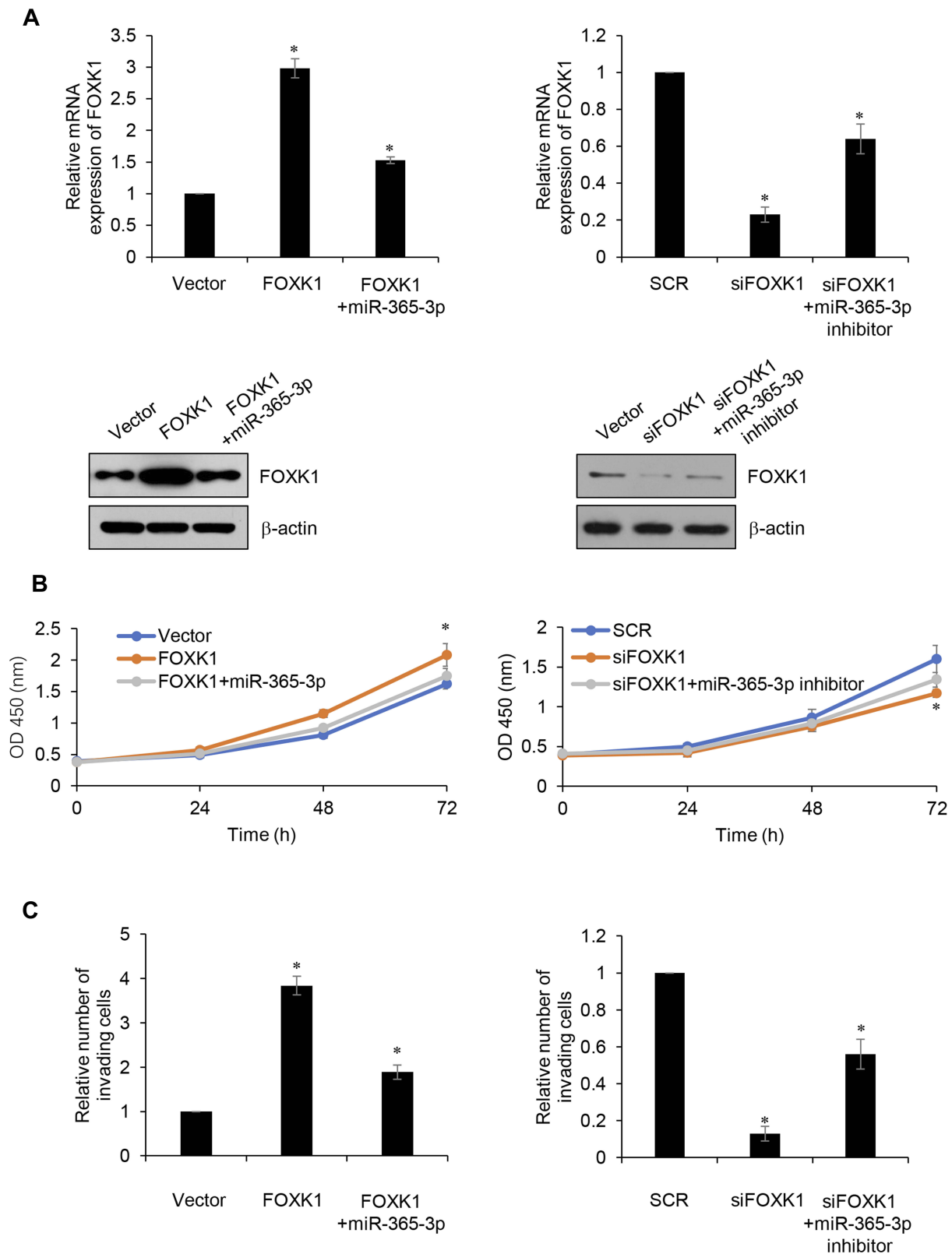


Figure 6 Up-reComment: Dear TS, please replace fig 6 with attached fig 6regulation of miR-365-3p reverses the effects of FOXK1 on proliferation and invasion in breast cancer cells. **(A)** MDA-MB-231 cells were co-transfected with FLAG-FOXK1 plasmid or/and miR-365-3p mimic or co-transfecting siFOXK1 or/and miR-365-3p inhibitor; the expression of FOXK1 was identified using RT-qPCR and Western blotting analyses. * $p < 0.05$ vs vector or SCR. **(B)** After transfection, CCK-8 assay was carried out to determine cell proliferation. * $p < 0.05$ vs vector or SCR. **(C)** After transfection, Transwell assay was performed to determine cell invasion. * $p < 0.05$ vs vector or SCR.

miR-365-3p Reverses the Effects of FOXC1 on Proliferation and Invasion in Breast Cancer Cells

To further investigate whether miR-365-3p was involved in FOXC1-mediated malignant phenotypes of breast cancer cells, MDA-MB-231 cells were co-transfected with pcDNA3.1-FOXC1 plasmids or/and miR-365-3p mimics or co-transfecting siFOXC1 or/and miR-365-3p inhibitor. The mRNA and protein levels of FOXC1 were established using RT-qPCR and Western blotting analyses, respectively (Figure 6A). The results of CCK-8 and Transwell invasion assay demonstrated that up-regulation of miR-365-3p reversed the effects of FOXC1 on proliferation and invasion in MDA-MB-231 cells, and inhibition of miR-365-3p reversed the effects of FOXC1 inhibition on proliferation and invasion in MDA-MB-231 cells, suggesting that miR-365-3p is indeed involved in FOXC1-mediated malignant phenotypes of breast cancer cells (Figure 6B and C).

Discussion

In the present study, FOXC1 expression was up-regulated in breast cancer tissues and cell lines. The expression of FOXC1 was associated with tumor size, TNM stage and metastasis. Moreover, the results of survival curve revealed that high FOXC1 expression predicted a poor prognosis, thus suggesting that FOXC1 may serve as a therapeutic target in breast cancer. In addition, CCK-8, colony formation and cell cycle assays demonstrated that ectopic expression of FOXC1 in breast cancer cells significantly promotes cell proliferation through facilitating G₁/S phase transition. Moreover, we found that FOXC1 also reduces cell migration and invasion through the regulation of EMT. EMT is a critical step in cancer metastasis,^{27,28} epithelial cells change phenotype and obtain the characteristics of mesenchymal cells, gaining the ability to migrate and contribute to tumor metastasis.²⁹ During EMT process, the expression of several proteins is alteration, such as epithelial marker E-cadherin, mesenchymal markers N-cadherin and vimentin, EMT-associated transcription factors TWIST, Snail and Slug.³⁰ Multiple studies have demonstrated that FOXC1 acts as a critical mediator of EMT in certain tumors, such as gastric cancer,³¹ colorectal cancer³² and prostate cancer,³³ indicating that FOXC1 regulates the EMT process and may serve as the core protein during EMT in human cancers.

MicroRNAs (miRNAs) are a class of 22-nucleotide non-coding RNAs encoded by endogenous genes.¹⁹ MiRNAs

display abnormal expression and functions in multiple kinds of malignancies, which act as different kinds of tumor-related genes.^{34,35} miR-365-3p has been proven to act as a potential inhibitor in different types of cancer. For example, miR-365-3p inhibits the progression of cutaneous squamous cell carcinoma through regulating cyclin-dependent kinase 6 and 4,³⁶ and miR-365-3p also suppresses tumor growth, proliferation and metastasis in malignant myeloma through targeting NRP.³⁷ Consistent with a previous report, we found that miR-365-3p is down-regulated in breast cancer, and regulates cell proliferation, migration and invasion through targeting FOXC1.

However, there are still present several limitations to our work. First, how FOXC1 regulates proliferation need to be further investigated. RNA-seq and ChIP-seq can perform to reveal the down-stream of FOXC1 in breast cancer. Moreover, whether FOXC1 facilitates tumor proliferation and metastasis needs to be verity in vivo.

In conclusion, our study provides evidence that FOXC1 is up-regulated in breast cancer and serve as an oncogene. Abnormal expression of FOXC1 can promote cell proliferation, migration and invasion. miR-365-3p has been found to down-regulate in breast cancer, here, our work supports that FOXC1 is a target of miR-365-3p, miR-365-3p-FOXC1 axis may as a potential therapeutic strategy for breast cancer.

Disclosure

The authors report no conflicts of interest in this work.

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