

# Circular RNA\_0074027 participates in cell proliferation, apoptosis and metastasis of colorectal cancer cells through regulation of miR-525-3p

GANG XIONG, JUN ZHANG, YICHAO ZHANG, XIAO PANG, BIAO WANG and YONGCHUAN ZHANG

Department of General Surgery, Dazhou Central Hospital, Dazhou, Sichuan 635000, P.R. China

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**Abstract.** The present study aimed to elucidate the biological function of circular RNAs (circRNA) 0074027 in colorectal cancer (CRC). The expression of circRNA-0074027 in CRC tissues and cells was determined by reverse transcription-quantitative PCR. The *in vitro* experiments, including Cell Counting Kit-8 (CCK-8) assay, 5-Ethynyl-2'-deoxyuridine assay, flow cytometry and Transwell assay, were applied to evaluate cell proliferation, apoptosis and metastasis ability respectively following downregulation of circRNA-0074027. The correlation between circRNA-0074027 and micro (mi)RNA-525-3p was determined via dual-luciferase reporter assay. Finally, western blotting was used to explore the possible regulatory mechanism. CircRNA-0074027 was upregulated in CRC tissues, while miR-525-3p expression was reduced. In addition, patients with CRC and circRNA-0074027 overexpression were more likely to have low tumor differentiation, lymph node metastasis and advanced TMN stage. Deletion of circRNA-0074027 could suppress cell proliferation and metastasis through upregulating p53 expression and forbidding epithelial-mesenchymal transition signaling pathway. The addition of miRNA-525-3p inhibitors could reverse the anti-tumor effects induced by the deletion of circRNA-0074027. The downregulation of circRNA\_0074027 inhibited tumor progression via sponging miR-525-3p, which could be a promising treatment bio-marker for CRC.

## Introduction

Colorectal cancer (CRC) is one of the commonest malignant tumors worldwide, with high mortality and morbidity (1,2). Due to a lack of early symptoms and effective screening strategies, patients are commonly diagnosed with advanced

CRC (2). Although there has been major progress in molecular target therapy and immunotherapy (3,4), the overall survival time for patients with CRC is still not satisfactory. Therefore, it is essential to identify an effective bio-marker for CRC and explore its potential regulatory mechanism.

Circular RNAs (circRNAs), a novel type of non-coding RNA, are characterized as closed ring structure without 5' caps or 3' poly-A tails (5,6). Emerging studies have shown that circRNAs may serve a function in the regulation of physiological and pathological processes via serving as microRNA (miRNA) sponges (7-9). A number of circRNAs are believed to serve a function as a tumor promoter or a tumor inhibitor in various cancer progression, including liver, ovarian and lung cancer (10-12). For example, silencing of circRNA-0060428 can inhibit cell growth in osteosarcoma by regulating the miR-375/RPBJ axis (13). CircRNA-0003645 can sponge miR-1229 and then promote hepatocellular carcinoma progression (14). However, the biological role of circRNAs in the development of CRC remains to be elucidated.

CircRNA-0074027 is a newly discovered competing endogenous (ce)RNA, which is oriented from chr5:134363423-134369964. The biological role of circRNA-0074027 in malignant transfection has been explored (15,16). Gao *et al* (16) reported that circRNA-0074027 is upregulated in lung cancer and its overexpression can promote cell growth and metastasis ability via regulation of the miR-185-3p/bromodomain-containing protein 4/MAPK-activating death domain protein axis. In addition, Qian *et al* (15) demonstrated that the deletion of circRNA-0074027 can inhibit glioblastoma cell proliferation and metastasis by regulating the miR-518a-5p/IL17RD signaling pathway. However, it remains to be fully elucidated whether circRNA-0074027 serves as a tumor promoter in CRC progression.

The present study detected the expression of circRNA-0074027 in CRC tissue samples and further investigated the role of circRNA-0074027 in oncogenesis. *In vitro* experiments were performed to identify the cell apoptosis rate, cell proliferation and metastasis ability. In addition, miR-525-3p was identified as a downstream target gene of circRNA-0074027 by bioinformatics analysis and the relationship was further confirmed by dual-luciferase reporter assays. Taken together, circRNA-0074027 is a promising bio-marker in CRC.

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**Correspondence to:** Dr Yongchuan Zhang, Department of General Surgery, Dazhou Central Hospital, 56 Nanyuemiao Street, Dazhou, Sichuan 635000, P.R. China  
E-mail: xiong123456g@126.com

**Key words:** circular RNA-0074027, microRNA-525-3p, colorectal cancer, cell proliferation, cell metastasis

## Materials and methods

**Patient tissue samples.** A total of 60 paired CRC tissue and normal tissue samples were obtained from the patients with CRC who underwent surgery at Dazhou Central hospital between January 2014 and January 2018. The inclusion criteria were: i) Age >18 and <80 years; ii) written informed consent; and iii) primary CRC confirmed by pathological examination. The exclusion criteria were: i) Patients with other malignant diseases and ii) patients with previous neoadjuvant chemotherapy or radiotherapy. All samples were preserved in liquid nitrogen following histological confirmation by experienced pathologists. Written informed consent was sought from the participants before the samples were obtained. The present study was approved by the Ethical Committees of Dazhou Central Hospital (approval no. KY2020-086-01).

**Cell culture.** Normal epithelial cells (FHC) and CRC cancer cells (Caco-2, LoVo, SW480, HCT-116 and HCT-8), as well as 293T cell lines were acquired from American Type Culture Collection. All cell lines were incubated in 10% fetal bovine serum (FBS; Thermo Fisher Scientific, Inc.) supplemented with DMEM medium (Thermo Fisher Scientific, Inc.) under standard culture conditions (5% CO<sub>2</sub> and 37°C).

**RNA transfection.** Small interfering (si) RNA against circRNA-0074027 (si-circRNA-0074027-1 and si-circRNA-0074027-2) and its paired control (si-Control), as well as miR-525-3p inhibitors and its negative control (miR-525-3p NC) were acquired from Shanghai GenePharma Co., Ltd. Lipofectamine® 3000 (Invitrogen; Thermo Fisher Scientific, Inc.) was used for cell transfection. In brief, Lipofectamine® 3000 was incubated with miRNA inhibitor at a concentration of 100 nmol/l or siRNAs at a concentration of 200 nmol/l for 20 min at room temperature and the complex was then added to each well of a 6-well plate. Following transfection for 24 h, the subsequent experiments were performed. The sequences were: si-circRNA-0074027-1 (5'-GCGTGC TAAGCACCTGGCGCA-3'), circRNA-0074027-2 (5'-GTG CTAAGCACCTGGCGCAGG-3') and miR-525-3p inhibitor (5'-CGCUCUAAAGGGAAGCGCCUUC-3').

**Reverse transcription-quantitative (RT-q) PCR.** According to the manufacturer's protocols, TRIzol® (Thermo Fisher Scientific, Inc.) was used for total RNA extraction from tissue and cell samples. Subsequently, total RNA (1 µg) was transcribed to cDNA via using PrimeScript RT Master Mix (Takara Biotechnology Co., Ltd.). The temperature and duration of RT were: 37°C for 30 sec, followed by 85°C for 5 sec and 4°C for 10 min. The cDNA (10 ng) was subjected to qPCR (Bio-Rad Laboratories, Inc.) using SYBR Premix Ex Taq II kit (Takara Biotechnology Co., Ltd.), and the reaction volume was 10 µl. The thermocycling conditions were: 95°C for 30 sec, followed by 40 cycles of 95°C for 5 sec and 60°C for 40 sec. GAPDH was used as reference gene and relative mRNA expression changes were calculated by 2<sup>-ΔΔC<sub>q</sub></sup> method (17). The primer sequences used are shown in Table I.

**Cell proliferation assay.** A CCK-8 kit (Dojindo Molecular Technologies, Inc.) was used to assess the cell viability at

different time points. A 5-Ethynyl-2'-deoxyuridine (EdU) cell proliferation kit (Guangzhou RiboBio Co., Ltd.) was used to identify the EdU positive rate of CRC cells.

**Flow cytometry.** An Annexin V-FITC/PI apoptosis detection kit (Nanjing KeyGen Biotech Co., Ltd.) was used for the detection of the cell apoptosis rate. According to the manufacturer's protocols, the transfected cells were collected and suspended with binding buffer. Then, Annexin V-FITC and propidium iodide were added to the buffer. Subsequently, the mixture was shielded from light and incubated at room temperature for 15 min. Subsequently, the samples were detected by flow cytometer (BD FACSCelesta™ Flow Cytometer; BD Biosciences), and early plus late apoptotic cells were analyzed using FlowJo software (FlowJo LLC).

**Transwell assay.** For the cell migration assay, the transfected cells were suspended in DMEM without FBS, added into the upper Transwell chambers (0.8 µm; Corning, Inc.) and incubated for 48 h. The lower chamber was filled with DMEM supplemented with 20% FBS. The migrated and invasive CRC cells were fixed with 4% methanol for 10 min, stained with crystal violet for 10 min at room temperature, and finally captured with a light microscope. For the cell invasion assay, the filter membranes were precoated with Matrigel® (Corning, Inc.) at 37°C for 30 min, and the other steps were performed as described in the migration assay.

**Western blotting.** The un-transfected and transfected cells were lysed using RIPA buffer (Beyotime Institute of Biotechnology), followed by homogenization at 4°C for 10 min. Protein concentration was evaluated with BCA Protein Assay kit (Beyotime Institute of Biotechnology). Equal amounts (20 µg) of proteins were loaded and subjected to 10% SDS-PAGE electrophoresis, followed by transferred onto polyvinylidene fluoride membranes. The membranes were blocked with 10% bovine serum albumin (Beijing Solarbio Science & Technology Co., Ltd.) for 1 h at room temperature. Subsequently, the membranes were incubated with primary antibodies: p-53 (cat. no. ab32389), Bcl-2 (cat. no. ab32124), Bax (cat. no. ab32503), E-cadherin (cat. no. ab11512), N-cadherin (cat. no. ab245117), Vimentin (cat. no. ab16700) and GAPDH (cat. no. ab9485; all 1:1,000 dilution; Abcam) overnight at 4°C. Subsequently, proteins were incubated with the goat anti-rabbit IgG H&L (HRP; cat. no. ab6721; 1:2,000 dilution; Abcam) for 1 h at room temperature. Finally, the protein bands were then analyzed with super sensitive ECL luminescence reagent (Dalian Meilun Biology Technology Co., Ltd.). The protein expression levels were detected using a chemi-luminescence detection system with Quantity One software (v3.0; Sigma-Aldrich; Merck KGaA).

**Prediction of downstream molecules regulated by circRNA-0074027.** A publicly available bioinformatics algorithm (Starbase 2.0) was utilized to predict the downstream miRNAs of circRNA-0074027 (9).

**Dual-luciferase reporter assays.** The wild-type (WT) and mutant (MUT) circRNA-0074027 fragments were constructed and inserted into the pmirGLO vector (Ybscience). The WT

Table I. Sequences of oligomers and primers used in the present study.

Name	Sequence (5'-3')
CircRNA-0074027 forward	GATTTCCCGACCCCGTACAA
CircRNA-0074027 reverse	GGGGTGTCTCTGAGATGGACC
miR-525-3p forward	GGAAGGCGCTTCCCTTT
miR-525-3p reverse	GTTGTGGTTGGTTGGTTTGT
GAPDH forward	CCTTCCGTGTCCCCACT
GAPDH reverse	GCCTGCTTCACCACCTTC
circRNA, circular RNA; miR, microRNA.	

and MUT plasmids, as well as miR-525-3p inhibitor and miR-525-3p NC, were co-transfected into 293T cells using Lipofectamine®3000 (Invitrogen; Thermo Fisher Scientific, Inc.). After 24 h, the firefly and *Renilla* luciferase activities were measured using a dual-luciferase reporter assay system (Promega Corporation) according to the manufacturer's instructions.

**Statistical analysis.** All experiments were repeated at least three times. The data are shown as the mean  $\pm$  standard deviation. Comparisons between two groups were performed using Student's t-test. In addition, the samples were divided into a high and a low expression group according to the median value of circRNA-0074027 expression and  $\chi^2$  test was performed to analyze the relationship between the circRNA-0074027 expression and clinicopathological characterization of patients with CRC. The data among three or more groups were performed using one-way ANOVA followed by Fisher's least significant difference post hoc test. The correlation between circRNA-0074027 and miR-525-3p were analyzed by Pearson's correlation. Statistical analyses were performed using SPSS 20.0 (IBM Corp.).  $P < 0.05$  was considered to indicate a statistically significant difference.

## Results

**CircRNA-0074027 expression is increased in CRC tissue samples.** Results comparing circRNA-0074027 expression levels in CRC tissues samples and samples obtained from adjacent normal tissues revealed that there was an overexpression of circRNA-0074027 in the CRC tissues (Fig. 1A). Subsequently, it was discovered that among 60 paired tissue samples, almost 73.3% (44/60) demonstrated higher circRNA-0074027 expression in CRC tissues (Fig. 1B). Next, these 60 patients with CRC were divided into two groups, including circRNA-0074027 high and low expression group, based on the median value of circRNA-0074027 expression. According to the results of statistical analysis, it was found that the circRNA-0074027 expression was closely related to the differential status ( $P=0.001$ ), N stage ( $P=0.004$ ), vascular invasion ( $P=0.016$ ), tumor size ( $P<0.001$ ) and TNM stage ( $P=0.02$ ; Table II). Similarly, the data revealed that patients with CRC and circRNA\_0047027 overexpression were more

Table II. Correlation between the clinicopathological data and circRNA-0074027 expression of in colorectal cancer (n=60).

Characteristics	n	circRNA-0074027 expression		P-value
		High	Low	
Sex				
Male	27	12	15	0.586
Female	33	17	16	
Age				
<60 years	25	12	13	0.879
$\geq 60$ years	35	18	17	
Differential status				
Moderate/well	20	13	7	<b>0.001</b>
Undifferentiated/poorly	40	9	31	
Vascular invasion				
Negative	22	14	8	<b>0.016</b>
Positive	38	12	26	
Tumor size				
$\leq 5$ cm	25	19	6	<b>&lt;0.001</b>
$> 5$ cm	35	8	27	
N stage				
N0	27	19	8	<b>0.004</b>
N1-N3	33	11	22	
TNM stage				
I-II	24	16	8	<b>0.02</b>
III-IV	36	13	23	
Tumor location				
Left colon cancer	20	8	12	0.625
Right colon cancer	22	10	12	
Rectal cancer	18	10	8	

Significant differences are presented in bold. circRNA, circular RNA.

likely to have low tumor differentiation (Fig. 1C), lymph node metastasis (Fig. 1D) and advanced TMN stage (Fig. 1E). In summary, circRNA-0074027 might have a role as a tumor inhibitor role in CRC progression.

**Knockdown of circRNA-0074027 suppresses CRC cell growth.** Since the expression of circRNA-0074027 was reduced in CRC tissues, its expression was further detected in normal epithelial cells and five different CRC cell lines. When compared with normal epithelial cell lines (FHC), the expression of circRNA-0074027 was higher in CRC cell lines, especially in Caco-2 and LoVo cells (Fig. 2A). Therefore, the Caco-2 and LoVo cells were selected in the present study for future *in vitro* experiments. The expression of circRNA-0074027 in Caco-2 and LoVo cells was knocked down using RNA transfection (Fig. 2B and C). The results of CCK-8 assay displayed that silencing of circRNA-0074027 could markedly inhibit the cell

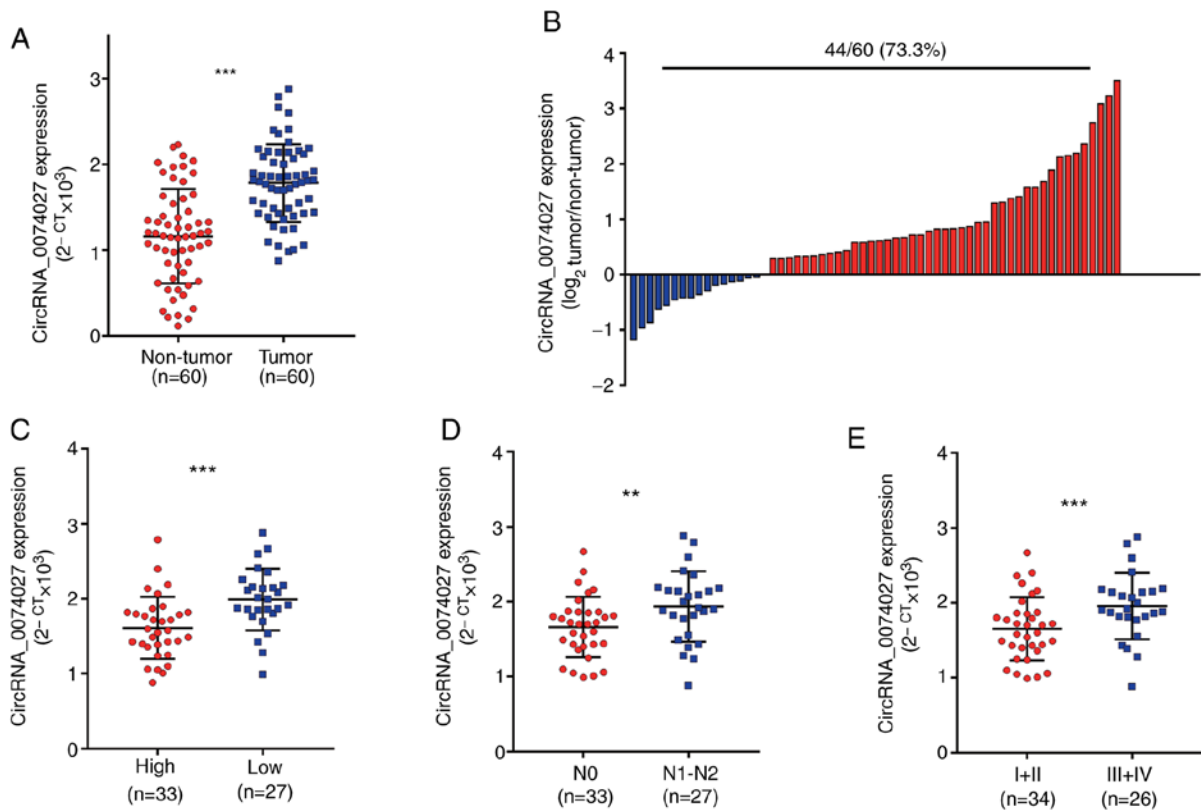


Figure 1. Expression of circRNA-0074027 is enhanced in CRC. (A) The expression of circRNA-0074027 in CRC tissues. (B) CircRNA-0074027 is upregulated (73.3%; 44/60) in CRC. (C) Relationship between circRNA-0074027 expression and tumor differentiation. (D) The relationship between circRNA-0074027 expression and lymph node metastasis. (E) The relationship between circRNA-0074027 expression and TMN stage. \*\* $P < 0.01$ , \*\*\* $P < 0.001$ . circRNA, circular RNA; CRC, colorectal cancer.

growth rate in both of HGC-27 and AGS cells (Fig. 2D and F). Consistent with the previous results, the EdU assay also indicated that there were fewer EdU-positive Caco-2 cells in siRNA-circRNA-0074027-1 and siRNA-circRNA-0074027-2 group than that in si-Control group (Fig. 2E). Additionally, EdU-positive LoVo cells in siRNA-circRNA-0074027-1 and siRNA-circRNA-0074027-2 group were fewer compared with the si-Control group (Fig. 2G). In total, silencing of circRNA-0074027 could suppress cell growth rate in CRC.

**Knockdown of circRNA-0074027 increases cell apoptosis rate via activating the p53 pathway.** As circRNA-0074027 upregulation could suppress cell proliferation ability in CRC, the changes of apoptosis level with low expression of circRNA-0074027 required further research. Therefore, in current study, the cell apoptosis rate was detected by flow cytometry experiments. As shown in Fig. 3A, the total cell apoptosis rate of Caco-2 cells in siRNA-circRNA-0074027-1 and siRNA-circRNA-0074027-2 group were higher compared with the si-Control group. Similarly, silencing of circRNA-0074027 could induce more apoptosis in LoVo cells compared with the si-Control group (Fig. 3B).

Western blotting was performed to identify the potential mechanism of the downregulation of circRNA-0074027 on cell apoptosis. The downregulation of circRNA-0074027 could significantly increase p53 and Bax proteins expression, while reducing the expression of anti-apoptosis protein Bcl-2 compared with the si-Control group (Fig. 3C and D). In all,

these data consistently displayed that the downregulation of circRNA-0074027 could significantly promote cell apoptosis via the regulation of p53 signaling pathway in CRC.

**Knockdown of circRNA-0074027 inhibits cell metastasis via interfering with the epithelial to mesenchymal transition (EMT) signaling pathway.** Statistical analysis of data revealed that circRNA-0074027 expression was associated with lymph node metastasis in CRC. Therefore, Transwell assay was performed to determine the involvement of circRNA-0074027 in CRC cell metastasis. As shown in Fig. 4A, the number of cells migrating and invading decreased markedly following downregulation of the expression of circRNA-0074027 in Caco-2 cells compared with si-Control group. Fig. 4B demonstrates that the silencing of circRNA-0074027 could significantly suppress cell migration and invasion ability in LoVo cells compared with the control group. The results of western blot analysis demonstrated that there was an increase of epithelial-like phenotype protein (E-cadherin) and a decrease of mesenchymal phenotype proteins (N-cadherin and vimentin) when circRNA-0074027 was downregulated (Fig. 4C and D). Taken together, these data suggested that knockdown of circRNA-0074027 suppresses CRC cell metastasis ability by inhibiting the EMT pathway.

**miR-525-3p is the target gene of circRNA-0074027.** Previous studies (18-20) have revealed that circRNAs may act as miRNA sponges in regulation of CRC progression. In the present

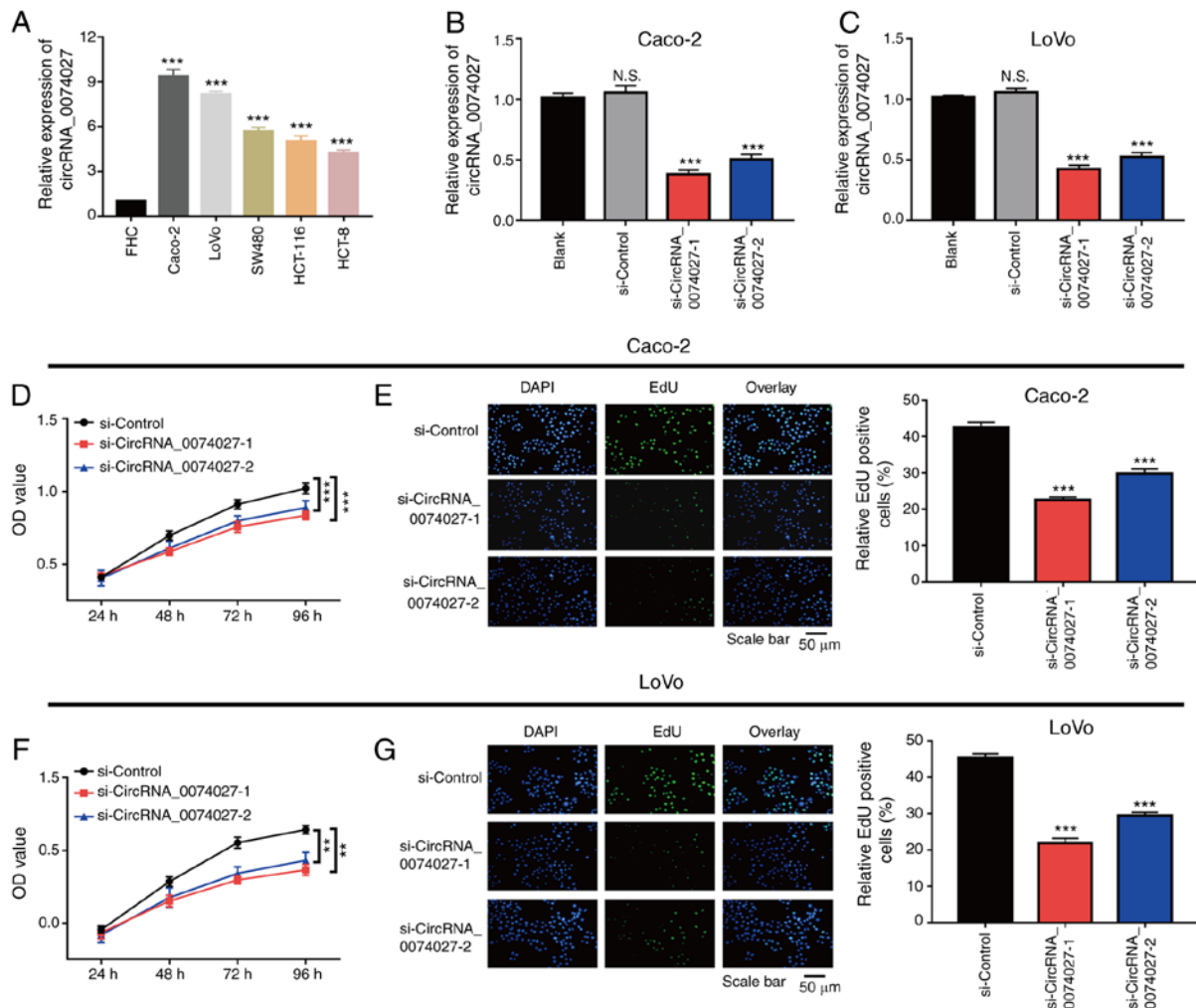


Figure 2. Downregulation of circRNA\_100395 suppresses CRC cell proliferation. (A) The expression of circRNA-0074027 in CRC cells. Successful construction of circRNA-0074027-downregulated (B) Caco-2 and (C) LoVo cells. The cell proliferation ability of (D) Caco-2 and (F) LoVo cells upon inhibiting circRNA-0074027 expression. 5-Ethynyl-2'-deoxyuridine assay of (E) Caco-2 and (G) LoVo cells upon inhibiting circRNA-0074027 expression. Every experiment was repeated three times. \*\* $P < 0.01$ , \*\*\* $P < 0.001$ . circRNA, circular RNA; CRC, colorectal cancer; N.S., no significance; si, short interfering.

study, miR-525-3p was proposed as the downstream gene of circRNA-0074027 based on the results of bioinformatics algorithms (Starbase.2). Subsequently, we discovered that the low-expression of miR-525-3p accounted for 80.0% (48/60) of the CRC tissue samples (Fig. 5A). The potential binding site between circRNA-0074027 and miR-525-3p was predicted as shown in Fig. 5B. Similarly, the results of RT-qPCR demonstrated that miR-525-3p was significantly downregulated in CRC tissue samples (Fig. 5C). Pearson correlation analysis revealed that the circRNA-0074027 expression was negatively correlated with the expression of miR-525-3p in CRC (Fig. 5D,  $r = -0.5483$ ,  $P < 0.0001$ ). To further explore their association, luciferase reporter assay was performed and the results indicated that miR-525-3p mimics could markedly decrease the luciferase activity of wild-type circ\_0074027, but with no effect on MUT circ\_0074027 (Fig. 5E). The results of RT-qPCR indicated that miR-525-3p inhibitor could significantly downregulate the expression of miR-525-3p in Caco-2 cells, when compared with the control group (Fig. 5F). In addition, RT-qPCR assay was performed to determine the expression of miR-525-3p in Caco-2 cells. Notably, both si-Circ\_0074027-1 and si-Circ\_0074027-2 groups demonstrated higher expression

of miR-525-3p in comparison with si-Control group, with the si-Circ\_0074027-1 group showing the highest expression (Fig. 5G). Thus si-circRNA-0074027-1 Caco-2 cells were suggested as ideal for future rescue experiments. To elucidate the potential regulatory role of miR-525-3p, the miR-525-3p inhibitor was used to decrease the intra-cellular miR-525-3p expression in Circ\_0074027 downregulated Caco-2 cells and this was verified with the RT-qPCR assay (Fig. 5H). These data consistently revealed that miR-525-3p would be direct target gene of circRNA-0074027 in CRC.

**Knockdown of miR-525-3p expression reverses the anti-tumor effects induced by the overexpression of circRNA-0074027 in CRC.** To explore the regulatory role of miR-525-3p in the regulation of CRC progression, miR-525-3p inhibitors were added to inhibit the expression of miR-525-3p in circRNA-0074027-downregulated Caco-2 cells. As the results of CCK-8 and EdU assay demonstrated, the effects of circRNA-0074027 downregulation on suppression of cell proliferation could be rescued by adding miR-525-3p inhibitors (Fig. 6A and B). Addition of miR-525-3p inhibitors may also reduce the cell apoptosis rate in the si-circRNA-0074027-1



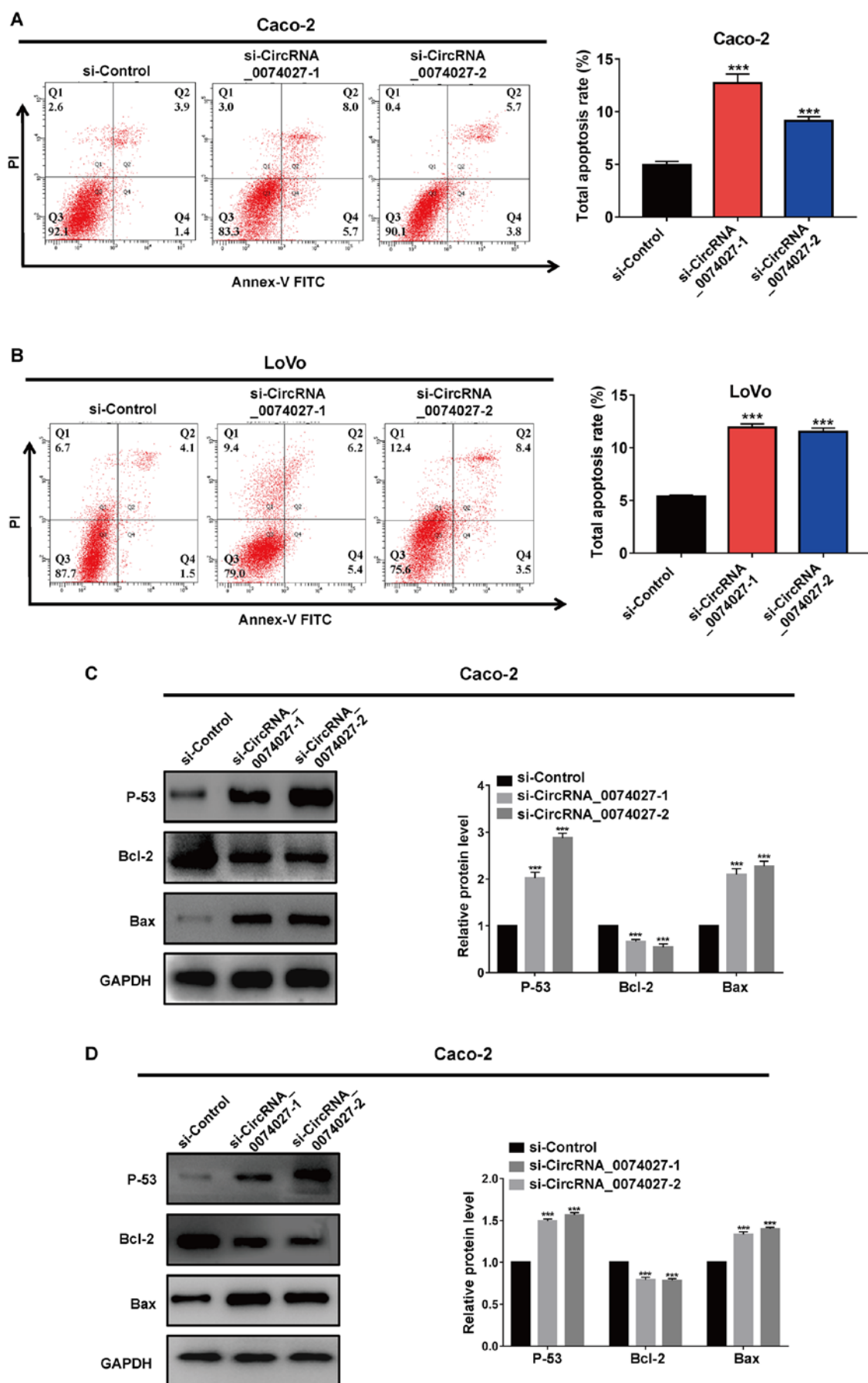


Figure 3. Downregulation of circRNA-0074027 promotes cell apoptosis in CRC. The cell apoptosis images (left) and statistical analysis (right) of (A) Caco-2 and (B) LoVo cells upon circRNA-0074027 downregulation. The total apoptosis rate includes cells in right quadrants (Q2 and Q4). Western blotting images of p53, Bcl-2 and Bax protein expression in (C) Caco-2 and (D) LoVo cells with circRNA-0074027 downregulation. Every experiment was repeated three times. \*\*\* $P < 0.001$ . circRNA, circular RNA; CRC, colorectal cancer; si, short interfering.

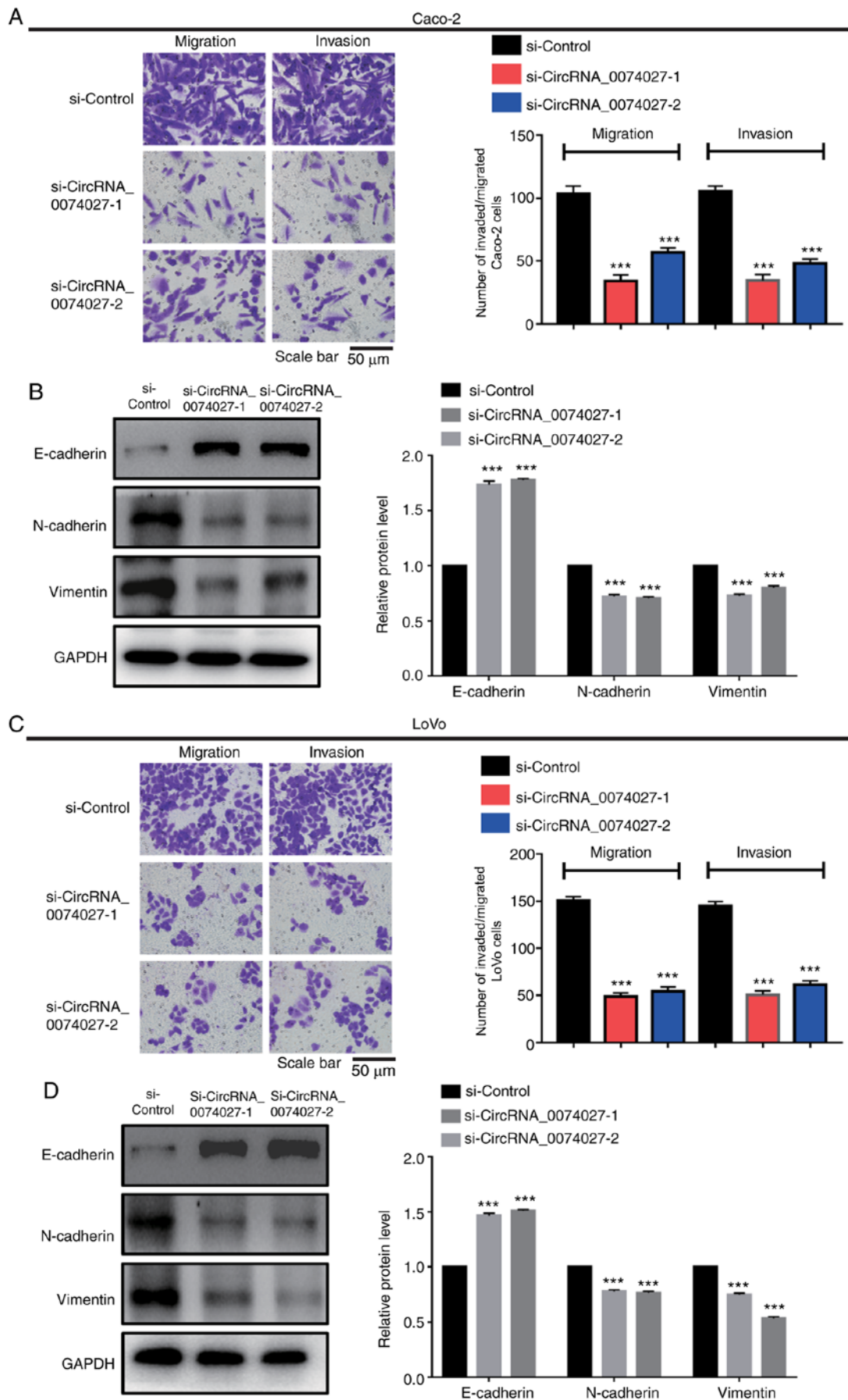


Figure 4. Downregulation of circRNA-0074027 inhibits cell metastasis in CRC. The cell migration and invasion images (left) and statistics analysis (right) of (A) Caco-2 and (B) LoVo cells upon downregulation of circRNA-0074027. Western blotting images of EMT signaling pathway related proteins expression in (C) Caco-2 and (D) LoVo cells when circRNA-0074027 is downregulated. Every experiment was repeated three times. \*\*\* $P$ <0.001. circRNA, circular RNA; CRC, colorectal cancer; si, short interfering.

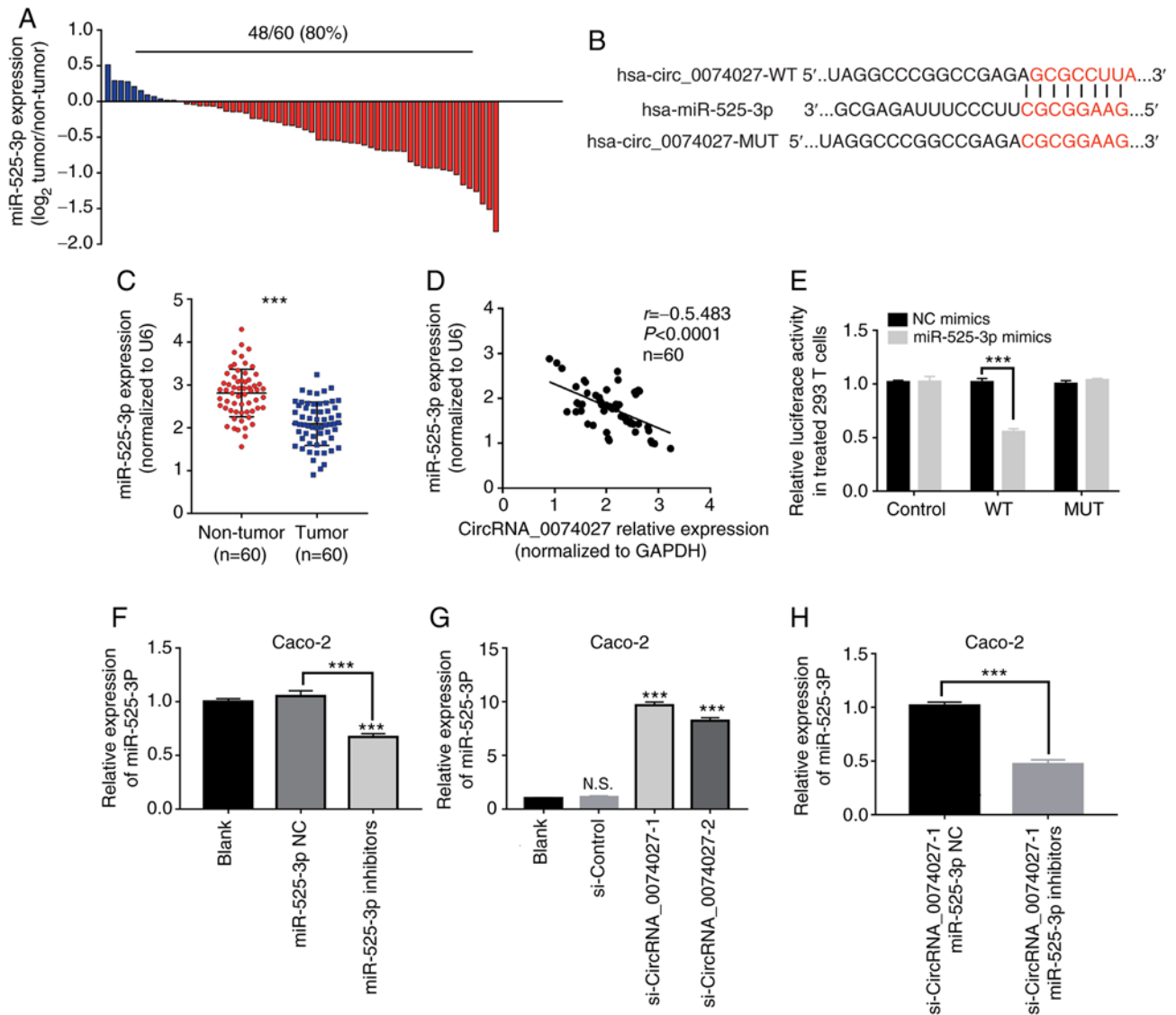


Figure 5. miR-525-3p is a downstream target gene of circRNA-0074027. (A) miR-525-3p is downregulated in CRC tissues. (B) The predicted 3'UTR binding regions of circRNA-0074027 on miR-525-3p. (C) The expression of miR-525-3p in CRC tissues. (D) The Pearson correlation analysis between the miR-525-3p and circRNA-0074027 expression. (E) Relative luciferase activity in 293T cells following co-transfection. (F) The expression level of miR-525-3p in Caco-2 cells following treatment with miR-525-3p inhibitors, as detected by RT-qPCR. (G) The expression of miR-525-3p in circRNA\_007402-downregulated Caco-2 cells. (H) The expression of miR-525-3p in circRNA\_007402-downregulated Caco-2 cells following treatment with miR-525-3p inhibitors. \*\*\* $P < 0.001$ . miR, microRNA; circRNA, circular RNA; CRC, colorectal cancer; RT-qPCR, reverse transcription-quantitative PCR; N.S., no significance; si, short interfering; WT, wild-type; MUT, mutant; NC, negative control.

Caco-2 cells (Fig. 6C). Similarly, the western blotting results also revealed that when incubating with miR-525-3p inhibitors could lead to increased expression of p53 and Bax proteins but decrease Bcl-2 protein expression in si-circRNA-00074027-1 Caco-2 cells (Fig. 6D).

An increase in migrated and invasive Caco-2 cells were observed in si-circRNA-0074027-1 groups following the addition of miR-525-3p inhibitors (Fig. 6E). Furthermore, the EMT signaling pathway-related markers in CRC cells were also determined following adding miR-525-3p inhibitor. The addition of miR-525-3p could lead to decreased expression of E-cadherin proteins and an increased expression of N-cadherin and Vimentin proteins in circRNA-0074027-low expression CRC cells (Fig. 6F). Therefore, these data revealed that down-regulation of circRNA-0074027 could exert anti-tumor effects through sponging miR-525-3p in CRC.

## Discussion

CRC is one of the most prevalent malignancies in the world (1,2). The main strategies for CRC treatment include surgery, radiotherapy and chemotherapy, which have shown positive therapeutic effects in early-stage patients. However, due to the shortage of effective biomarkers, most patients are diagnosed with advanced CRC and thus have short survival periods (1). Therefore, the present study was designed to identify effective diagnosis and therapeutic bio-markers and determine their potential mechanism.

It is known that circRNAs are evolutionally conserved and more stable than other non-coding RNAs, suggesting that circRNAs might be the most suitable bio-markers for human diseases (21-23). Several studies (15,16,24) indicate that circRNA-0074027 is an important regulatory factor in the



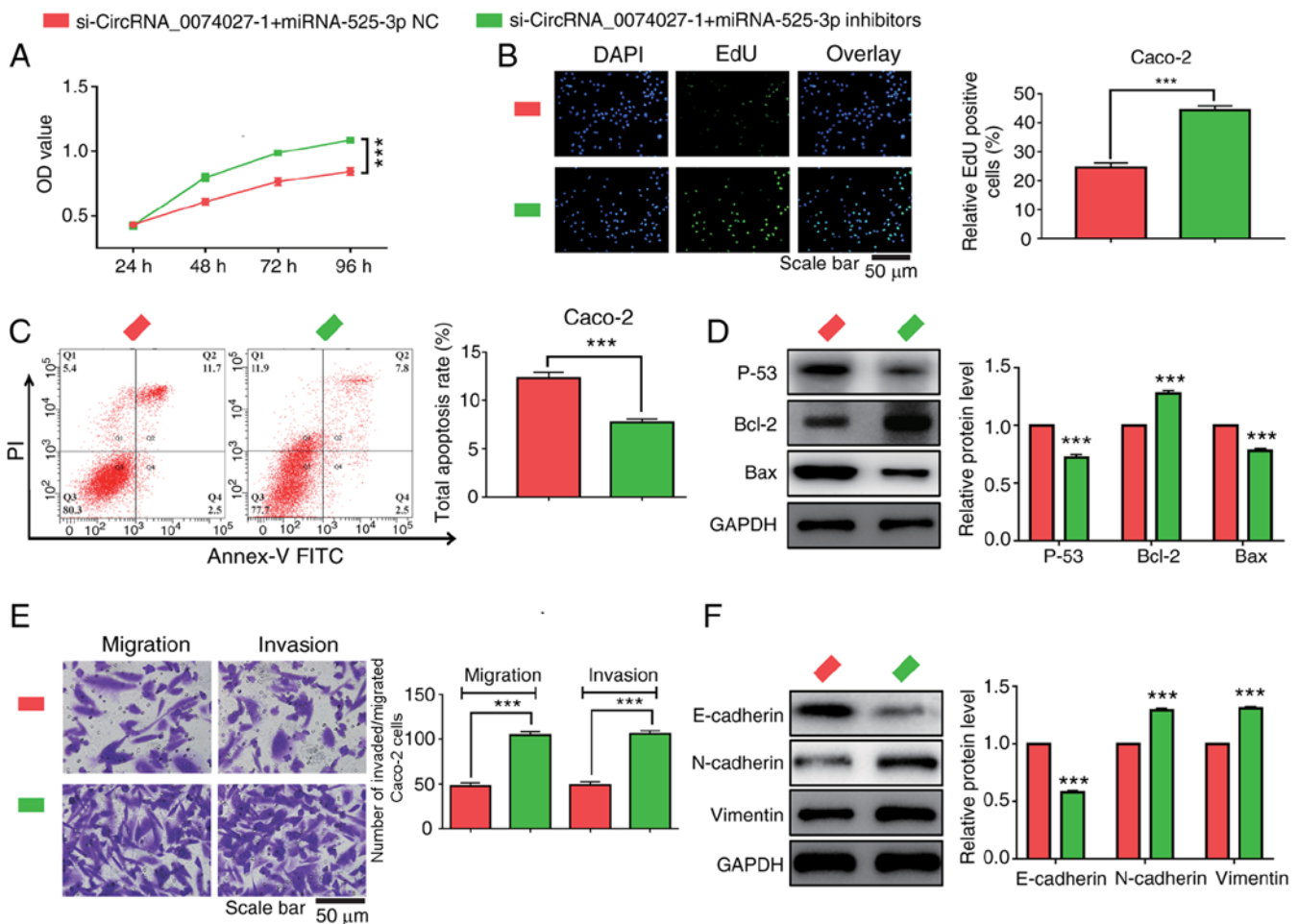


Figure 6. Downregulation of miR-525-3p rescues anti-tumor effects induced by depletion of circRNA-0074027. (A) Cell proliferation analysis of circRNA-0074027-downregulated Caco-2 cells. (B) 5-Ethynyl-2'-deoxyuridine assay of circRNA-0074027-downregulated Caco-2 cells. (C) Cell apoptosis rate of circRNA-0074027-downregulated Caco-2 cells. (D) Expression of p53, Bax and Bcl-2 proteins in circRNA-0074027-downregulated Caco-2 cells following addition of miR-135a-5p. (E) Cell migration and invasion of circRNA-0074027-downregulated Caco-2 cells. (F) Expression of epithelial to mesenchymal transition signaling pathway-related proteins in circRNA-0074027-downregulated Caco-2 cells following addition of miR-135a-5p. Every experiment was repeated three times. \*\*\* $P < 0.001$ . miR, microRNA; circRNA, circular RNA; OD, optical density.

progression of various types of cancer, including glioblastoma and non-small-cell lung cancer. However, no study, to the best of the authors' knowledge, has been performed on the regulatory role of circRNA-0074027 in CRC progression. In the current study, RT-qPCR was performed to detect the expression of circRNA-0074027 in CRC tissue and adjacent normal tissue samples. The results indicated that circRNA-0074027 expression was upregulated in CRC tissues compared with the normal tissues. According to the results of statistical analysis, patients with CRC and circRNA-0074027 overexpression were more likely to have poor differential status, larger tumor size, advanced TNM stage, advanced N stage and vascular invasion. These results demonstrated that circRNA-0074027 might function as a tumor promoter in the development of CRC.

To clarify the biological role of circRNA-0074027 in CRC progression, circRNA-0074027-downregulated CRC cells were constructed. Results revealed that the deletion of circRNA-0074027 could markedly inhibit cell growth and promote cell apoptosis. However, it remains unclear how circRNA-0074027 participated in the regulation of cell proliferation and apoptosis in CRC. Previous studies report that p53 is a pivotal anti-oncogene in cancer progression and its dysfunction

might contribute to the rapid cell proliferation and decreased DNA repair capacity (25,26). circRNA ZNF609 is shown to promote cell apoptosis rate via upregulating p53 expression in CRC (27). Su *et al* (28) showed that circRNA\_0055538 can exert an anti-tumor effect by activating the p53/caspase signaling pathway. The present study also demonstrated that knockdown of circRNA-0074027 increased p53 and Bax proteins expression and decreased Bcl-2 protein expression. Therefore, the silencing of circRNA-0074027 could activate the p53-mediated pathway, which contributes to the suppression of cell proliferation and promotion of cell apoptosis in CRC.

The present study discovered that the patients with CRC and circRNA-0074027 overexpression had advanced N stage, thus it was hypothesized that circRNA-0074027 participated in the regulation of cell migration and invasion ability in CRC. The data of the present study demonstrated that the deletion of circRNA-0074027 could markedly inhibit the cell metastasis ability in CRC, but the potential regulatory mechanism remains to be elucidated. EMT is a significant biological process, in which the epithelial (E) cells transition to a mesenchymal (M) phenotype (29-31). Previous studies indicate that EMT signaling pathway serves an important role in the

development of malignant tumors (32-34). Wang *et al* (35) revealed that the circRNA circP4HB can promote cell aggressiveness and metastasis of non-small cell lung carcinoma via sponging miR-133a-5p. In addition, circRNA circPTPRA can sponge miR-96-5p, thereby inhibiting cell metastasis ability in non-small cell lung carcinomas cells via regulating the EMT pathway (36). The results of the present study were consistent with these previous studies and revealed that the downregulation of circRNA-0074027 led to increased expression of epithelial-like phenotype protein (E-cadherin) and decrease expression of mesenchymal phenotype proteins (N-cadherin and vimentin). Taken together, the present study hypothesized that downregulation of circRNA-0074027 mediated inhibition of cell migration and invasion in CRC and this might be caused by the inactivation of EMT signaling pathway.

The primary function of circRNAs is miRNA sponging, which contributes to the carcinogenesis and progression of various types of cancer (8,37). For example, circRNA-MAN2B2 had been identified as a tumor promotor in hepatocellular carcinoma progression via sponging miRNA-217 (38). Another study indicates that circFOXK2 can sponge miR-942, followed by enhanced cell proliferation, migration and invasion ability in pancreatic ductal adenocarcinoma (39). The present study proposed miR-525-3p as the target gene of circRNA-0074027 in CRC progression. The results of RT-qPCR revealed that miR-525-3p was downregulated in CRC tissues in comparison with normal tissue samples. The Pearson correlation analysis showed that circRNA-0074027 expression was negatively correlated with the expression of miR-525-3p, suggesting that miR-525-3p might be the downstream target gene of circRNA-0074027. The association between the circRNA-0074027 and miR-525-3p was confirmed by dual-luciferase reporter assays. Previous studies identify that miR-525-3p may function as a significant regulatory factor in the development of various types of cancer, including liver cancer (40) and Hodgkin lymphoma (41). Pang *et al* (40) determined that miR-525-3p can enhance the cell migration and invasion of liver cancer via modulating ZNF395 expression. The results of the current study demonstrated that the addition of miR-525-3p could reverse the anti-tumor effects induced by the silencing of circRNA-0074027. Therefore, these data suggested that circRNA-0074027 could sponge miR-525-3p, thereby participating in the regulation of cell proliferation, cell metastasis and cell apoptosis in CRC. However, the present study has some limitations. First, the molecular targets downstream of the circRNA-0074027/miR-525-3p axis were not fully elucidated. Second, the present study was performed *in vitro* and there is a need to carry it out *in vivo* to confirm the biological function of circRNA-0074027. Third, the sample size and the follow-up duration was insufficient. In the future, a large sample size and a longer follow-up duration should be employed to validate the diagnostic and prognostic significance of circRNA-0074027.

CircRNA-0074027 might function as a tumor promoter in CRC, as the patients with CRC and overexpression were more likely to have poor prognosis. In addition, the *in vitro* experiments indicated that silencing of circRNA-0074027 could directly regulate the function of miR-525-3p and then lead to the suppression of cell proliferation and metastasis ability via interfering with the p53/EMT signaling pathway.

In conclusion, circRNA-0074027 could act as a promising therapeutic bio-marker for CRC.

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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

GX and YZ conceived and designed the study. GX, JZ and YZ performed the experiments. GX, XP and BW were responsible for the analysis and interpretation of data. XP, BW and YZ wrote the manuscript. GX and YZ were responsible for confirming the authenticity of the data. All authors read and approved the final manuscript. This manuscript was revised by all authors.

## Ethics approval and consent to participate

Written informed consent was sought from the participants before the samples were obtained. The present study was approved by the Ethical Committees of Dazhou Central Hospital (approval no. KY2020-086-01).

## Patient consent for publication

Not applicable.

## Competing interests

The authors declare that they have no competing interests.

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