

# miR-141 Promotes Colon Cancer Cell Proliferation by Targeted *PHLPP2* Expression Inhibition

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**Objective:** Colon cancer (CC) is the third most common cancer with a high rate of incidence and mortality. Therefore, it is highly necessary to explore novel targets of CC.

**Methods:** The miRNA-seq and RNA-seq data of CC were accessed from the TCGA database. Differential analysis was performed using the “edgeR” package to identify differentially expressed miRNAs (DE\_miRNAs). The downstream target genes of the target miRNA were then predicted by miRNA target prediction databases to identify the target mRNA. Normal colon cell line CCD-18Co and CC cell lines HCT-116, HT-29, SW620 and SW480 were chosen, and qRT-PCR was conducted to detect *miR-141* expression in these cell lines. qRT-PCR and Western blot were carried out to determine *PHLPP2* mRNA and protein expression, respectively. Dual-luciferase reporter gene assay was performed to verify the targeting relationship between *miR-141* and *PHLPP2* 3'UTR. CCK-8 assay and colony formation assay were carried out to detect cell proliferation. Meanwhile, tumor xenograft model in nude mice was constructed to assess CC cell tumorigenic ability in vivo.

**Results:** *miR-141* was markedly up-regulated in CC tissue. CC cell proliferation and in vivo tumorigenic ability were suppressed by *miR-141* silencing but promoted by *miR-141* over-expression. *PHLPP2* was significantly down-regulated in cancer tissue. Dual-luciferase reporter gene assay indicated that *miR-141* could bind to *PHLPP2* 3'UTR. *PHLPP2* expression was noticeably elevated upon *miR-141* deficiency but significantly inhibited upon *miR-141* over-expression. CCK-8 and colony formation assay suggested that *miR-141* facilitated CC cell proliferation by silencing *PHLPP2*.

**Conclusion:** *miR-141* promotes CC cell proliferation by targeted silencing *PHLPP2*.

**Keywords:** *miR-141*, *PHLPP2*, colon cancer, proliferation

## Introduction

Colon cancer (CC) is one of the major causes leading to tumor-related deaths and its incidence rate is rising in many countries.<sup>1</sup> According to Surveillance, Epidemiology, and End Results Program of the USA, it was estimated that CC made up 8.3% of the total cancer cases and 8.4% of the total cancer-related deaths in 2019 (Available at: <https://seer.cancer.gov/statfacts/html/colorect.html>; cited 5 July 2019). Although the overall rate of incidence and mortality has been reduced by the progress made in screening and treatment methods, CC remains a burden and it is predicted that CC patients will increase by 60% worldwide by 2030.<sup>2</sup> As a result, it is highly necessary to find novel molecular markers.

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MicroRNAs (miRNAs) are evolutionarily conservative endogenous non-coding RNA molecules about 22 nucleotides in length<sup>3</sup> that regulate gene expression by binding to mRNAs at the 3'-untranslated region (3'UTR).<sup>4</sup> Research suggests that miRNAs play a crucial role in various biological processes, such as cell proliferation,<sup>5</sup> differentiation,<sup>6</sup> invasion,<sup>7</sup> apoptosis<sup>8</sup> and metabolism.<sup>4</sup> Besides, another evidence demonstrates that dysregulation of miRNAs is related to the occurrence and development of various human cancers.<sup>9</sup> For example, *miR-101-5p* has ectopic expression in breast cancer and can weaken the aggressive phenotypes of BrCa cells, such as proliferation, migration and invasion.<sup>10</sup> While *miR-302a* is poorly expressed in colorectal cancer (CRC) and suppresses CRC metastasis and cetuximab resistance.<sup>11</sup> *miR-141* is a member of *miR-200* family reported to be differentially expressed in multiple human malignant tumors.<sup>12</sup> For instance, *miR-141* is up-regulated in ovarian cancer<sup>13</sup> and down-regulated in hepatocellular carcinoma (HCC) and prostate cancer,<sup>14,15</sup> but the specific regulatory mechanism of *miR-141* in various cancers remains unclear.

*PHLPP2* (PH domain leucine-rich-repeats protein phosphatase 2) is able to regulate the activity of AGC kinase including Akt4. It is indispensable to keep *PHLPP2* expression in balance for pathological prevention as the change of *PHLPP2* steady-state level is connected with various diseases, including diabetes, hepatic steatosis and cancers.<sup>16,17</sup> Recently, many studies have unveiled that *PHLPP2* is ubiquitously lost in various cancers and plays an important role in multiple biological processes, such as cancer cell proliferation, metastasis, autophagy and apoptosis.<sup>18-23</sup> Nonetheless, the diagnostic and prognostic significance of *PHLPP2* in CC have not been fully researched.

In this study, we investigated *miR-141* expression in CC cells and validated its effect on CC cell proliferation and tumorigenic ability. Additionally, we also studied the regulatory mechanism of *miR-141* and *PHLPP2* in CC cells. Our study will shed light on the mechanism of CC tumorigenesis and verify that *miR-141* is a potential therapeutic target for CC treatment.

## Materials and Methods

### Bioinformatics Analysis

The miRNA-seq and RNA-seq data of CC were accessed from the TCGA-COAD dataset (<https://portal.gdc.cancer.gov/>) and subjected to differential analysis using the "edgeR" package.  $|\log_{2}FC| > 2$  and  $p$  value  $< 0.05$  were set as

the threshold to screen out differentially expressed genes (DEGs). TargetScan ([http://www.targetscan.org/vert\\_72/](http://www.targetscan.org/vert_72/)), miRDB (<http://mirdb.org/>) and miRTarBase (<http://mirtarbase.mbc.nctu.edu.tw/php/index.php>) databases were employed for target prediction to validate the candidate mRNAs, which were then intersected with the differentially expressed mRNAs (DE\_mRNAs) in the TCGA-COAD dataset. Subsequently, the mRNA which showed a marked effect on patient's prognosis was chosen.

### Cell Culture

Normal colon cell line CCD-18Co (ATCC<sup>®</sup> CRL-1459<sup>TM</sup>) and CC cell lines HCT-116 (ATCC<sup>®</sup> CCL-247), HT-29 (ATCC<sup>®</sup> HTB-38), SW620 (ATCC<sup>®</sup> CCL-227<sup>TM</sup>) and SW480 (ATCC<sup>®</sup> CCL-228<sup>TM</sup>) were all purchased from the American Type Culture Collection (ATCC; Rockville, MD, USA). All the cells were kept in a humidified atmosphere of 5% CO<sub>2</sub> at 37°C, among which CCD-18Co cell line was cultured in Eagle's Minimum Essential Medium (EMEM; Catalog No. 30-2003) containing 10% fetal bovine serum (FBS; Gibco, Grand Island, NY, USA), HCT-116 and HT-29 cell lines were grown in McCoy's 5a Medium Modified (Catalog No. 30-2007) containing 10% FBS, while SW620 and SW480 cell lines were maintained in Leibovitz's L-15 Medium (Catalog No. 30-2008) containing 10% FBS. The mediums were replaced every two or three days depending on cell growth until cells were cultured to be 80-90% confluent. Thereafter, cells were used for subculture and then harvested when reaching the logarithmic phase.

### Cell Transfection

Inhibitor NC, miR-141 inhibitor, mimic NC, miR-141 mimic, oe-NC and oe-PHLPP2 ordered from GenePharma (Shanghai, China) were transfected into cells ( $1 \times 10^6$  cells per well in a 6-well plate) by Lipofectamine 2000 (Invitrogen; Thermo Fisher Scientific, Inc.) following the manufacturer's protocol. After transfection, cells were cultured for additional 6 h at 37°C and 5% CO<sub>2</sub>, after which the mediums were changed. All the cells were collected after 48 h of transfection.

### Real-Time Quantitative PCR (qRT-PCR)

Total RNA was extracted from CC tissue and cells using Trizol reagent (16,096,020; Thermo Fisher Scientific, NY, USA) according to the manufacturer's instructions. Then, 5 µg RNA was transcribed into complementary DNA (cDNA) by using the cDNA Reverse Transcription Kit

(K1622; Fermentas Inc., Ontario, CA, USA). qRT-PCR was performed with the TaqMan MicroRNA Assay and TaqMan® Universal PCR Master Mix under the following thermocycler conditions: 95°C for 2 min, and 45 cycles of 95°C for 15 s and 60°C for 45 s, followed by 72°C for 45 s. Each measurement was normalized to U6 to ensure comparable amounts of miRNA in all wells.

According to the TaqMan Gene Expression Assays protocol (Applied Biosystems, Foster City, CA, USA), qRT-PCR was conducted to assess mRNA under the following thermocycler conditions: 95°C for 10 min, and 35 cycles of 95°C for 15 s and 60°C for 30 s, followed by 72°C for 45 s. *GAPDH* was used as an internal reference. Three repeated wells were set for each treatment. The primers were ordered from TAKARA (Beijing, China) and listed in Table 1. The quantitative value was expressed using the  $2^{-\Delta\Delta Ct}$  method:  $\Delta\Delta Ct = \Delta Ct$  (experiment group) -  $\Delta Ct$  (control group), where  $\Delta Ct = Ct$  (target gene) -  $Ct$  (internal reference).<sup>24</sup>

## Western Blot

Total proteins were extracted from cells after transfection and the concentration of proteins was measured by bicinchoninic acid (BCA) protein assay kit (Thermo Fisher Scientific, USA). Protein samples were separated on 10% sodium dodecyl sulphate-polyacrylamide gel electrophoresis (SDS-PAGE) and transferred onto nitrocellulose membranes (ZY-160FP, SHZYSW, Shanghai, China). After being blocked with 5% non-fat milk at room temperature for 1 h, the membranes were oscillated in 3% BSA in an oscillator for 1 h. The membranes were washed with Tris-buffered saline with Tween (TBST; Solarbio, Beijing, China) in triplicate with 10 min each time and then incubated with diluted primary rabbit polyclonal antibodies *PHLPP2* (ab71973, 1:2000, abcam, UK) and *GAPDH* (ab9485, 1:2500, abcam, UK) overnight at 4°C. The membranes were washed with TBST in triplicate with 10 min each time. Subsequently, goat anti-rabbit immunoglobulin G (IgG) H&L (Alexa Fluor® 790) secondary antibody (ab205718, 1:5000, abcam, UK) was added onto the membranes and incubated for 1 h. The membranes were washed

with TBST in triplicate with 10 min each time, followed by development with electrochemiluminescence (ECL) reagent (ECL808-25, Biomiga, USA) for 1 min at room temperature. Images of the protein bands were observed and captured under an optical luminometer (GE, USA). Subsequently, analysis of relative protein expression was performed with ImageProPlus6.0 (MediaCybernetics, USA) gray scale scanning software. *GAPDH* was used as an internal reference. The experiment was repeated three times.

## Cell Counting Kit-8 (CCK-8) Assay

HCT-116 and SW480 cells were digested and resuspended after transfection and were then seeded in 96-well plates at  $1 \times 10^5$  cells/mL (100  $\mu$ L/well) and cultured overnight. Subsequently, at 0 h, 24 h, 48 h, 72 h and 96 h, each well was added with 10  $\mu$ L CCK-8 solution (Beyotime, Shanghai, China) and cells were incubated for additional 4 h. At last, the absorbance of each well at 450 nm was detected by using a microplate reader.

## Colony Formation Assay in vitro

Soft agar plate was made in 6-well plates, with the bottom layer being 0.5% agar mixed with serum-free medium. Firstly, cells were seeded in 35-mm tissue cultivation dishes for 24 h and then transfected with different reagents. After cells were digested with trypsin,  $1 \times 10^3$  cells and 0.35% agar were mixed in the cell medium containing 10% FBS and seeded into the agar plate as the top agar layer. Subsequently, cells were cultured in an incubator at 37°C for 3 weeks. The qualified colonies were stained with 0.1% crystal violet, washed extensively with phosphate-buffered saline (PBS) and counted.

## Dual-Luciferase Reporter Gene Assay

The wild-type (WT) *PHLPP2* 3'UTR that could bind with miR-141 and the mutant (MUT) *PHLPP2* 3'UTR formed by site-directed mutagenesis method were, respectively, cloned into psiCheck2 luciferase vector, contributing to the establishment of *PHLPP2*-WT and *PHLPP2*-MUT vectors. *PHLPP2*-WT/*PHLPP2*-MUT vectors were co-transfected with mimic

**Table 1** Primers for qRT-PCR

Genes	Forward (5'→3')	Reverse (5'→3')
<i>miR-141</i>	GTCCATCTCCATCAGTACAGGTTG	AGCCATCTTACTCACAGAGTGTG
<i>U6</i>	GCTTGCTTCACACACACATA	AAAAAACATCGACTCADG
<i>PHLPP2</i>	AGGTTCCCTGAGCATCTCTTC	GTTCAGGCCCTTCAGTTGAG
<i>GAPDH</i>	TGCACCACCAACTGCTTAGC	GGCATGCACTGTGGTCATGAG

NC/*miR-141* mimic into HCT-116 and SW480 cells. The luciferase activities were assessed using the Dual-Luciferase Reporter Assay Kit (Promega, Madison, WI, USA). After 48 h of incubation, the cells were lysed using 1×Passive lysis buffer and Dual-Luciferase Reporter Assay Kit (Promega) was used to evaluate the firefly luciferase activities. Firefly luciferase activity was normalized to Renilla luciferase activity. The experiment was performed in triplicate.

## Tumor Formation in Nude Mice

Twelve male BALB/C nude mice (4–6 weeks old, weighing 18–24 g) derived from Hunan SJA Laboratory Animal (Hunan, China) were fed in a specific pathogen-free (SPF) environment. The mice were randomly divided into the antagomir NC group and the *miR-141* antagomir group with 6 mice in each group. Lipofectamine 2000 reagent (Invitrogen) was employed to transfect antagomir NC and *miR-141* antagomir into CC cell line SW480. The stably transfected SW480 cells were obtained to prepare cell suspension at  $1 \times 10^6$  cells/mL, which was subcutaneously injected into nude mice. Tumor volume was monitored once a week. The long diameter (A) and short diameter (B) of the xenograft tumor were measured by a vernier caliper each week. Tumor volume was calculated with a formula of:  $\text{Volume} = (A \times B^2) / 2$ . At the end of the 5th week, mice were given a euthanasia with tumors being isolated and then fixed in 4% paraformaldehyde. Tumor tissues were frozen and stored in liquid nitrogen for further study 4 weeks later. All the animals being experimented were used for

medical research and this experiment was approved by the Ethics Committee of Laboratory Animal.

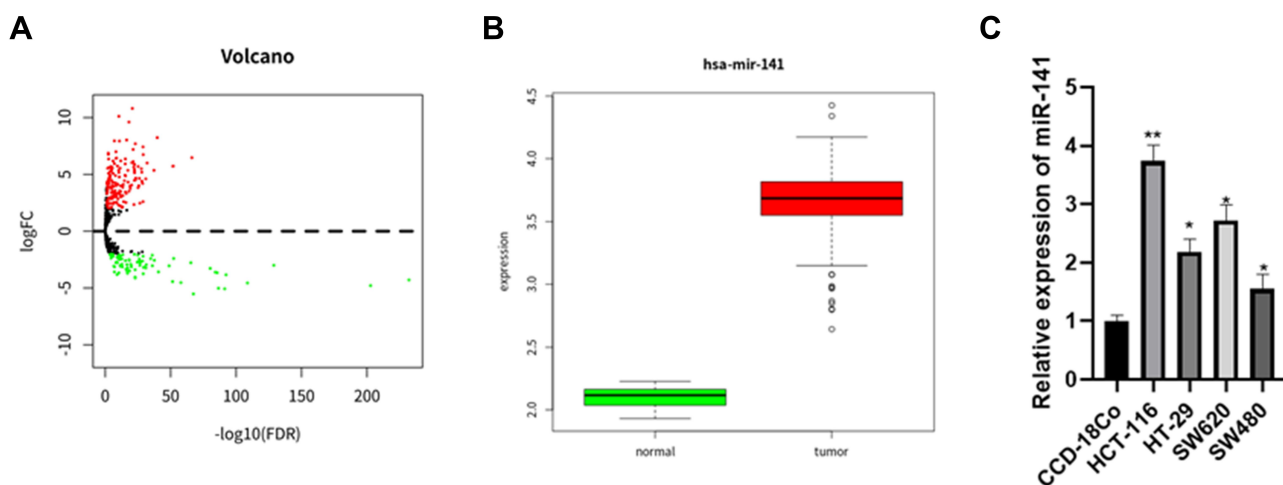
## Statistical Analysis

All statistical analyses were performed using SPSS 21.0 (SPSS, Inc., Chicago, IL, USA) software. All the data were expressed as mean  $\pm$  standard deviation (SD), and differences between two groups were analyzed by Student's *t*-test.  $p < 0.05$  was considered statistically significant and  $p < 0.01$  was considered highly statistically significant.

## Results

### *miR-141* is Highly Expressed in CC Cells

Normal samples ( $n=8$ ) and CC tissue samples ( $n=445$ ) were used for analysis of differentially expressed miRNAs (DE\_miRNAs), and 271 DE\_miRNAs in total were obtained (Figure 1A), among which *miR-141* was markedly up-regulated in cancer tissue (Figure 1B). A study validated that *miR-141* is able to serve as an oncogene,<sup>25</sup> and it is abnormally expressed in CC and involved in various cell processes, including epithelial-mesenchymal transition (EMT), proliferation, migration, invasion and drug resistance.<sup>26</sup> We chose normal colon cell line CCD-18Co and CC cell lines HCT-116, HT-29, SW620 and SW480, in which *miR-141* expression was detected via qRT-PCR. The results suggested that compared with CCD-18Co, *miR-141* was highly expressed in CC cells (Figure 1C). HCT-116 and SW480 cell lines with



**Figure 1** *miR-141* is highly expressed in CC cells. (A) Volcano plot of DE\_miRNAs in the TCGA-COAD dataset. Red and green represent markedly up-regulated and down-regulated DE\_miRNAs, respectively; (B) Relative expression of *miR-141* in normal and tumor samples in the TCGA-COAD dataset; (C) *miR-141* expression in normal colon cell line and CC cell lines were detected using qRT-PCR; \*  $p < 0.05$ , \*\*  $p < 0.01$ . The experiment was repeated in triplicate.

the highest and the lowest miR-141 expression were chosen for follow-up experiments.

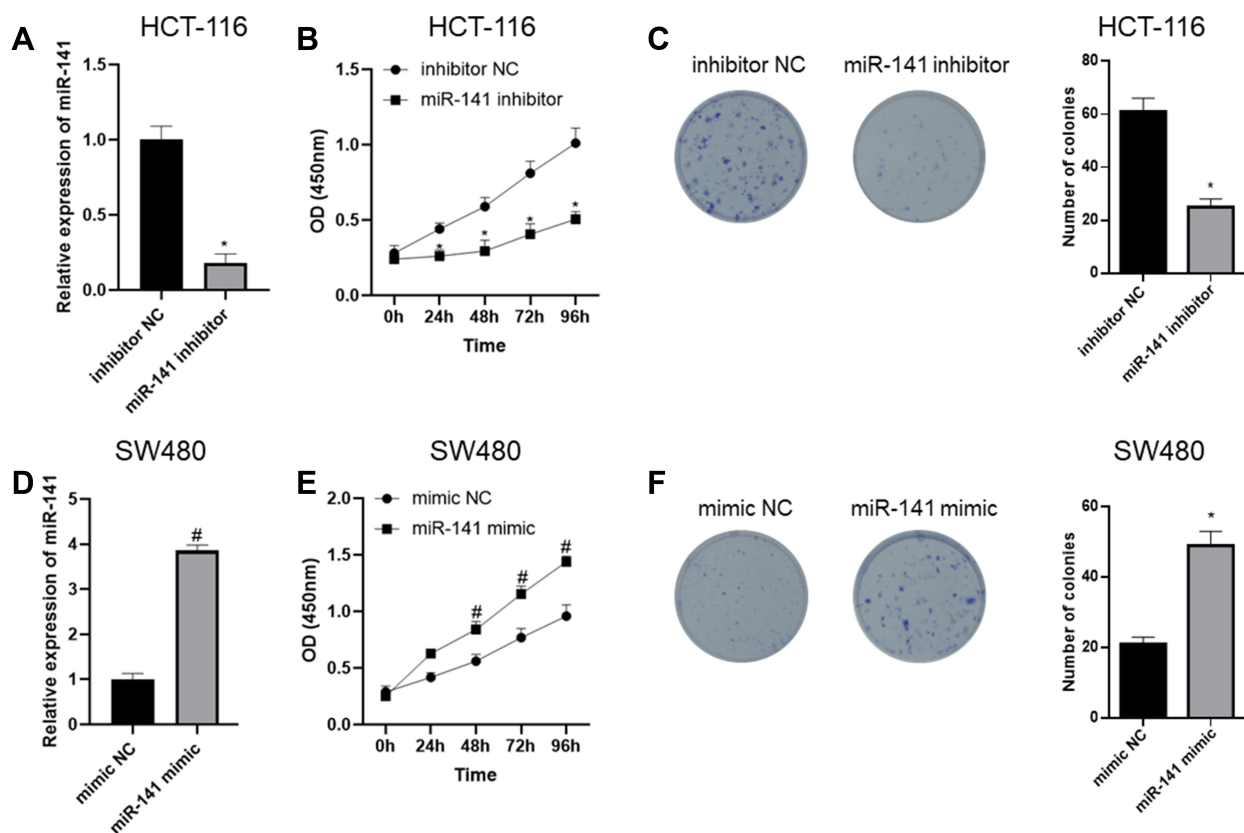
## CC Cell Proliferation is Suppressed by miR-141 Silencing but Promoted by miR-141 Over-Expression

In order to further verify the mechanism by which miR-141 regulates CC cell proliferation, we firstly silenced *miR-141* in HCT-116 cell line and detected *miR-141* expression via qRT-PCR. The results showed that compared with the inhibitor NC group, miR-141 expression in the *miR-141* inhibitor group was significantly reduced (Figure 2A). Cell proliferation was assessed by CCK-8 assay and colony formation assay, the results of which indicated that compared with the inhibitor NC group, cell proliferation in the *miR-141* inhibitor group was noticeably decreased (Figure 2B and C). Meanwhile, we over-expressed *miR-141* in SW480 cell line and determined the expression of *miR-141* through qRT-PCR and found that

compared with the mimic NC group, miR-141 expression in the *miR-141* mimic group was markedly increased (Figure 2D). We further detected cell proliferation by CCK-8 and colony formation assay, discovering that compared with the mimic NC group, cell proliferation in the *miR-141* mimic group was prominently raised (Figure 2E and F). Collectively, the above results demonstrated that CC cell proliferation was inhibited by *miR-141* silencing but fostered by miR-141 over-expression.

## miR-141 Represses Tumor Formation of CC Cells

After stably transfected cell lines were constructed, they were subcutaneously injected into nude mice to construct xenograft tumor model in nude mice so as to verify the effect of *miR-141* on the in vivo tumorigenic ability of CC cells. Firstly, qRT-PCR was carried out to detect *miR-141* expression in transfected cells, which revealed that compared with the antagomir NC group, the expression of *miR-141* in the *miR-141* antagomir group was markedly



**Figure 2** CC cell proliferation is suppressed by *miR-141* silencing but promoted by *miR-141* over-expression. (A) qRT-PCR was done to detect *miR-141* expression after *miR-141* was silenced in HCT-116 cells; (B) CCK-8 assay was performed to detect cell viability upon *miR-141* silencing; (C) colony formation assay was carried out to detect cell proliferation upon *miR-141* silencing; (D) qRT-PCR was done to detect *miR-141* expression after *miR-141* was overexpressed in SW480 cells; (E) CCK-8 assay was performed to detect cell viability upon *miR-141* overexpression; (F) colony formation assay in vitro was carried out to detect cell proliferation upon *miR-141* over-expression; \*Represents comparison with inhibitor NC,  $p < 0.05$ , #Represents comparison with mimic NC,  $p < 0.05$ . The experiment was repeated in triplicate.



reduced (Figure 3A). The detection of cell tumor growth speed each week out of five weeks after subcutaneous injection indicated that compared with the antagomir NC group, tumor formation speed in the *miR-141* antagomir was significantly decreased (Figure 3B). All mice were sacrificed five weeks later, and the tumors were taken out and weighed, which unveiled that compared with the antagomir NC group, tumor weight in the *miR-141* antagomir was noticeably reduced (Figure 3C and D). Taken together, *miR-141* suppressed CC cell tumor formation in vivo.

## *miR-141* Targeted Silences *PHLPP2* in CC Cells

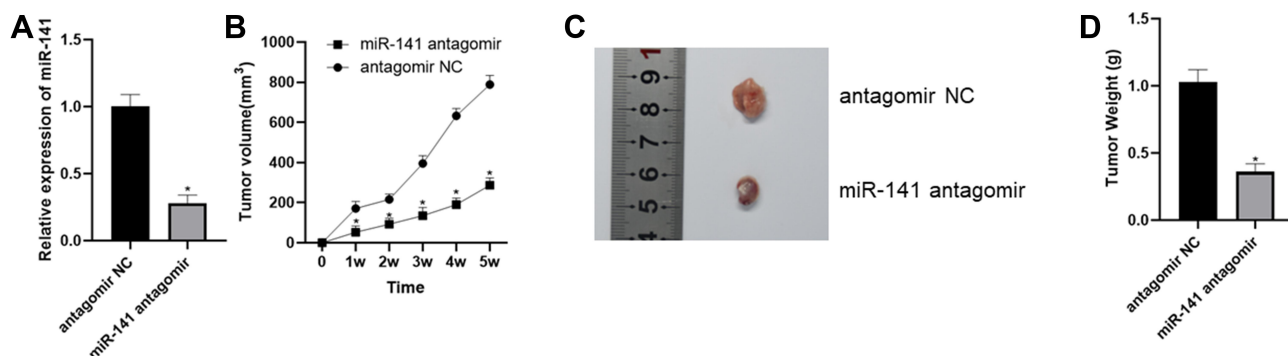
In order to further detect the functional mechanism by which *miR-141* regulates CC cell proliferation, firstly we performed differential analysis of mRNAs by using 41 normal samples and 462 cancer samples and obtained 2042 DE\_mRNAs (Figure 4A). Next, TargetScan, miRDB and miRTarBase databases were used to predict the target genes of *miR-141*, which were then intersected with down-regulated DE\_mRNAs in the TCGA-COAD dataset, and 2 DE\_mRNAs which had the binding sites with *miR-141* were obtained (Figure 4B), among which *PHLPP2* was markedly down-regulated in cancer tissues (Figure 4C) and led to a poor prognosis in patients (Figure 4D). A study showed that low *PHLPP2* expression promotes cell proliferation.<sup>18</sup> Consequently, we speculated that *miR-141* was likely to regulate CC cell proliferation by targeting *PHLPP2*.

To validate our speculation, we conducted qRT-PCR and Western blot to detect *PHLPP2* mRNA and protein expression in normal colon cell line CCD-18Co and CC cell lines HCT-116, HT-29, SW620 and SW480, which unveiled that compared with CCD-18Co, *PHLPP2* was poorly expressed in CC cells (Figure 4E and F). *miR-141* was silenced and over-

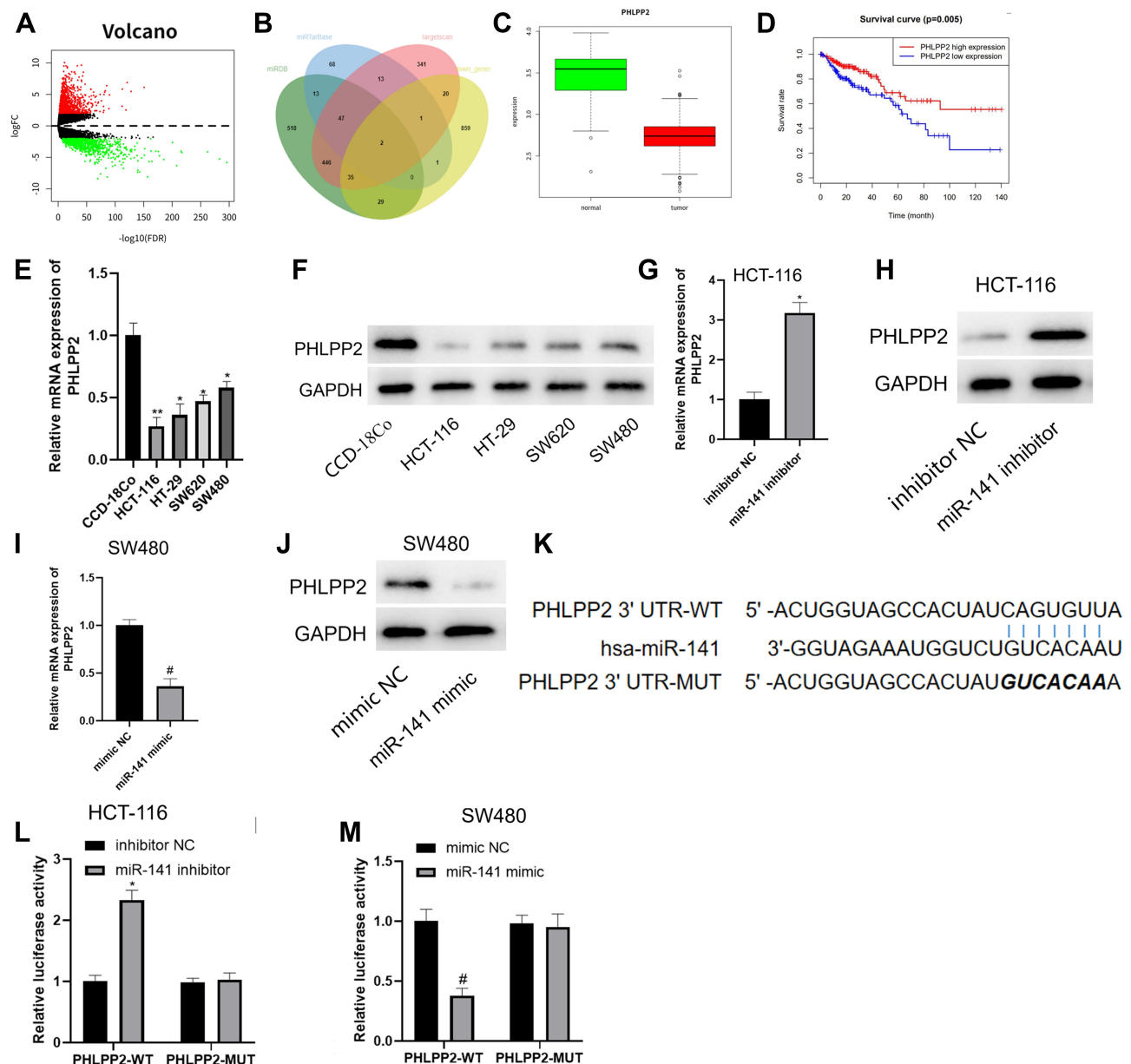
expressed in CC cell line HCT-116 and SW480, respectively, and *PHLPP2* mRNA and protein expression were detected by qRT-PCR and Western blot. The results suggested that compared with the inhibitor NC group, *PHLPP2* expression in the *miR-141* inhibitor group was markedly increased (Figure 4G and H); compared with the mimic NC group, *PHLPP2* expression in the *miR-141* mimic group was significantly decreased (Figure 4I and J). *miR-141* was predicted to bind with *PHLPP2* 3'UTR through TargetScan website (Figure 4K). Dual-luciferase reporter gene assay demonstrated that the luciferase activity of *PHLPP2*-WT could be markedly enhanced by silencing *miR-141* (Figure 4L) but weakened by over-expressing *miR-141* (Figure 4M), whereas *PHLPP2*-MUT was not noticeably affected by silencing or over-expressing *miR-141*. Collectively, *miR-141* targeted to down-regulate *PHLPP2* in CC cells.

## *miR-141* Facilitates CC Cell Proliferation by Silencing *PHLPP2*

In order to clarify the functional mechanism by which *miR-141* regulates CC cell proliferation by targeting *PHLPP2*, we simultaneously over-expressed *miR-141* and *PHLPP2* in CC cell line SW480. Firstly, qRT-PCR and Western blot were carried out to detect the expression of *miR-141* and *PHLPP2*, which indicated that compared with the mimic NC +oe-NC group, *miR-141* expression was markedly increased while *PHLPP2* expression was significantly reduced in the *miR-141* mimic+oe-NC group, and *PHLPP2* expression was remarkably raised in the mimic NC+oe-*PHLPP2* group; whereas compared with the *miR-141* mimic+oe-NC group, *PHLPP2* expression was noticeably elevated in the *miR-141* mimic+oe-*PHLPP2* group (Figure 5A and B). CCK-8 and colony formation assay were performed to detect cell proliferation, which revealed that compared with the mimic NC+oe-



**Figure 3** *miR-141* suppresses CC cell tumor formation in vivo. (A) qRT-PCR was carried out to detect *miR-141* expression in transfected cells; (B) detection of cell tumor growth speed each week out of five weeks after subcutaneous injection; (C) picture of xenograft tumors; (D) statistics of tumor weight; \* $p < 0.05$ . N=6 nude mice.

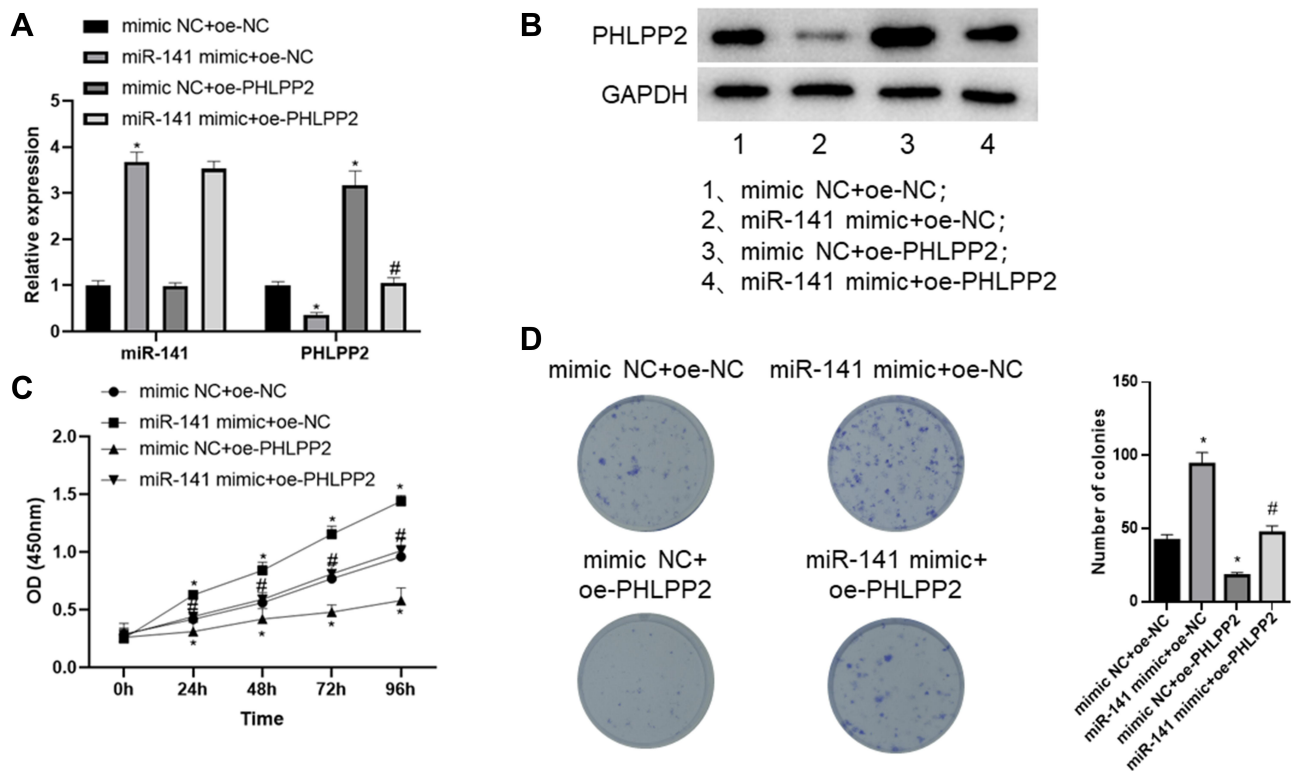


**Figure 4** *miR-141* targeted silences *PHLPP2* in CC cells. (A) Volcano plot of DE\_mRNAs in the TCGA-COAD dataset. Red and green represent markedly up-regulated and down-regulated DE\_mRNAs, respectively; (B) Venn diagram of down-regulated DE\_mRNAs and the predicted target mRNAs of *miR-141*; (C) relative expression of *PHLPP2* in normal and tumor samples in the TCGA-COAD dataset; (D) survival analysis of *PHLPP2* in the TCGA-COAD dataset; (E) qRT-PCR and (F) Western blot were conducted to detect *PHLPP2* mRNA and protein expression in normal colon cell line and CC cell lines; (G, H) *miR-141* was silenced in CC cell line HCT-116 and *PHLPP2* mRNA and protein expression were detected by (G) qRT-PCR and (H) Western blot; (I, J) *miR-141* was over-expressed in CC cell line SW480 and *PHLPP2* mRNA and protein expression were detected by (I) qRT-PCR and (J) Western blot; (K) *miR-141* was predicted to bind with *PHLPP2* 3'UTR through TargetScan website; (L, M) dual-luciferase reporter gene assay was performed; *miR-141* was silenced and over-expressed in CC cell line HCT-116 and SW480, respectively; \* in (I) represents comparison with CCD-18Co,  $p < 0.05$ , \*\*  $p < 0.01$ ; \* in (G) and (L) represents comparison with the inhibitor NC group,  $p < 0.05$ ; # in (I) and (M) represents comparison with the mimic NC group,  $p < 0.05$ . The experiment was repeated in triplicate.

NC group, cell proliferation was markedly increased in the *miR-141* mimic+oe-NC group but decreased in the mimic NC +oe-PHLPP2 group; whereas compared with the *miR-141* mimic+oe-NC group, cell proliferation was remarkably reduced in the *miR-141* mimic+oe-PHLPP2 group (Figure 5C and D). Taken together, *miR-141* fostered CC cell proliferation by silencing *PHLPP2*.

## Conclusion

*miR-141* is a member of miR-200 family and has been reported to be associated with various cancers.<sup>27</sup> For example, *miR-141* is down-regulated in HCC,<sup>14</sup> pancreatic ductal adenocarcinoma<sup>28</sup> and renal cell carcinoma,<sup>29</sup> in which it plays a role as a potential tumor suppressor gene. However, *miR-141* has been found to act as an oncogene in



**Figure 5** *miR-141* facilitates CC cell proliferation by silencing *PHLPP2*. (A) qRT-PCR was conducted to detect *miR-141* and *PHLPP2* mRNA expression; (B) Western blot was performed to detect *PHLPP2* protein expression; (C) CCK-8 assay was carried out to assess cell viability; (D) colony formation assay was conducted to evaluate cell proliferation; \*Represents comparison with the mimic NC+oe-NC group,  $p < 0.05$ ; #Represents comparison with the miR-141 mimic+oe-NC group,  $p < 0.05$ . The experiment was repeated in triplicate.

nasopharyngeal carcinoma, suggesting that it plays dual effects on different cancers.<sup>3</sup> Earlier Liu et al<sup>30</sup> validated that *miR-141* expression is higher than that of normal tissue in non-small cell lung cancer. However, there have been no further reports on the functional mechanism of *miR-141* in CC. In this study, we found that *miR-141* was highly expressed in CC tissue by bioinformatics analysis, which was further verified by the detection of *miR-141* expression in CC cells and normal colon cells using qRT-PCR. Additionally, by conducting CCK-8 assay, colony formation assay and tumor formation in nude mice experiment, we validated that *miR-141* could promote CC tumor growth and unveiled that it played a carcinogenic role in CC.

*PHLPP2* has been found to be down-regulated in various cancers<sup>31,32</sup> and is able to suppress the invasion of bladder cancer by facilitating the autophagy and degradation of MMP2 protein.<sup>33</sup> Besides, *PHLPP1* is capable of inhibiting the metastasis of melanoma by repressing AKT2 activation.<sup>22</sup> However, the regulatory function of *PHLPP2* in CC needs further research. In this study, we further investigated the molecular mechanism by which *miR-141* promoted CC cell proliferation. We discovered that

*PHLPP2* was a theoretical target gene of *miR-141* through prediction using publicly available bioinformatics websites, which was subsequently validated by dual-luciferase reporter assay. Meanwhile, we also found that *PHLPP2* could reverse miR-141-induced CC cell proliferation as revealed by CCK-8 and colony formation assay. Collectively, *miR-141* negatively regulated *PHLPP2* expression in CC cells by directly targeting *PHLPP2* 3'UTR so as to foster CC cell proliferation.

In all, this study verified that miR-141 was up-regulated in CC and facilitated CC cell proliferation by targeted inhibiting the expression of *PHLPP2*. Therefore, *miR-141* is likely to be a potential therapeutic target of CC treatment.

## Highlights

1. *miR-141* targeted to regulate *PHLPP2* in colon cancer;
2. Colon cancer cell proliferation and in vivo tumorigenic ability could be suppressed by *miR-141* down-regulation but promoted by *miR-141* over-expression;
3. *miR-141* fostered colon cancer cell proliferation by silencing *PHLPP2*.



## Ethics approval

This study was conducted in accordance with Guide for the care and use of laboratory animals and was approved by the Institutional Review Boards of Zhejiang Jinhua Guangfu Hospital.

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There is no funding to report.

## Disclosure

The authors declare that they have no potential conflicts of interest for this work.

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