

Downregulation of miR-216a-5p by long noncoding RNA PVT1 suppresses colorectal cancer progression via modulation of YBX1 expression

This article was published in the following Dove Press journal:
Cancer Management and Research

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Purpose: Increasing evidence has demonstrated that microRNAs (miRNAs) are closely related to the occurrence and development of tumors. MiR-216a-5p, located at 2p16.1, has been shown to suppress proliferation of cancerous cells. However, its expression and function in colorectal cancer (CRC) remain unclear.

Materials and methods: The significance of miR-216a-5p in CRC was studied by analyzing miR-216a-5p expression in CRC tissues and its association with clinicopathological parameters. CRC cells, stably overexpressing miR-216a-5p, were evaluated for cell proliferation and metastasis using cell counting kit-8 (CCK-8) and transwell assay methods. Epithelial-mesenchymal transition (EMT) pathway was analyzed by Western blotting. Bioinformatics, quantitative real-time polymerase chain reaction (RT-qPCR), and luciferase reporter assay were performed to define the regulation of PVT1/miR-216a-5p/Y Box Binding Protein 1 (YBX1) axis in CRC.

Results: The expression of miR-216a-5p was found to be significantly downregulated in CRC and was correlated with the various stages and differentiation degree of the tumors. Moreover, the overexpression of miR-216a-5p could significantly inhibit the tumor growth, metastasis, and EMT progression in CRC. Furthermore, the expression of miR-216a-5p was negatively correlated with the expression of PVT1, and PVT1 could reverse tumor suppressive effect of miR-216a-5p in CRC cells. Finally, YBX1 might be the key target of PVT1/miR-216a-5p axis in CRC.

Conclusion: Downregulation of miR-216a-5p by PVT1 could suppress CRC progression via modulating YBX1 expression.

Keywords: colorectal cancer, PVT1, miR-216a-5p, YBX1, epithelial-mesenchymal transition, metastasis

Introduction

Colorectal cancer (CRC) is one of the most common malignant tumors worldwide, with mortality rate just lower than that of lung cancer, as estimated by the recent global cancer statistics.^{1,2} Despite the rapid development of various therapeutic approaches such as endoscopic, laparotomic or laparoscopic removal of tumor, chemotherapy, radiotherapy, or immunotherapy, the prognosis of patients with CRC remains unsatisfactory. Therefore, a detailed study of the underlying mechanisms of CRC progression is required to develop more precise diagnostic methods and effective treatments.^{3,4}

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The emerging data in the past decade have established the complex structure and function of noncoding RNAs (ncRNAs) and have altered the perception of ncRNAs from “junk” to “functional regulatory molecules”. Long non-coding RNAs (lncRNAs) and microRNAs (miRNAs) regulate numerous physiological and pathophysiological processes within cells.⁵ MiRNAs are 20–25 nucleotides long and regulate gene expression post-transcriptionally by binding to the 3'-untranslated region of target mRNAs, resulting in RNA degradation or translational inhibition.⁶ Deregulation of miR-216a-5p has been reported in the development of prostate cancer,⁷ hepatocellular carcinoma,⁸ and so on. Its role as an oncogene or tumor suppressor in the complex cellular signaling network varies with cancer types and its target genes. For example, miR-216a-5p significantly suppresses cellular proliferation and/or viability in pancreatic and prostate cancers, whereas it promotes tumor progression in renal and cervical cancers. However, the expression and effects of miR-216a-5p in CRC have not been defined.

In the present study, RT-qPCR was performed to analyze the expression of miR-216a-5p in CRC tissues and its correlation with the clinicopathological parameters of CRC patients. We further explored the underlying mechanism causing the downregulation of miR-216a-5p and its role in cellular proliferation, metastasis and epithelial–mesenchymal transition (EMT) process via in vitro assays and bioinformatics analysis.

Materials and methods

Patients tissue samples

CRC and adjacent normal tissues were collected from 70 patients who were admitted to the Affiliated Cancer Hospital & Institute of Guangzhou Medical University from 2012 to 2014. The patients were not treated with radio- or chemical therapy before collection of the tissues. The pathological diagnosis of tissue samples was performed by veteran pathologists. All patients signed the informed consent before participating in this research. This study was approved by the medical ethics committee of Affiliated Cancer Hospital & Institute of Guangzhou Medical University and conducted in accordance with the Declaration of Helsinki (6th revision, 2008).

Cell culture

The intestinal normal cell line (FBS), and five CRC cell lines (LoVo, SW480, HT-29, HCT-116, and Caco-2) were

acquired from ATCC (Shanghai, China). The McCoy's 5a Medium (Gibco, Grand Island, NY, USA) with 10% fetal bovine serum (Gibco, Australia origin) was used to culture these cell lines.

RNA transfection

Hsa-miR-216a-5p and has-PVT1 cDNA plasmid (pcDNA3.1) were developed by GENESSEED (Guangzhou, China). The lentiviruses expressing either miR-216a-5p (OE-miR-216a-5p) or PVT1 (OE-PVT1) and the respective negative controls (OE-vector) were constructed by GENE (Shanghai, China). The LoVo and HCT-116 cells were seeded and transfected with OE-miR-216a-5p or OE-PVT1 lentiviruses with 5 µg/mL polybrene to improve the transfection efficiency. The multiplicity of infection (MOI) is known to be 50 for LoVo cells, whereas 30 for HCT-116 cells. Seven days later, stably transfected cells were collected for further study. The sequences of miR-216a-5p mimics are shown in Table 1.

Quantitative real-time polymerase chain reaction

Total RNA from the frozen tissues and cell lines were extracted by RNAiso Plus reagent (TaKaRa, Dalian, China), followed by reverse transcription using PrimeScript RT Master Mix (TaKaRa), according to the manufacturer's instruction. Next, the SYBR Premix Ex Taq II Kit (TaKaRa) was used to determine the expression of target genes using StepOnePlus system (Applied Biosystems, CA, USA). $2^{-\Delta\Delta C_t}$ method was applied to detect the fold changes. The primers sequences are listed in Table 1.

Table 1 Sequences of oligomers and primers used in the study

Name	Sequence (5'-3')
miR-216a-5p mimics-F'	UAAUCUCAGCUGGCAACUGUGA
miR-216a-5p mimics-R'	ACAGUUGCCAGCUGAGAUUAAU
NC mimics-F'	UUCUCCGAACGUGUCACGUTT
NC mimics-R'	ACGUGACACGUUCGGAGAATT
PVT1-F'	CAGCACTCTGGACGGAC
PVT1-R'	CAACAGGAGAAGCAAACA
YBX1-F'	GGAGTTTGATGTTGTTGAAGGA
YBX1-R'	AACGGAACACCACCAGGAC
miR-216a-5p-RT	CTCAACTGGTGTCTGCTGGA
miR-216a-5p-F'	TGTCGCAAATCT CTGCAGG
miR-216a-5p-R'	CAGAGCAGGGTCCGAGGTA
β -actin forward	CGC TCT CTG CTC CTG TTC
β -actin reverse	ATC CGT TGA CTC CGA CCT TCA C

Cell counting kit 8 assay

The OE-miR-216a-5p or OE-PVT1 cells (LoVo and HCT-116) were plated into 96-well plates at the concentration of 5×10^3 cells/well. Next, CCK-8 (Donjindo, Kumamoto, Japan) was added to the wells according to the manufacturer's instructions. Optical density was measured using microplate reader (Bio-Rad, Hercules, CA, USA) to the cell viability.

Cell migration assay

The transwell assays were performed to analyze cellular migration and invasion abilities. First, 500 μ L McCoy's 5a Medium with 10% fetal bovine serum was added into the lower chambers, followed by the addition of 200 μ L serum-free McCoy's 5a medium containing 5×10^4 cells into the upper chambers. After 24 hrs, non-migrating and non-invading cells were removed from the upper surface of the membranes. Attached cells were stained in 0.5% crystal violet, and the cell migration and invasion ability were evaluated through the digital imaging of the cells.

Western blot analysis

Total protein was extracted from cells, and its concentration was analyzed through the BCA protein Assays Kit (Thermo Fisher Scientific; Shanghai, China). Twenty micrograms of each protein samples were separated on SDS-PAGE gel and transferred onto the polyvinylidene fluoride (PVDF) membranes. Next, the PVDF membranes were incubated with 10% BSA, followed by primary and secondary antibody incubations. Finally, the protein bands were visualized by GeneSnap using SynGene system.

Dual-luciferase reporter assays

Based on publicly available bioinformatic algorithms (Starbase), miR-216a-5p is predicted as a downstream target of PVT1. To address this issue, the 293T cells were seeded in 24-well plates at the concentration of 5×10^4 cells/well. Subsequently, cells were transfected with miR-216a-5p mimics or negative control (NC), with pmirGLO-PVT1-WT (wild type) or pmirGLO-PVT1-MUT (mutant) plasmid. After 48 hrs, cells were lysed in passive lysis buffer, and the relative luciferase activity was calculated with the dual-luciferase reporter assay (Promega, Madison, WI, USA). The firefly luminescence was normalized to Renilla luminescence. According to Starbase algorithm, miR-216a-5p is also predicted to directly regulate the YBX1 expression.

Similarly, to address this point, 293T cells were transfected with miR-216a-5p mimics or miR-216a-5p NC, along with pmirGLO-YBX1-WT or pmirGLO-YBX1-MUT plasmid, followed by dual-luciferase reporter assay.

Statistics

All data in this study are shown as mean \pm standard deviation (SD). In addition, all statistical data were analyzed in the IBM SPSS Statistics 20.0 (IBM, IL, USA). The chi-square test, two-tailed Student's *t*-test, Mann-Whitney U test, analysis of variance (ANOVA) and Kruskal-Wallis test were applied to evaluate the data. The Kaplan-Meier method and log-rank test were performed to analyze the survival data, significant difference was considered at $P < 0.05$.

Results

Downregulation of miR-216a-5p in CRC

In order to evaluate the correlation between the miR-216a-5p expression and the clinicopathological characteristics in CRC patients, miR-216a-5p expression was estimated in cancerous and adjacent normal tissues. MiR-216a-5p was significantly downregulated in cancerous tissue as compared to adjacent normal tissues (Figure 1A). Moreover, low-expression of miR-216a-5p accounted for 77.14% (54/70) of CRC specimens (Figure 1B). Further, low expression of miR-216a-5p in CRC patients was correlated to have higher T (Figure 1E) and N stages (Figure 1F) and shorter disease-free survival (DFS) (Figure 1G). Although no significant correlation was observed between differential expression of miR-216a-5p among patients and their tumor size (Figure 1D), and its differentiation stages (Figure 1C), these results showed that miR-216a-5p can be a promising prognostic biomarker for CRC.

Furthermore, CRC patients were divided into two groups; a high-expression group and low-expression group, based on the median expression of the miR-216a-5p. Consistent with the earlier data, the expression of miR-216a-5p was closely related to the lymphatic metastasis ($P=0.014$) and T stage ($P=0.029$) (Table 2). Furthermore, Cox analyses showed that lymphatic metastasis ($P=0.020$) and miR-216a-5p expression ($P=0.005$) are independent risk factors for DFS of patients with CRC (Table 3). Together, these results demonstrated that miR-216a-5p expression is negatively related with the poor prognosis of CRC patients.

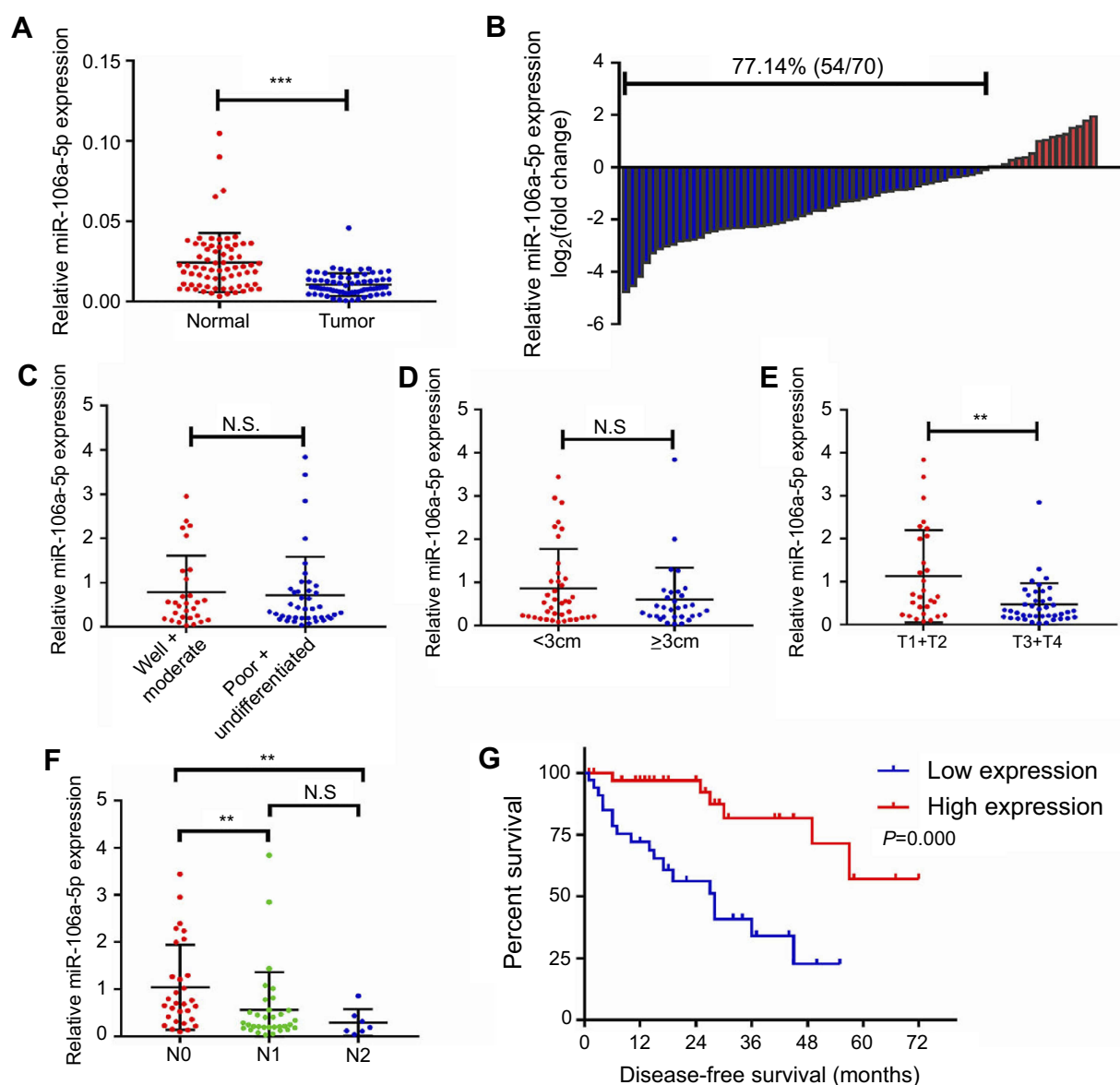


Figure 1 The expression profile of miR-216a-5p in CRC tissues and its correlation with clinicopathological parameters of CRC patients. **(A)** RT-qPCR results showing significant downregulation of miR-216a-5p in CRC tissues than in adjacent normal tissues. **(B)** Waterfall plot showing relatively low expression of miR-216a-5p in 77.14% (54/70) patients; the data are represented as Log₂ (T/N). Scatter plots showing correlation analysis between miR-216a-5p expression and **(C)** tumor differentiation, **(D)** tumor size, **(E)** T stage, and **(F)** N stage of the tumor. **(G)** Kaplan-Meier plot showing the correlation between miR-216a-5p expression and duration of disease-free survival (DFS) in patients. Note that each dot represents a patient. ** $P < 0.01$, *** $P < 0.001$, N.S. means "not significant."

MiR-216a-5p overexpression suppresses the CRC cells proliferation in vitro

The gain-of-function experiments were performed to further demonstrate the biological role of miR-216a-5p in CRC. The RT-qPCR results showed a significantly lower endogenous expression of miR-216a-5p in CRC cell lines (HT-29, LoVo, SW480, HCT-116, and Caco-2) when compared to a normal intestinal normal cell line (FHC), with lowest expression in HCT-116 and LoVo cells (Figure 2A). Next, we stably overexpressed miR-216a-5p in HCT-116

and LoVo cells (OE-miR-216a-5p) via lentiviral transfection, and expression of miR-216a-5p was compared with a negative control group (OE-vector) as well as with a naive group (non-transfected cells). As shown in Figure 2B and C, OE-miR-216a-5p cells showed a significantly increased expression of miR-216a-5p when compared to OE-vector and naive cells. Subsequently, CCK-8 assays were performed to evaluate the in vitro proliferation abilities of OE-miR-216a-5p cell lines. The results showed that the miR-216a-5p upregulation effectively suppresses the

Table 2 Correlation between miR-216a-5p expression and clinicopathological characteristics of colorectal cancer patients

Clinical parameter		MiR-216a-5p		P-value
		Low (n=35)	High (n=35)	
Gender	Male	17 (42.5%)	23 (57.5%)	0.147
	Female	18 (60.0%)	12 (40.0%)	
Age	<60	18 (50.0%)	18 (50.0%)	1.000
	≥60	17 (50.0%)	17 (50.0%)	
Differentiation	Well+moderate	13 (44.8%)	16 (55.2%)	0.467
	Poor+undifferentiated	22 (53.7%)	19 (46.3%)	
Tumor size (cm)	<3	18 (46.2%)	21 (53.8%)	0.470
	≥3	17 (54.8%)	14 (45.2%)	
T stage	T1 T2	10 (34.5%)	19 (65.5%)	0.029
	T3+T4	25 (61.0%)	16 (39.0%)	
Lymphatic metastasis	N0	9 (30.0%)	21 (70.0%)	0.014
	N1	21 (63.6%)	12 (36.4%)	
	N2	5 (71.4%)	2 (28.6%)	

Table 3 Cox analyses of disease-free survival in colorectal cancer patients

Clinical parameters	Univariate				Multivariate			
	HR	95% CI		P	HR	95% CI		P
Gender (male vs female)	1.511	0.676	3.379	0.315				
Age (<60 vs ≥60)	0.411	0.174	0.970	0.042	0.538	0.211	1.373	0.195
Differentiation (well+moderate vs poor+undifferentiated)	2.671	1.056	6.755	0.038	1.355	0.497	3.694	0.552
Tumor size (cm) (<3.0 vs ≥3.0)	1.990	0.881	4.495	0.098				
T stage (T1+T2 vs T3+T4)	3.433	1.278	9.220	0.014	2.051	0.732	5.748	0.172
Lymphatic metastasis (N0, N1, and N2)	4.238	2.075	8.656	0.000	2.990	1.188	7.525	0.020
MiR-216a-5p expression (low vs high)	0.178	0.065	0.485	0.001	0.223	0.079	0.630	0.005

proliferation of LoVo and HCT-116 cells (Figure 2D and E).

Overexpression of miR-216a-5p inhibits cell metastasis via silencing the EMT pathway in CRC cell lines

Given that the expression level of miR-216a-5p was correlated to the lymph node metastasis, we hypothesized a role of miR-216a-5p in the regulation of CRC metastasis. Therefore, Transwell assays were performed to evaluate the effect of miR-216a-5p on cellular migration and invasion capabilities. The OE-miR-216a-5p cells showed significantly reduced cellular migration (Figure 3A) and (Figure 3B) invasion in HCT-116 and LoVo cells. To study the potential mechanism of miR-216a-5p as a tumor-suppressor miRNA, the protein expression of EMT-related markers was analyzed. Interestingly, OE-miR-216a-5p cells showed an increase

in the expression of E-cadherin (an epithelial marker) and a lower expression of N-cadherin, Vimentin, and Snail (the mesenchymal markers), when compared to OE-vector (Figure 3C). Therefore, we concluded that miR-216a-5p inhibits metastasis and EMT progression in CRC.

MiR-216a-5p is a direct target of PVT1 in CRC

Next, based on Starbase prediction, we addressed the possibility of miR-216a-5p acting as a downstream target of PVT1. To test this, we first evaluated the expression pattern of PVT1 in CRC patients. As shown in Figure 4A, PVT1 was overexpressed in 83.33% (25/30) patients with CRC. Consistently, the expression level of PVT1 was significantly higher in CRC tissues as compared to adjacent normal tissues (Figure 4B). Further, a negative correlation was observed between the expression levels of miR-216a-5p

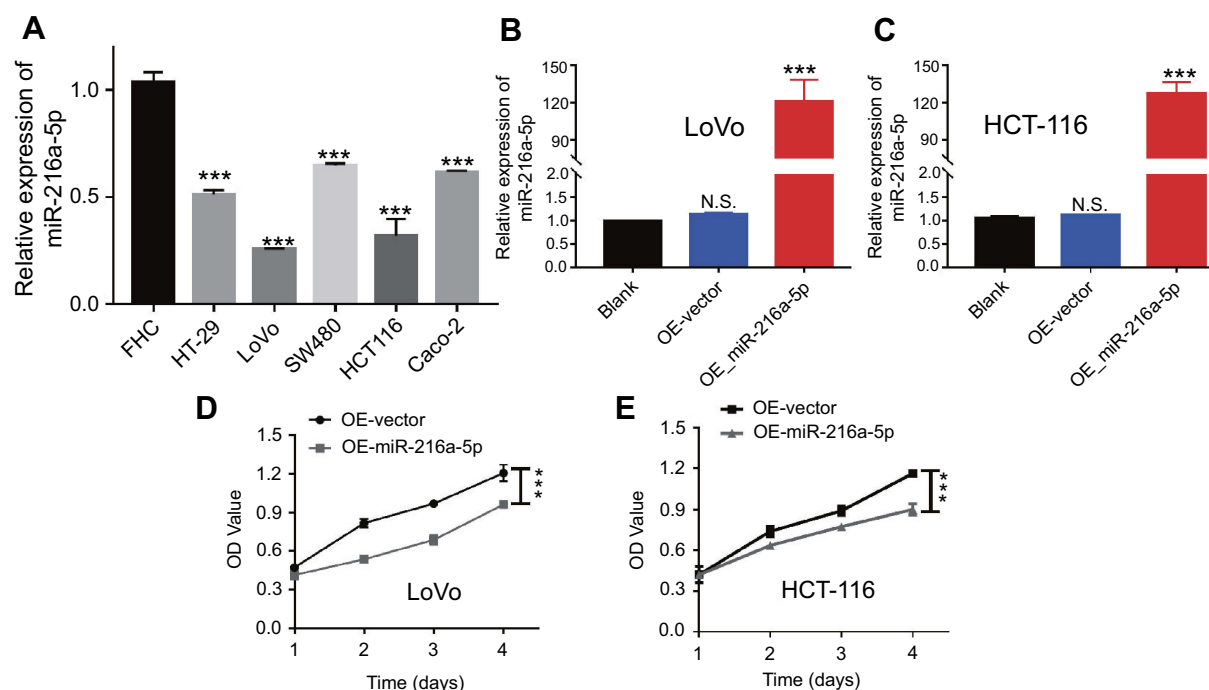


Figure 2 Overexpression of miR-216a-5p suppresses the proliferation of CRC cells in vitro. **(A)** Bar diagram showing the relatively low expression of miR-216a-5p in CRC cell lines (HT-29, LoVo, SW480, HCT-116, and Caco-2) as compared to normal intestinal cell line (FHC). **(B and C)** RT-qPCR results showing the overexpression of miR-216a-5p in stable OE-miR-216a-5p LoVo and HCT-116 cell lines, as compared to OE-vector cells. **(D and E)** CCK-8 assay showing the effect of miR-216a-5p upregulation on proliferation of LoVo and HCT-116 cells. All experiments were repeated at least three times. *** $P < 0.001$, N.S. means "not significant."

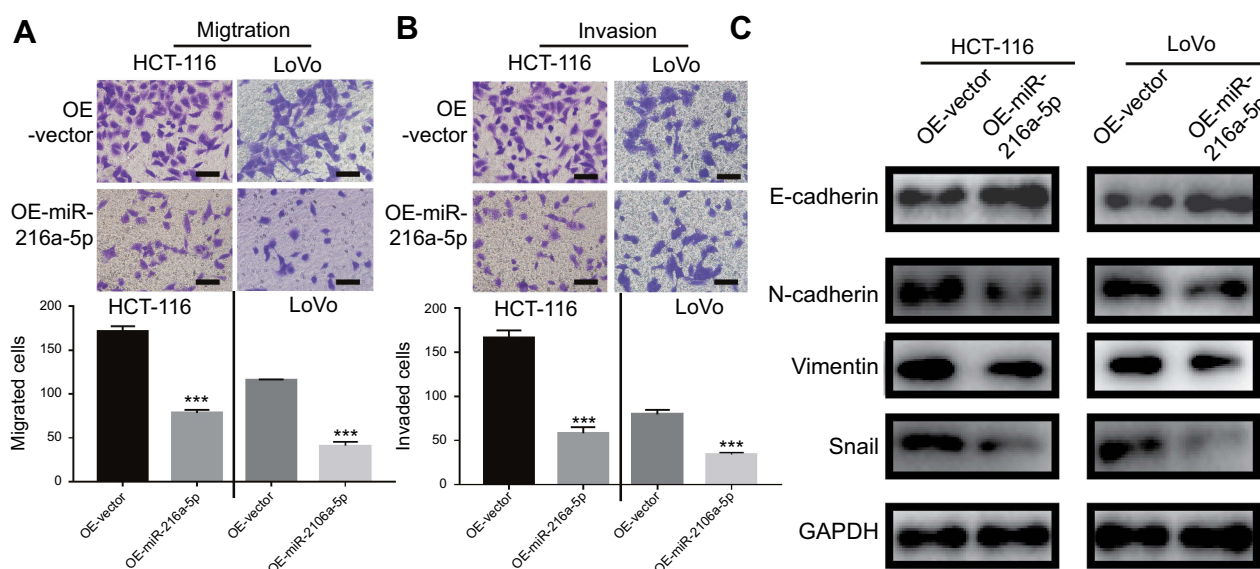


Figure 3 MiR-216a-5p suppresses migration, invasion, and EMT in CRC cell lines. Transwell assay showing reduced **(A)** cell migration and **(B)** cell invasion upon overexpression of miR-216a-5p in HCT-116 and LoVo cell. **(C)** Immunoblots showing increased and decreased expression of E-cadherin (epithelial marker) and N-cadherin/Vimentin/Snail (mesenchymal markers) respectively, upon overexpression of miR-216a-5p in HCT-116 and LoVo cells. *** $P < 0.001$.

and PVT1 (Figure 4C, $r = -0.3048$, $P = 0.00016$). These results showed that PVT1 and miR-216a-5p were highly associated with CRC.

To further explore their relationship, we predicted a miR-216a-5p binding site at 3'UTR region of PVT1

(Figure 4D). Luciferase tagged PVT1-wild type (pmirGLO-PVT1-WT) and PVT1-mutant type (pmirGLO-PVT1-MT) 293T cells were constructed and proceeded for luciferase assays. Luciferase activity was dramatically decreased in pmirGLO-PVT1-WT as

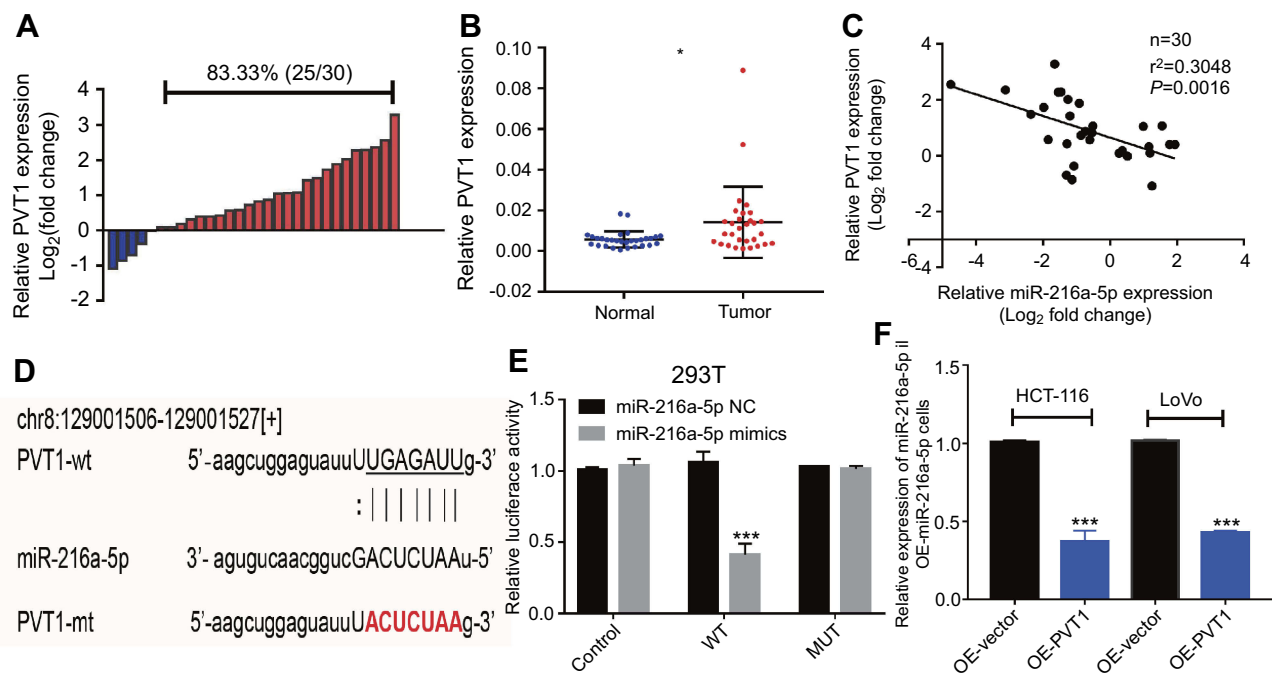


Figure 4 MiR-216a-5p is a direct target of lncRNA PVT1 in CRC. (A) Waterfall plot showing relatively high expression of PVT1 in 83.33% (25/30) patients; the data are represented as Log₂ (T/N). (B) Scatter plot showing higher expression of PVT1 in CRC tissues than that of adjacent normal tissues, evaluated by RT-qPCR. (C) Correlation analysis between the expression level of miR-216a-5p and PVT1 in 30 paired CRC tissues. (D) Illustration is showing the predicted sequence of the miR-216a-5p binding site on 3'UTR of PVT1 mRNA. Sequence of the mutated binding site in PVT1 is also represented. (E) The relative luciferase activity pmirGLO-PVT1-WT or pmirGLO-PVT1-MUT 293T cells, upon transfection with miR-216a-5p mimics or negative control (miR-216a-5p NC). (F) RT-qPCR data showing the relative expression of miR-216a-5p in OE-miR-216a-5p cells, upon overexpression of PVT1. **P*<0.05, ****P*<0.001.

compared to pmirGLO-PVT1-MT cells when treated with miR-216a-5p mimics (Figure 4E). Furthermore, we successfully constructed the PVT1-overexpressed (OE-PVT1) HCT-116 and LoVo cell lines via lentiviral transfection (Figure S1). Interestingly, miR-216a-5p expression was found to be dramatically downregulated in OE-PVT1 cells as compared to the control group (Figure 4F). Therefore, we concluded that PVT1 might be an upstream regulator of miR-216a-5p in CRC.

Upregulation of PVT1 rescues the anti-proliferative phenotype of OE- miR-216a-5p cells

Because the high expression of miR-216a-5p could inhibit proliferation, metastasis, and EMT progression in CRC, its tumor suppressive function might be reversed by the upregulation of PVT1, as miR-216a-5p is predicted as a direct target of PVT1. To confirm this hypothesis, proliferation, metastasis, and EMT progression of OE-miR-216a-5p cells were evaluated under overexpression of PVT1. According to CCK-8 assay, OE-miR-216a-5p showed a higher cell proliferation when transfected with OE-PVT1 as compared

to OE-vector (Figure 5A and B). Consistently, migration and invasion of cells were increased after transfection with OE-PVT1 as compared to OE-vector (Figure 5C and D). Similar rescue effect was observed when expression of EMT markers was analyzed (Figure 5E and F).

YBX1 might be the key target of PVT1/ miR-216a-5p axis in CRC

In order to find the downstream target of PVT1/miR-216a-5p axis in CRC, we shortlisted only those 200 target genes of miR-216a-5p, which were predicted by both TargetScan and miRanda databases (Figure 6A). Next, we shortlisted some genes that are co-expressed with PVT1, using cBioPortal (<http://www.cbioportal.org>). By drawing a Venn diagram, we found 10 overlapping genes (Figure S2) that were suspected to be the potential targets for PVT1/miR-216a-5p axis; further analyses were performed (Figure 6B). Next, protein-protein interaction (PPI) network analysis was performed on the selected 10 genes using STRING. YBX1 was found to be located at the central position, which indicates that YBX1 might play the key role downstream of PVT1/miR-216a-5p axis (Figure 6C).

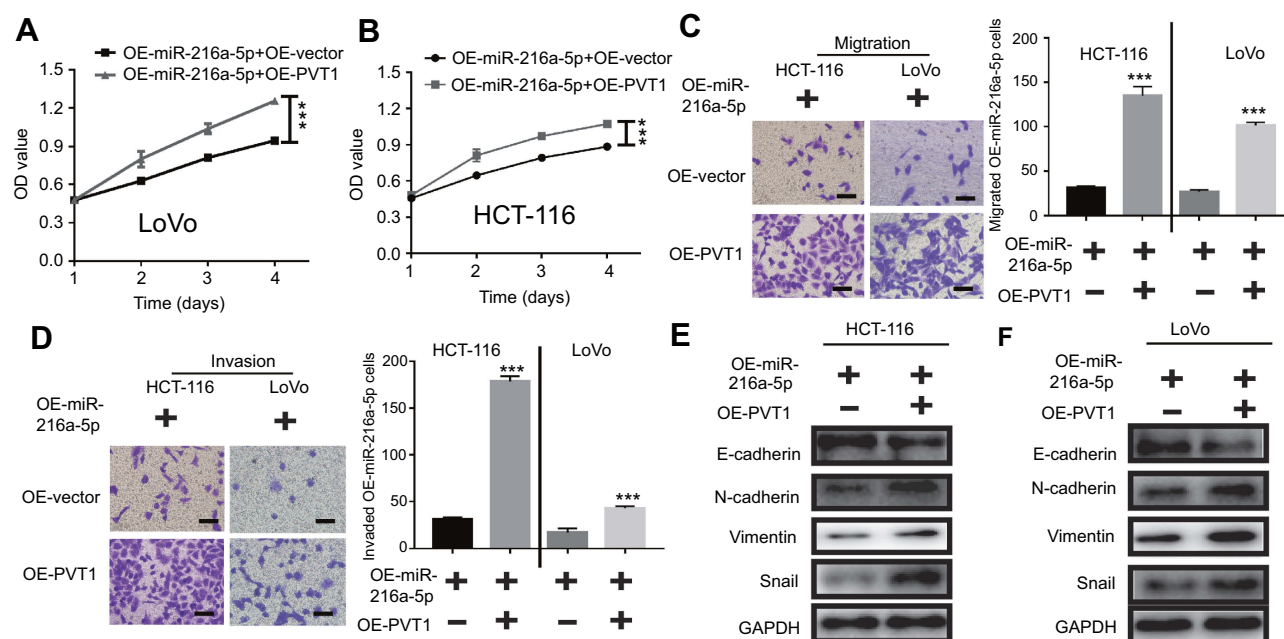


Figure 5 Upregulation of PVT1 reverses the anti-proliferative effect of miR-216a-5p overexpressed (OE- miR-216a-5p) CRC cells. **(A and B)** CCK-8 assay showing the effect of PVT1 upregulation on proliferation of OE-miR-216a-5p LoVo and HCT-116 cells. Transwell assay showing increased **(C)** cell migration and **(D)** cell invasion upon overexpression of PVT1 in OE-miR-216a-5p cells. **(E)** Immunoblots showing decreased and increased expression of E-cadherin (epithelial marker) and N-cadherin/Vimentin/Snail (mesenchyme markers) respectively, upon overexpression of PVT1 in OE-miR-216a-5p cells. *** $P < 0.001$.

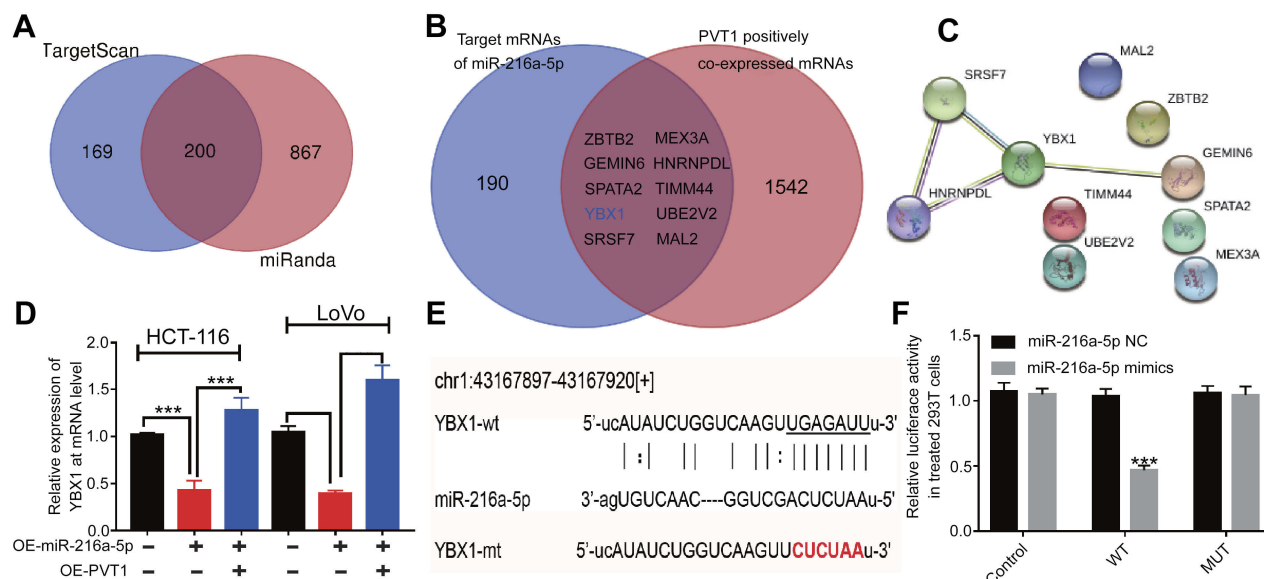


Figure 6 YBX1 might be the key target of PVT1/miR-216a-5p axis in CRC. **(A)** Predicted target genes of miR-216a-5p from TargetScan and miRanda databases, with 200 common predicted targets. **(B)** Venn diagram representing the potential 10 target genes, regulated by both PVT1 and miR-216a-5p. **(C)** Protein-protein interaction networks of the 10 target genes regulated by PVT1/miR-216a-5p axis using STRING database. **(D)** RT-qPCR data showing the relative expression of YBX1 mRNA in miR-216a-5p and/or PVT1 upregulated CRC cells. **(E)** Illustration showing the predicted sequence of the miR-216a-5p binding site on 3'UTR of YBX1 mRNA. Sequence of the mutated binding site in YBX1 is also represented. **(F)** The relative luciferase activity pmirGLO-YBX1-WT or pmirGLO-YBX1-MUT 293T cells, upon transfection with miR-216a-5p mimics or negative control (miR-216a-5p NC). *** $P < 0.001$.

To further explore the involvement of YBX1, its mRNA level was estimated in OE-PVT1 and OE-miR-216a-5p cells. The results showed that the expression of YBX1 was found to be downregulated in OE-miR-216a-

5p cells, and this effect was reversed by the overexpression of PVT1 (Figure 6D). We also predicted the binding regions of miR-216a-5p on 3'UTR of YBX1 mRNA and constructed pmirGLO-YBX1-WT and pmirGLO-YBX1-

MT 293T cells (Figure 6E). As shown in Figure 6F, the luciferase activity of pmirGLO-YBX1-WT cells was dramatically reduced in the presence of miR-216a-5p mimics, which remains unchanged in pmirGLO-YBX1-MT cells (Figure 6F). Therefore, we concluded that YBX1 might be the key downstream regulatory target of PVT1/miR-216a-5p axis in CRC.

Discussion

The rates of colorectal cancer are rising equally among women and men,^{1,9} and despite the availability of advanced laparoscopic techniques and molecular target therapies, the 5-year survival rate is still unsatisfactory.¹⁰ Thus, identification of novel biomarkers for diagnosis and prognosis of CRC is urgently needed. In recent years, a plethora of evidences suggest that miRNAs play a significant role in the development of tumors.^{11–13} The dysfunction of miRNAs interferes with gene expression at the post-transcriptional level, resulting in deregulation of tumor growth, metastasis, and drug resistance.^{14,15} For example, Guo et al have demonstrated that miRNA-532-3p can inhibit the gastric cancer cell proliferation via suppressing the expression of Rab3IP.¹⁶ MiR-216a-5p has been reported as an inhibitor for several malignant tumors such as hepatocellular and thyroid carcinoma.^{17,18} However, the exact function of miR-216a-5p in CRC remains largely unknown. Therefore, in the present study, we attempted to investigate the function and underlying mechanism of miR-216a-5p in CRC.

We first demonstrated that the miR-216a-5p was down-regulated in the CRC tissues. Next, the correlation analysis between the miR-216a-5p expression and the clinicopathologic characteristics of CRC patients indicated that the patients with lower expression of miR-216a-5p had poor prognosis. Consistent with the earlier findings, *in vitro* analyses showed that miR-216a-5p overexpression could significantly inhibit the proliferation, cellular migration, and invasion capabilities of CRC cell lines. Together, we suggest that miR-216a-5p suppresses cancer progression and might be a promising biomarker for the treatment of CRC. However, the potential mechanism of miR-216a-5p as a tumor suppressor remains unidentified.

The transition of cells from epithelial to mesenchymal phenotype (EMT) enhances the migratory and invasive properties of cancerous cells and cause metastasis of malignant tumor.^{19,20} MiRNAs are reported to regulate EMT mediated metastasis.^{21–23} MiR-320 has been shown to suppress EMT by regulating the expression of KLF5

and HIF-1 α , in turn suppressing the migration and invasion of gastric cancer.²⁴ Further, Pan et al have reported that miR-4429 inhibits CDK6 expression and suppress tumor progression of clear cell renal cell carcinoma by interfering with the EMT pathway.²⁵ In our correlation analysis, a significant association of miR-216a-5p expression with tumor stage and lymphatic metastasis was observed. Furthermore, the overexpression of miR-216a-5p leads to simultaneous upregulation of E-cadherin and downregulation of N-cadherin, Vimentin, and Snail expression. Taken them together, we propose that miR-216a-5p exerts anti-metastatic role in CRC by deregulating the EMT pathway.

In recent years, a lot of attention has been directed toward long non-coding RNAs (lncRNAs), due to their significant role in the development of tumors.^{26–28} It has been established that lncRNAs suppress the function of miRNAs through molecular sponging.²⁹ A novel plasmacytoma variant translocation 1 (PVT1) lncRNA, located at 8q24.21, has been shown as an important regulator of CRC.^{30,31} Consistent with the previous reports, we demonstrated that PVT1 is upregulated in the CRC tissues. Additionally, we showed a negative correlation between PVT1 and miR-216a-5p expression levels, which was confirmed by the dual-luciferase reporter assays. Further, anti-tumor effects on proliferation, migration, invasion, and EMT process of miR-216a-5p were rescued in the cells with PVT1 overexpression. Therefore, we concluded that PVT1 might be an upstream regulator of miR-216a-5p in CRC.

Y-box binding protein 1 (YBX1) is a multifunctional oncoprotein that deregulates numerous genes involved in cell proliferation, survival, drug resistance, chromatin destabilization, and metastasis of various cancers.³² Upregulation of YBX1 in CRC is demonstrated associated with local recurrence; its expression in tumor tissues was an independent prognostic factor for overall survival of CRC patients.³³ Furthermore, YBX1 is reported as a key inducer of partial EMT and tumorigenicity by elevating the secretion of angiogenic factors such as TGF- β , CSF-1, NGF, VGF, ADAM9, and ADAM17.³⁴ Another study has suggested that overexpression of YBX1 in lung adenocarcinoma cells promotes TGF- β 1-induced EMT via upregulation of the phosphorylation level of Akt.³⁵ In our study, using TargetScan, miRanda, and cBioPortal databases, we predicted YBX1 as a downstream target of PVT1/miR-216a-5p axis in CRC. Subsequently, the results of the luciferase reporter assay validated the relationship

between the YBX1 and the PVT1/miR-216a-5p axis, where the luciferase activity was significantly lower in pmirGLO-YBX1-WT cells as compared to pmirGLO-YBX1-MT, when treated with miR-216a-5p mimics. Therefore, we propose that miR-216a-5p inhibits tumor growth and metastasis by directly regulating YBX1 expression.

Conclusion

In summary, our results revealed that the overexpression of miR-216a-5p might silence YBX1 expression, thus resulting in inhibition of cell proliferation, metastasis, and the EMT pathway of CRC cells. Furthermore, we suggest that PVT1 upregulation can reverse the anti-tumor effect of miR-216a-5p on CRC cells. Therefore, miR-216a-5p might be a promising prognostic and therapy biomarker for CRC.

Availability of data and materials

The datasets used during the present study are available from the corresponding author upon reasonable request.

Acknowledgments

We would like to thank Editage for their English language editing service. This research was supported by Guangzhou science and technology planning project (201804010077).

Disclosure

The authors declare that they have no competing interests in this work.

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Supplementary materials

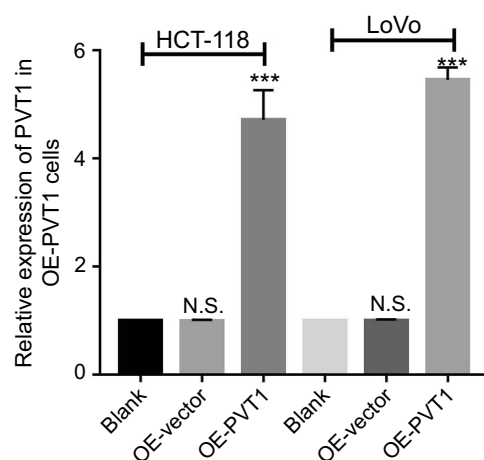


Figure S1 The overexpression efficiency of PVT1 by a lentivirus-based method in LoVo and HCT-116 cell lines.

Note: *** $P < 0.001$, N.S. means not significant.

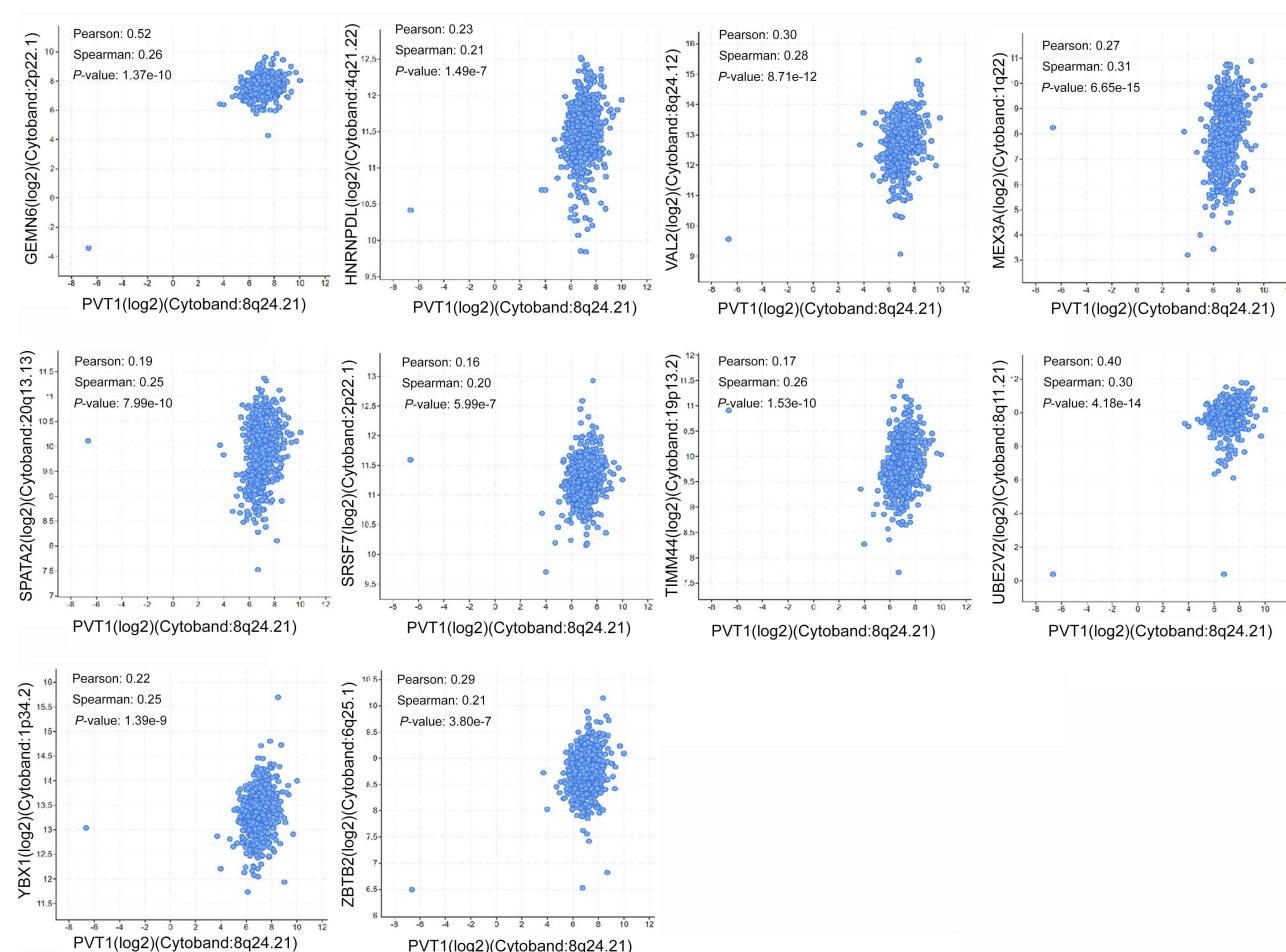


Figure S2 The potential 10 target genes that regulated by PVT1/miR-216a-5p axis in CRC.

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