

PTPN9 promotes cell proliferation and invasion in Eca109 cells and is negatively regulated by microRNA-126

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Abstract. Protein tyrosine phosphatase non-receptor type 9 (PTPN9), also named PTP-MEG2, is an important member of the protein tyrosine phosphatase family that is involved in variety of human diseases. However, the role of PTPN9 in esophageal squamous cell carcinoma (ESCC) remains to be established. The present evaluated the potential effect and underlying mechanism of action of PTPN9 in ESCC. Immunohistochemistry was performed to detect PTPN9 protein expression in 84 ESCC tumor specimens and 30 normal esophageal tissues. The association between positive expression of PTPN9 and clinicopathological features and prognosis was analyzed. The prognostic role of PTPN9 was further investigated using multivariate regression analysis. PTPN9-small interfering RNA and microRNA (miR-126)-mimics were transfected into Eca109 cells to construct PTPN9 silencing and an miR-126 ectopic expression cell model. Reverse transcription-quantitative polymerase chain reaction, western blot analysis, cell counting kit-8, Transwell assays and flow cytometry were used to investigate the role of PTPN9 in the process of ESCC progression and its potential downstream signaling pathway. Immunohistochemical analysis revealed that PTPN9 was upregulated in ESCC tumor specimens compared with normal esophageal tissues. The χ^2 test indicated that positive expression of PTPN9 was correlated with tumor node metastasis stage, tumor classification and node classification. Patients with PTPN9 positive expression had shorter survival time, compared with those that were PTPN9 negative. Multivariate regression analysis with the Cox proportional hazards regression model revealed that PTPN9 expression was a prognostic factor of overall survival for patients with ESCC.

Using RNA interference, the present study demonstrated that knockdown of PTPN9 significantly suppressed cell proliferation and invasion in Eca109. Additionally, it was hypothesized that miR-126, described as a tumor suppressor in ESCC, may act at least in part via its inhibition of PTPN9 at the post-transcriptional level. To the best of our knowledge, this is the first study to demonstrate that PTPN9 is overexpressed in ESCC and associated with poor survival, and may therefore be important in the pathogenesis of ESCC.

Introduction

Globally, esophageal cancer is a highly common malignancy associated with significant mortality, with ~6.5 billion new cases and 400,200 mortalities in 2012 (1). In China, it has been ranked as the fifth and fourth highest tumor type for incidence and mortality, respectively (2). Esophageal squamous cell carcinoma (ESCC) remains the major histological subtype in China, accounting for 90% of newly diagnosed patients (3). When accompanied by low socioeconomic status, active tobacco and alcohol abuse, malnutrition, pulmonary comorbidities and secondary malignancies, the long-term survival rate is poor and requires improvement (4). Despite the improvements in therapeutic strategies, including surgical techniques and induction therapy, prior to or following surgery, biomarkers for tailored multimodal treatment with increased efficacy are required.

Protein tyrosine phosphatase non-receptor type 9 (PTPN9), also termed PTP-MEG2, is highly expressed in the brain, leukocytes and endocrine cells and is a cytoplasmic phosphatase that is hyperactivated in erythroid progenitors (5). PTPN9 is usually present in the cytoplasm and cell membrane of the majority of cells (5,6). It is unique among the protein tyrosine phosphatases due to its N-terminal Sec 14p homology domain, which is able to activate the enzyme of the phosphatase domain of PTPN9 via binding phosphoinositides (7,8). PTPN9 is involved in numerous cellular processes, including cell proliferation, differentiation and migration through the regulation of signaling pathways (9). For example, it promotes secretory vesicle fusion (10), mediates insulin signaling in hepatocytes (11,12), inhibits breast cancer cell growth (13-15) and regulates endothelial cell function (16). However, the role of PTPN9 in ESCC remains to be established.

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MicroRNAs (miRNAs) are a class of highly conserved, non-coding, small single-stranded RNAs of 22-25 nucleotides, which have been reported to serve important roles in a number of types of human disease (17,18). Due to their sequences being complementary to specific sequences in transcripts, miRNAs negatively regulate gene expression by altering mRNA abundance at the post-transcriptional level, as well as allowing for transcription modification. Located at chromosome 9q34.3, an intron of EGF-like domain-containing protein 7 (EGFL7), miRNA (miR)-126 is associated with various human tumors (19). Commonly exhibiting tumor suppressive properties, miR-126 is expressed at a low level in numerous human malignancies, including lung cancer (20), colon cancer (21), breast cancer (22), osteosarcoma (23) and gastric cancer (24). Repression of cancer cell proliferation (25), migration (26) and invasion (27) mediated by miR-126 can be achieved via targeting specific oncogenes, including phosphoinositide 3-kinase, V-Ki-ras2 Kirsten rat sarcoma viral oncogene, EGFL7 and vascular endothelial growth factor (VEGF). In addition, reduced levels of miR-126 are an effective predictor of poor survival in patients with cancer (20,28). A previous study indicated that miR-126 was able to negatively regulate erythropoietic development by targeting PTPN9 (29). However, the role of PTPN9 and the association between miR-126 and PTPN9 in ESCC has, to the best of our knowledge, never been investigated. Therefore, the present study aimed to investigate their potential roles in the development of ESCC.

Materials and methods

Patients and specimens. In total, 84 patients with ESCC were included in the present study. The inclusion criteria were as follows: i) Diagnosis of ESCC with histopathological identification; ii) no treatment received prior to surgery; iii) 7th edition of Union for International Cancer Control tumor-node-metastasis (TNM) classification system (30) was used for staging the tumor; iv) limited or extended surgical history including esophagectomy at the Department of Thoracic Surgery, Henan Tumor Hospital (Zhengzhou, China) between January 2008 and December 2009; v) detailed clinical records were available with follow-up duration extending until May 10th, 2014; vi) tumor specimens were frozen in liquid nitrogen and stored at -80°C following surgical resection, in order to perform immunohistochemistry (IHC). Specimens were obtained with the informed consent of patients. The present study was approved by the Ethics Committee of Henan Tumor Hospital. Post-surgical follow-up was conducted every 3 months for the first 2 years, every 6 months between the 3rd and 6th year, or until patient mortality. The overall survival rate was calculated as the period between surgery and the date of mortality or final observation.

Cell line and cell culture. Obtained from the Cell Bank of the Tumor Hospital of the Chinese Academy of Medical Sciences (Beijing, China), the Eca109 human ESCC cell line was cultured in RPMI-1640 supplemented with 10% fetal bovine serum (FBS; both from Thermo Fisher Scientific, Inc., Waltham, MA, USA) and 100 U/ml penicillin-streptomycin at 37°C in an atmosphere containing 5% CO₂. Cells at the logarithmic growth phase were used for subsequent experiments.

RNA oligonucleotides and cell transfection. For the upregulation of miR-126 expression or the knockdown of PTPN9, RNA oligonucleotides were synthesized by Shanghai GenePharma, Ltd. (Shanghai, China). The sequences of the miR-126 mimics were as follows: Forward, 5'-UCGUACCGUGAGUAAUAAUGCG-3' and reverse, 5'-CAUUAUUACUCACGGUACGAU-3'. The sequences of the PTPN9-siRNA were as follows: Forward, 5'-GUGGACAGUUCAGUACAAUTT-3' and reverse, 5'-AUUGUACUGAACUGUCCACTT-3'. The mock miRNA control sequences were forward, 5'-UUCUCCGAACGUGUCACGUTT-3' and reverse, 5'-ACGUGACACGUUCGGAGATT-3'. Eca109 cells were treated with miR-126 mimics, mock, PTPN9-siRNA or equal amount of PBS (non-treated group) using Lipofectamine[®] 2000 (Invitrogen; Thermo Fisher Scientific, Inc.) in OptiMEM I Reduced Serum media (Thermo Fisher Scientific, Inc.) at a final concentration of 100 pmol/10⁶ cells in all experiments. Cells were incubated with the complexes for 6 h prior to replacement of the medium. A fluorescence microscope was used to evaluate the transfection efficiency.

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR). Total RNA was extracted using TRIzol[®] Reagent (Invitrogen; Thermo Fisher Scientific, Inc.). Reverse transcription and real-time PCR for miRNAs were conducted using the SYBR Green Hairpin-it[™] miRNAs qPCR quantitation kit (Shanghai GenePharma, Ltd.) in 20 μl reaction mixtures, and the reaction was performed on an ABI 7500 real-time PCR system (Applied Biosystems; Thermo Fisher Scientific, Inc.). The DNA was denatured at 95°C for 3 min, followed by 40 amplification cycles consisting of 95°C for 12 sec and 62°C for 40 sec. The primers for quantitative PCR were as follows: hsa-miR-126 forward, 5'-ACAGTTCTCTCGTACCGTGAGTAAT-3'; reverse, 5'-AAAGGTTGATCTGCTCTCTCTCTC-3'; human RNU6 forward, 5'-ATTGGAACGATACAGAGAGATT-3'; reverse, 5'-GGAACGTTACGAATTTG-3'. The gene expression threshold cycle values of miRNAs were calculated by normalizing to the level of human U6 small nuclear RNA in the reaction.

For the analysis of mRNA, the SYBR Green RT-qPCR kit (Shanghai GenePharma, Inc.) was used for cDNA synthesis and qPCR according to the manufacturer's protocol. β-actin was used as an internal control and oligo (dT) was the common primer for reverse transcription. The sequences of the primers used for qPCR are as follows: PTPN9 forward, 5'-ATGTGCTCCGTGCCATAGAATTG-3'; reverse, 5'-GAGGATCTGAGAACGAAGAGGTTCC-3'; β-actin forward, 5'-TCTGGCACCACACCTTCTAC-3'; reverse, 5'-GATAGCACAGCC TGGATAGC-3'. The thermocycler settings were the same as described previously. The 2^{-ΔΔC_q} method (31) was used for data analysis.

IHC and western blot analysis. Immunohistochemistry was performed as previously described (32). Rabbit anti-human PTPN9 antibody (dilution, 1:50; cat. no., sc-130859; Santa Cruz Biotechnology, Inc., USA) was used as the primary antibody, followed by a peroxidase-conjugated goat anti-rabbit secondary antibody (dilution, 1:200; cat. no., AB10058; Sangon Biotech Co., Ltd., Shanghai, China). Any intensity of cell membrane and cytoplasmic staining was considered a positive stain for PTPN9. Positive expression of protein was considered if the

Table I. Expression of PTPN9 protein in patients with ESCC.

Group	No. of patients	PTPN9 expression		P-value
		Positive (%)	Negative (%)	
Tumor tissue	84	31 (36.9)	53 (63.1)	0.041
Normal tissue	30	5 (16.7)	25 (83.3)	

PTPN9, protein tyrosine phosphatase, non-receptor type 9; ESCC, esophageal squamous cell carcinoma

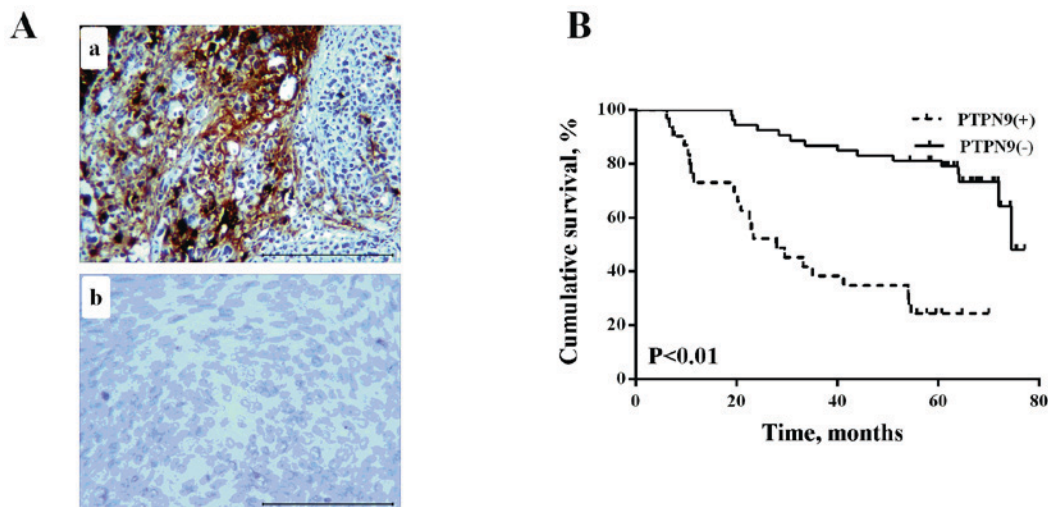


Figure 1. PTPN9 protein expression in ESCC tissues compared with normal esophageal tissues and the association with prognosis in patients with ESCC. (A) Representative immunostaining images of PTPN9 in ESCC specimens and normal esophageal tissues (magnification, x200). a, ESCC specimens; b, normal esophagus tissues. Scale bar, 100 μ m. (B) Kaplan-Meier survival analysis of patients with ESCC. Patients with PTPN9-positive tissues had a shorter survival time compared with those with PTPN9-negative tissues ($P < 0.01$). PTPN9, protein tyrosine phosphatase, non-receptor type 9; ESCC, esophageal squamous cell carcinoma.

percentage of stained cells was $\geq 10\%$. IHC results were further evaluated at high-power magnification (x200) once regions containing positive immunoreactivity were identified with low-power magnification (x40) using an optical microscope (BX41; Olympus Corporation, Tokyo, Japan). The section was examined by two pathologists individually.

Protein lysates were lysed on ice in cold radioimmuno-precipitation assay buffer containing a protease inhibitor cocktail (Pierce; Thermo Fisher Scientific, Inc.), according to the manufacturer's protocol. The protein concentration was determined using a bicinchoninic acid assay with a BCA protein assay kit (Beyotime Institute of Biotechnology, Haimen, China), equivalent amounts of protein (34 μ g) were separated by SDS-PAGE on a 10% gel and blotted onto polyvinylidene fluoride membrane (Bio-Rad Laboratories, Inc., Hercules, CA, USA). Following blocking with 5% non-fat milk in Tris-buffered saline containing 0.1% Tween-20 for 2 h at room temperature, the membranes were probed with primary antibodies against PTPN9, as previously described (dilution, 1:200), and β -actin (dilution, 1:5,000; cat. no., ab8227; Abcam, Cambridge, UK) at 4°C overnight. The bands were visualized using an enhanced chemiluminescence western blotting substrate (Thermo Fisher Scientific, Inc.), following incubation with the previously described secondary antibody (dilution,

1:5,000) for 2 h at room temperature. The internal control β -actin was used for normalization. Protein bands were quantified using FluorChem FC3 AlphaView software (version 2.0; ProteinSimple, San Jose, CA, USA).

Prediction of miRNA targets. PicTar (<http://pictar.mdc-berlin.de/>), Targetscan (<http://www.targetscan.org/>) and Microcosm Targets (<http://www.ebi.ac.uk/enright-srv/microcosm/htdocs/targets/v5/>) were searched to predict potential targets for miR-126. The 3'-untranslated region of PTPN9 mRNA (RefSeq NM_002833) was identified as containing a putative miRNA-126 binding site.

Cell proliferation assay. The miR-126 mimic, mock control and PTPN9-siRNA were transfected into Eca109 cells at a concentration of 100 pmol/ 10^6 cells. A total of 24 h later, following trypsinization, cells were counted and seeded into 96-well plates (5×10^3 cells/well) in quadruplicate. Cell proliferation was monitored at 5, 24, 48, 72 and 96 h following transfection using the Cell Counting Kit-8 (Dojindo Molecular Technologies, Inc., Kumamoto, Japan), according to the manufacturer's protocol. The number of cells was evaluated through measurement of absorbance at 450 nm using a Wellsan MK-3 (Labsystems Dragon, Helsinki, Finland).

Table II. Association between PTPN9 Expression and clinicopathological features in patients with ESCC.

Features	No of patients	PTPN9 expression		P-value ^a
		Positive, n=31 (%)	Negative, n=53 (%)	
Age, years				
≤60	49	16 (32.7)	33 (67.3)	0.339
>60	35	15 (42.9)	20 (57.1)	
Sex				
Male	58	21 (36.2)	37 (63.8)	0.834
Female	26	10 (38.5)	16 (61.5)	
Tumor location				
Upper	12	5 (41.7)	7 (58.3)	0.418
Middle	58	23 (39.7)	35 (60.3)	
Lower	14	3 (21.4)	11 (78.6)	
Maximum tumor size				
≤3	51	22 (43.1)	29 (56.9)	0.141
>3	33	9 (27.3)	24 (72.7)	
Differentiation				
Poor	28	15 (53.6)	13 (46.4)	0.058
Moderate	37	12 (32.4)	25 (67.6)	
Well	19	4 (21.1)	15 (78.9)	
T classification				
T1+T2	39	9 (23.1)	30 (76.9)	0.014
≥T3	45	22 (48.9)	23 (51.1)	
N classification				
Yes	27	16 (59.3)	11 (40.7)	0.003
No	57	15 (26.3)	42 (73.7)	
TNM stage				
I+II	64	17 (26.6)	47 (73.4)	0.001
>III	20	14 (70.0)	6 (30.0)	
Postoperative chemotherapy				
Yes	34	11 (32.4)	23 (67.6)	0.476
No	50	20 (40.0)	30 (60.0)	
Postoperative recurrence				
Yes	9	4 (44.4)	5 (55.6)	0.896
No	75	27 (36.0)	48 (64.0)	

^aStatistical analysis was performed using χ^2 analysis. PTPN9, protein tyrosine phosphatase, non-receptor type 9; ESCC, esophageal squamous cell carcinoma.

Transwell cell invasion assay. Transwell inserts coated with Matrigel (Corning Incorporated, Corning, NY, USA) were used for a cell invasion assay. Cells transfected with miR-126 mimics, mock controls and PTPN9-siRNA for 24 h were detached and resuspended in serum-free medium. A 200 μ l suspension containing 5×10^4 cells was added to the upper insert. RPMI-1640 containing 20% FBS was added to the lower wells in the 6-well cell culture plate as a chemoattractant. Following a 24-h incubation, cells were fixed in 4% paraformaldehyde for 15 min and stained with 1% crystal violet for 10 min at room temperature. The cells on the upper surface of filter were wiped off with a cotton swab and invaded cells on

the lower surface of the membrane were visualized with an optical microscope in 5 fields of view at x200 magnification.

Cell apoptosis analysis. At 48 h following transfection, cells were washed and resuspended in binding buffer at a final concentration of 1×10^6 cells/ml, and 100 μ l cell suspension was incubated with 10 μ l Annexin V-fluorescein isothiocyanate (FITC) and 5 μ l propidium iodide (PI) in the dark for 15 min using the Annexin V-FITC kit (Miltenyi Biotec GmbH, Bergisch Gladbach, Germany), according to the manufacturer's protocol. The stained cells were analyzed using flow cytometry (FACSCalibur™; BD Biosciences, Franklin Lakes,

Table III. Multivariate analysis of PTPN9 expression status with regard to OS in patients with ESCC.

Variables	OS				
	β	SE	Wald	HR (95% CI)	P-value
Differentiation	-0.734	0.285	6.610	0.480 (0.274-0.840)	0.010
TNM stage	1.536	0.367	17.487	4.646 (2.261-9.543)	0.001
PTPN9, positive vs. negative	1.087	0.388	7.867	2.967 (1.388-6.342)	0.005
Postoperative chemotherapy	1.102	0.511	4.464	3.010 (1.105-8.196)	0.031
Postoperative recurrence	-1.465	0.432	11.497	0.231 (0.099-0.539)	0.001

SE, standard error; HR, hazard ratio; CI, confidence interval; OS, overall survival; TNM, tumor-node-metastasis; PTPN9, protein tyrosine phosphatase, non-receptor type 9; ESCC, esophageal squamous cell carcinoma.

NJ, USA). Annexin V-FITC-positive and PI-negative staining was indicative of cells undergoing early apoptosis, and the percentage of apoptotic cells of each group was compared.

Statistical analysis. Quantitative data were expressed as the mean \pm standard deviation and were statistically analyzed by t-test. Pearson's χ^2 test was used to analyze the association between PTPN9 expression levels and the clinicopathological features of ESCC specimens. Survival curves were calculated using the Kaplan-Meier method, and compared using the Log-rank test. The influence of each clinicopathological parameter on survival was assessed through multivariate regression analysis with the Cox proportional hazards regression model. A Wald test was used to test the association of each variable in Table II with overall survival. All data were analyzed using SPSS software (17.0; SPSS, Inc., Chicago, IL, USA). $P < 0.05$ was considered to indicate a statistically significant difference.

Results

PTPN9 overexpression in ESCC specimens. A total of 84 ESCC specimens were obtained for an IHC assay to analyze the expression of PTPN9 in ESCC. The data revealed that positive PTPN9 staining was observed in 36.9% (31/84) of tumor tissues, and in 16.7% of normal specimens (5/30; Table I), suggesting that the positive rate of PTPN9 in tumor specimens was increased compared with healthy tissues (Fig. 1A).

PTPN9 protein expression and clinicopathological features of ESCC. The association between PTPN9 expression levels and patient clinicopathological features is presented in Table II. According to the expression levels of PTPN9 protein, clinical data were divided into two groups, the PTPN9-positive and PTPN9-negative staining group. As presented in Table II, PTPN9 was overexpressed in patients with increased T classification ($P = 0.014$), N classification ($P = 0.003$) and TNM stage ($P < 0.001$). No significant association was detected between PTPN9 expression levels and other clinicopathological features, including patient age, sex and tumor location (Table II).

PTPN9 expression is negatively correlated with the survival time of patients with ESCC. The median follow-up time was

60.7 months (range, 6.1-77.17), Kaplan-Meier analysis survival curves revealed that patients with PTPN9-negative expression had a longer overall survival compared with PTPN9-positive group (log-rank test, $P < 0.01$; Fig. 1B). The results of multivariate Cox proportional hazards regression analysis demonstrated that PTPN9 protein expression, differentiation, TNM stage, postoperative chemotherapy and postoperative recurrence were independent prognostic markers for patients with ESCC (Table III).

PTPN9 protein expression level, and not mRNA level, is altered with the upregulation of miR-126 expression. As PTPN9 was identified as a putative target for miR-126 (Fig. 2A), it was assessed whether miR-126 regulates endogenous PTPN9 expression by transfecting an miR-126 mimic into Eca109 cells. miR-126 and PTPN9 mRNA expression levels were assessed with RT-qPCR. Following transfection, the expression level of miR-126 was ~ 9 -fold higher compared with the NC group (Fig. 2B). A decreasing trend in PTPN9 mRNA expression level in cells transfected with miR-126 mimics was observed; however, it was not statistically significant (Fig. 2C).

The expression of PTPN9 protein level was also observed in Eca109 cells transfected with miR-126 mimics. An increase of miR-126 level was significantly associated with a decrease in PTPN9 protein expression level as determined by western blot ($P < 0.05$; Fig. 2D). No significant differences were observed in the expression of PTPN9 mRNA, miR-126 and PTPN9 protein expression between the mock and NC groups (Fig. 2B, C and D). Taken together, these results indicated that PTPN9 was targeted by miR-126 and the expression of PTPN9 in Eca109 cells was negatively regulated by miR-126.

PTPN9 depletion inhibits cell proliferation and invasion in Eca109 cells, but has no effect on apoptosis. PTPN9-siRNA was used for inhibiting the expression of PTPN9 to assess its effect on ESCC cells. In the PTPN9-siRNA group, the expression level of PTPN9 was $\sim 50\%$ lower compared with the NC group, as detected using RT-qPCR (Fig. 2C). CCK-8 and Transwell assays as well as flow cytometry were used to examine the impacts of PTPN9 on cell proliferation, invasion and apoptosis in Eca109 cells. The results revealed that the knockdown of PTPN9 significantly inhibited cell proliferation at 48 ($P < 0.05$), 72 ($P < 0.01$) and 96 h ($P < 0.01$) after transfection

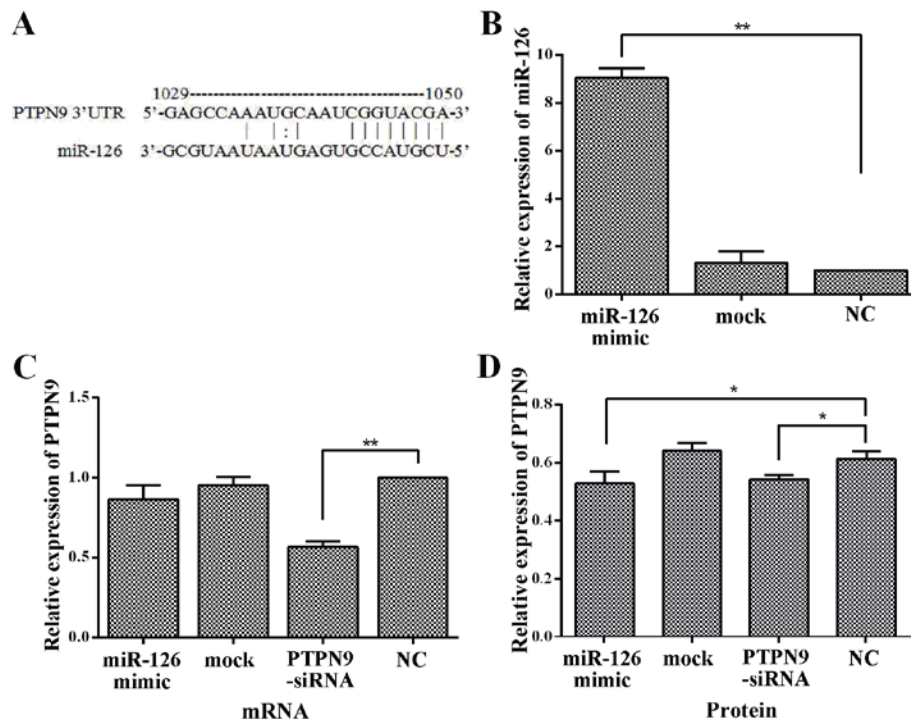


Figure 2. PTPN9 as a potential target for miR-126. (A) Predicted complementary sequences between miR-126 and PTPN9. (B) Detection of the relative expression of miR-126 in Eca109 cells following transfection with miRNA-126 mimics and mock controls. (C) Reverse transcription-quantitative polymerase chain reaction analysis of PTPN9 mRNA in Eca109 cells following transfection with PTPN9-siRNA, mock control or miR-126 mimics. $^{**}P < 0.01$. (D) Quantification of the western blot analysis of PTPN9 protein levels in Eca109 cells following transfection with miR-126 mimics, mock control or PTPN9-siRNA. $^{*}P < 0.05$. PTPN9, protein tyrosine phosphatase, non-receptor type 9; miR-126, microRNA-126; NC, negative control; siRNA, small interfering RNA.

(Fig. 3A). A decrease in the number of cells migrating through the Matrigel was also observed (Fig. 3B and C). However, no significant difference in apoptotic rate was identified between any two groups (Fig. 3D and E). These results indicated that PTPN9 may serve an important role in the cell proliferation and invasion of ESCC.

Discussion

Esophageal cancer is a global health challenge, with 5-year overall survival ranging between 15 and 35% (33). Therefore, novel molecular markers used to predict the progression and prognosis of ESCC are required. In the last few years, a number of studies have revealed that a variety of human diseases are associated with aberrant expression of PTPN9 (34,35). However, the role of PTPN9 in ESCC remains to be elucidated. The present study revealed that the protein expression rate of PTPN9 is increased in ESCC specimens compared with healthy esophageal tissues using IHC analysis. Furthermore, the protein levels of PTPN9 were associated with various clinicopathological parameters, including T classification, N classification and TNM stage. Multivariate Cox regression analyses revealed that PTPN9 is an independent prognostic predictor for patients with ESCC.

A previous study indicated that knockdown of PTPN9 in the liver of diabetic mice was able to lead to insulin sensitization and normalization of hyperglycemia (12). In addition, it has been demonstrated that PTPN9 is able to negatively regulate the VEGF-induced cell signal through inhibition of the phosphorylation of VEGF receptor 2 on Tyr1175 in endothelial

cells (16). Furthermore, it has been reported that PTPN9 is able to inhibit ErbB2 and epidermal growth factor receptor (EGFR) signaling by dephosphorylating ErbB2/EGFR to impair growth and invasiveness in breast cancer cells (13), supported by the results of Du *et al* (15). In the present study, PTPN9 was observed to be overexpressed in ESCC specimens, compared with healthy esophageal tissues, indicating that PTPN9 may be an oncogene during the development of ESCC. To evaluate this hypothesis, the PTPN9 depletion cell model was constructed following transfection of PTPN9-siRNA into the Eca109 ESCC cell line. As expected, cell proliferation and invasion was significantly suppressed upon the knockdown of PTPN9.

A number of previous studies have demonstrated that aberrant expression of miRNAs serves a key role in the tumorigenic process (26,36). Therefore, an improved understanding of the underlying mechanisms of these non-coding RNAs may improve technologies for the diagnosis and treatment of human diseases. A previous study reported that miR-126 was able to regulate PTPN9 in the hematopoietic differentiation of human embryonic stem cells at the post-transcriptional level (29). Frequently described as a tumor suppressor in a number of studies, miR-126 was observed to be downregulated in ESCC tissues and cell lines (32,37-39). However, the interaction between miR-126 and PTPN9 in the process of ESCC remains to be established. In the present study, using the miRNA target prediction program, a putative miR-126 binding site was identified within the 3'UTR of PTPN9, suggesting that PTPN9 may be a target of miR-126. The present study also revealed that ectopic expression of miR-126 reduced the levels of PTPN9,

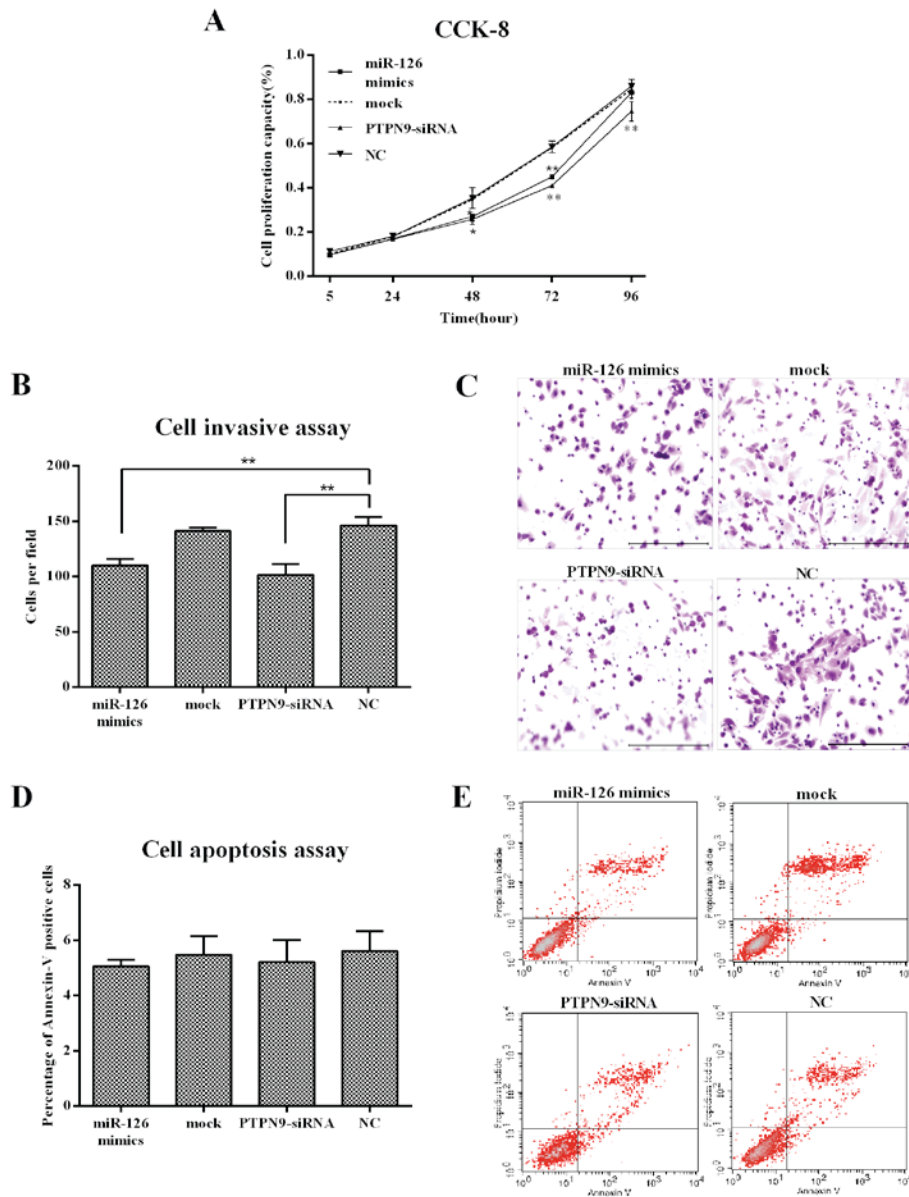


Figure 3. Knockdown of PTPN9 inhibited ESCC cell proliferation and invasion, but not apoptosis. (A) CCK-8 analysis of Eca109 cell growth following transfection with PTPN9-siRNA, mock control or miR-126 mimics. * $P < 0.05$, ** $P < 0.01$ compared with NC. (B) Quantification of invasion data. ** $P < 0.01$. (C) Transwell analysis of Eca109 cell invasion following transfection with PTPN9-siRNA, mock control or miR-126 mimics (stained with crystal violet). Scale bar, 100 μm . (D) Quantification of apoptosis data. (E) Apoptotic analysis of Eca109 cells using flow cytometry following transfection with PTPN9-siRNA, mock control and miR-126 mimics. miR-126, microRNA-126; ESCC, esophageal squamous cell carcinoma; PTPN9, protein tyrosine phosphatase, non-receptor type 9; CCK-8, cell counting kit-8; siRNA, small interfering RNA; NC, negative control; UTR, untranslated region.

and inhibited the growth and invasion of Eca109 cells, similar to the results from the PTPN9-siRNA group. These results indicate that miR-126 may be an important component of the signal pathway induced by PTPN9 in ESCC.

However, the present study has certain limitations. The sample size of patients with ESCC was relatively small; therefore, studies with a larger sample size may be conducted in order to further identify the prognostic impact of each clinicopathological factor on survival time at various clinical stages. In addition, a dual-luciferase reporter assay may be used to examine the direct targets of PTPN9.

In conclusion, the present study demonstrated that PTPN9 expression levels are associated with T classification,

N classification and TNM stage, and may represent a useful prognostic marker for patients with ESCC. Through being regulated by miR-126, downregulation of PTPN9 may inhibit growth and decrease the invasive capacity of ESCC cells, indicating that PTPN9 may be a promising molecular therapeutic target for ESCC in the future.

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