Insulin receptor substrate-1 and Golgi phosphoprotein 3 are downstream targets of miR-126 in esophageal squamous cell carcinoma

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Abstract. Esophageal squamous cell carcinoma (ESCC) is a common histologic subtype in China. It has been suggested that abnormal expression of microRNAs (miRNAs) is associated with carcinogenesis. We investigated miR-126 expression and its potential targets in ESCC. The expression of miR-126 was detected in cancerous and paired paracancer tissues from 102 patients with ESCC. Target analysis of miR-126 was predicted using online tools. The effect of miR-126 expression on target proteins was assessed using miR-126 mimics or miR-126 inhibitors in ESCC cell lines. In addition, the impact of miR-126 on cell proliferation, apoptosis, migration and invasion was detected in ESCC cell lines. The expression of miR-126 was significantly lower in ESCC tissues, which was associated with tumor differentiation, lymph node metastasis, tumor in-depth and TNM stage. Insulin receptor substrate-1 (IRS-1) and Golgi phosphoprotein 3 (GOLPH3) were overexpressed in ESCC. Overexpression of IRS-1 was associated with cell differentiation, whereas GOLPH3 was related to lymph node metastasis, tumor invasion in-depth and TNM stage in ESCC patients. miR-126 mimics downregulated the expression of IRS-1 and GOLPH3 protein and suppressed the proliferation, migration and invasion of ESCC cells, whereas miR-126 inhibitors led to the opposite results. miR-126 suppressed the proliferation, migration and invasion of ESCC cells, and acted as a tumor suppressor in the carcinogenesis of ESCC. IRS-1 and GOLPH3 are downstream targets of miR-126 at the post-transcriptional level in ESCC.

Introduction

Esophageal cancer is the eighth most common cancer worldwide and the sixth most common cause of cancer-related mortality (1,2). Although Barrett's adenocarcinoma is the most rapidly increasing cancer in Western countries, esophageal squamous cell carcinoma (ESCC) remains dominant in East Asia (3). Often diagnosed at later stages, the prognosis of affected patients is unsatisfactory despite the development of therapeutic options such as surgery, chemotherapy and radiotherapy (4). Consequently, there is a great need for ESCC biomarkers to allow a tailored multimodal approach with increased efficacy. Nevertheless, efforts to identify molecular markers in association with the pathogenesis of ESCC have, to date, been unsuccessful (5).

Mature microRNAs (miRNAs) are a class of small, well-conserved, non-coding RNA molecules that silence gene expression usually by interfering with mRNA stability or protein translation (6,7). The function is performed by identifying 3'-UTRs (untranslated regions) of target mRNAs with conserved complementarities to the seed (nucleotides 2-7) of the miRNA. Up to 30% of human protein-coding genes may be regulated by miRNAs (8,9). miRNAs are involved in biological and pathological processes including cell differentiation, proliferation, apoptosis and metabolism, and they are emerging as highly tissue-specific biomarkers with potential clinical application for defining cancer types and origins. miRNAs can either function as oncogenes or tumor suppressors (10,11).

It has been reported that miR-126 is located on chromosome 9q34.3 within the host gene encoding for epidermal growth factor like-7 (EGF-L-7), an endothelial cell-derived, secreted inhibitor of smooth muscle cell migration and a regulator of blood vessel formation (12,13). Vasculature is required for the expansion of tumor masses, as inhibition of new vessel formation prevents tumor growth. Studies have shown that the expression of miR-126 is downregulated in cancers such as non-small cell lung cancer and colon cancer (14,15), suggesting that the downregulation of miR-126 is significantly related to the occurrence and development of...
cancer. However, studies have also reported that aberrant overexpression of miR-126 contributes to carcinogenesis, suggesting that miR-126 acts as an oncogene in gastric cancer and oral squamous cell carcinoma (16,17). Therefore, miR-126 may have diverse roles in tumor carcinogenesis. To date, some genes have been identified as miR-126 target genes, including sex determining region Y-box 2 (SOX2), Crk, vascular endothelial growth factor (VEGF), matrix metalloproteinase-9 (MMP-9), insulin receptor substrate-1 (IRS-1), SLC7A5, TOM1, and CD97/G-protein-coupled receptor (GPCR) (16,18-21). Bioinformatics has also shown that the 3′-UTR of genes such as IRS-1, FBXO33, PTPN9, PLK2, and Golgi phosphoprotein 3 (GOLPH3) contains a putative binding site for miR-126.

However, the functional role and mechanism of miR-126 in regulating tumorigenesis of ESCC is still not fully understood. To identify miRNAs which can be specifically expressed and exert distinct biological actions in ESCC, we investigated miR-126 expression and its functional role in ESCC. miR-126 expression was measured in ESCC tissue specimens and paired paracancer tissues, and the relationship between the expression level of miR-126 and the clinicopathological characteristics were analyzed. Further functional studies of miR-126 showed that it could inhibit ESCC cell growth, migration and invasion. IRS-1 and GOLPH3 are potential downstream targets of miR-126 in ESCC.

Materials and methods

Tissue specimens and cancer cell lines. Specimens of 102 ESCC tissues and paired paracancer tissues were collected from Henan Cancer Hospital between July 2012 and July 2013. Clinicopathological data of patients were collected. None of the patients received preoperative antitumor therapy. All specimens were collected using two methods: one was snap frozen in liquid nitrogen immediately after surgery and stored at -80°C for further miRNA detection, the other was formalin-fixed for histopathological assay. The clinicopathological information of the ESCC patients is shown in Table I. All patients provided informed consent under a protocol reviewed and approved by the Institutional Review Board, Henan Cancer Hospital.

The human ESCC cell line Eca9706 was purchased from the cell bank of the tumor hospital of the Chinese Academy of Medical Sciences/Biological Detection Center (Beijing, China). The cells were cultured in RPMI-1640 (Thermo, Waltham MA, USA) supplemented with 10% fetal bovine serum (FBS; Thermo) and penicillin/streptomycin in a 5% CO₂ humidified incubator at 37°C (normal conditions for cell culture).

RNA isolation and real-time reverse transcription polymerase chain reaction (qRT-PCR). Total-RNA in cancer tissues or cell lines was isolated using QIAzol lysis reagent and miRNeasy mini kit (Qiagen, Düsseldorf, Germany), according to the manufacturer’s instructions. Total-RNA was isolated 48 h after cell transfection. Quantitation of total-RNA was carried out using spectrophotometry. All samples had an excellent 260/280 ratio. RNAs were reverse-transcribed into cDNAs using miScript Reverse Transcription kit (Qiagen) and stored at -20°C for immediate or further use. The qRT-PCR of miRNA and mRNA expression was detected using miScript SYBR-Green PCR kit (Qiagen) on an ABI 7500 detector (Applied Biosystems). RUN6 was used as internal control according to the protocol. miR-126 miScript Primer assay (no. MS00003430), RNU6 miScript Primer assay (no. MS00033740), IRS-1 QuantiTect Primer assay (QT00074144), GOLPH3 QuantiTect Primer assay (no. QT00061320), and β-actin QuantiTect Primer assay (no. QT00095431) for miRNA or mRNA detection were commercially provided by Qiagen. Relative gene expression determinations were made using the comparative ΔΔCt method (2^ΔΔCt). Patients were divided into two groups (high or low miR-126 expression) for further analysis. The high miR-126 expression group was defined as an expression level of miR-126 in tumor tissue higher than the paired paracancer tissue.

Online prediction of miRNA targets. To predict potential targets of miR-126, we searched PicTar (http://www.picTar.org), miRGenTargets (http://microrna.gr) and TargetScanHuman (http://www.targetscan.org/vert_61/). The 3′-UTR of IRS-1 mRNA (RefSeq 005544) and GOLPH3 mRNA (RefSeq 022130) may have putative miRNA-126 binding sites.

Immunohistochemistry. To explore expression of IRS-1 and GOLPH3 proteins in ESCC, immunohistochemistry experiments were carried out on the collected 3-5 µm tissue slices. After deparaffinization and antigen retrieval, slices were incubated at 4°C overnight with rabbit polyclonal GOLPH3 antibody (1:100; Abcam, Hong Kong), or rabbit polyclonal IRS-1 antibody (1:50; Santa Cruz Biotechnology, Inc., Santa Cruz, CA, USA; sc-559). After washing with phosphate-buffered saline (PBS) three times, the slides were incubated with biotinylated secondary antibody (1:80; goat anti-rabbit IgG, ready-to-use; Dako, Denmark) for 1 h at 37°C, washing with PBS three times and chromogenic use of 3,3′-diaminobenzidine (DAB) for 5 min. Counterstaining was carried out with hematoxylin. The immunohistochemically stained tissue sections were reviewed and scored separately by two pathologists blinded to the clinical information. Any intensity of cell membrane, endosome or cytoplasmatic staining was considered a positive stain for GOLPH3, whereas cytoplasmic or nuclear staining was considered positive for IRS-1. Increased expression of proteins was considered if the percentage of stained cells was ≥10%.

Transfection of miR-126 mimics or inhibitor into ESCC cells. Eca9706 cells were seeded into 24-well plates and transfected when the cells reached 60% confluence. Syn-hsa-miR-126-3p miScript miRNA Mimic (no. MSY0000445), anti-hsa-miR-126-3p miScript miRNA Inhibitor (no. MIN0000445), and miScript miRNA Inhibitor Negative Control (mock sequence, no. 1027271), commercially provided by Qiagen, were transfected into Eca9706 cells. siRNAs were allowed to form transfection complexes with HiPerFect Transfection Reagent (Qiagen) in serum-free Opti-MEM, according to the manufacturer’s protocol. All groups were performed in triplicate. Cells were collected for RNA or protein assay at different time points after transfection.

For the invasion and apoptosis assays, cells were seeded in 6-well plates and the transfected cells were suspended using tyrode; a hemocytometer was used for cell counting.
Western blot analysis. Transfected cells were lysed in RIPA buffer (Sigma, St. Louis, MO, USA) with protease inhibitors. Proteins were resolved in 10% SDS-PAGE gel and electro transferred onto a PVDF membrane, blocked with 5% milk, and then the membrane was incubated with a primary antibody at 4°C overnight (rabbit polyclonal GOLPH3 antibody, 1:500; Abcam, or rabbit polyclonal IRS-1 antibody; 1:500; Santa Cruz). After washing with Tris-buffered saline 3 times, corresponding horseradish peroxidase-conjugated secondary antibody (1:2,000; goat anti-rabbit IgG; Dako) was used to incubate the membrane at room temperature for 2 h. The membrane was then washed 3 times and the blots were visualized using Western Lightning Ultra (PerkinElmer, Waltham, MA, USA). Bands were quantified using FluorChem FC3 Software® (ProteinSimple, San Francisco, CA, USA) for band intensity and β-actin was used as internal control.

Cell proliferation assay. The cells transfected with 50 nM miR-126 mimics, inhibitors or mock sequence were initially seeded at a density of 5.0x10^3 cells/well in 96-well plates with siRNAs. After 6-8 h, non-adherent cells were removed by washing with PBS. The remaining adherent cells were cultured under normal conditions, and this time point was set as the start point. Cell counting kit-8 (CCK-8) analyses were performed at 5, 24, 48, 72 or 96 h for cell proliferation observation. Absorbance was detected at 490 nm in an enzyme microplate reader (Wellscan MK-3; Labsystems Dragon, Finland).

Table I. Expression of miR-126, IRS-1 protein and GOLPH3 protein in ESCC patients.

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<th>P-value</th>
<th>IRS-1-positive</th>
<th>P-value</th>
<th>GOLPH3-positive</th>
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Relationship between expression levels of miR-126, IRS-1 protein, GOLPH3 protein and clinicopathological characteristics in 102 patients with ESCC analyzed using the Fisher's exact test. miR-126 high is defined as an expression level of miR-126 in tumor tissue higher than the paired paracancerous tissue. IRS-1- or GOLPH3-positive are defined as the percentage of immunoreactive cells ≥10%.
Let al: IRS-1 AND GOLPH3 ARE DOWNSTREAM TARGETS OF miR-126

**Cell apoptosis assay.** Cells were suspended in 0.25% trypsin without EDTA and harvested at 48 h after transfection. After washing twice with PBS, the cell concentration was adjusted to 1x10^6 using Annexin V-FITC binding buffer. Then, 100 µl of the suspended cells were stained with 5 µl of Annexin V for 30 min and 10 µl of propidium iodide (4Abio, Beijing, China) for 15 min in the dark at room temperature. After washing with PBS, 400 µl of PBS was added to the samples and analysis was performed using a flow cytometer (BD Biosciences, San Jose, CA, USA). The percentage of Annexin V-positive cells was recorded for analysis.

**Cell migration and invasion assay.** The Transwell invasion system (8.0 µm polycarbonate membrane, 6.5 mm insert in a 6-well plate; Corning, Lowell, MA, USA) with or without Matrigel was used for the invasion or migration assay. After transfection for 24 h, cells were harvested using 0.25% trypsin-EDTA (Thermo). For the migration assay, 2x10^6 cells were added into the upper compartment of the Transwell chamber. For the invasion assay, 1x10^5 cells were plated on chambers preloaded with Matrigel. In both assays, all cells were suspended in 100 µl of RPMI-1640 without serum, 1,000 µl RPMI-1640 supplemented with 10% FBS was added in the bottom compartment of the chamber. After 12 h of incubation, Matrigel, dead cells and non-migrated or non-invaded cells on the upper surface of the membrane were removed with a cotton swab. The membranes were fixed with 4% paraformaldehyde and stained with 1% crystal purple. The invasive cells that were stuck to the lower surface of the membrane were then counted.

**Statistical analysis.** Statistical analysis was performed using SPSS 17.0 software (SPSS, Inc., Chicago, IL, USA). Data obtained from qRT-PCR were log transformed. Differences in miR-126 expressions between ESCC and paired paracancer tissues were analyzed using non-parametric test. The two-sided Fisher's exact test was used to test the relationship between miR-126, IRS-1 and GOLPH3 expression and clinicopathological variables. Quantitative data expressed as means ± standard deviation and differences between groups were calculated with the t-test. All results were considered statistically significant when P<0.05.

**Results**

**Downregulation of miR-126 expression and association with lymph node metastasis, tumor in-depth or cell differentiation in ESCC.** To investigate the expression of miR-126 in ESCC, we first evaluated the status of miR-126 in samples of ESCC and paracancer tissues. Lower expression of miR-126 was found in 66 (64.7%) patients. The expression of miR-126 in ESCC samples was on average decreased ~68.36% of that in paracancer samples (P<0.05; Fig. 1A and B). To further investigate the relationship between the expression level of miR-126 and clinicopathological characteristics, all patients were divided into two groups. Lymph node metastasis, tumor in-depth, cell differentiation and TNM stage were all associated with miR-126 expression. Greater possibility of lymph node metastasis was observed in the miR-126 low expression group. There were no statistically significant associations between miR-126 expression and other clinicopathological parameters, such as gender, age, tumor location, alcoholism or smoking (P>0.05; Table I).

**IRS-1 and GOLPH3 are potential targets of miR-126 predicted online.** Since miR-126 was closely related to clinicopathological characteristics in ESCC, we further analyzed its function and molecular mechanism in the carcinogenesis of ESCC. We predicted targets of miR-126 in humans using the online tools miRGenTargets, TargetScanHuman and PicTar. Following computational prediction, the proteins IRS-1, FBXO33, PLK2, PTPN9, and GOLPH3 were analyzed. We reviewed published reports that focused on individual miRNAs of the proteins. Among them, IRS-1 has been reported as a target of miR-126 in colon cancer, whereas GOLPH3 shows overexpression in ESCC and is associated with poor prognosis. Therefore IRS-1 and GOLPH3 were selected as targets of our research (Fig. 2A).

**IRS-1 and GOLPH3 are overexpressed in human ESCC.** To verify our prediction, we first studied the relationship between the two proteins in ESCC. IRS-1 and GOLPH3 were determined using immunohistochemistry in ESCC tissues and the paracancer tissues respectively. The expression of IRS-1
protein was significantly higher in ESCC samples compared with paracancer tissues, and the same results were found for GOLPH3 expression levels (P<0.01; Fig. 2B).

We also analyzed the relationship between IRS-1 or GOLPH3 protein and clinical characteristics. Positive expression of IRS-1 was associated with cell differentiation but no obvious association with any other clinical characteristics. Meanwhile, GOLPH3 was closely related to lymph node metastasis, tumor invasion in-depth and TNM stage (Table I).

**IRS-1 and GOLPH3 protein expression after up/down regulation of miR-126.** To further explore the impact of aberrant expression of miR-126 on IRS-1 and GOLPH3, we transfected miR-126 mimics, inhibitors or mock sequence into Eca9706 cells. miR-126 levels increased on an average 5.17-fold (P<0.01) after transfection with miR-126 mimics and decreased 0.37-fold (P<0.01) when transfected with miR-126 inhibitors. There was no obvious change in cells transfected with the mock sequence (Fig. 3A).

Initially, relative mRNA levels of IRS-1 and GOLPH3 in Eca9706 cells were detected. Then, 50 nM of miR-126 mimics, inhibitors or mock sequence were transfected into Eca9706 cells. After transfection for 48 h, we detected the mRNA levels of IRS-1 and GOLPH3 in these cells. In cells transfected with miR-126 mimics, altered relative mRNA levels of IRS-1 (1.05-fold) and GOLPH3 (0.94-fold) had no statistical difference compared with non-treated controls. Downregulation of miR-126 in Eca9706 cells showed no significant difference in IRS-1 (0.92-fold) and GOLPH3 (0.92-fold) mRNA levels, and there were also no differences in the mock sequence group (P>0.05; Fig. 3B).

We then analyzed the relationship between miR-126 and IRS-1 or GOLPH3 protein. The expression level of IRS-1 protein decreased 0.71-fold (P<0.05) compared with the non-treated control, and GOLPH3 protein decreased 0.66-fold (P<0.01) when transfected with miR-126 mimics. In addition, when transfected with miR-126 inhibitors, IRS-1 protein expression increased 2.01-fold (P<0.01) and GOLPH3 protein increased 2.14-fold (P<0.01) compared with the non-treated controls, and there were no differences between the mock sequence group and the non-treated controls (Fig. 3C and D).

**miR-126 suppresses cell proliferation but has no effect on apoptosis in ESCC cells.** To further characterize the functional role of miR-126 in ESCC, we observed the effect of miR-126 on the proliferation and apoptosis of ESCC cells. After transfection with miR-126 mimics, cell proliferation was suppressed after 48 h (0.433±0.054 vs. 0.514±0.061, P<0.05) and 72 h (0.680±0.038 vs. 0.943±0.070, P<0.01). In addition, transfection with miR-126 inhibitors showed an improvement in cell proliferation at 48 h (0.584±0.088 vs. 0.514±0.061, P<0.05) and 72 h (1.130±0.098 vs. 0.943±0.070, P<0.01); no
alteration of cell proliferation occurred in the mock sequence group (P>0.05; Fig. 4A) compared with the non-treated group. However, cell apoptosis, which was measured using flow cytometry at 48 h after transfection, showed no significant differences between all three groups compared to that in the non-treated controls (P>0.05, Fig. 4B and C).

miR-126 regulates cell migration and invasion. To explore the correlation between miR-126 expression and migration or invasion potential in human ESCC cell lines, we carried out the Transwell assay. To reduce the bias of proliferation suppression of miR-126, the incubation time was controlled for 24 h. In the migration assay, the number of migrated cells was significantly decreased in the group transfected with miR-126 mimics compared to the non-treated controls (43.87±7.59 vs. 67.77±10.57, P<0.01), whereas the number of cells increased in the miR-126 inhibitor group (89.59±12.16 vs. 67.77±10.57, P<0.01, Fig. 5A and B). In the invasion assay, cell invasiveness increased significantly when transfected with the miR-126 inhibitor (48.21±10.21 vs. 27.03±8.36, P<0.01), whereas over-expression of miR-126 significantly suppressed Eca9706 cell invasion (14.44±6.14 vs. 27.03±8.36, P<0.01); results in the mock sequence group were similar to the non-treated controls (P>0.05, Fig. 5A).

Discussion

miRNAs have important roles in tumorigenesis by various mechanisms. miRNAs can act as tumor suppressors depending on the targeting of specific suppressor genes or oncogenes, or they can alter tumors epithelial-mesenchymal transitions to promote tumor invasion and metastasis, and they can also regulate the expression of metastasis-associated genes in tumors (22,23). Thus, the aberrant expression of tumor-suppressive miRNAs may contribute to human carcinogenesis. For example, downregulation of miR-7 can promote metastasis of gastric cancer by targeting the insulin-like growth factor-1 receptor, and the low level of miR-210 is associated with the carcinogenesis of ESCC (24,25). The expression of miR-126 is downregulated in many cancers, such as lung, breast and liver cancer (14,26), suggesting that the suppression of miR-126 was significantly related to the occurrence and development of cancer. The results of the present study confirmed that miRNA-126 was downregulated in ESCC compared with paracancer tissues, indicating that miR-126 may be associated with the carcinogenesis of ESCC. We therefore analyzed the relationship between downregulation of miR-126 and the clinicopathological characteristics of ESCC patients. We found that downregulation of miR-126 was associated
Figure 4. Effect of miR-126 on cell proliferation and apoptosis in the Eca9706 cell line. (A) Cell proliferation significantly increased 48 and 72 h after transfecting miR-126 inhibitors, whereas upregulation of miR-126 led to a decrease of cell proliferation. (B and C) miR-126 on cell apoptosis. Neither the miR-126 mimics nor the inhibitors caused a statistical change in cell apoptosis. *P<0.05, **P<0.01.

Figure 5. Effect of miR-126 on cell invasion and migration in vitro. (A) Results of cell invasion and migration assay. After incubation for 12 h, the invaded or migrated cells that penetrated to the lower surface of the membrane were counted from 5 randomly selected visual fields. (B) In the migration assay, downregulation of miR-126 promoted cell migration, whereas upregulation of miR-126 caused a decrease of migrated cells. **P<0.01.
with tumor cell differentiation, lymph nodes metastasis and tumor invasion in-depth in ESCC patients, which indicated that miR-126 acts as an oncogene in ESCC and participates in tumor invasiveness and metastasis. This result is similar to that found in colon and lung cancer (18,20).

To further explore the molecular mechanism of miR-126 in suppressing ESCC development and invasion, we predicted IRS-1 and GOLPH3 as novel targets of miR-126. The insulin receptor substrates are cytoplasmic signaling adaptor proteins that function as intermediates of the insulin receptor and insulin-like growth factor receptor (IGF-1R), which are involved in cell growth and survival. As a member of the IRS family, IRS-1 has been a hotspot and is widely expressed. Tyrosine-phosphorylated IRS-1 bind proteins containing Src homology 2 (SH2) domains, which activate mitogen-activated protein kinase (MAPK) by effector cascades such as the PI3K pathway (27). Porter et al (28) reported that IRS-1 was highly expressed in localized breast tumors and that high IRS-1 levels can be a good indicator of the effectiveness of specific types of chemotherapy in breast cancer, but were not related to tumor aggressiveness. Several studies have also shown that IRS-1 forced expression of VEGF expression via transcriptional activation and promoted cell proliferation in human colon cancer (29,30). The results of the present study showed that IRS-1 was overexpressed in ESCC tissues and was associated with cell differentiation. Forced expression of miR-126 downregulated the expression level of IRS-1 protein in Eca9706 cells. Therefore, we suggest that IRS-1 is a direct target of miR-126 in human ESCC.

The endocytic protein, GOLPH3, is a highly conserved 34 kDa protein which was initially identified by proteomic characterization of the Golgi apparatus. It was suggested an enhanced activation of mTOR signaling represents a molecular basis of the oncogenic activity of GOLPH3 (31). Several studies have reported that the increased expression of GOLPH3 is closely associated with the clinicopathological characteristics of cancers, such as prostate, lung, ovary and gastric cancer (31-33). Wang et al (34) showed that GOLPH3 was overexpressed in ESCC and was positively associated with clinical stage, TNM classification, histological differentiation and vital status. The present study indicates that GOLPH3 is highly expressed in ESCC and is related to tumor invasion in-depth, lymph nodes metastasis and TNM stage, which is the same as the above study. However, in our study, expression of GOLPH3 was not related to histological differentiation. Although there are few differences, results from the present study and previous studies suggest that GOLPH3, as well as IRS-1, are targets of miR-126 in ESCC, indicating that miR-126 can have different targets and roles across malignancies. In the present study, the mRNA levels of IRS-1 and GOLPH3 did not change significantly with miR-126 levels (when up/downregulated) in ESCC cell lines. This indicated that IRS-1 and GOLPH3 were negatively regulated by miR-126 at the post-transcriptional level in ESCC.

Crawford et al (18) indicated that miR-126 inhibited the proliferation of the human lung cancer cell line A549 both in vitro and in vivo. Li et al (35) found that miR-126 suppresses the proliferation and invasion of the human colon cancer cell lines HCT116, SW620 and HT-29. However, the functions of miR-126 in esophageal cancer progression remain unresolved.

In the present study, the overexpression of miR-126 suppressed cell proliferation in the ESCC cell line Eca9706. Moreover, the Transwell experiment also showed that miR-126 can inhibit ESCC cell migration and invasion in vitro. Collectively, these results suggest that miR-126 may function as a tumor suppressor by inhibiting cell proliferation, suppressing migration and invasion, which, in turn, affect multiple clinicopathological characteristics of cancer such as lymph node metastasis and depth of invasion. At the same time, this is the first report indicating the suppressor action of miR-126 in ESCC.

In conclusion, as a tumor suppressor, miR-126 expression decreased in ESCC and might have a tumor suppression role via the regulation of IRS-1 and GOLPH3. It can suppress cell proliferation, migration and invasion. miR-126 correlates with lymph node metastasis, tumor invasion in-depth and TNM stage of ESCC, and could serve as a potential marker and therapeutic target in ESCC.

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