STMN1, a prognostic predictor of esophageal squamous cell carcinoma, is a marker of the activation of the PI3K pathway

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Abstract. The esophageal squamous cell carcinoma (ESCC) subtype with STMN1 overexpression has a high likelihood of lymphatic metastatic recurrence. However, the underlying mechanism remains to be further elucidated. We assessed the expression level of STMN1 and PTEN in 96 pN0 ESCC patient tissues using immunohistochemistry and western blot analysis. Then, the association between STMN1 overexpression and postoperative lymphatic metastatic recurrence was evaluated. In addition, the relationship between STMN1 and PTEN was also assessed. The results showed that STMN1 expression was significantly higher in tumor tissues (P=0.013). STMN1 overexpression was related to tumor length (P=0.003) and depth of invasion (P=0.019). In addition, STMN1 overexpression was significantly associated with postoperative lymphatic metastatic recurrence rate (P=0.024). Furthermore, in laboratory experiments, STMN1 expression was stably silenced using lentiviral vector delivery of shRNA in Eca109 and EC9706 cell lines to assess the functional effect of STMN1 in vitro. The results indicated that stable silencing of STMN1 expression significantly inhibited cell proliferation, migration and invasion. Moreover, we inactivated the PI3K pathway in ESCC cell lines with the PI3K inhibitor LY294002 and then detected STMN1 expression by western blot analysis. STMN1 levels were robustly reduced consistent with the downregulation of p-Akt (S473) by PI3K pathway inhibition. STMN1 can act as a marker to quantitatively measure the activation of the PI3K pathway and stratify patients accordingly.

Introduction

Esophageal cancer is one of the most common upper gastrointestinal tract malignant neoplasms, and is the sixth leading cause of cancer-related mortality worldwide (1). Esophageal squamous cell carcinoma (ESCC) accounts for more than 70% of esophageal cancers worldwide and is the main pathological type of all esophageal cancer in the Chinese population (2-4). Recently, despite advances in the systemic treatment of ESCC, the prognosis is far from satisfactory (5). In addition, the prognosis of ESCC varies widely between individuals. To stratify patients and develop personalized treatment, it is of clinical importance to elucidate the molecular mechanisms involved in the development and progression of ESCC.

Stathmin 1 (STMN1) is highly conserved and plays a critical role in the assembly and disassembly of the mitotic spindle, which is necessary in the final stage of cell division, and its mutation may lead to uncontrolled cell proliferation (6-8). This role in cell cycle regulation may classify STMN1 as an oncoprotein. It was reported that STMN1 overexpression correlates with invasion and metastasis in many human malignancies, such as lymphoma, ovarian, prostate, breast and lung cancer (9). In addition, STMN1 overexpression was also detected in ESCC. Although, a previous study has shown that STMN1 overexpression predicts high risk for lymphatic metastatic recurrence and a poor prognosis in pN0 ESCC patients (10), the underlying mechanism remains unclear. A series of recent studies suggest that malfunction of signaling pathways plays an important role in promoting proliferation and invasion of ESCC cells and is associated with poor ESCC prognosis. Activation of the oncogenic phosphatidylinositol 3-kinase (PI3K) pathway is frequent in solid tumors (11-13). Multiple cellular processes that are critical for tumorigenesis, such as cell proliferation, apoptosis, migration, glucose metabolism and angiogenesis, are regulated by PI3K signaling (14,15). Meanwhile, the PI3K/Akt pathway is negatively regulated by the tumor suppressor gene phosphatase and tensin homolog deleted on chromosome 10 (PTEN) (16). It is well documented that activation of PTEN/PI3K pathway signaling is a biological marker of poor prognoses in breast, prostate and bladder carcinoma (17) and it is involved in the cisplatin resistance of ESCC cells (18,19). There may be a relationship between STMN1 acting as an oncoprotein and activation of the PI3K pathway.

In the present study, we investigated the correlation between the expression of STMN1 and the prognosis of pN0 ESCC patients. Moreover, we employed laboratory experiments to detect the functional effect of STMN1 on cellular
ability related to tumor metastasis in vitro. To determine the possible underlying mechanisms of high lymphatic metastatic recurrence rate in ESCC patients with the STMN1 overexpression subtype, we performed laboratory research on the PI3K pathway to explore the possible relationships between STMN1 and the regulatory proteins involved in the PI3K pathway.

Materials and methods

Ethics statement. The study protocol was approved by the Research Ethics Committee of Shandong Provincial Hospital affiliated to Shandong University (protocol no. 2017-204). All participants provided their written informed consent for use of the tissues and data analysis.

Patients and specimens. Thirty paired samples of frozen ESCC tissues and corresponding healthy esophageal mucosa (CHEM, >5 cm from the margin of ESCC) were harvested from surgical specimens in our department from January 2016 to May 2016. In addition, from December 2011 through December 2012, 113 patients with mid-thoracic ESCC who had undergone an Ivor Lewis esophagectomy with two-field lymph node dissection in our department were retrospectively studied. Patients did not receive chemotherapy or radiotherapy before surgery and all of them underwent a complete tumor resection. In addition, individuals enrolled in the present study were all restaged with stage pN0 according to postoperative pathology (American Joint Committee on Cancer Staging Manual, 7th edition). Among them, 17 patients were lost to follow-up. The remaining eligible 96 patients were enrolled in this study, and the detailed characteristics of the 96 patients are listed in Table I.

Immunohistochemistry of tissue specimens. The STMN1 and PTEN expression levels were detected by immunohistochemistry using a streptavidin-peroxidase (SP) method according to a previously published procedure (20). Rabbit anti-STMN1 (cat. no. ab52906) and anti-PTEN polyclonal (cat. no. ab170941) antibodies were respectively diluted at 1:150 and 1:50 (Abcam, Cambridge, MA, USA), and the secondary biotinylated antibody kit was purchased from Beijing ZSGB Biotechnology (Beijing, China).

All sections were examined by two independent pathologists who were blinded to the clinical data. The immunohistochemical score (IHS) was identified by combining the proportion score (percentage of positively stained cells) with the staining intensity score. For STMN1 expression, the proportion score ranges were as follows: 0 (<5%), 1 (5-24%), 2 (25-49%), 3 (50-74%) and 4 (>75%) while the staining intensity was scored as 0 (negative), 1 (weak), 2 (moderate) and 3 (strong). The IHS of each case was generated by multiplying the proportion score and staining intensity score. Cases with an IHS ≥4 were considered to have STMN1-positive expression (10). For PTEN expression, the quantity score ranges were as follows: 0 (<5%), 1 (5-25%), 2 (26-50%) and 3 (>50%). The staining intensity was scored as: 0 (absent), 1 (weak staining), 2 (moderate staining) and 3 (strong staining). The total score was classified into negative expression (from 0 to 2) and positive expression (from 3 to 9) (21).

Cell culture, treatment and transfection. The human ESCC cell lines (Eca109 and EC9706) were purchased from the Cell Bank of Shanghai Institute in China (Shanghai, China). All cells were maintained in RPMI-1640 medium supplemented with 10% fetal bovine serum (FBS) and 1% penicillin/streptomycin (both from HyClone Laboratories, Inc., Logan, UT, USA). These cells were grown at 37°C in an atmosphere of 5% CO2.

The PI3K-inhibitor LY294002 was purchased from Selleck Chemicals (Houston, TX, USA). Cells were cultured in LY294002 (20 µM) for 4 days to inhibit the activation of the PI3K pathway.

Human short hairpin RNA (shRNA) was synthesized and packaged in Open Biosystems (Thermo Fisher Scientific, Inc., Waltham, MA, USA). The targeting sequence of STMN1 was 5'-TTATAGGCTTCATTTTG-3' and 5'-TTATTAACC ATTAAGTCC-3' (22). Eca109 and EC9706 cell lines were transfected with the lentivirus-mediated shRNA according to the manufacturer's instructions. The multiplicity of infection (MOI) was 20 for Eca109 and 30 for EC9706 cells. A normal control (NC) shRNA was used as a blank control. Puromycin at a concentration of 5 µg/ml was used to select the transfected cells. qRT-PCR and western blot analysis were used to determine the transfection efficiency.

RNA extraction and qRT-PCR. Total RNA was extracted from the cells using TRIzol reagent (Invitrogen, Carlsbad, CA, USA). The procedures for RNA extraction and qRT-PCR were
detailed as described in our previous study (20). The sequences of primers used were as follows: STMN1 forward 5’-AGA ATA CAC TGC CTG TCG CTT G-3’ and reverse 5’-AGG CAC GCT T C T C C A G T T -3’; β-actin forward 5’-TGG AGA AAA TCT GGc ACC AC-3’ and reverse 5’-GGT CTC AAA CAT GAT CTG G-3’ (22). The relative expression levels were normalized to endogenous β-actin expression.

Cell proliferation assay. In the cell proliferation assay using a Cell Counting Kit-8 (CCK-8; Dojindo Laboratories, Kymamoto, Japan), viable cells were seeded into 96-well tissue culture plates such that there were 3,000 cells in a final volume of 100 µl/well. Every 24 h, 10 µl of CCK-8 solution was added to each well and then the plate was incubated for 4 h at 37˚C. The viable cells were identified by absorbance measurements at 450 nm using a microplate reader (Bio-Rad Laboratories). The experiment was performed in triplicate.

Migration and invasion assays. For the Transwell migration and invasion assays, cells were pre-cultured in serum-free medium for 24 h. In addition, 1.5x10^3 cells in 200 µl serum-free medium were seeded into upper chambers of a 24-well Transwell apparatus (8-µm pore size; Merck Millipore, Darmstadt, Germany). Six hundred microliters of medium with 15% FBS was added to the lower chambers. After incubation for 24 h, cells remaining in the upper chambers were removed by scraping. Cells that had migrated through the membrane were fixed and stained with hematoxylin and eosin (H&E). Then, average numbers of cells per visual field were counted under a light microscope (Leica DM 4000B; Leica Microsystems, Wetzlar, Germany).

For the invasion assay, 40 µl of Matrigel (BD Biosciences, San Jose, CA, USA) was diluted 1:4 in serum-free medium and used to pre-coat the upper chambers of the Transwell apparatus and left to solidify for 1 h. Then, pre-cultured cells in 200 µl of serum-free medium were added to the upper chambers. The remaining procedures were the same as the migration assay, except the duration of incubation was 48 h.

Clonogenic assay. Transfected cells which were trypsinized to generate a single cell suspension were seeded in 6-well plates at 500 cells/well. After 14 days, the number of colonies that were stained with crystal violet and contained at least 50 cells was counted. The colony survival fraction was calculated for each treatment.

Statistical methods. The quantitative data are expressed as the mean ± standard deviation (SD). Differences among multiple groups were analyzed using two-tailed Student’s t-tests using GraphPad Prism 5 software (GraphPad Software, Inc., La Jolla, CA, USA). The Mann-Whitney U test was chosen to identify the differences in STMN1 expression as detected by immunohistochemistry. A Chi-square test was employed to analyze the correlations between the STMN1 overexpression and clinicopathological factors of the pN0 ESCC cases.

Table I. Correlations between STMN1 expression and the clinicopathological factors of the pN0 ESCC cases.

<table>
<thead>
<tr>
<th>Clinical characteristics</th>
<th>Patients (N=96)</th>
<th>STMN1 expression</th>
<th>P-valuea</th>
<th>Recurrence rate (%)</th>
<th>P-valueb</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>(-) (n=43)</td>
<td>(+) (n=53)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>74</td>
<td>33</td>
<td>41</td>
<td>0.943</td>
<td>28 (38.9)</td>
</tr>
<tr>
<td>Female</td>
<td>22</td>
<td>10</td>
<td>12</td>
<td></td>
<td>9 (40.9)</td>
</tr>
<tr>
<td>Age (years)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;50</td>
<td>36</td>
<td>16</td>
<td>20</td>
<td>0.958</td>
<td>12 (33.3)</td>
</tr>
<tr>
<td>≥50</td>
<td>60</td>
<td>27</td>
<td>33</td>
<td></td>
<td>25 (41.7)</td>
</tr>
<tr>
<td>Tumor length (cm)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;3</td>
<td>44</td>
<td>27</td>
<td>17</td>
<td>0.003</td>
<td>14 (31.8)</td>
</tr>
<tr>
<td>≥3</td>
<td>52</td>
<td>16</td>
<td>36</td>
<td></td>
<td>26 (50)</td>
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<tr>
<td>Differentiation</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Well + moderate</td>
<td>64</td>
<td>26</td>
<td>38</td>
<td>0.658</td>
<td>21 (32.8)</td>
</tr>
<tr>
<td>Poor</td>
<td>32</td>
<td>17</td>
<td>15</td>
<td></td>
<td>16 (50)</td>
</tr>
<tr>
<td>pT</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>T1</td>
<td>8</td>
<td>5</td>
<td>3</td>
<td>0.019</td>
<td>0 (0)</td>
</tr>
<tr>
<td>T2</td>
<td>37</td>
<td>22</td>
<td>15</td>
<td></td>
<td>9 (24.3)</td>
</tr>
<tr>
<td>T3</td>
<td>51</td>
<td>16</td>
<td>35</td>
<td></td>
<td>28 (54.9)</td>
</tr>
<tr>
<td>PTEN expression</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>(-)</td>
<td>51</td>
<td>15</td>
<td>36</td>
<td>0.001</td>
<td>25 (49.0)</td>
</tr>
<tr>
<td>(+)</td>
<td>45</td>
<td>28</td>
<td>17</td>
<td></td>
<td>12 (26.7)</td>
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</table>

*aChi-square test; bP-value log-rank test.
and the clinicopathological factors, and those between STMN1 expression and PTEN expression. A univariate analysis was performed by modeling Kaplan-Meier survival curves. The log-rank test was used to calculate the survival rate. A multivariate analysis was carried out using the Cox proportional hazards model. Differences were considered significant when P<0.05. The statistical data were obtained using the SPSS software package (SPSS 17.0; SPSS, Inc., Chicago, IL, USA).

Results

The expression of STMN1 in ESCC tissues and CHEM. STMN1 protein expression levels in 30 pairs of tissue specimens (ESCC tissue and CHEM) were investigated by IHC and western blot analysis. The IHC results showed that the positive expression of STMN1 was detected as a yellow or brownish yellow stain in the cytoplasm (Fig. 1). The IHC staining demonstrated that the expression level of STMN1 in ESCC was significantly higher than that in CHEM (P=0.013; Fig. 2). STMN1 overexpression was found in 16 cases (53.3%) of ESCC tissues and 4 cases (13.3%) of CHEM. Moreover, we randomly selected 8 pairs of tissue specimens (ESCC and CHEM) to confirm the STMN1 protein level by western blot analysis (Fig. 3A). Higher STMN1 protein expression was identified in tumor tissues (STMN1/β-actin: 0.76±0.10 vs. 0.49±0.18, P=0.0021; Fig. 3B). The results corresponded to those of the immunohistochemistry analysis.

STMN1 expression is correlated with clinical characteristics and with PTEN expression. The clinicopathological data of the 96 eligible patients with mid-thoracic ESCC were retrospectively studied. ESCC tissues from 53 patients were identified with STMN1 overexpression. The diagnostic sensitivity was 55.2% (53/96) (Table I).

By immunohistochemistry, positive expression of PTEN was detected mainly in the cytoplasm (Fig. 4) and rarely in the nuclei (only 4 stained heterogeneously). Forty-five tumor samples (46.9%) showed positive expression of PTEN, whereas 51 (53.1%) tumor samples showed negative expression of PTEN.

The correlation between STMN1 expression and clinicopathological features is shown in Table I. The Chi-square analysis indicated that STMN1 overexpression was significantly associated with tumor length (P=0.003) and depth of invasion (P=0.019). In addition, STMN1 expression was inversely correlated with PTEN expression (P=0.001). No other clinicopathological parameter was associated with STMN1 overexpression.

STMN1 expression is correlated with the overall survival and lymphatic metastatic recurrence. Through the follow-up, a first lymph node metastatic recurrence within 3 years was
identified in 37 cases (38.5%), in which STMN1 overexpression was detected in 26 patients (70.3%). In the group with low STMN1 expression, the 3-year lymphatic metastatic recurrence rate was only 25.6% (11/43), whereas in the group with STMN1 overexpression, this rate reached 49.1% (26/53). The Kaplan-Meier analysis and the log-rank test showed that patients with STMN1 overexpression had a significantly higher lymphatic metastatic recurrence rate (Fig. 5A; P=0.024) and a significantly worse overall survival rate (Fig. 5B; P=0.030).

In addition, T stage (P=0.002), differentiation degree (P=0.016) and PTEN levels (P=0.026) were also indicated to be associated with lymphatic recurrence in pN0 ESCC patients by univariate analysis (Table I). The Cox proportional hazards model was performed to identify factors involved in lymphatic recurrence of pN0 ESCC patients. The Cox multivariate regression analysis revealed that the T stage (P=0.019) and the tumor differentiation degree (P=0.060) were both independent prognostic factors for pN0 ESCC (Table II).

Table II. Cox regression analysis for risk factors of 3-year lymphatic metastatic recurrence in pN0 ESCC patients.

<table>
<thead>
<tr>
<th></th>
<th>B</th>
<th>SE</th>
<th>Wald</th>
<th>P-value</th>
<th>HR</th>
<th>95% CI</th>
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<tbody>
<tr>
<td>Sex</td>
<td>0.422</td>
<td>0.422</td>
<td>1.000</td>
<td>0.317</td>
<td>0.656</td>
<td>0.287-1.500</td>
</tr>
<tr>
<td>Age (years)</td>
<td>0.022</td>
<td>0.373</td>
<td>0.003</td>
<td>0.954</td>
<td>1.022</td>
<td>0.492-2.125</td>
</tr>
<tr>
<td>Tumor length</td>
<td>-0.122</td>
<td>0.391</td>
<td>0.097</td>
<td>0.755</td>
<td>0.885</td>
<td>0.411-1.906</td>
</tr>
<tr>
<td>T stage</td>
<td>0.980</td>
<td>0.419</td>
<td>5.461</td>
<td>0.019</td>
<td>2.664</td>
<td>1.171-6.060</td>
</tr>
<tr>
<td>Differentiation</td>
<td>0.676</td>
<td>0.359</td>
<td>3.542</td>
<td>0.060</td>
<td>1.967</td>
<td>0.972-3.977</td>
</tr>
<tr>
<td>STMN1 expression</td>
<td>0.450</td>
<td>0.391</td>
<td>1.325</td>
<td>0.250</td>
<td>1.568</td>
<td>0.729-3.374</td>
</tr>
<tr>
<td>PTEN expression</td>
<td>0.143</td>
<td>0.395</td>
<td>0.131</td>
<td>0.717</td>
<td>0.867</td>
<td>0.399-1.880</td>
</tr>
</tbody>
</table>

B, regression coefficient; SE, standard error; Wald, Wald value; HR, hazard ratio; CI, confidence interval.

Figure 4. Representative immunohistochemical staining of PTEN in esophageal squamous cell carcinoma (ESCC) tissue and corresponding healthy esophageal mucosa (CHEM) (original magnification, x400).

Figure 5. Kaplan-Meier analysis and log-rank test of STMN1 expression for lymphatic metastatic recurrence (A) and overall survival (B).
Expression of STMN1 in cell lines. qRT-PCR was performed to evaluate the expression of STMN1 in four esophageal cancer cell lines: Eca109, KYSE150, TE-1 and EC9706. Eca109 and EC9706 showed relatively high expression of STMN1 (Fig. 6). Immunocytochemistry was used to verify the result of qRT-PCR. By immunocytochemistry, the positive expression of STMN1 protein showed a yellow or brownish yellow stain in the cytoplasm of tumor cells (Fig. 7). Strong immunoreactivity of STMN1 protein was detected in the cytoplasm of Eca109 and EC9706 cells. Therefore, Eca109 and EC9706 were selected as the candidate cell line for shRNA transfection.

Lentiviral-mediated shRNA silencing of STMN1 gene expression. After transfection, western blot analysis was used to confirm the efficacy. The results showed that the expression of the STMN1 protein was significantly downregulated in the lentiviral-mediated STMN1 shRNA-transfected Eca109 and EC9706 cells (P<0.001; Fig. 8). There was no significant difference in STMN1 expression between the NC shRNA (control) group and the untransfected group. These results showed that stable transfection of STMN1 shRNA can effectively and specifically silence STMN1 gene expression.

Stable silencing of STMN1 inhibits cell proliferation. The CCK-8 assay showed that the cell growth rate of Eca109 and EC9706 cells transfected with STMN1 shRNA was significantly lower compared to those cells transfected with NC shRNA. The OD450 values of the Eca109 and EC9706 cells transfected with STMN1 shRNA were significantly decreased at 24, 48 and 72 h (P<0.01; Fig. 9A and B). The clonogenic assay (Fig. 9C) showed that the colony numbers of Eca109 and EC9706 cells transfected with STMN1 shRNA were significantly less than those transfected with NC shRNA (P<0.05; Fig. 9D and E). These results indicated that stable silencing of STMN1 may inhibit the cell proliferation.

Stable silencing of STMN1 inhibits cell migration and invasion. Migration and invasion assays were used to test the effect of STMN1 on cell motility and invasion. The migration assay showed that the number of migrated cells in the STMN1 shRNA group were significantly decreased in the lower chamber compared with cells in the NC shRNA group (P<0.0001; Fig. 10A). At the same time, the number of invaded cells in the STMN1 shRNA group was also decreased compared to that in NC shRNA group in the invasion assay (P<0.0001; Fig. 10B and C). These results indicated that stable silencing of STMN1 may inhibit the invasive and metastatic ability of ESCC cells.

STMN1 is PI3K pathway-regulated in ESCC cells in vitro. To identify the possible relationship between STMN1 and...
the activation of the PI3K pathway, we detected the status of p-Akt (S473) to identify the activation of the PI3K pathway in STMN1 shRNA Eca109 cells. Western blot analysis showed that there was no obvious change in p-Akt expression after STMN1 was silenced (Fig. 11A).

Then, we treated Eca109 and EC9706 cells with LY294002 (diluted in DMSO) to inhibit the activation of the PI3K pathway and then detect STMN1 expression by western blot analysis. The results showed that STMN1 levels were robustly reduced, consistent with the downregulation of p-Akt (S473) by PI3K...
pathway inhibition both in the Eca109 and EC9706 cell line (Fig. 11B).

**Discussion**

Currently, postoperative adjuvant therapy is not recommended in pN0 ESCC after radical operation according to the NCCN guidelines. However, multiple ESCC patients in the pN0 stage tend to relapse after R0 resection, and the common type of relapse is lymph node metastatic recurrence. With surgery alone, the 5-year survival rate of ESCC patients in stage IB and IIA is 62 and 55%, respectively (23,24). The prognosis of ESCC is far from satisfactory even in pN0 stage. Also, it is of clinical importance to predict lymphatic metastatic recurrence early and select those candidates to undergo postoperative adjuvant therapy.

STMN1 is a ubiquitously expressed phosphoprotein. It plays a critical role in the assembly and disassembly of the mitotic spindle, which is necessary in the final stage of cell division. In addition, it regulates cell cycle progression by influencing the dynamics of microtubules (25,26). Its role in regulating the cell cycle makes STMN1 act as an oncoprotein. Overexpression of STMN1 in human cancer was reported to be associated with malignancy and poor prognosis (27). It was demonstrated that STMN1 is overexpressed in ESCC, and STMN1 overexpression predicts a high risk for lymphatic metastatic recurrence in pN0 ESCC patients (10). However, the functional effect of STMN1 in vitro was not elucidated. It is well known that the abilities of primary tumor cells to invade and propagate are vital factors that are involved in tumor metastasis (28).

In the present study, we used shRNA to silence the expression of STMN1 in ESCC cell lines to study the effect of STMN1 on cellular ability related to tumor metastasis in vitro. Eca109 and EC9706 cells were chosen to analyze the effect of STMN1 suppression by shRNA due to their higher expression of STMN1. After stable transfection, western blot analysis showed that expression of STMN1 in Eca109 and EC9706 was stably suppressed. The CCK-8 assay showed that suppressing STMN1 expression significantly inhibited the growth rates of the Eca109 and EC9706 cells. Migration and invasion assays confirmed that knockdown of STMN1 significantly inhibited the abilities of motility and invasion in Eca109 and EC9706 cells. We concluded that the STMN1 expression was associated with the ability for metastasis in vitro in ESCC.

Despite the fact that STMN1 has proven to act as a tumor marker that is capable of discriminating ESCC patients with good or poor outcomes, deciphering the biological basis of why it is predictive remains a significant challenge. The activation of the oncogenic PI3K pathway is frequent in solid tumors. In addition, it was estimated that aberrant PI3K pathway signaling is present in more than 30% of human cancers (29). The PI3K pathway is involved in many aspects of tumor biology: cell transformation, growth, proliferation, migration, protection from apoptosis, genomic instability, angiogenesis and metastasis (29,30). Akt, a small family of serine/threonine protein kinases, is the end-point of the PI3K pathway. Activated Akt can phosphorylate a number of downstream substrates to regulate the above cellular processes (31). Conversely, the lipid phosphatase PTEN can dephosphorylate the 3'-end of phosphatidylinositol so that it can attenuate Akt activation and negatively regulate the PI3K pathway (32).

The activation of the oncogenic PI3K pathway is frequent in ESCC, and STMN1 overexpression predicts a high risk for lymphatic metastatic recurrence in pN0 ESCC patients. However, the functional effect of STMN1 in vitro was not elucidated. It is well known that the abilities of primary tumor cells to invade and propagate are vital factors that are involved in tumor metastasis (28). In the present study, we used shRNA to silence the expression of STMN1 in ESCC cell lines to study the effect of STMN1 on cellular ability related to tumor metastasis in vitro.

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Figure 11. (A) Bands of p-Akt and β-actin in STMN1 shRNA-transfected Eca109 cells. (B) Expression of STMN1 in Eca109 and EC9706 cells treated with DMSO (negative control) or the PI3K pathway inhibitor LY294002. LY, LY294002.
act as a marker to quantitatively measure the activation of the PI3K pathway and stratify patients accordingly.

The limitation of the present study is that the functional effect of STMN1 on the cellular ability related to tumor metastasis was determined only in vitro, and the in vivo effect was not included. In future research, we will enroll xenograft tumor models in nude mice to study the functional effect of STMN1 in ESCC in vivo.

In conclusion, STMN1 overexpression was significantly associated with lymphatic metastatic recurrence in pN0 ESCC patients. STMN1 levels are regulated by the PI3K pathway, and STMN1 can act as a surrogate marker of PI3K pathway signaling related to tumor recurrence. STMN1 may be clinically useful to select patients for PI3K pathway-targeted therapy and to monitor the therapeutic efficacy in ESCC.

Acknowledgements

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References