Overexpressed PKMYT1 promotes tumor progression and associates with poor survival in esophageal squamous cell carcinoma

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Background: Esophageal squamous cell carcinoma (ESCC) is one of the most common malignant tumors worldwide and the 5-year overall survival rate remains poor. Protein kinase, membrane-associated tyrosine/threonine (PKMYT1) is overexpressed in several cancers and participates in tumor progression. However, the mechanism of PKMYT1 in ESCC is unclear.

Purpose: The objective of our study was to demonstrate the expression and role of PKMYT1 in ESCC.

Patients and methods: We detected the expression of PKMYT1 in ESCC patients and analysed the correlation with overall survival time and disease-free survival time. Then we detected PKMYT1 expression in ESCC cell lines and immortalized human esophageal epithelial cell line. Down-regulated PKMYT1 was carried out in KYSE70 and KYSE450 cells to investigate the mechanism of PKMYT1 in ESCC cells.

Results: PKMYT1 was up-regulated in tumor tissues and ESCC cell lines, and higher expression of PKMYT1 correlated with poorer overall survival in ESCC patients. Besides, in ESCC cell lines KYSE70 and KYSE450, knocking down PKMYT1 allowed more cells to skip G2/M checkpoint to complete mitosis, which promoted cell apoptosis, inhibited cell proliferation, and prevented the EMT phenotype in vitro. Meanwhile, we also observed that down-regulated PKMYT1 in ESCC cells suppressed AKT/mTOR signaling pathway. These results demonstrated PKMYT1 may act as an oncogene in ESCC.

Conclusion: PKMYT1 plays an crucial role in ESCC progression, downregulated PKMYT1 might inhibit the development of ESCC by AKT/mTOR signaling pathway, and might be a novel target in the treatment of ESCC.

Keywords: ESCC, PKMYT1, AKT, EMT, cell apoptosis

Introduction

Esophageal carcinoma (EC) is the ninth common carcinoma and the sixth leading cause of cancer-related death all over the world.1–3 The histological type of EC includes esophageal squamous cell carcinoma (ESCC) and esophageal adenocarcinoma, with ESCC accounting for the majority of EC patients in China.4–6 With the evolution of the multidisciplinary treatment, including novel surgical approaches, chemotherapy, radiotherapy, and molecular-targeted therapy, the 5-year overall survival rate remains poor.7 Thus, it is necessary and urgent to explore the pathogenesis of ESCC which might provide new insights into the treatment of ESCC.

Protein kinase, membrane-associated tyrosine/threonine (PKMYT1) is a member of the Wee family of protein kinases. PKMYT1 acts as a negative regulator of the cell cycle by inactivating the CDK1-cyclinB complex through phosphorylation of Tyr14/Tyr15 and...
preventing cell cycle entering into mitosis at the G2/M transition. In cell cycle, there will be a large number of DNA mismatches and DNA defect when DNA is duplicated. Normal cells repair DNA at G1/S checkpoint, whereas tumor cells repair DNA and shun immature cells entry into mitosis to prevent mitotic catastrophe at G2/M checkpoint because they have abrogated G1/S checkpoint due to the mutation of P53. Therefore, PKMYT1 is more essential for tumor cells than normal cells. Nowadays, a new tendency for cancer therapy is to keep cell cycle with unrepaired DNA damage in premature mitosis, which induces apoptotic or non-apoptotic cell death. Previous studies have demonstrated that upregulated PKMYT1 plays an important role in the progression of these malignancies such as hepatic carcinoma, colon cancer, glioblastoma. A high frequency of TP53 mutations has been found in ESCC patients, but how PKMYT1 participates in the pathogenesis and development of ESCC have not been fully investigated. In this study, we found that PKMYT1 was highly expressed in ESCC cell lines and ESCC patients, and high PKMYT1 expression induced poorer prognosis in patients with ESCC. Then, we investigated its biological functions in ESCC cells. Our results indicated that PKMYT1 might be a potential target in the treatment of ESCC.

Materials and methods

Patients

Totally, 60 ESCC patients tissues including paired tumor tissues and normal adjacent tissues (>5 cm from tumor) were collected in the First Affiliated Hospital of Zhengzhou University from February 2018 to October 2018. Fresh tissues were immediately frozen at −80°C for RNA extraction. Besides, we collected histopathological sections of 104 patients who underwent radical resection of esophageal cancer from 2014 to 2017 to perform immunohistochemistry. All patients in this subject had none preoperative treatments including chemotherapy and radiotherapy. This study was approved by the Ethics Committee of the First Affiliated Hospital of Zhengzhou University and all patients signed written informed consent. And all these assays were conducted in accordance with the Declaration of Helsinki.

Expression profiling and interactive analyses (GEPIA) database and Kaplan–Meier plotter database

GEPIA is a new Bioinformatics tool for analyzing the expression of mRNA by The Cancer Genome Atlas (TCGA) and the Genotype-Tissue Expression (GTEx) projects (http://gepia.cancer-pku.cn/). We analyzed the mRNA expression of PKMYT1 in ESCC tumor tissues and normal tissues, then we also explored the correlation between PKMYT1 expression and TWIST, Ki67, and AKT expression in GEPIA. Kaplan–Meier Plotter was performed to analyze the correlation between the expression of PKMYT1 and survival time in breast cancer, lung cancer, and gastric cancer, and this database had 54,675 genes and 10,461 cancer samples, which included 5143 breast, 1186 ovarian, 2437 lung, and 1065 gastric cancer samples (http://kmplot.com/analysis/).

Cell lines and cell culture

Both ESCC cell lines (KYSE450, KYSE70) and immortalized human esophageal epithelial cell line HET-1A were purchased from Cell Bank of the Chinese Academy of Sciences (Shanghai, China). All ESCC cells and HET-1A cells were cultured in RPMI1640 medium (Hyclone, SH30809.01B, Logan, UT, USA) supplemented with 10% fetal bovine serum (Hyclone, FBS, Logan, UT, USA), 100 units/mL penicillin, and 100 µg/mL streptomycin. All cells were maintained in humidified atmosphere with 5% carbon dioxide at 37°C.

siRNA and cell transfection

siRNAs targeting PKMYT1 were synthesized by GenePharma (Shanghai, China), the sequence of siRNAs was listed in Table 1. Lipofectamine 3000 (Invitrogen, Lipofectamine 3000, Carlsbad, CA, USA) was used for cell transfection according to the manufacturer’s protocol. We transfected ESCC cells in a six-hole plate, and 48 hrs after transfection, we collected ESCC cells for further assays.

RNA extraction, reverse transcription, and quantitative real-time PCR (qRT-PCR)

All RNA was isolated from cells or tissues using Trizol (Invitrogen, TRIzol RNA Isolation Reagents, Carlsbad, CA, USA) according to the manufacturer’s protocol. Whole cDNA was synthesized from 1 µg RNA using the Primescript™ RT reagent Kit (Takara, Primescript™ RT reagent Kit, Beijing, China). The level of PKMYT1 mRNA was assessed by qRT-PCR using FastStart Essential DNA Green Master (Roche, Penzberg, Upper Bavaria, Germany) according to the manufacturer’s instructions. Glyceraldehyde 3-phosphate dehydrogenase was used for normalization of data and these data were analyzed by 2−ΔΔCT. The primers sequences (Sangon Biotech, Shanghai, China) for the RT-PCR analyses are listed in Table 2.
Cell cycle
After the transfection, $1 \times 10^6$ were collected for flow cytometry, cell cycle detection kit (KeyGEN BioTECH, KGA512, Nanjing, Jiangsu Province, China) was used in this assay according to the manufacturer’s protocol. All data were analyzed by Modfit software.

CCK8
CCK8 assay was measured by Cell Counting Kit-8 (Dojindo, CCK-8, Shanghai, China). Transfected ESCC cells were seeded in 96-well plate for 1000 cells per well. Cells were cultured for 24, 48, 72, 96 hrs, and 10 μL of CCK-8 was added into each well. The absorbance was read at 450 nm (Thermo Scientific, Multiskan MK3, Waltham, MA, USA) after incubation for 4 hrs at 37°C.

Colony formation
In the colony formation assay, 2000 cells were plated in six-well plate and cultured for 2 weeks at 37°C in humidified atmosphere with 5% carbon dioxide. Four percent paraformaldehyde was used to fix cell colonies, and cell colonies were stained in 0.5% crystal violet for 20 mins. Then the cell colonies were counted and analyzed.

Cell apoptosis
One hundred thousand ESCC cells were plated into 24-well plates and harvested 24 hrs later. Cell apoptosis was detected using propidium iodide/Annexin-V-FITC (Biolegend, San Diego, CA, USA), then flow cytometry was performed, and Flowjo software (Tree Star Software, San Carlos, CA, USA) was used to analyze the results. All the assays were performed in triplicate.

Migration and invasion
The migration assay was performed by using 8 μm transwell chamber (Corning Costar corporation, Corning, NewYork, NY, USA). $1 \times 10^5$ ESCC cells were plated into the upper chamber with serum-free medium, and medium containing 10% fetal bovine serum was added into the lower chamber. After incubating for 48 hrs, upper chamber cells were scraped off and the migration cells were fixed in 4% paraformaldehyde, and then stained in 0.5% crystal violet. The migration cells were counted under a microscope (Olympus Corporation, IX73, Tokyo, Japan). Matrigel (Corning Life Science, C-Matrigel, NewYork, NY, USA) was used to measure the invasion ability of ESCC cells, 50 μL matrigel was plated in the upper chamber, the other procedures were the same as migration assay.

Wound healing
When there was 80–90% confluence in six-well plates, 200 μL pipette was used to scratch the cells, and then PBS was used to remove floating cells and detached cells. Serum-free medium was added into plates, and photos were photographed using a microscope in 0, 12, and 24 hrs to analyze cell migration.

Western blot
All the protein was collected by using RIPA lysis buffer (Beyotime Biotechnology, P0013, Shanghai, China). Protein was quantified using Coomassie blue staining in microplate spectrophotometer (Thermo Scientific, Multiskan MK3, Waltham, MA, USA) at 594 nm. Identical amount of protein was separated into 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis, and then the gels were transferred onto nitrocellulose membrane (GE Healthcare Life Sciences, Amersham™ Protran™ 0.45 NC, Chicago, IL, USA) after

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**Table 1 Sequences of siRNAs**

<table>
<thead>
<tr>
<th>Name</th>
<th>Sequence of siRNA (5′–3′)</th>
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<tr>
<td>PKMYT1-868</td>
<td>CUCUGGGAUGUCUCAATTUUGAAGACCCUCUCGGGUAGTTT</td>
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<tr>
<td>PKMYT1-1396</td>
<td>UCUGGUAUGUGCAUGCAATTUUGUCAGGCACUUCACAGGATT</td>
</tr>
<tr>
<td>PKMYT1-NC</td>
<td>UUCUGGAACGUGUCAGGUTACGUGACAGUGCGAGAATT</td>
</tr>
</tbody>
</table>

Abbreviation: PKMYT1, protein kinase, membrane associated tyrosine/threonine.

**Table 2 Primers used for RT-qPCR**

<table>
<thead>
<tr>
<th>Genes</th>
<th>Primers sequences</th>
</tr>
</thead>
<tbody>
<tr>
<td>PKMYT1</td>
<td></td>
</tr>
<tr>
<td>Forward</td>
<td>5′-CATGGAACGCTTACGGAGGT-3′</td>
</tr>
<tr>
<td>Reverse</td>
<td>5′-ACATGGAACGCTTACGGAGGT-3′</td>
</tr>
<tr>
<td>GAPDH</td>
<td></td>
</tr>
<tr>
<td>Forward</td>
<td>5′-GGGT GTGAA CATGGAAGT-3′</td>
</tr>
<tr>
<td>Reverse</td>
<td>5′-GGCATG GACTGTGGTCATGA-3′</td>
</tr>
</tbody>
</table>

Abbreviations: PKMYT1, protein kinase, membrane associated tyrosine/threonine; GAPDH, glyceraldehyde 3-phosphate dehydrogenase.
electrophoresis. Membranes were blocked by 5% skim milk for 2 hrs. Then, the membranes were incubated with primary antibodies (E-cadherin (ECAD), N-cadherin (NCAD), Vimentin (VIM), and β-actin were purchased from Proteintech group, Wuhan, China; PKMYT1, AKT, mTOR, S6, Phospho-AKT, Phospho-mTOR, Phospho-S6, and Twist1 were purchased from Cell Signaling Technology, Danvers, MA, USA) in 5% bovine serum albumin overnight at 4°C. After rewarming to room temperature for 1 hr, the membranes were incubated with secondary antibody for 1 hr at 37°C. We used enhanced chemiluminescence (Beyotime Biotechnology, Shanghai, China) to detect the blots with ChemiDoc MP (Bio-Rad Laboratories, Hercules, CA, USA). Images were taken under a fluorescence microscope and analyzed by ImageJ software (National Institutes of Health, Bethesda, MD, USA).

**Immunofluorescence**

Two thousand cells were seeded into 24-well plates, and then fixed in 4% paraformaldehyde and permeabilized by 0.5% Triton X-100. Five percent goat serum was used to block cells for 1 hr, and cells were incubated with primary antibodies (ECAD, NCAD, VIM) overnight at 4°C. Then, the cells were incubated with secondary antibodies (Biolegend, San Diego, CA, USA) for 1 hr, and then stained with 4’,6-diamidino-2-phenylindole (DAPI) (Biolegend, San Diego, CA, USA). Images were taken under a fluorescence microscope and analyzed by ImageJ software (National Institutes of Health, Bethesda, MD, USA).

**Immunohistochemistry (IHC)**

Patients tissues were fixed with formalin and embedded in paraffin. After dewaxing and hydration, tissue sections were incubated with antigen retrieval solution. Then, the tissue sections were treated with 3% H2O2 and 5% goat serum. Sections were incubated with primary antibody (PKMYT1, 1:50) at 4°C overnight. After rewarming to room temperature for 1 hr, the sections were marked with secondary biotinylated antibody, and diaminobenzidine was used as a chromogen. Then, we used hematoxylin for nuclear counterstain. Subsequently, the sections were observed by microscope and all images were at 200× magnification and assessed by two pathologists independently using a semi-quantitative immunoreactivity score (IRS). According to the proportion of positive tumor cells examined, the tissues were scored as 0 (no positive tumor cells), 1 (0–25%), 2 (26–50%), 3 (51–75%), and 4 (76–100%). Staining intensity was scored as 0 (no staining), 1 (weak staining), 2 (intermediate staining), or 3 (strong staining). IRS was evaluated as staining intensity × proportion of positive tumor cells that ranging between 0 and 12. Patients with a total score<4 were considered low PKMYT1 expression and total score ≥4 was considered as high PKMYT1 expression. We divided these patients into two groups, high PKMYT1 group (n=50), and low PKMYT1 group (N=54).

**Statistical analysis**

Statistical analysis was performed by using SPSS 22.0 (IBM Corporation, Armonk, NY, USA) program. The results of RT-PCR, apoptosis, migration, and invasion were analyzed with Student’s t-test. Chi-square tests were conducted for analyzing the associations of PKMYT1 and clinicopathological data. The overall survival rate and disease-free survival rate were analyzed by Kaplan–Meier curve with the log-rank test method. Cox regression models were used to analyze the clinicopathological data. P<0.05 was considered to indicate statistically significant differences.

**Results**

**Expression of PKMYT1 in ESCC and its association with clinical-pathological features**

To investigate the expression of PKMYT1 in ESCC patients, RT-PCR and IHC assays were carried out with ESCC patients’ paired tissues. The results indicated that the expression levels of PKMYT1 in tumor tissues were higher than the one in the paired normal adjacent tissues (Figure 1A and B). We then analyzed the expression of PKMYT1 in ESCC cell lines and HET-1A cells by RT-PCR, showing that PKMYT1 expression in ESCC cells was significantly higher than HET-1 cells (Figure 3A). These results demonstrated that PKMYT1 was up-regulated in ESCC. Then, we analyzed the correlation between PKMYT1 and clinicopathological parameters of patients. The results revealed that the protein expression of PKMYT1 in ESCC tissues was associated with lymph nodes metastasis (P=0.018), tumor differentiation (P=0.018), and pathological stage (P=0.014). However, there were no correlations between PKMYT1 with tumor invasion depth, age, cigarette or alcohol intake, and gender (Table 3).

**PKMYT1 with clinical outcomes in patients with ESCC**

We used Kaplan–Meier method and log-rank test to assess whether the expression of PKMYT1 was related to the
prognosis of ESCC patients. The results of IHC demonstrated that high PKMYT1 expression was significantly related to poor overall survival in patients with ESCC (Figure 1C and D), and high PKMYT1 expression also related to shorter disease-free time (Figure 1E). Then, we used Cox proportional hazards model with univariate and multivariate analysis to verify whether PKMYT1 can be used as an independent risk factor, these results indicated that high PKMYT1 expression was an independent risk factor \( (P=0.014) \) in patients with ESCC (Table 4). All these results indicated that PKMYT1 associated with the prognosis of patients with ESCC.

Also, we performed in-silico analysis of microarray gene expression data using an online survival analysis tool (data from http://kmplot.com/analysis/) to assess the prognostic effect of PKMYT1 in lung cancer, breast cancer, and gastric cancer, showing that high expression of PKMYT1 was related to poor prognosis (Figure 2A). Moreover, using a web server for cancer and normal gene GEPIA, we found that PKMYT1 expression was significantly higher in ESCC tumor tissues than in normal tissues (Figure 2B). The expression of PKMYT1 associated with some important genes of ESCC, such as Ki67, TWIST,\(^{23,24}\) AKT (Figure 2C).\(^{25,26}\)

**Effect of PKMYT1 knockdown on cell cycle, cell proliferation, and cell apoptosis**

To further explore the function of PKMYT1, KYSE70, KYSE450 cell lines were chosen for the subsequent experiments. Two different siRNAs were designed and transfected into those ESCC cells to knockdown PKMYT1 in ESCC cells (Figure 3B). Forty-eight hours after the transfection, cell cycle...
assays indicated that knocked down PKMYT1 expression reduced the G2/M phase in cell cycle (Figure 3C), and CCK-8 assays showed that knockdown of PKMYT1 significantly suppressed cell proliferation in KYSE450 cells compared with negative control group (Figure 3D). This observation was further supported by reduced numbers of colony formation upon PKMYT1 knockdown (Figure 3E). Additionally, we observed that the negative control group had

Table 3 The correlation between PKMYT1 expression and clinicopathological factors in ESCC patients

<table>
<thead>
<tr>
<th>Clinic factors</th>
<th>All cases</th>
<th>PKMYT1 expression</th>
<th>P-value</th>
</tr>
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<tbody>
<tr>
<td></td>
<td></td>
<td>High (n=50)</td>
<td>Low (n=54)</td>
</tr>
<tr>
<td>Age (years)</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>&lt;65</td>
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<td>28</td>
<td>24</td>
</tr>
<tr>
<td>≥65</td>
<td>52</td>
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<td>30</td>
</tr>
<tr>
<td>Gender</td>
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</tr>
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<tr>
<td>No</td>
<td>51</td>
<td>22</td>
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<tr>
<td>Tumor invasion depth</td>
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</tr>
<tr>
<td>T1+T2</td>
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<td>23</td>
<td>31</td>
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<tr>
<td>T3+T4</td>
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<td>27</td>
<td>23</td>
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<td>Lymph nodes metastasis</td>
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<td>Positive</td>
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<td>34</td>
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<tr>
<td>III-IV</td>
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</table>

Note: The bold values mean that the P-value<0.05.
Abbreviations: ESCC, esophageal squamous cell carcinoma; PKMYT1, protein kinase, membrane associated tyrosine/threonine.

Table 4 Univariate and multivariate Cox regression analysis of the relative risk of death according to the expression of PKMYT1

<table>
<thead>
<tr>
<th>Variables</th>
<th>Univariate analysis</th>
<th>Multivariate analysis</th>
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<tr>
<td></td>
<td>HR(95% CI)</td>
<td>P-value</td>
</tr>
<tr>
<td>Age</td>
<td>1.769(1.083–2.889)</td>
<td>0.023</td>
</tr>
<tr>
<td>Gender</td>
<td>0.706(0.420–1.187)</td>
<td>0.189</td>
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<tr>
<td>Tumor invasion depth</td>
<td>1.544(0.950–2.510)</td>
<td>0.079</td>
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<tr>
<td>Differentiation grade</td>
<td>3.309(1.870–5.857)</td>
<td>0.000</td>
</tr>
<tr>
<td>Lymph nodes metastasis</td>
<td>8.966(5.037–15.959)</td>
<td>0.000</td>
</tr>
<tr>
<td>TNM stage</td>
<td>7.828(4.445–13.785)</td>
<td>0.000</td>
</tr>
<tr>
<td>PKMYT1 expression</td>
<td>2.376(1.453–3.883)</td>
<td>0.001</td>
</tr>
</tbody>
</table>

Note: The bold values mean that the P-value<0.05.
Abbreviation: PKMYT1, protein kinase, membrane associated tyrosine/threonine.
lower apoptosis rate than siRNA groups in KYSE450 cells (Figure 3F), and Western blot analysis indicated that the expression of Bcl-2 was decreased, but the expression of Bax and cleaved caspase3 was increased (Figure 3G). Similar results were obtained when performing these analyses in KYSE70 cells following PKMYT1 knockdown. All these assays showed that inhibited the expression of PKMYT1 promoted ESCC cell apoptosis and reduced the proliferation of ESCC cells.

Effect of PKMYT1 knockdown on invasion and migration and PKMYT1 implication in establishing the mesenchymal phenotype ESCC cells

Transwell assays were used to investigate the effects of knockdown PKMYT1 on the invasion and migration capability in ESCC cells. The transwell assays showed that there were fewer cells migrated in siRNA group than negative group in KYSE450 and KYSE70 cells (Figure 4A and B), indicating that PKMYT1 knockdown could inhibit the migration and invasion capacities of ESCC cells. Similarly, the wound healing assays revealed that the siRNA groups had less migrated cell proportions compared to the control groups in both ESCC cell lines (Figure 4C and D). All these results indicated that PKMYT1 might involve in metastasis of esophageal cancer. To further investigate how PKMYT1 affect metastasis capability of ESCC cells, we detected epithelial-mesenchymal transition (EMT) phenotype, an important pathway in tumor progression and metastasis. The necessary markers of EMT phenotype NCAD, ECAD, and VIM were detected by immunofluorescence. In KYSE450 cells, the protein level of ECAD was upregulated while NCAD, and VIM levels were reduced in the siRNA groups. Parallel results were observed in KYSE70 cells (Figure 4E). Western blot was also used to analyze the protein levels of these EMT markers, and we found that siRNA groups had higher ECAD expression, while expressions of NCAD, VIM, and TWIST were lower in siRNA groups (Figure 4F). All these results revealed that PKMYT1 could accelerate ESCC cells migration and invasion by EMT.
Down-regulated PKMYT1 suppressed Akt/mTOR signaling pathway in ESCC cells

To investigate the further mechanisms of how down-regulated PKMYT1 inhibited ESCC cells proliferation and migration, we detected the expressions of Akt/mTOR signaling pathway in ESCC cells by Western blot. Western blot confirmed that down-regulated PKMYT1 significantly decreased the expressions of p-Akt, p-mTOR, and p-S6 (Figure 5), suggesting down-regulated PKMYT1 might inhibit Akt/mTOR signaling pathway in ESCC cells and thus weakened the basic biological functions in ESCC cells in vitro.

Discussion

ESCC is one of the most malignant tumors with high recurrence and high mortality.5,6 Due to the unique anatomical structure of esophagus, metastasis in early-stage ESCC patients and recurrence after surgery are common, which result in the lower 5-year overall survival than the ones in other gastrointestinal cancers.27 Hence, it is necessary to explore molecular mechanism and discover novel therapeutic targets for the ESCC treatment.

PKMYT1 plays a negative role in the G2/M phase of cell cycle and has a great impact on the normal mitosis of tumor cells.9,12 However, to our knowledge, there is no report on its expression and specific role in ESCC. In this study, we found that PKMYT1 was highly expressed in ESCC patients and ESCC cell lines. To investigate the potential role of PKMYT1 in ESCC, we down-regulated PKMYT1 expression in vitro by siRNA and we found that it could inhibit the proliferation of ESCC cells. This is in line with previous findings in other tumors, such as hepatocellular carcinoma and colorectal
We hypothesized that aberrant PKMYT1 expression was involved in abnormal proliferation and tumorigenesis in ESCC. Recent research has shown that abnormal apoptosis is associated with tumor initiation and progress of ESCC.28 The key regulator in apoptosis-related pathway is Bcl-2 family, which has two representative molecules Bcl-2 and Bax, and the Bax/Bcl-2 ratio determines whether cells enter an apoptotic state or not. Our assays demonstrated that down-regulated PKMYT1 induced higher rate of cell apoptosis. Furthermore, we also found that the expression ratio of the apoptosis-related proteins Bcl-2/Bax decreased by Western blot. In cancer cells, G2 checkpoint is important for DNA repair because of the defective G1 checkpoint mechanism which is caused by a p53 mutation.10,11 We speculated that down-regulated PKMYT1 in ESCC cells might result in DNA-deficient in ESCC cells, thereby contributing to a large amount of apoptosis. Therefore, we consider that PKMYT1 inhibition might play an anti-tumor role in ESCC cells by inducing apoptosis, which might be a potential therapy in the treatment of ESCC.31

The characteristics of invasion and metastasis of malignant tumors are the main factors related to tumor-specific death.32,33 The EMT is one of the most important events associated with tumor metastasis, and EMT is closely related to disease progression and tumor metastasis of ESCC.34,35 The biomarkers of EMT changed during tumor progression and metastasis, such as the downregulation of ECAD and upregulation of NCAD, VIM, and TWIST.36,37 Substantial evidence has indicated the importance of EMT in cancer proliferation, metastasis, and chemoresistance which makes it a potential therapeutic target for tumor treatment.38 In our assays, we detected decreased migration and invasion abilities in ESCC cell lines after PKMYT1 knockdown. Moreover, PKMYT1
Thus, PKMYT1 may participate in EMT of ESCC cells. Several studies indicated AKT/mTOR signaling pathway is a crucial target in the tumor progression of ESCC. The down-regulated of PKMYT1 expression in ESCC cells could inhibit cell proliferation, promote cell apoptosis, and restrict the EMT phenotype in vitro. Besides, the PKMYT1 downregulation also suppressed the Akt/mTOR/S6 signaling pathway in ESCC cells. These results demonstrated that PKMYT1 acted as an oncogene that might promote tumor progression by the AKT/mTOR pathway, and might be a potential target in ESCC treatment.

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Disclosure
The authors report no conflicts of interest in this work.

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