TOPK promotes metastasis of esophageal squamous cell carcinoma by activating the Src/GSK3β/STAT3 signaling pathway via γ-catenin

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Abstract

Background: Esophageal squamous cell carcinoma (ESCC) is a fatal disease with poor prognosis. The predominant reason for ESCC-related death is distal metastasis. A comprehensive understanding of the molecular mechanism underlying metastasis is needed for improving patient prognosis. T-LAK cell-originated protein kinase (TOPK) is a MAPKK-like kinase, which plays a vital role in various physiological and pathophysiological processes. However, the role of TOPK in ESCC metastasis is unclear.

Methods: Tissue array was used to evaluate the correlation between TOPK expression and ESCC lymph node metastasis. Wound healing assay, transwell assay, and lung metastasis mice model were used to examine the role of TOPK in the migration of ESCC cells in vitro and in vivo. Protein kinase array, mass spectrometry (MS), and molecular modeling were used to examine the pathways and direct target proteins of TOPK that are involved in ESCC metastasis. Additionally, immunofluorescence and western blotting analyses were performed to verify these findings.

Results: The enhanced expression of TOPK was correlated with lymph node metastasis in the ESCC tissues. TOPK knockdown or treatment with the TOPK inhibitor (HI-TOPK-032) decreased the invasion and migration of ESCC cells in vitro. HI-TOPK-032 also inhibited the lung metastasis in ESCC cell xenograft in vivo model. Moreover, TOPK promoted the invasion of ESCC cells by activating the Src/GSK3β/STAT3 and ERK signaling pathways via γ-catenin.

Conclusion: The findings of this study reveal that TOPK is involved in ESCC metastasis and promoted the ESCC cell mobility by activating the Src/GSK3β/STAT3 and ERK signaling pathways. This indicated that TOPK may be a potential molecular therapeutic target for ESCC metastasis.

Keywords: Esophageal squamous cell carcinoma, TOPK, Tumor metastasis, Src/GSK3β/STAT3, γ-Catenin

Background

Globally, esophageal cancer is the sixth leading cause of cancer-related death [1]. Esophageal cancer is classified as adenocarcinoma (EAC) or squamous cell carcinoma (ESCC) based on the histological examination [2]. In recent decades, there has been a drastic increase in EAC incidence in western countries. ESCC is the main histological type, which accounts for about 80% of esophageal cancer cases in China. Most patients with ESCC are diagnosed at advanced stages of the disease. The patients with ESCC exhibit a low 5-year survival rate (10–15%), which is mainly due to the lack of effective drugs [2]. Moreover, ESCC frequently metastasizes to the liver, lungs, regional lymph nodes, bone, and adrenal glands in the advanced stage [3].

Several wide-spread genomic alterations have been detected in ESCC using high-throughput sequencing technologies [2, 4]. There are 79 key genes that promote...
the development of ESCC. These genes are involved in 12 signal transduction pathways and are associated with genome stability, cell survival, and cell fate in ESCC. Recent studies revealed that the aberrant activation of various signaling pathways, such as PI3KCA and NOTCH1 pathways promote the development of ESCC and contribute to several processes of ESCC, such as metastasis and proliferation [2]. Therefore, there is an urgent need to identify an effective target in these pathways for the clinical treatment of ESCC. Additionally, the molecular mechanisms underlying ESCC metastasis must be elucidated. LAK cell–originated protein kinase (TOPK) is a serine-threonine kinase, which is involved in various physiological and pathological processes, such as cell cycle [5], apoptosis [6], cancer cell proliferation [7], and cell invasion [8]. Several studies have indicated that TOPK is an oncogenic kinase that is highly expressed in several types of human cancers, such as lung, colon, and ovary cancers. TOPK expression is positively correlated with the aggressive phenotype of these tumors [9, 10]. However, the expression and function of TOPK in ESCC metastasis are unclear. Hence, it is important to elucidate the underlying mechanism of TOPK in ESCC.

Methods

Chemical reagents

Tris, NaCl, and sodium dodecyl sulfate (SDS) used for molecular biology experiments and buffer preparation were purchased from Sigma-Aldrich. The following antibodies for western blotting analysis were purchased from Cell Signaling Technology: anti-Stat3 (#9139), anti-p-Stat3 (#9145), anti-Src (#2109), anti-p-Src (#12432), anti-Erk (#4696), anti-p-Erk (#4370), anti-GSK3β (#5676), and anti-p-GSK3β (#9327) antibodies. The anti-TOPK (ab236872), anti-p-TOPK (ab250790), and anti-β-catenin (ab32572) antibodies were purchased from Abcam. The antibodies against β-actin (sc-47,778) and tubulin (sc-73,242) were purchased from Santa Cruz Biotechnology. The human phospho-kinase arrays were obtained from R&D Systems. RPMI-1640, Dulbecco’s modified Eagle’s medium (DMEM), and minimum essential medium (MEM) were purchased from Bi (Bioind, Israel). Fetal bovine serum (FBS) was purchased from Gibico. The TOPK inhibitor, HI-TOPK-032 was a gift from Prof Zigang Dong. The stock solution of HI-TOPK-032 (50 mM) was prepared in dimethyl sulfoxide (DMSO) and stored at −20 °C. The working solution of HI-TOPK-032 was prepared by diluting the stock solution to 1000 times and used for in vitro treatment.

Human ESCC tissue array

The tissue microarray was purchased from Shanghai Outdo Biotech Co., Ltd. The information on clinical stages (stage I-IV) (AJCC 7.0) and TNM score was obtained from the patient clinical data. The tissue microarray comprises 10 paired ESCC and adjacent normal tissues with lymph node-negative tissue (N0), 19 N1 ESCC tissues, 10 N2 ESCC tissues, 8 N3 ESCC tissues with lymph node-positive tissue. The expression of TOPK in the ESCC metastasis tissue was evaluated by immunohistochemical staining. The tissue microarray was washed and incubated with the secondary antibody. The proteins were detected using DAB. The positive cells were graded under a microscope (20 X) and analyzed using the Image J software.

Cell culture

The following ESCC cell lines were purchased from Shanghai Xinyu Biological Technology Co., Ltd. (Shanghai, China): KYSE450 (XY-H600), KYSE510 (AD0604), KYSE410 (AD0602), KYSE70 (XY-H601), and KYSE30 (AD0601). These cells have been authenticated by STR technology of GENEWIZ Co., Ltd. (Suzhou, China) and no mycoplasma contamination. The KYSE450 cell line was cultured in DMEM supplemented with 10% FBS. The other ESCC cell lines were cultured in RPMI-1640 medium supplemented with 10% FBS, 100 μg/mL penicillin, and 100 μ/mL streptomycin at 37 °C and 5% CO2.

Western blot

The cells were lysed in the RIPA buffer [50 mM Tris-HCl (pH 7.4), 150 mM NaCl, 1 mM EDTA, 1% NP-40, 0.1% SDS, 0.5% sodium deoxycholate, 1 mM Na3VO4, and complete EDTA-free protease inhibitor (Roche)] for 20 min on ice. The cells were centrifuged and the supernatant was subjected to SDS-polyacrylamide gel electrophoresis (SDS-PAGE) using 10% gel. The resolved proteins were blotted onto a polyvinylidene difluoride membrane (Millipore, USA). The proteins were developed using the enhanced chemiluminescence (ECL) western blotting detection reagent (GE, USA) and analyzed using the LAS-1000UVmini image analyzer (GE, USA).

 Lentiviral infections

The lentiviral TOPK constructs, shTOPK#1 (ATTAGT GCATAAGAGAAGGTT) and shTOPK#2 (GTCTGT GTCTTGCTATGGAAT) were gifts from Prof Zigang Dong. The lentiviruses were engineered by co-transfecting the shRNA-expressing vector with pSPAX2 and pMD2.G constructs into the 293 T cells using jetPrime reagent. The viral supernatant was harvested and used to infect the cells along with 8 μ/mL polybrene. The cells were selected using 2 μg/mL puromycin to establish the stable KYSE510 and KYSE30 TOPK knockdown cell lines.

Wound healing assay

For wound healing assays, the shMock-, shTOPK#1-, and shTOPK#2-transfected KYSE510 and KYSE30 cells
were seeded on 6-well plates containing culture medium at 80% confluency and incubated for 24 h. The confluent monolayer was scratched using a fine pipette tip. The cells were washed twice with phosphate-buffered saline (PBS pH 8.0). The cell debris was removed and the cells were incubated with the growth medium containing 1% FBS. The migration of cells to the injured blank area was evaluated by capturing the images at 0, 24, and 48 h.

Transwell migration assay
Transwell migration assay was performed using the transwell inserts for a 24-well plate. The upper chamber membrane (8 μm; Millipore) was coated with 100 μL of 200 μg/mL Matrigel (BD Biosciences, Franklin Lakes, NJ, USA) at 37 °C for 1 h before adding the cells. The shMock-,
shTOPK#1-,
and shTOPK#2-transfected KYSE510 and KYSE30 cells were plated in culture medium without serum or growth factors in the upper chamber of 1 × 10^5 cells. The lower chamber contained medium supplemented with 20% FBS, which served as a chemo-attractant. After incubating for 24 h, the cells that did not migrate or invade through the pores were removed using a cotton swab. The cells on the lower surface of the membrane were fixed with methanol for 10 min and stained with 0.1% crystal violet. The cells were then counted in 5 randomly selected microscopic fields (×100) in each well.

Protein kinase array
The phosphorylated proteins were analyzed using the human phospho-kinase arrays, following the manufacturer’s instructions. Briefly, the cell lysates (500 μg) of shMock-,
shTOPK#1-,
and shTOPK#2-transfected KYSE510 cells were collected and incubated with each array at 4 °C overnight on a rocking platform shaker. The cell lysate was removed the next day and the arrays were washed thrice with washing buffer. The arrays were incubated with the secondary antibody for 2 h at room temperature (25 °C) and washed thrice with washing buffer. The array was incubated with horseradish peroxidase (HRP) solution on a rocking shaker for 30 min. The array was then washed thrice with washing buffer and the protein spots were visualized using a chemiluminescence detection kit. The density of each duplicated array spot was assessed using the Image J software (v.1.37v, NIH). The density of the sample was calculated after subtracting the background density and the density of negative control (PBS). The expression level in the shTOPK#1- and shTOPK#2-transfected cells was normalized (indicated as a value of 1) to that of the shMock-transfected cells.

Surface plasmon resonance (SPR)
The glutathione-S-transferase (GST)-tagged TOPK proteins were purified from BL21 bacteria using the glutathione-sepharose 4B beads. The anti-GST antibody was immobilized onto the CM5 chip by amine coupling, following the manufacturer’s instructions (GE). The recombinant TOPK (Signalchem, T14-10G-10) and purified GST-tagged TOPK protein (> 90%) were captured by the anti-GST antibody on the CM5 chip. The KYSE510 cell lysate was introduced into the CM5 chip at a flow rate of 10 μL/min for 5 min with running buffer (1X kinase buffer, pH7.2) for one cycle and the captured protein was recycled. The control group did not contain recombinant TOPK or GST-TOPK protein. The SPR procedure was performed at Bicore T200 (GE, USA).

Nanoflow liquid chromatography-electrospray ionization-tandem mass spectrometry (LC-ESI-MS/MS)
The captured proteins bound to the recombinant TOPK, GST-TOPK (TOPK protein was purified by GST tag), and negative group (KYSE510 lysis buffer) in the SPR were subjected to mass spectrometry (MS). The samples were subjected to reduction by incubation with dithiothreitol (DTT) (final concentration 10 mmol/L) for 1 h at 37 °C. Next, the samples were subjected to alkyla-tion by incubation with iodoacetamide (final concentration 30 mmol/L) at room temperature for 45 min. Finally, the samples were diluted with 25 mmol/L ammonium hydrogen carbonate buffer to ensure that the urea concentration was below 2 mol/L and incubated with sequencing grade trypsin at a mass ratio of 1:50 at 37 °C for 14 h. The LCS-ESI.MS/MS analysis was performed using freeze-dried samples dissolved in 60% acetonitrile (ACN) containing 0.1% formic acid. The sample was passed through a R3 self-loading desalting column conditioned with 99.9% acetonitrile (ACN) and 0.1% trifluoroacetic acid (TFA) twice. Next, the samples were subjected to gradient elution as follows: 0.1% TFA, 1% TFA, with 30% ACN, 60% ACN, 30% ACN. The MS/MS conditions were as follows: mobile phase, solution A (0.1% formic acid, 99.9% H2O), solution B (0.1% formic acid, 99.9% ACN); sample volume, 10 mL; continuous gradient elution; flow rate, 200 nL/min. The analytical column gradient elution settings for the LTQ Orbitrap elite and the MS parameters were as follows: setting mode, cation; collision model, collision-induced dissociation (CID); resolution, 120,000; mass range, 350–1800, before top 15 ionic strength. The data were analyzed using the Thermo Proteome Discoverer software (1.4.0.288) with the following settings: databases, Mascot and UniProt; enzyme, trypsin-P; maximum missed cleavage, 2; precursor mass tolerance, 10 ppm; fragment mass tolerance, 0.5 Da; fixed modifications, carbamidomethyl (c); variable modification, Gln > pyro-Glu (N-term q), acetyl (protein N-term), oxidation (M), Phospho (st), and Phospho (y); peptide confidence, high false discovery rate (FDR) < 0.01.
Molecular modeling
To confirm whether TOPK binds to γ-catenin, in silico docking was performed using the Schrödinger Suite 2016 software. The TOPK and γ-catenin crystal structures were obtained from the Protein Data Bank [11] (PDB ID: 5F0A and 3IFQ, respectively) and prepared according to the standard procedures of the Protein Preparation Wizard (Schrödinger Suite 2012). The protein-protein docking of TOPK and γ-catenin was performed using the protein docking server with the interactive molecular graphics program [12]. The best configuration was selected to represent the binding mode.

Immunofluorescence assay
Immunofluorescence assay was performed using the primary antibodies against TOPK and γ-catenin. The samples were then incubated with the fluorescein-conjugated affinity pure goat anti-mouse IgG or rhodamine (TRITC)-conjugated affinity pure goat anti-rabbit IgG secondary antibodies (1:2000) for 1 h. The samples were mounted on coverslips along with 4′, 6-diamidino-2-phenylindole (DAPI) (Santa Cruz TM). The images were captured and analyzed using high-throughput confocal microscopy (IN Cell Analyzer 6000, GE).

Co-immunoprecipitation
For co-immunoprecipitation studies, the HEK293 cells were transfected with 4 μg pcDNA3-HA-TOPK and/or pDEST-myc-γ-catenin using jetPRIME reagent (Polyplus-transfection® SA, France) for 72 h. The protein was isolated in CHAPS buffer (as previously described) supplemented with protease inhibitor cocktail (Roche). The lysates were centrifuged at 14500 g for 30 min. Next, 500 μg protein in 500 μL was incubated with anti-hemagglutinin (HA) antibody overnight at 4 °C. The samples were then incubated with secondary antibodies (sc-2004, Santa Cruz) immobilized on A/G agarose (40 μL) for 4 h at 4 °C. The collected protein complexes were washed thrice with cold PBS and eluted by boiling in loading buffer at 95 °C, followed by incubation on ice for 2 min. The myc protein was resolved by SDS-PAGE and analyzed by western blotting.

Lung metastasis in ESCC cell xenograft mouse models
The stable GFP-KYSE510 cells were established by transfecting the pcDNA3.1-green fluorescent protein (GFP) vector and screened using G418. The GFP signal of KYSE510 cells was evaluated using the IVIS® Lumina III In Vivo Imaging System. Next, the GFP-KYSE510 cells (2 × 10⁶ cells/mL) were injected into the tail vein of BALB/c nude mice, which were purchased from Vital River, Beijing, China. After two weeks, these mice were divided into vehicle and treatment groups. The vehicle group was treated with 5% DMSO-PBS (n = 12, 6 females and 6 males), while the treatment group was injected with HI-TOPK-032 (n = 13, 7 females and 6 males). The mice in the treatment group were treated with HI-TOPK-032 was dissolved in 5% DMSO-PBS and was given to each mouse in treatment group once daily at 10 mg/kg (i.p., 100 μL per mouse). Four to five mice were kept in a pathogen-free environment with light controlled rooms (12 h cycles) and provided with food and water ad libitum. After two weeks, the mice were subjected to in vivo imaging. The fluorescence of GFP was evaluated as described previously [13]. The mice were euthanized by CO₂ inhalation. The lung tumor tissues were used to evaluate the changes in the signaling pathways. This study was approved by the Ethics Committee of Basic Medical College of Zhengzhou University.

Results
TOPK was positively correlated with lymph node metastasis of ESCC patients
To investigate the clinical significance of TOPK in ESCC metastasis, the expression of TOPK was analyzed in the tissue samples of 49 patients with ESCC with or without the lymph node metastasis. The expression of TOPK was detected mainly in the cytoplasm and/or cell nucleus. The TOPK expression varied depending on the lymph node metastasis type (Fig. 1a). The TOPK expression level in the lymph node metastasis groups (N1 and N2–3 groups) was significantly (p < 0.001) higher than that in the no lymph node metastasis group (N0 group). This indicated a positive correlation between TOPK expression and lymph node metastasis (Fig. 1b). Additionally, the expression of TOPK varied in different ESCC cell lines. The expression levels of TOPK in the KYSE510, KYSE140, and KYSE30 cells were higher than those in the KYSE450 and KYSE70 cells (Fig. 1c-d).

TOPK knockdown attenuated the migration and invasion of ESCC cells
The TOPK knockdown stable cell lines were established by transfecting shTOPK#1 and shTOPK#2 into the KYSE510 (Fig. 2a) and KYSE30 (Fig. 2b) cells. The wound healing assay revealed that TOPK knockdown inhibited the migration of KYSE510 (Fig. 2c) and KYSE30 cells (Fig. 2d). The transwell migration assay indicated that TOPK knockdown
decreased the invasion of KYSE510 (Fig. 2e) and KYSE30 (Fig. 2f) cells. The role of TOPK in the metastasis of ESCC was evaluated by treating the KYSE510 and KYSE30 cells with HI-TOPK-032, which inhibits the TOPK activity. Treatment with HI-TOPK-032 inhibited the migration of KYSE510 (Fig. 3a) and KYSE30 (Fig. 3b) cells. Moreover, HI-TOPK-032 dose-dependently decreased the invasion and migration of KYSE510 (Fig. 3c) and KYSE30 (Fig. 3d) cells after treatment for 24 h. These data suggested that TOPK was involved in ESCC cell mobility.

Src/GSK3β/STAT3 and ERK signaling pathways were strongly inhibited by knocking down TOPK or by HI-TOPK-032 treatment

The human phospho-kinase array (ARY003B, R&D) was used to examine the signaling pathway that mediates the TOPK-regulated ESCC metastasis. The effect of TOPK knockdown in the KYSE510 cells on the signal transduction pathways was evaluated using the human phospho-kinase array. Several signaling pathways were affected in the shTOPK#1- and shTOPK#2-transfected KYSE510 cells when compared with the shMock-transfected KYSE510 cells (Fig. 4a). The phosphorylation level of STAT3 Y705 markedly decreased to 0.35 and 0.11 in the shTOPK#1- and shTOPK#2-transfected KYSE510 cells, respectively. The phosphorylation levels of Src Y419, GSK3β kinase S21/S9, and ERK1/2 T202/Y204 in the shTOPK#1-transfected KYSE510 cells decreased to 0.71, 0.65, and 0.81, whereas those in the shTOPK#2-transfected cells decreased to 0.65, 0.66, and 0.79, respectively (Fig. 4a). These data indicated that TOPK might mediate the crosstalk between different signaling pathways and that TOPK played a complex role in ESCC metastasis. To further verify the phospho-kinase array results, the same cell lysates were subjected to western blotting analysis. The western blotting analysis revealed that the phosphorylation of STAT3 Y705, GSK3β S21, Src Y419, and ERK1/2 T202/Y204 was strongly inhibited upon TOPK knockdown in the KYSE510 and KYSE30 cells (Fig. 4b), which concurred with the array data. Next, the KYSE510 and KYSE30 cells were treated with different doses of HI-TOPK-032 for 24 h. The western blotting analysis revealed that HI-TOPK-032 dose-dependently inhibited the p-TOPK, STAT3, Src/GSK3β, and ERK signaling pathways (Fig. 4c). These findings indicated that TOPK may promote
the invasion and migration of ESCC cells by activating several signaling pathways, such as the Src/GSK3β/STAT3 and ERK signaling pathways. This indicated that TOPK may mediate the crosstalk between various signaling pathways during ESCC metastasis.

**Fig. 2** Knocking down TOPK attenuates the migration and invasion of esophageal squamous cell carcinoma (ESCC) cells. Knocking down TOPK expression in the KYSE510 (a) and KYSE30 cells (b) by lentiviral infection. Cell migration was evaluated at 0, 24, and 48 h in the stable TOPK knockdown cells (shTOPK-transfected KYSE510 (c) and KYSE30 cells (d)) by wound healing migration assay. The invasion of KYSE510 (e) and KYSE30 (f) cells was measured by transwell migration assay after knocking down TOPK. ***p < 0.001 vs shMock group

TOPK interacts with multiple proteins and binds with γ-catenin to enhance the invasion and migration of ESCC cell lines

SPR and MS were performed to screen the potential TOPK target proteins. The proteins with a score higher...
than 25 in MS analysis were chosen for further analysis. There were 53 and 39 proteins in the recombinant TOPK and GST-TOPK protein groups, respectively (Fig. 5a). Among these proteins, 39, 25, and 14 proteins bound to the recombinant TOPK, GST-TOPK, and both recombinant TOPK and GST-TOPK, respectively. Among the 14 proteins that bound to both recombinant TOPK and GST-TOPK, plakophilin-1, dermcidin, EEF1A had scores of 81.95, 65.91, and 41.6, respectively, and were associated with cell migration and invasion (Fig. 5b). Among the migration-related proteins, γ-catenin had the highest score (184.85) (Fig. 6a). Molecular modeling was also performed to understand the detailed binding interactions between TOPK and γ-catenin (Fig. 6b). The binding of TOPK to γ-catenin was evaluated by transfecting pcDNA3-HA-TOPK and/or pDEST-myc-γ-catenin into the HEK293 cell for 72 h. Immunoprecipitation analysis using the anti-HA antibody revealed that γ-catenin interacted with TOPK (Fig. 6c). Immunofluorescence analysis indicated that TOPK knockdown induced the expression of γ-catenin in the KYSE510 and KYSE30 cells (Fig. 6d). Furthermore, HI-TOPK-032 dose-dependently increased the expression of γ-
catenin in the KYSE30 cells (Fig. 6e). These data demonstrated that TOPK interacts with several proteins, including γ-catenin, plakophilin-1, dermcidin, and EEF1A to promote the migration and invasion of ESCC cells.

The TOPK inhibitor, HI-TOPK-032 suppressed ESCC lung metastasis in a KYSE510 cell xenograft mouse model

Next, we evaluated the effect of TOPK on ESCC metastasis in vivo. Treating the athymic nude xenograft mouse model with HI-TOPK-032 suppressed lung metastasis without affecting the bodyweight in vivo (Fig. 7a-c). The western blotting analysis revealed that treatment with HI-TOPK-032 inhibited the expression of p-TOPK and the activation of Src, GSK3β, and ERK signaling pathways (Fig. 7d). This indicated that HI-TOPK-032 exerted an inhibitory effect on the metastasis of KYSE510 cells in the xenograft model by suppressing the Src, GSK3β, and ERK signal transduction pathways in vivo.

Discussion

Metastasis is a complex multi-stage process and is the most lethal hallmark of ESCC [14]. Tumor metastasis involves enhanced cell motility. The cancer cells are transmitted from the primary tumor to the surrounding tissues and distant organs through the lymphatic and blood vessels. Clinically, tumor metastasis is mostly incurable and is the primary cause of cancer-related morbidity and mortality. Over 50% of ESCC-associated
deaths are caused by distal metastasis, which is associated with poor prognosis, to the lymph nodes, lung, liver, bones, adrenal glands, and brain [15]. Therefore, identification of critical regulators that drive invasion and metastasis in ESCC will facilitate the development of new treatment strategies.

TOPK was first discovered in the lymphokine-activated killer T cells and is a branch of the MEK1/2 and MKK7 phylogenetic tree [16, 17]. Several studies have revealed a correlation between TOPK expression and poor prognosis in numerous cancers, such as leukemia [18], breast cancer [19, 20], ovarian cancer [21], cervical cancer [22], lung cancer [23], colon cancer [24], and oral cancer [9]. The enhanced expression of TOPK was reported to be associated with tumor aggressiveness, invasion, and metastasis [25]. Recent studies indicated that TOPK is a potential cancer-specific biomarker and a potential therapeutic target for cancer that can be used without markedly affecting the normal tissues [26]. However, there are no systemic studies that have evaluated the TOPK function in ESCC metastasis. In this study, the expression of TOPK in the N1–2 and N3 ESCC tissues was higher than that in the N0 ESCC tissues. The knockdown of TOPK inhibited the migration and invasion of ESCC cells. Additionally, treatment with HI-TOPK-032 inhibited the motility of ESCC cells in vitro and lung metastasis in vivo. These data indicated that TOPK overexpression was associated with ESCC metastasis and that the TOPK expression level may be a potential biomarker for ESCC. Therefore, it is important to explore the TOPK as a therapeutic target for ESCC.

Various studies indicated that the abnormal expression of TOPK in many cancers triggered aberrant activation of different signaling pathways, such as the ERK1/2, p38, c-Jun, and NF (nuclear factor)-κB signaling pathways [27]. In this study, the ERK signaling pathway was activated by TOPK, which concurred...
with the results of other studies. The results of this study also indicated that TOPK may mediate the crosstalk between several signaling pathways during ESCC metastasis, such as Src, GSK3β, and STAT3 signaling pathways. Shegan Gao et al. [28] reported that enhanced GSK3β expression was associated with higher metastasis rates and poor prognosis of ESCC. Additionally, the study demonstrated that STAT3 was a target of GSK3β using cancer phospho-antibody array and that enhanced GSK3β expression promoted ESCC progression through STAT3 in vitro and in vivo. These results indicated that GSK3β-STAT3 signaling could be a potential therapeutic target for ESCC treatment. However, Src was also overexpressed and functionally relevant to the progression of human ESCC. This indicated that Src might be a useful...
molecular target for ESCC prognosis and treatment [29]. Moreover, some studies demonstrated that Src-mediated Y216 GSK-3β phosphorylation and activation increased prostate cancer cell motility, proliferation, micrometastasis, and progression [30]. This suggested that GSK-3β activity can be modulated by Src kinases and that the pharmacological inhibition of Src-GSK-3β pathway may be a useful therapeutic strategy for prostate cancer. In this study, TOPK knockdown decreased the phosphorylation levels of Src, GSK3β, STAT3, and ERK proteins in the ESCC tissues. To the best of our knowledge, this is the first study to report the regulation of Src/GSK3β/STAT3 signaling pathway by TOPK, which is involved in the invasion process of ESCC in vitro and in vivo. Thus, TOPK may be vital for promoting the progression of ESCC. Therefore, TOPK as a therapeutic target may have unexpected effects by inhibiting other signaling pathways in ESCC.

To identify the direct target of TOPK that mediates the progression of ESCC, the proteins that bind to the purified TOPK protein and recombinant TOPK in the KYSE510 cell lysates were screened by SPR and MS. Several proteins bind to the recombinant TOPK, such as plakophilin-1, dermcidin, and EEF1A proteins. Among these TOPK-binding proteins, γ-catenin had the highest score. Therefore, we focused on evaluating the correlation between TOPK and γ-catenin. γ-catenin (also known as junction plakoglobin) is a desmosomal protein and may play a central role in desmosomes organization. Additionally, γ-catenin is required for the effective intermediate filament anchorage to desmosomes. γ-catenin was reported to be involved in cell invasion and metastasis of human oral squamous cell cancer, lung cancer, and bladder cancer [31]. Some studies have reported that γ-catenin functions as a tumor suppressor in ESCC and that the γ-catenin expression levels affected the overall survival. This indicated that low γ-catenin was associated with lymph node metastasis and may serve as a prognostic marker in human ESCC [32]. The results of this study demonstrated that TOPK knockdown or treatment with HI-TOPK-032 rescued the expression of γ-catenin in the ESCC cells. Moreover, TOPK directly binds to γ-catenin, which results in the dissociation of desmoplakin. Therefore, this suggested that
downregulating TOPK may increase and stabilize γ-catenin and subsequently decrease the cell adhesion and motility.

Similar to its homolog β-catenin, γ-catenin is also reported to be involved in several cell signaling cascades during cell motility, cell proliferation, and apoptosis [33]. All these processes involve the interaction of cytoplasmic γ-catenin with various kinases and the inhibition of Wnt – β-catenin signaling by nuclear γ-catenin. The silencing of γ-catenin but not desmoplakin is reported to activate the p38-dependent keratin filament collapse and cell dissociation [34]. However, the signaling mechanisms underlying the tumor suppressor activity of γ-catenin in ESCC cells are not well understood. Recent studies demonstrated that γ-catenin knockout cells exhibited enhanced activation of Src signal pathways and that γ-catenin regulated cell motility through distinct Src and Rho GTPase-dependent pathways [35]. Therefore, decreased γ-catenin may be the primary reason for activating the Src signal pathways during ESCC metastasis.

The following model illustrates the promotion of ESCC cell metastasis by TOPK (Fig. 8): TOPK binds γ-catenin and the deregulated γ-catenin activates the Src/GSK3β/STAT3 signaling pathway. TOPK also activates the ERK and Src/GSK3β/STAT3 signal pathway. These two activated pathways increase the cancer cell invasion and mobility and subsequently promote ESCC metastasis. Further, the TOPK inhibitor, HI-TOPK-032 inhibits the activated Src/GSK3β/STAT3 and ERK signaling pathways resulting in decreased ESCC metastasis.

**Conclusion**

TOPK is involved in ESCC metastasis and is a potential molecular therapeutic target for ESCC. This study demonstrated that TOPK promotes the metastasis of ESCC cells by activating the Src/GSK3β/STAT3 signal pathway via γ-catenin.
Abbreviations
ESC C: Esophageal squamous cell carcinoma; IHC: immunohistochemical staining; MS: mass spectrometry; SEM: standard error of mean; SPR: surface plasmon resonance; STAT3: signal transducer and activator of transcription 3; TOPK: T-LAK cell-originated protein kinase

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Authors’ contributions
KL, ZD, and YJ conceived of the study and participated in its design and coordination and drafted the manuscript. JZ1 (Jiang Zhang) and JZ2 (Jimin Zhao) participated in the design of the study and helped to correct the drafts. ZL carried out the animal studies. XC performed the immunoassays, HC and YQ performed molecular modeling experiments. All authors read and approved the final manuscript.

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Availability of data and materials
All data generated or analyzed during this study are included in this published article.

Ethics approval and consent to participate
The study was approved and supervised by the research ethics committee of Zhengzhou University, Zhengzhou, China. All animal experiments were performed in accordance with the animal experimental guidelines of Zhengzhou University.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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