Sphingosine kinase 1 promotes the metastasis of colorectal cancer by inducing the epithelial-mesenchymal transition mediated by the FAK/AKT/MMPs axis

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Abstract. It was demonstrated that Sphingosine kinase 1 (SphK1) promotes tumor progression and confers the malignancy phenotype of colorectal cancer by activating the focal adhesion kinase (FAK) pathway. However, further clarification is required to determine if SphK1 promotes the metastasis of colorectal cancer by inducing epithelial-mesenchymal transition (EMT), and its mechanisms have not been fully elucidated. Immunohistochemistry staining was used to detect protein expression in normal colonic mucosa tissues and colorectal cancer tissues. Cells were transfected to overexpress SphK1, downregulate SphK1 or downregulate FAK. An MTT assay was used to detect the drug toxicity to cells. Transwell and wound healing assays were used to detect cell migration ability. Reverse transcription-polymerase chain reaction and western blot analysis were used to detect the expression of mRNA and protein, respectively. Scanning electron microscopy was used to observe the microvilli and pseudopodia of the cells. The analysis of protein expression in 114 human colorectal cancer tissues demonstrated that the expressions of SphK1, FAK, phosphorylated (p)-FAK, E-cadherin and vimentin were associated with the metastasis of colorectal cancer. Furthermore, the patients with colorectal cancer with SphK1-positive cancer demonstrated poorer prognosis compared with SphK1-negative cancer. FAK knockdown and SphK1 knockdown of human colon cancer RKO cells inhibited the EMT and migrational potency, along with the expression of p-FAK, p-protein kinase B (AKT) and matrix metalloproteinase (MMP)2/9. In contrast, SphK1 overexpression promoted EMT, migrational potency, and the expression of p-FAK, p-AKT and MMP2/9 in HT29 cells. Additionally, the EMT and migrational potency of SphK1-overexpressing HT29 cells was suppressed by a FAK inhibitor, and the expression of p-FAK, p-AKT and MMP2/9 was suppressed by blocking the FAK pathway. In conclusion, SphK1 promoted the migration and metastasis of colon cancer by inducing EMT mediated by the FAK/AKT/MMPs axis.

Introduction

Poor prognosis and a lack of effective therapeutic strategies pose principal challenges for the treatment of colorectal cancer (1). There is an urgent requirement to identify a better therapeutic target in the treatment of colorectal cancer.

Sphingosine kinase 1 (SphK1) is involved in the regulation of cellular behaviors. Accumulating evidence suggested that the activation of SphK1 contributes to tumor growth, neovascularization, metastasis and drug resistance (2). SphK1 is overexpressed in multiple types of human cancer tissues, including colorectal cancer tissues (3). Furthermore, migrational potency of cancer cell was improved by the overexpression of SphK1 and decreased by the knockdown of SphK1 (4). Our previous study demonstrated that the expression of SphK1 in primary colorectal cancer tissues was significantly increased compared with matched normal tissues (5). A further previous study suggested that the migrational potency of colon cancer LOVO cells was enhanced by the overexpression of SphK1, and inhibited by suppression of SphK1 via short hairpin (sh)RNA transfection (6). These results suggested that SphK1 may serve an important role in promoting the migration and metastasis of colorectal cancer. However, the precise molecular mechanism still requires investigation.

Emerging evidence suggested an association between the development of metastasis and epithelial-mesenchymal transition (EMT) processes in cancer (7). EMT is defined as the process of epithelial cell transformations towards

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the mesenchymal phenotype that results in metastasis (8). Accordingly, epithelial tumor cells were associated with the reactivation of EMT (9). EMT was characterized by a downregulation of epithelial markers, including E-cadherin (10), an upregulation of mesenchymal markers, including vimentin (11) and fibronectin (12), and the production of matrix-degrading enzymes, including matrix metalloproteinase (MMP)2 and MMP9 (13,14). The migrational potency of cells was facilitated once the EMT process was reactivated in epithelial tumor cells, including colon cancer cells (15). Previous studies demonstrated that focal adhesion kinase (FAK), phosphoinositide 3 kinase (PI3K)/protein kinase B (AKT) and MMP2/9 were involved in mediating the EMT process (16-19); however, the regulatory mechanism has not yet been fully understood.

It was confirmed that SphK1 promoted tumor progression and advanced malignancy phenotype of colon cancer by regulating the FAK pathway and upregulating the production of MMP2/9 (4). Recent studies suggested that the FAK pathway was one of the most important factors in regulating EMT (20,21). Furthermore, tyrosine phosphorylation of p85 subunits of PI3K was modulated by FAK (22), whereas, PI3K/AKT affected MMP2/9 secretion in cancer cells (23). Therefore, in the present study, the hypothesis that SphK1 promotes the migration and metastasis of colorectal cancer by EMT induction mediated by the FAK/AKT/MMPs axis was investigated.

Materials and methods

Tissue specimens. Tissues samples from patients were collected at the First Affiliated Hospital of Guangxi Medical University (Guangxi, China) from March 2013 to October 2013 (24). There were 71 male and 43 female patients included in the present study, all of whom were newly diagnosed and had not received any chemotherapy or radiotherapy previously, and the age of the patients ranged between 25 and 83 (56.5±13.47) years old (24). Based on the National Comprehensive Cancer Network classification, 19 patients were diagnosed with stage I CRC, 44 with stage II CRC, 30 with stage III CRC and 21 with stage IV CRC (24). The study was approved by the Institutional Ethics Committee of Guangxi Medical University under full consideration of the Helsinki declaration of human rights. Written informed consent was obtained from all patients.

Immunohistochemistry staining. Immunohistochemistry staining was performed on deparaffinized 4-µm thick sections according to the manufacturer’s protocol. Endogenous peroxidase activity was blocked with 3% hydrogen peroxide at room temperature for 10 min. Subsequently, the sections were blocked with 10% normal goat serum (Beyotime Institute of Biotechnology, Haimen, China) in PBS for 1 h, incubated with primary antibody for overnight 4°C, and finally incubated with undiluted horseradish peroxidase-conjugated secondary antibody for 30 min at 37°C. The secondary antibody was the B reagent in an immunohistochemistry kit (Zhongshan Golden Bridge Biotechnology, Beijing, China; cat. no. PV-9000). Visualization of the immunohistochemical reaction was performed using DAB as the chromogen. The slides were evaluated by two pathologists independently. Protein expression was examined by using five random fields on each slide at using a light microscope (magnification, x100; PH50-3A43L-PL optical microscope; Phmias2008 Computer Image Analysis System; Phoenix Optical Co., Ltd., Jiangxi, China). Polyclonal antibodies against SphK1 (cat. no. 10670-1-AP; 1:400) and FAK (cat. no. 12636-1-AP; 1:800) were from ProteinTech Group, Inc. (Chicago, IL, USA). Polyclonal antibodies against phosphorylated (p)-FAK (Tyr397; cat. no. 8556P; 1:1,200) were from Abcam (Cambridge, UK). Polyclonal antibodies against E-cadherin (cat. no. bs-1519R; 200) and vimentin (cat. no. bs-8533R; 1:200) were from BIOSS (Beijing, China). The degree of staining was determined by semi-quantitative evaluation and categorized by the extent and intensity of staining as follows: i) The percentage of positively stained cells <5% was scored as 0, 5-25% scored as 1, 26-50% scored as 2, >50% scored as 3; and ii) The intensity of staining was scored as: 0, achromatic; 1, light yellow; 2, yellow; and 3, brown. The two scores were added for the final score: 0-2, negative (-) and ≥3, positive (+).

Reverse transcription (RT)-quantitative (q)PCR analysis. RNA isolation was performed using the Total RNA Extraction kit (Tiangen Biotech Co., Ltd., Beijing, China), according to the manufacturer's protocol. cDNA synthesis was performed using the RT kit (Takara Bio, Inc., Otsu, Japan). RT was performed using 2.0 µl 5X gDNA Eraser Buffer, 1.0 µl gDNA Eraser, 1 µg Total RNA and RNase Free dH2O to a total volume of 10 µl. The reaction conditions were 42°C for 2 min and rapidly cooled to 4°C. Subsequently, 10.0 µl reaction solution, 1.0 µl PrimeScript RT Enzyme Mix 1, 1.0 µl RT Primer Mix, 4.0 µl 5X PrimeScript Buffer 2 and 4.0 µl RNase Free dH2O were used. The reaction conditions were 37°C for 15 min, 85°C for 5 sec, rapidly cooled to 4°C and stored at -20°C. A fluorescence-based qPCR method was performed using 2 µl cDNA, 10 µl SYBR Green (Takara Bio, Inc.), 0.6 µl PCR forward primer (10 µM), 0.6 µl PCR reverse primer (10 µM) and 6.8 µl dH2O in a 20 µl PCR reaction volume.
The RT-qPCR reaction was run on a StepOne Real-Time PCR system (Applied Biosystems; Thermo Fisher Scientific, Inc.). The cycling parameters were as follows: Denaturing at 95°C for 30 sec, 40 cycles at denaturing at 95°C for 5 sec, primer annealing at 60°C for 34 sec and extension temperature at 95°C for 15 sec; final extension at 60°C for 1 min and final denaturing at 95°C for 15 sec. Gene expression levels were determined using the 2^(-ΔΔCq) method (25). The primers were from Takara Bio, Inc. The sequences were as follows: GAPDH, forward: 5'-GCA CCG TCA AGG CTT AGA AC-3'; reverse: 5'-TGG TGA AGA CGC CAG TGG-3'; SphK1, forward: 5'-GGC TTT ATT GCT GCT TTT A-3'; reverse: 5'-AGG AAG GTG CCC AGA GTG AA-3'; FAK, forward: 5'-CAA CCA CCT GGG CCA GTA TTA TC-3'; reverse: 5'-CCA TAG CAG GCC ACA TGC TTAT-3'; vimentin, forward: 5'-TGA CAT TGA GAT TGC CAC CTA CAG-3'; reverse: 5'-TCA ACC GTC TTA ATC AGA AGT GTTC-3'; E-cadherin, forward: 5'-GAG TGC CCA CCT ACA ACT TTG AGAA-3', reverse: 5'-GCC TGT CGG TGA GAT TGG TT-3'; fibronectin, forward: 5'-GAC CAC ATC GAG CCG ATG TCA TG-3', reverse: 5'-GTC TCT TGG CAG CTG ACT CC-3'.

Western blot analysis. Total proteins were extracted using radioimmunoprecipitation buffer (Beyotime Institute of Biotechnology). Protein concentrations were measured by a bicinchoninic acid assay (Solarbio Biotech Co., Ltd., Beijing, China) according to the manufacturer’s protocol. A total of 30 µg of protein from each sample was separated via 12% SDS-PAGE (Beyotime Institute of Biotechnology) for 1 h at 100 V, and subsequently transferred onto nitrocellulose membranes. Samples were blocked with 5% non-fat milk in Tris-buffered saline with Tween-20 (Solarbio Biotech Ltd., Tokyo, Japan) and S4800 (Hitachi, Japan) for 1 h at room temperature. The membranes were incubated overnight at 4˚C with primary antibodies diluted in Tris-buffered saline with Tween-20 containing 5% non-fat milk. The membranes were washed with Tris-buffered saline, followed by incubation with anti-rabbit (cat. no. 60329-11; 1:10,000) secondary antibodies purchased from ProteinTech Group, Inc. IRDy™ 680RD goat anti-rabbit (cat. no. 60329-11; 1:10,000) secondary antibodies were from LI-COR Biosciences.

Cells viability. Cells (1x10^5) were seeded into a 96-well plate. Following treatment with the FAK pathway inhibitor (PF-562271) at concentrations of 0, 2.5, 5 and 10 µM for 24, 48 and 72 h, the cells were incubated in fresh medium containing MTT solution (2 mg/ml) and were incubated for 4 h at 37°C in the dark. Following removal of the MTT solution, 150 µl dimethyl sulfoxide was added to dissolve the formazan crystals and the absorbance (A) at 450 nm was measured using a fully automated microplate reader (ELx800™; BioTek Instruments, Inc., Winooski, VT, USA). Cell viability (%) = A450 (sample)/A450 (control) x100.

Transwell and wound healing assays. The cells (1x10^5) were cultured in fresh serum-free medium in the upper chambers of Transwell plates. The lower chambers contained fresh medium with 10% FBS. After 48 h of incubation at 37°C, cells remaining on the upper surface of the membrane were removed. The cells that migrated to the lower membrane surface were fixed with 4% paraformaldehyde (Beyotime Institute of Biotechnology) at 4°C for 5 min and stained with 0.1% crystal violet solution (Beyotime Institute of Biotechnology) at 37°C for 20 min. Migrating cells were counted in five random fields under a light microscope (Nikon Corporation, Tokyo, Japan; magnification, x200). The migrated cells were counted with Nikon microscope software (version 4.0; Nikon Corporation). Cells were seeded into a 6-well cell culture plate. When the cells reached 80-90% confluency, the cells were scratched with a pipette tip (1 mm). Images were acquired after cells were cultured in medium for 0, 24 and 48 h. The migration distance was measured with the microscope software (Nikon Corporation).

Scanning electron microscope analysis. Cells were grown on cover glass slides placed in 6-well plates. Subsequent to being treated, cells were fixed with 3% glutaraldehyde at 4°C for 2 h, gradient ethanol dehydration: 50% for 10 min one time and 70% for 10 min one time. Gold was used for sputter coating. Images of RKO cells and HT29 cells were obtained using the scanning electron microscope of VEGA3, TESCAN (Brno-Kohoutovice, Czech Republic) and S4800 (Hitachi, Ltd., Tokyo, Japan).

Statistical analysis. Each experiment was performed at least three times. Statistical analysis was performed using SPSS 20.0 software (IBM, Corp., Armonk, NY, USA). The significance between SphK1, FAK, p-FAK, E-cadherin, vimentin and clinicopathological characteristics of the patients was assessed with the χ² test. Survival curves were compared using the log-rank test. The PCR was analyzed by the Mann-Whitney U test. Statistical significance was determined by unpaired Student's t-test. Analysis of variance was performed, followed by the Least Significant Difference's test for multiple comparisons. P<0.05 was considered to indicate a statistically significant difference.

Results

Expression of SphK1, FAK, p-FAK, E-cadherin and vimentin in colorectal cancer tissues and their clinical significance. Immunohistochemistry staining demonstrated that the expression density of SphK1, FAK, p-FAK and vimentin in colorectal cancer tissues was higher compared with
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normal colonic mucosa tissues, whereas, the expression of E-cadherin was lower compared with normal colonic mucosa tissues (Fig. 1A; Tables I and II). Furthermore, the expression of SphK1, FAK, p-FAK and vimentin in metastatic cancer tissues were higher compared with non-metastatic cancer tissues, whereas, the expression of E-cadherin was lower (Fig. 1A; Tables I and II). As presented in Tables III and IV, the positive scores of SphK1, FAK, p-FAK and vimentin expression in advanced tumors (stage III and stage IV) with lymph nodes and distant metastases of colorectal cancer tissues were higher compared with those identified in less advanced tumors (stage I and stage II) without-lymph nodes and distant metastases. However, the positive rate of E-cadherin was lower in advanced tumors. There was a significant difference in E-cadherin and vimentin expression between different infiltration depths in colorectal cancer tissues; however, there was no significant difference in SphK1, FAK and p-FAK expression. These results suggested that the expression of SphK1, FAK, p-FAK, E-cadherin and vimentin was associated with the malignant invasion and metastasis of colorectal cancer.

Association between SphK1 expression and survival of patients with colorectal cancer. Patients with colorectal cancer with SphK1-positive cancer cells had a significantly lower survival rate compared with patients with SphK1-negative (Fig. 1B; P=0.0169). The results suggested that the prognosis of patients with colorectal cancer with SphK1-positive tumor was poorer. Therefore, SphK1 may be used as a prognostic indicator for patients with colorectal cancer.

Suppression of FAK inhibits the cell migrational potency, EMT, and the expression of p-AKT and MMPs in RKO cells. Our previous study demonstrated that the relative protein expression of p-FAK was 0.93±0.02 in Caco2 cells, 0.71±0.03

Table I. Expression of SphK1, FAK and p-FAK in tissues assessed by immunohistochemistry staining.

| Colonic samples                | SphK1 | FAK | p-FAK |
|-------------------------------|-------|-----|-------|
| Normal colonic mucosa tissues | 50    | 44  | 6     | 0.001 |
| Non-metastatic cancer tissues | 63    | 27  | 36    | 0.001 |
| Metastatic cancer tissues     | 51    | 9   | 42    |       |

Table II. Expression of E-cadherin and vimentin in tissues assessed by immunohistochemistry staining.

| Colonic samples                | E-cadherin | Vimentin |
|-------------------------------|------------|----------|
| Normal colonic mucosa tissues | 50         | 9        | 41      | 0.001 |
| Non-metastatic cancer tissues | 63         | 15       | 48      |       |
| Metastatic cancer tissues     | 51         | 27       | 24      |       |

Figure 1. Expression of SphK1, FAK, p-FAK, E-cadherin and vimentin in colorectal cancer. (A) Immunohistochemical staining of SphK1, FAK, p-FAK, E-cadherin and vimentin proteins in normal colonic mucosa tissues, non-metastatic cancer tissues and metastatic cancer tissue samples (magnification, x100). (B) Survival rate of patients with SphK1-positive and SphK1-negative colorectal cancer. P=0.0169 vs. SphK1-negative. SphK1, Sphingosine kinase 1; FAK, focal adhesion kinase; p, phosphorylated.
in HT29 cells, 0.96±0.01 in RKO cells and 0.80±0.02 in HCT116 cells (26). The protein expression of p-FAK in RKO cells was the highest. Therefore, RKO cells were selected for FAK shRNA stable transfection and the FAK and p-FAK expression were successfully suppressed (Fig. 2A -C). The expression of p-AKT, MMP2/9, vimentin and fibronectin was decreased with the suppression of FAK, whereas, E-cadherin was increased, and no noticeable alteration occurred for AKT expression (Fig. 3A and B). The microvilli and pseudopodia of FAK knockdown RKO cells [FAK(-)-RKO] were decreased compared with transfection negative control (NC) and control (Con) RKO cells (Fig. 3C). These results suggested that the EMT process was inhibited by suppression of FAK by decreasing p-AKT and MMP2/9. The migrational potency of cells may be facilitated once the EMT process is reactivated in colorectal cancer cells (15). Therefore, the migrational potency of RKO cells was significantly suppressed by FAK knockdown (Fig. 4A and B; P<0.05).

Table III. Clinicopathological characteristics of the patients with colorectal cancer and SphK1, FAK and p-FAK expression in the colorectal cancer tissues.

| Pathologic feature                        | SphK1 | FAK    | p-FAK   |
|------------------------------------------|-------|--------|---------|
|                                         | n     | -      | +       | P-value | n     | -      | +       | P-value | n     | -      | +       | P-value |
| Infiltration depth                       |       |        |         |         |       |        |         |         |       |        |         |         |
| Mucosa and superficial muscular layer    | 21    | 10     | 11      | 0.080   | 9     | 12     | 16      | 0.181   | 10    | 11     | 0.701   |
| Deep muscular layer and below            | 93    | 26     | 67      | 0.004   | 26    | 67     | 26      | 0.001   | 40    | 53     | 0.001   |
| TNM staging                              |       |        |         |         |       |        |         |         |       |        |         |         |
| I/II stage                               | 63    | 27     | 36      | 0.004   | 29    | 34     | 37      | 0.001   | 37    | 31     | 0.006   |
| III/IV stage                             | 51    | 9      | 42      | 0.023   | 6     | 45     | 13      | 0.001   | 13    | 33     | 0.001   |
| Lymphatic metastasis                     |       |        |         |         |       |        |         |         |       |        |         |         |
| -                                        | 68    | 27     | 41      | 0.003   | 29    | 39     | 37      | 0.004   | 48    | 45     | 0.001   |
| +                                        | 46    | 9      | 37      | 0.003   | 6     | 40     | 13      | 0.001   | 45    | 33     | 0.001   |
| Distant metastasis                       |       |        |         |         |       |        |         |         |       |        |         |         |
| -                                        | 93    | 35     | 58      | 0.003   | 34    | 59     | 48      | 0.004   | 48    | 45     | 0.001   |
| +                                        | 21    | 1      | 20      | 0.004   | 1     | 20     | 2       | 0.004   | 19    | 17     | 0.001   |

SphK1, Sphingosine kinase 1; FAK, focal adhesion kinase; p, phosphorylated.

Table IV. Clinicopathological characteristics of the patients with colorectal cancer and E-cadherin, vimentin expression in the colorectal cancer tissues.

| Pathologic feature                        | E-cadherin | Vimentin |
|------------------------------------------|------------|----------|
|                                         | n          | -        | +        | P-value | n          | -        | +        | P-value |
| Infiltration depth                       |            |          |          |         |            |          |          |         |
| Mucosa and superficial muscular layer    | 21          | 3        | 18       | 0.018   | 16         | 5        | 0.013    |
| Deep muscular layer and below            | 93          | 39       | 54       |          | 43         | 50       |          |
| TNM staging                              |            |          |          |         |            |          |          |         |
| I/II stage                               | 63          | 15       | 48       | 0.001   | 40         | 23       | 0.005    |
| III/IV stage                             | 51          | 27       | 24       |          | 19         | 32       |          |
| Lymphatic metastasis                     |            |          |          |         |            |          |          |         |
| -                                        | 68          | 16       | 52       | 0.001   | 41         | 27       | 0.027    |
| +                                        | 46          | 26       | 20       |          | 18         | 28       |          |
| Distant metastasis                       |            |          |          |         |            |          |          |         |
| -                                        | 93          | 30       | 63       | 0.033   | 55         | 38       | 0.001    |
| +                                        | 21          | 12       | 9        |          | 4          | 17       |          |

TNM, tumor, node and metastasis.
SphK1 knockdown inhibits the cell migrational potency, EMT, and the expression of p-FAK, p-AKT and MMPs in RKO cells. RKO cells were used for SphK1 shRNA stable transfection. SphK1 mRNA and protein expressions were successfully suppressed in the transfected RKO cells (Fig. 2A and B). The expression of p-FAK, p-AKT and MMP2/9 was decreased with the suppression of SphK1 in RKO cells. However, there were no noticeable alterations of FAK and AKT expression levels (Figs. 2C and 3A-C). SphK1 knockdown was accompanied by an increased expression of E-cadherin and a decrease in the expression of vimentin and fibronectin (Fig. 3A and B). The microvilli and pseudopodia of SphK1 knockdown RKO cells [SphK1(-)-RKO] were decreased compared with NC and Con RKO cells (Fig. 3C). These results suggested that the EMT was repressed with the suppression of SphK1 by decreasing p-FAK, p-AKT and MMP2/9. The migrational potency of cells was facilitated once the EMT process was reactivated in colorectal cancer cells (17). Therefore, the migrational potency of RKO cells was additionally significantly decreased with the knockdown of SphK1 (Fig. 4; P<0.05).

SphK1 overexpression strengthens cell migrational potency, EMT, and the expression of p-FAK, p-AKT and MMPs in HT29 cells. Our previous study demonstrated that the relative protein expression of SphK1 was 0.96±0.02 in Caco2 cells, 0.61±0.07 in HT29 cells, 0.92±0.05 in RKO cells and 0.97±0.02 in HCT116 cells (22). The protein expression of SphK1 in HT29 cells was the lowest. HT29 cells were used for stable transfection of the SphK1 overexpressing vector (Fig. 5). SphK1 mRNA and protein expression levels were significantly increased in the transfected cells (Fig. 5A and C; P<0.05). In contrast to the cells with knockdown of SphK1, the negatively transfected control (NC) and Con-HT29 cells, the expression levels of p-FAK, p-AKT and MMP2/9 were significantly increased in SphK1-overexpressing HT29 cells [SphK1(+)‑HT29] (Fig. 5B and C; P<0.05). However, there were no statistical differences in FAK and AKT expression levels among SphK1(+)‑HT29 cells, NC-HT29 cells and Con-HT29 cells (Fig. 5A and B; P>0.05). Compared with NC-HT29 and Con-HT29 cells, the expression of E-cadherin decreased, whereas, vimentin and fibronectin increased in SphK1(+)‑HT29 cells (Fig. 5B and C), along with the microvilli and pseudopodia (Fig. 5D). These results suggested the EMT was induced by the overexpression of SphK1 by increasing p-FAK, p-AKT and MMP2/9. The migrational potency of cells was facilitated once the EMT process was reactivated in colorectal cancer cells (13). The migrational potency of the cells was additionally increased with the upregulation of SphK1 (Fig. 5E).
Figure 3. FAK knockdown or SphK1 knockdown inhibits the epithelial-mesenchymal transition of RKO cells mediated by the FAK/AKT/MMPs axis. (A) RT-PCR analysis for the target molecules. GAPDH was used as a reference and the Con-RKO was set to 1. (B) Western blot analysis. GAPDH was used as an internal control. (C) Microvilli and pseudopodia of Con, NC, FAK(-) and SphK1(-) RKO cells under scanning electron microscopy (magnification, x5.00k). All data are presented as the mean ± standard deviation of three independent experiments. FAK, focal adhesion kinase; SphK1, Sphingosine kinase 1; AKT, protein kinase B; MMP, matrix metalloproteinase; Con, control; NC, negative control; p, phosphorylated.

Figure 4. FAK knockdown or SphK1 knockdown decreases the migrational potency of RKO cells. (A) Cell migrational potency demonstrated by a wound healing assay (magnification, x100) and analysis. (B) Cell migrational potency demonstrated by a Transwell assay (magnification, x200) and analysis. All data are presented as the mean ± standard deviation of three independent experiments. FAK, focal adhesion kinase; SphK1, Sphingosine kinase 1; Con, control; NC, negative control.
Cell migrational potency, EMT, and the expression of p-FAK, p-AKT and MMPs in HT29 cells. As demonstrated in Fig. 6A, PF562271 (inhibitor of the FAK pathway) suppressed the proliferation of HT29 cells in a time- and dose-dependent manner. The protein expression of FAK was decreased when HT29 cells were treated with PF562271 at concentrations of 2.5 and 5 µM for 48 h. The protein expression of p-FAK was
significantly decreased when HT29 cells were treated with PF562271 at a concentration of 5 µM for 48 h (P<0.05). The expression of FAK, p-FAK, p-AKT and MMP2/9 was decreased when SphK1(+)‑HT29 cells were treated with PF562271 at a concentration of 5 µM for 48 h [PF‑SphK1(+)‑HT29]. However, there was no noticeable alteration in the expression level of AKT (Fig. 6B and C). In addition, the expression of E-cadherin was increased, whereas, vimentin and fibronectin expression levels were decreased in SphK1(+)‑HT29 cells treated with PF562271 (Fig. 6B and C). Furthermore, the microvilli, pseudopodia and migrational potency of PF‑SphK1(+)‑HT29 cells were suppressed by treatment with PF562271 (Fig. 6D and E). These results suggested that the FAK pathway inhibitor suppressed the EMT and migrational potency induced by SphK1 overexpression in HT29 cells.

**Discussion**

SphK1 was identified to be overexpressed in various types of cancer cells and contributed to the development, progression,
metastasis and chemo-resistance of cancer (2-6). Advanced invasion and metastasis are the key factors affecting cancer prognosis, and the leading causes of mortality in patients with cancer (27). The present study not only confirmed that the expression of SphK1 was increased in colorectal cancer cells; however, additionally demonstrated that it contributes to the metastasis and shorter survival time of patients with colorectal cancer. However, the molecular mechanism of SphK1 promoting the migration and metastasis of colon cancer cells requires further study.

EMT is considered a process of basic transformation during embryonic development. During EMT, epithelial cells may further boost polarity and transform into mesenchymal type cells capable of advanced migration under specific physiological and pathological conditions (28). In a previous study, the cancer cell migrational potency and the morphological transformation of cells microvilli and pseudopodia were enhanced when cells underwent EMT-associated alterations (29). In the present study, it was confirmed that the expression of E-cadherin decreased and vimentin increased in metastatic colorectal cancer tissues and SphK1-overexpressed colon cancer cells. The migrational potency was enhanced with the overexpression of SphK1, along with the microvilli and pseudopodia. E-cadherin is an adherens junction protein that forms homophilic intercellular contacts in epithelial cells; its downregulation is commonly observed in EMT cancer cells (30). Fibronectin serves an important role in the regulation of epithelial cell adhesion, allowing transition of cell-cell contacts to cell-extracellular matrix interactions during EMT (31). Vimentin contributes to EMT cancer cell mechanics by mediating cytoskeletal organization and focal adhesion maturation (32). It additionally promotes the cell intermediate filament status, altering from a keratin-rich network to a vimentin-rich network connecting to focal adhesions (32). Therefore, SphK1 contributed to the metastasis of colon cancer by inducing EMT.

FAK is a non-receptor tyrosine kinase that mediates integrin signaling from the sites of connection to the extracellular membrane termed focal adhesions. FAK mediates essential cellular processes, including growth, proliferation, adhesion, migration and survival through its functions as a molecular scaffold and as a kinase (33). It was observed that FAK promoted malignancy by regulating tumorigenic and migrational potency (34). Inhibition of FAK caused less mesenchymal-like characteristics and decreased the mobility and migrational potency of Hep2 cells and mesenchymal triple negative breast cancer cells (16). The results of the present study suggested that blocking FAK/p-FAK (Tyr397) suppressed the expression of fibronectin and vimentin, and enhanced E-cadherin in colon cancer cells. Cell migrational potency was inhibited by FAK knockdown. Furthermore, it was demonstrated in the present study and in our previous study that SphK1 promoted the migrational potency of colon cancer by regulating the FAK pathway (4). Induced by SphK1 overexpression, the EMT and migrational potency were suppressed by inhibition of the FAK pathway. These results demonstrated that SphK1 promoted the migrational potency of colon cancer by inducing EMT, which was mediated by FAK/p-FAK (Tyr397).

A previous study identified that SphK1 enhanced the migrational potency of non-small cell lung cancer cells by activating the AKT pathway (35). The expression of p-AKT in colon cancer cells was enhanced with the upregulation of SphK1 and suppressed with its downregulation. Notably, the expression of AKT/p-AKT, induced by the upregulation of SphK1 may be suppressed by the inhibitor of the FAK pathway. The PI3K/AKT pathway was involved in the regulation of cell mobility via activation of FAK, and associated phosphorylation of pS8 subunits of tyrosine of PI3K in human cancer cells (22). PI3K, an upstream activator of AKT (36), activated AKT by promoting the phosphorylation of the serine phosphorylation site (Ser473) of AKT (37). AKT was frequently dysregulated in tumors and served a pivotal role in tumor metastasis (38). Therefore, SphK1 promoted the migration and metastasis of colorectal cancer by inducing EMT, which was mediated by the FAK/AKT pathway.

It has been suggested that SphK1 expression promoted the secretion of MMP-2/9 and urokinase-type plasminogen activator (39). The indicated molecular mechanisms are important for the regulation of malignant behavior, including invasiveness, in colon cancer (39). The expression of MMP-2/9 was suppressed with the downregulation of SphK1, p-FAK and p-AKT. The PI3K/AKT/nuclear factor (NF)-xB pathway was involved in the regulation of MMPs (39). AKT regulates MMP2/9 gene expression by promoting p65- and p52-DNA-binding activities of NF-xB (40). Furthermore, MMP2/9 are crucial for the remodeling of the extracellular matrix (41). These results suggested that SphK1 promoted the migration and metastasis of colorectal cancer by inducing EMT, which was mediated by the FAK/AKT/MMPs axis.

In summary, SphK1 promoted the metastasis of colorectal cancer through induction of the EMT, which was mediated by the FAK/AKT/MMPs axis. The association of SphK1 with EMT requires further examination as a potential therapeutic target in the treatment of colorectal cancer.

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Availability of data and materials

All data generated or analyzed during the current study are included in this published article.

Authors’ contributions

CYX and SQL performed the studies and drafted the manuscript. WHW, ZHF and SWH collected the clinical data and tissue samples. CYX and MBQ performed the statistical analysis. SQL and JAH designed the study and helped draft the manuscript. All authors read and approved the final manuscript.
Ethics approval and consent to participate

This present study was approved by the Ethics Committee of the First Affiliated Hospital of Guangxi Medical University (Guangxi, China). Written informed consent was obtained from all patients.

Patient consent for publication

Consent for the publication of the clinical and pathological data was obtained from all patients who were involved in the present study.

Competing interests

The authors declare that they have no competing interests.

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