Knockdown of KLK12 inhibits viability and induces apoptosis in human colorectal cancer HT-29 cell line

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Abstract. Kallikrein-related peptidase 12 (KLK12) is overexpressed in cancer tissues including gastric, breast and prostate cancer. However, the role of KLK12 in colorectal cancer is not fully understood. In the present study, the level of KLK12 was determined by performing reverse transcription-polymerase chain reaction (RT-qPCR) in colorectal cancer tissues and cell lines. Lipofectamine® 2000 was used to transfect HT-29 cells to overexpress and knockdown KLK12. Cell viability, migration, invasion and apoptosis were detected by MTT, wound healing, Transwell and flow cytometry assays, respectively. The mRNA and protein expression levels of EMT-associated proteins, apoptosis-associated proteins, phosphorylated adenosine monophosphate-activated protein kinase (p-AMPK) and phosphorylated mammalian target of rapamycin (p-mTOR) were determined by RT-qPCR and western blot analysis. It was identified that the KLK12 mRNA levels were increased significantly in colorectal cancer tissues and cell lines. KLK12 small interfering RNA inhibited cell viability, migration and invasion. Furthermore, epithelial-mesenchymal transition (EMT)-associated proteins were altered by siKLK12. Cell apoptosis was induced by KLK12 downregulation, which was demonstrated by the changes in apoptosis-associated proteins; however, KLK12 overexpression produced the opposite effect. SiKLK12 enhanced the expression of p-AMPK and suppressed the expression of p-mTOR, while KLK12 overexpression had the opposite effect. Promotion of KLK12 overexpression-induced cell viability was reversed by 5-aminoimidazole-4-carboxamide ribonucleotide, an activator of the AMPK signaling pathway, and rapamycin, a specific inhibitor of the mTOR signaling pathway. Taken together, the results of the present study indicated that KLK12 was overexpressed in colorectal cancer and may regulate cell behavior, potentially via the AMPK and mTOR pathways.

Introduction

Colorectal cancer, which is one of the most common types of malignant tumor, has the third highest incidence in men and the second highest incidence in women globally; as a result, colorectal cancer is responsible for a high number of mortalities (1). Radical surgical resection is currently the only method for treating colorectal cancer (2). The primary cause of failure in colorectal cancer surgery is metastasis and tumor recurrence. To date, no effective medication has been identified to inhibit the progression of colorectal cancer. Therefore, the development of protective therapies to suppress the progression of colorectal cancer is necessary in treatment of colorectal cancer.

Epithelial mesenchymal transition (EMT) is a key mechanism in tumor metastasis (3,4). In tumor cells, it is a biological process in which tumor cells allow epithelial cells to transform into cells with mesenchymal phenotype characteristics. During the process, tumor epithelial cells lose their polarity and adhesion among cells is decreased. Through this, interstitial cell characteristics including high migration and invasion ability, anti-apoptosis and degradation of extracellular matrix are obtained (5,6). Therefore, EMT, which serves an important role in the process of tumor metastasis is also a critical condition for invasion and metastasis in colorectal cancer, (7).

As a class of serine/threonine protein kinases in eukaryotic cells, the primary function of adenosine monophosphate-activated protein kinases (AMPK) is to serve as a cellular energy sensor to modulate cellular responses to a low nutrient environment (8). A previous study has indicated that AMPK may regulate cell proliferation, growth and autophagy (9). The tumor suppressor gene serine/threonine kinase 11 may activate AMPK, while tumor suppressor gene TSC complex subunit 2
AMPK/TS c2/mTOR signaling pathway was involved in EcM molecules (17). It was also demonstrated that the mTOR pathway is activated in patients with colorectal cancer (12,13). Therefore, targeting AMPK and mTOR signal pathways may be a promising protective strategy for management of colorectal cancer.

As a class of serine proteases, human kallikrein-related peptidases (KLKs) are encoded by 15 different genes that localize to human chromosome 19q13.4 and have a highly conserved set of gene structures and protein sequences (14). KLKs have been demonstrated to serve important roles in tumor growth and metastasis and are considered as markers for various types of cancer (15). A previous study indicated that KLKs were closely associated with colorectal cancer, as it participated in tumor cell proliferation, apoptosis and prognosis (16). An additional study also identified that KLK promoted the invasion and metastasis of cancer cells by hydrolyzing certain macromolecular substances including the extracellular matrix (ECM), cell adhesion molecules (CAM), and importantly, KLK may degrade almost all ECM molecules (17). It was also demonstrated that the AMPK/TSC2/mTOR signaling pathway was involved in the process of tissue kallikrein to protecting the SH-SY5Y neuronal cell line against oxygen and glucose deprivation-induced injury (18). KLK12 is known as kallikrein 5, and a previous study has demonstrated that KLK12 mRNA levels were upregulated in cancer tissues including gastric, breast and prostate cancer (19). It has been suggested that KLK12 may become a novel tumor biomarker, as it participated in tumor cell proliferation, apoptosis and malignant changes in tissues (19). However, the mechanisms underlying the effect of KLK12 in human colorectal cancer remain unclear.

The present study was performed to investigate the role of KLK12 in cellular and animal colorectal cancer models, and to identify the potential underlying mechanisms. The results from the study increased understanding on the role of KLK12 in colorectal cancer.

Materials and methods

Human tissue samples. From May 2016 to May 2017, 45 patients who received radical colorectal cancer resection in The First Affiliated Hospital of Zhejiang Chinese Medical University (Hangzhou, China), were recruited to the study. Specimens were obtained immediately following surgical resection and used as a control. Sequences of the siRNAs used in the present study were as follows: KLK12 siRNA sense, 5'-AACAGUGUACGCAUGUATT-3'; KLK12 siRNA antisense, 5'-UACGUGGCUGUACUGUU GG-3'; negative siRNA sense, 5'-UUC UCGAAGUGUACGUTT-3'; negative siRNA antisense, 5'-ACGUGACAGUUCCGAGATT-3'. Transfected cells were cultured in complete medium for 48 h prior to use in the invasion, migration and apoptosis experiments.

Immunohistochemistry (IHC). Tissues were fixed with 4% paraformaldehyde for 24 h at room temperature and embedded in paraffin. The paraffin-embedded block tissues were cut into 4 µm sections. The samples were dehydrated in a graded ethanol (100,95, 90, 80, 70 and 0%), and for antigen retrieval, deparaffinized sections were incubated in a microwave oven at 95°C for 5 min and cooled for 1 h at room temperature. Following washing with PBS for two times, the sections were treated with methanol containing 3% hydrogen peroxide for 10 min at room temperature and blocked with 5% goat serum (Beijing Solarbio Science & Technology Co., Ltd.) at room temperature for 20 min. Subsequently, anti-KLK12 antibody (cat. no. AF3095; R&D systems, Inc.) was added to the sections at 4°C and incubated for 24 h. Following washing with PBS three times, biotin-labeled goat anti-rabbit IgG secondary antibody (1:5,000, cat. no. ab97049, Abcam) was added to the sections and incubated for 10 min at room temperature. The sections were then stained by 0.05% DAB staining solution (Leica Microsystems, Inc.) at room temperature for 15 min and stained with 2 g/l hematoxylin (Beijing Solarbio Science & Technology Co., Ltd.) for 3 min at room temperature. The sections were then observed using an MF43 fluorescence microscope (Guangzhou Micro-shot Technology Co., Ltd.), with magnification of x100 and 200.

Cell culture. Colorectal cancer HT-29, LoVo, SW-480, SW-620, Caco2, HCT-116, HCT-15, RKO, LS174T and DLD-1 cell lines, and the normal colon CDD-18Co cell line were purchased from American Type Culture Collection. RPMI-1640 medium containing 2 mM L-glutamine, 10% fetal bovine serum (FBS), penicillin G (100 U/ml) and streptomycin (0.1 mg/ml) was used as cell culture medium and cultured with the cells at 37°C with 5% CO2. All culture reagents were purchased from Gibco; Thermo Fisher Scientific, Inc. HT-29 cells were inoculated and grown to 80% confluency prior to treatment with adequate concentrations of rapamycin (10 nM; cat. no. V900930; Sigma-Aldrich; Merek KGaA) and 5-amino-imidazole-4-carboxamide ribonucleotide (AICAR; 2 mM; cat. no. A9978; Sigma-Aldrich; Merek KGaA) for 90 min, and the cells were then transfected with KLK12 plasmid for 12, 24 and 48 h to detect cell viability.

Transfection. 50 nmol/l small interfering RNA targeted KLK12 (siKLK12) and negative siRNA (Shanghai GenePharma Co., Ltd.) were inserted into pLKO.1-TRc vector (Shanghai Institute of Biochemistry and Cell Biology, Chinese Academy of Sciences) to produce the recombinant plasmid. siRNA was used to transfect HT-29 cells using Lipofectamine™ 2000 (Invitrogen; Thermo Fisher Scientific, Inc.). The KLK12 coding region was inserted into the pLKO.1-TRC vector, and the HT-29 cells were transfected with KLK12 plasmid or empty vector. The cells that did not undergo transfection were used as a control. Sequences of the siRNAs used in the present study were as follows: KLK12 siRNA sense, 5'-AACAGUGUACGCAUGUATT-3'; KLK12 siRNA antisense, 5'-UACGUGGCUGUACUGUU GG-3'; negative siRNA sense, 5'-UUC UCGAAGUGUACGUTT-3'; negative siRNA antisense, 5'-ACGUGACAGUUCCGAGATT-3'. Transfected cells were cultured in complete medium for 48 h prior to use in the invasion, migration and apoptosis experiments.
Corporation), with the magnification of x200.

and captured using an inverted light microscope (Olympus glass slide, sealed with a neutral gum. Images were observed and dried with the bottom side up and then transferred to a membrane was removed using a small tweezers carefully were gently wiped off using a cotton swab. The microporous liquid was aspirated. The unmigrated cells in the upper layer with 0.1% crystal violet for 20 min at room temperature. Next, fixed in 100% methanol solution at 4˚C for 30 min and stained washed with calcium-free PBS twice, and the chamber was put under a 40-fold inverted microscope (Olympus corporation) to be observed, and images at the same position were captured twice.

Wound healing assay. After 48 h of cell culture transfection, the cell density of each group was ~90%. The cells were serum-starved for 24 h, and 200 µl sterile tip was used to scratch the cells in the well plate. The medium was discarded and the surface was gently washed 3 times with PBS to rinse off the exfoliated cells. After 0 and 24 h, the cell culture plate was placed under a 40-fold inverted microscope (Olympus Corporation) to be observed, and images at the same position were captured twice.

Transwell assay. Transwell chambers (8 µm; Corning Incorporated) was placed on a 24-well plate with a layer of 50 µl Matrigel (BD Biosciences) coated onto the Transwell chamber 37˚C for 30 min. Then, 6x10^4 transfected cells were cultured in serum-free medium for 12 h to eliminate the effects of the serum and then resuspended in RPMI-1640 medium without FBS. A total of 100 µl suspended cells were added to the Transwell chamber, while 400 µl RPMI-1640 medium containing 20% FBS was added to the lower chambers. The cells were cultured with 5% CO₂ for 24 h at 37°C in an incubator. The Transwell chamber was then removed, the culture solution in the Transwell plate was discarded and washed with calcium-free PBS twice, and the chamber was fixed in 100% methanol solution at 4°C for 30 min and stained with 0.1% crystal violet for 20 min at room temperature. Next, PBS was used to wash the chamber, and the upper chamber liquid was aspirated. The unmigrated cells in the upper layer were gently wiped off using a cotton swab. The microporous membrane was removed using a small tweezer carefully and dried with the bottom side up and then transferred to a glass slide, sealed with a neutral gum. Images were observed and captured using an inverted light microscope (Olympus Corporation), with the magnification of x200.

**MTT assay.** HT-29 cells were transfected with or without exposure to AICAR/rapamycin. The transfected cells were seeded in 96-well plates at a density of 5x10³/1. Following incubation at 37°C for 12, 24 and 48 h, 10 µl MTT solution (5 mg/ml) was added into the wells and the plate was incubated in an atmosphere containing 5% CO₂ at 37°C for 4 h. Next, a microplate reader was used to measure the absorbance at 450 nm. Cell viability was detected by MTT assay (Thermo Fisher Scientific, Inc.) according the manufacturer’s protocol.

**Apoptosis assay.** An Annexin V-fluorescein isothiocyanate (FITC)/propidium iodide (PI) apoptosis detection kit (BD Biosciences Medical Devices Shanghai Co., Ltd.) was used to detect HT-29 cells apoptosis. Colorectal cancer HT-29 cells (5x10⁴ cells per well) were seeded in 6-well plates until cell density reached 85%. siKLK12, KLK12 plasmids and the corresponding control plasmids were used to transfect HT-29 cells, which were then cultured in serum-free medium at 37°C for 5 h. After 48 h of culture, the cells were centrifuged at 4°C for 10 min, at the speed of 5,000 x g, and washed twice with PBS, and 100 µl 1X Annexin V Binding Buffer and 5 µl FITC-labeled Annexin V (20 µg/ml) were added to the cells and incubated at room temperature for 20 min. Next, the cells were added with 5 µl PI (50 µg/ml) for 5 min and 400 µl binding buffer (BD Biosciences). A flow cytometer (BD Biosciences Medical Devices Shanghai Co., Ltd.) was used to analyze cell apoptosis; among those cells, FITC/PI⁻ cells represented healthy living cells, FITC+/PI⁻ cells represented necrotic cells, FITC⁻/PI⁻ cells indicated early apoptotic cells, FITC⁻/PI⁻ cells represented late apoptotic cells.

**Reverse transcription quantitative polymerase chain reaction (RT-qPCR).** Following transfection, HT-29 cells were incubated in an incubator for 48 h, and total RNA from HT-29 cells (2x10⁴ cells/well in 6-well plates) and 50 mg of colorectal cancer and normal tissues were extracted using TRIzol® reagent (Thermo Fisher Scientific, Inc.) according to the manufacturer’s protocol. Nanodrop™ 2000 spectrophotometer (NanoDrop Technologies; Thermo Fisher Scientific, Inc.) was conducted to measure RNA quality; an A260/A280 ratio between 1.8-2.0 was required for the generation of cDNA. The oligo-dT or stem-loop reverse transcriptase primers (Takara Bio, Inc.) were used to obtain cDNA, and the RT thermocycler conditions were set at 42°C for 60 min, 70°C for 5 min and maintenance at 4°C. qPCR was performed with SYBR® Premix Ex Taq™ II (Takara Bio Inc.) using real-time PCR Detection System (ABI 7500; Thermo Fisher Scientific, Inc.). PCR reaction conditions were as follows: Pretreatment at 95°C for 10 min, followed by 40 cycles at 94°C for 15 sec, 60°C for 1 min and 60°C for 1 min, and preservation at 4°C. The 2^ΔΔCt method was used to quantify the data (21). The primers used in this assay are listed in Table I.

**Western blot analysis.** The HT-29 cells were treated with siKLK12, KLK12 and the corresponding control plasmid and

| Genes     | Forward       | Reverse       |
|-----------|---------------|---------------|
| KLK12     | GCCTCAACCTCTCCATCCTGTC | GCTGAAAGCTCCCCACAC |
| E-cadherin| CGAGAGCTACAGTTCAGG | GGTTGTCTCGAGGGAAAAATAGG |
| Vimentin  | TGCCGTGAAGCTGCTAACTA | CCAGAGGGAGTGAATCCAGGATTA |
| Snail     | TCGGAAAGCTAACTACAGGGA | AGATGAGCATGGTCAGGAG |
| MMP-2     | TTGATGGCAGATGGTCAAGT | TTGTCACGTGGCCGTACAGT |
| MMP-9     | GACGCAGACATCGTCTACCCA | CACAACTCCTCATCGTGCAA |
| GAPDH     | AACGTGTCAGTGGTTGGACCTG | AGTGGGTCGCTGTTTGAAGT |

KLK12, kallikrein-related peptide 12; E-cadherin, epithelial cadherin; MMP, matrix metalloproteinase.

Table I. Primers for reverse transcription quantitative polymerase chain reaction.
then cultured in an incubated for 48 h. Total proteins were extracted from HT-29 cells (2x10^5 cells/well in 6-well plates) and tissues (50 mg) by radioimmunoprecipitation assay lysis buffer (Cell Signaling Technology, Inc.). A BCA Protein Assay kit (Pierce; Thermo Fisher Scientific, Inc.) was applied to measure the concentrations of proteins, which were adjusted to a concentration of 6 µg/µl using 1X loading and DEPC water. The samples (5 µl) were separated by 10% SDS-PAGE gels and then transferred to polyvinylidene fluoride membrane (PVDF; EMD Millipore). Following blocking in 5% nonfat milk in PBST (0.1% Tween-20 in PBS) for 1 h at 37°C, the PVDF membranes were probed with the primary antibodies overnight at 4°C, followed washing 3 times with PBST. Next, the membrane was incubated with the secondary antibody (horseradish peroxidase-conjugated goat anti-mouse/rabbit IgG, 1:2,000; cat. nos. sc-516102/sc-2357; Santa Cruz Biotechnology, Inc.) at room temperature for 2 h and washed with PBST 3 times. An EZ-ECL kit (Biological Industries) was used to develop the membranes, and the gray values of the bands were analyzed and quantified using ImageJ v5.0 software (Bio-Rad Laboratories, Inc.). The antibodies used in the study were as follows: Anti-GAPDH (mouse; 1:1,000; cat. no. LS-B1625; LifeSpan BioSciences, Inc.); anti-epithelial (E)-cadherin (mouse; 1:1,000; cat. no. ab1416; Abcam); anti-vimentin (rabbit; 1:1,000; cat. no. ab92547; Abcam); anti-matrix metalloproteinase (MMP)-2 (rabbit; 1:1,000; cat. no. ab37150; Abcam); anti-MMP-9 (rabbit; 1:1,000; cat. no. ab73734; Abcam); anti-zinc finger protein SNAI1 (Snail; rabbit; 1:1,000; cat. no. 3879; Cell Signaling Technology, Inc.); anti-cleaved caspase-3 (rabbit; 1:1,000; cat. no. ab13847; Abcam); anti-Bax (rabbit; 1:1,000; cat. no. ab32503; Abcam); anti-Becl-2 (rabbit; 1:1,000; cat. no. ab32124; Abcam); anti-AMPK (mouse; 1:1,000; cat. no. ab2047; Abcam); anti-phosphorylated (p)-AMPK (rabbit; 1:1,000; cat. no. ab133448; Abcam); anti-mTOR (rabbit; 1:1,000; cat. no. ab32028; Abcam); and anti-p-mTOR (rabbit; 1:1,000; cat. no. ab84400; Abcam).

**Results**

**KLK12 is highly expressed in colorectal cancer tissues and cell lines.** In the present study, the expression levels of KLK12 in colorectal cancer tissues and adjacent normal tissues of cancer were initially explored using RT-qPCR. The results indicated that the KLK12 mRNA level was increased in the cancer tissues compared with the adjacent normal tissues (Fig. 1A). The IHC results indicated a large number of brown particles in the cancer tissues (Fig. 1B). Next, the KLK12 levels in the colorectal cancer cell lines and normal cells were determined using RT-qPCR; the results in the cell lines were consistent with those from the cancer tissues, in that the KLK12 mRNA levels were identified to be increased at different degrees in the majority of the colorectal cancer cell lines, with the exception of HCT-15 and RKO cells, compared with normal colorectal cells CCD-18Co (Fig. 1C). HT-29 cells exhibited the highest expression of KLK12; therefore, HT-29 cells were selected to be used in the following experiments.

**Knockdown of KLK12 inhibits HT-29 cells viability and promotes the cell apoptosis.** To further investigate the functional effects of KLK12 on colorectal cancer HT-29 cells, KLK12 expression was silenced by siRNA using Lipofectamine® 2000 transfection reagent, and RT-qPCR (Fig. 2A) and western blot analysis (Fig. 2B and C) were performed to confirm that the transfection was successful. Subsequently, HT-29 cells were seeded in a 96-well plate and transfected with small interfering RNA of KLK12 for 12, 24 and 48 h. The viability of HT-29 cells was analyzed using the MTT assay. It was identified that knockdown of KLK12 significantly decreased the viability of HT-29 cells at 24 and 48 h (Fig. 2D). Furthermore, cell apoptosis levels were quantified by flow cytometry using Annexin V and PI double staining, and it was demonstrated that KLK12 silencing noticeably increased the proportion of apoptotic HT-29 cells compared with the control and NC groups (Fig. 2E).

**Knockdown of KLK12 inhibits HT-29 cells migration and invasion.** Migration and invasion are typical biological characteristics of malignant tumors; therefore, invasion and migration were determined by wound healing and Transwell invasion assays, respectively. It was observed that KLK12 silencing for 24 h significantly decreased the migration rate in...
HT-29 cells (Fig. 3A and B). Concomitantly, cell invasion was inhibited following 24 h downregulation of KLK12 in HT-29 cells (Fig. 3C and D).

**Overexpression of KLK12 suppresses apoptosis in HT-29 cells.** The expression level of KLK12 was overexpressed by a KLK12 plasmid using Lipofectamine® 2000 transfection reagent, and RT-qPCR (Fig. 4A) and western blot analysis (Fig. 4B and C) were performed to confirm that the transfection was successful. Subsequently, HT-29 cells were seeded in a 6-well plate and transfected with the KLK12 plasmid for 12, 24 and 48 h. The viability of the HT-29 cells was analyzed using the MTT assay. KLK12 overexpression significantly increased the viability of HT-29 cells at 48 h (Fig. 4D). Furthermore, cell apoptosis levels were quantified by flow cytometry with Annexin V and PI double staining, and it was identified that KLK12 overexpression significantly decreased the proportion of apoptotic HT-29 cells compared with the control and mock groups (Fig. 4E).

**Upregulation of KLK12 promotes migration and invasion in HT-29 cells.** In the present study, invasion and migration were determined by wound healing and Transwell invasion assays, respectively. The results demonstrated that overexpressing KLK12 for 24 h significantly increased the migration rate of HT-29 cells (Fig. 5A and B), and that cell invasion was also increased following KLK12 upregulation for 24 h in the HT-29 cells (Fig. 5C and D).

**EMT-associated and apoptosis-associated proteins are altered in HT-29 cells treated with siKLK12/KLK12.** The effects of siKLK12/KLK12 on the EMT-associated proteins including E-cadherin, Vimentin, Snail, MMP-2 and MMP-9 were determined in HT-29 cells by performing RT-qPCR and western blot analysis. It was identified that treatment with KLK12 siRNA significantly upregulated the expression of E-cadherin and downregulated the expression levels of vimentin, Snail, MMP-2 and MMP-9 at mRNA levels (Fig. 6A) and protein levels (Fig. 6B and C). However, overexpression of KLK12 produced the opposite effect in these proteins, in comparison with the results from siKLK12 treatment (Fig. 6A-C). It was also identified that apoptosis was induced by siKLK12 treatment and inhibited by KLK12 overexpression from the results of flow cytometry; however, alterations to the apoptosis-associated proteins remained unclear. Therefore, the effects of KLK12 silencing and overexpression on the expression of apoptosis-association proteins including Bcl-2, Bax and cleaved caspase-3 was examined in the HT-29 cells by western blot analysis. The results indicated that KLK12 silencing induced apoptosis, as the expression of anti-apoptosis protein Bcl2 was significantly downregulated and the expression levels of pro-apoptosis proteins Bax and cleaved caspase-3 were upregulated, when compared with the control group (Fig. 6D and E), while overexpression of KLK12 elicited the opposite effect in these proteins, in comparison with the effects of siKLK12 treatment (Fig. 6D and E).

**AMPK and mTOR pathways are involved in HT-29 cells treated with siKLK12/KLK12.** The AMPK/mTOR signaling pathway serves an important role in tumor progression and metastasis. To explore the potential mechanism of KLK12 in the regulation of colorectal cancer cell migration and invasion, the present study examined the effects of KLK12 on AMPK/mTOR signaling. The levels of p-AMPK, AMPK, p-mTOR and mTOR in the HT-29 cells, which were transfected with siKLK12 and the KLK12 plasmid, were determined by western blot analysis. The results demonstrated that in HT-29 cells transfected with siKLK12, p-AMPK levels were significantly upregulated compared with control group; by contrast, the expression of p-mTOR was decreased (Fig. 7A and B). However, overexpression of KLK12...
resulted in the opposite effect in those proteins compared with siKLK12 treatment (Fig. 7A and B). In order to confirm the role of AMPK/mTOR in the progression of HT-29 cells, an activator of the AMPK signaling pathway, AICAR (2 mM), and a specific inhibitor of mTOR signaling pathway, rapamycin (10 nM), were incubated with HT-29 cells with or without the KLK12
Figure 5. Migration and invasion are promoted by KLK12 overexpression in HT-29 cells. (A and B) Migration rate of HT-29 cells treated with KLK12 plasmid for 0 and 24 h were (A) visualized and (B) quantified using a wound healing assay. (C and D) Invasion rate of HT-29 cells treated with KLK12 plasmid for 24 h were (C) visualized and (D) quantified using a Transwell assay. **P<0.01 vs. control group. ^^P<0.01 vs. mock group. KLK12, kallikrein-related peptidase 12; con, control.

Figure 6. Expression of epithelial-mesenchymal transition-associated and apoptosis-associated proteins is altered in HT-29 cells treated with siKLK12 or KLK12 overexpression plasmids. (A) The mRNA levels of E-cadherin, vimentin, Snail, MMP-2 and MMP-9 were analyzed by reverse transcription quantitative polymerase chain reaction. (B and C) The protein levels of E-cadherin, vimentin, Snail, MMP-2 and MMP-9 were analyzed by western blot analysis. (D and E) The protein levels of Bcl-2, Bax protein and cleaved caspase-3 were analyzed by western blot analysis. *P<0.05 and **P<0.01 vs. Nc group. ^P<0.05 and ^^P<0.01 vs. mock group. The ‘KLK12’ group represents the KLK12 overexpression group. KLK12, kallikrein-related peptidase 12; si, small interfering RNA; E-cadherin, epithelial cadherin; Snail, zinc finger protein SNAI1; MMP, matrix metalloproteinase; NC, negative control.
The viability of the HT-29 cells was analyzed using MTT assay. The data demonstrated that AIcAR reversed the effect of KLK12 overexpression on cell viability, and that AICAR had significant inhibitory effect on viability of HT-29 cells (Fig. 7c). Surprisingly, the results of rapamycin on the effect of KLK12 on HT-29 cell viability were consistent with AIcAR (Fig. 7d).

**Discussion**

Tissue or serum levels of KLK are examined individually or in small groups as a diagnostic or prognostic factor for determining different types of cancer (22, 23). In the colorectal cancer, KLK has been demonstrated to act as a diagnostic or prognostic factor (16). However, KLK12 has not been studied extensively in colorectal cancer. In the present study, KLK12 overexpression was demonstrated promoting the viability and inhibit apoptosis in the colorectal cancer-derived HT29 cell line via the activation of the AMPK/mTOR signaling pathways.

There have been multiple previous studies concerning KLK12 in different types of cancer: Li et al identified that KLK12 levels were significantly increased in gastric cancer cells compared with in GES-1 cells (24). In addition, analysis of a combined sample of 3,153 cases and 3,199 controls indicated that the KLK12 tag single nucleotide polymorphism rs3865443 was marginally and statistically correlated with the risk of developing prostate cancer (25). KLK12 expression at mRNA level was markedly increased in MKN-45 gastric cancer cells compared with normal mucosal cells and 2 other gastric cancer cell lines (26); the mRNA level of KLK12 observed in the present study was consistent with previous studies. It was also identified that KLK12 expression levels were increased in colorectal cancer tissues and colorectal cancer-derived cell lines compared with their corresponding controls. These
data demonstrated that the oncogenic potential of KLK12 in colorectal cancer was similar to that in other types of cancer. Metastasis is regarded as a major factor contributing to the poor prognosis of patients with colorectal cancer and is a typical feature of malignant tumors. Overexpression of KLK12 protein was significantly associated with lymph node metastasis, and the proliferation of gastric cancer MKN-45 cells was markedly decreased by the knockdown of KLK12 protein (26).

Transfection of AGS cells with KLK12 siRNA led to decreased cell proliferation (24). KLK12 efficiently cleaved human extracellular matrix proteins fibronectin and tenasin, both of which are involved in the regulation of endothelial cell adhesion and migration (27). In the present study, cell viability, migration and invasion were inhibited when the HT-29 cells were transfected with siKLK12. Concomitantly, E-cadherin expression was upregulated and vimentin, Snail, MMP-2 and MMP-9 expression were downregulated in siKLK12-transfected HT-29 cells. However, KLK12 overexpression elicited the opposite effect on those factors. Therefore, it was concluded that KLK12 may be involved in the process of EMT in colorectal cancer, at least in HT-29 cells.

Previous data supports the hypothesis that targeting apoptosis may be a promising and protective strategy for the management of colorectal cancer. MLK7-AS1 knockdown promoted CRC cell apoptosis in vitro (28). Lupeol-induced cellular apoptosis of both colorectal cancer cell lines, which increased p53 and decreased Bcl2 protein levels (29). In the present study, apoptosis levels and the expression levels of apoptosis-associated proteins were determined by flow cytometry and western blot analysis, respectively. It was identified that cell apoptosis was induced by the knockdown of KLK12 in HT-29 cells. In addition, the anti-apoptosis protein (Bcl-2) was expressed at low levels and the pro-apoptosis proteins (Bax and cleaved caspase-3) were expressed at high levels in the siKLK12 group, compared with the control group. However, KLK12 upregulation produced the opposite effect on those factors. These results suggested that the function of KLK12 as a carcinogenic factor in colorectal cancer may be realized by the inhibition of apoptosis, at least in HT-29 cells. However, the precise mechanisms underlying KLK12 activity in colorectal cancer required further investigation. At present, the mechanism by which KLK12 induced viability and metastasis of colorectal cancer is not fully understood. AMPK, a ubiquitous serine/threonine protein kinase, regulates tumorigenesis, development and chemical resistance through negative regulation of mTOR (30). The phosphorylation of AMPK and inhibition of downstream phosphorylation of mTOR in cholangiocarcinoma cells were activated by FYN proto-oncogene, Src family tyrosine kinase knockdown (31). In the present study, the functions of KLK12 on AMPK/mTOR signaling were explored, and it was identified that the phosphorylation of AMPK was activated by KLK12 in colorectal cancer. siKLK12 inhibited the viability and EMT process of HT-29 cells via regulation of the AMPK/mTOR signaling pathway. However, the method by which KLK12 functions in colorectal cancer animal models or in patients with colorectal cancer remains unclear. The present study also demonstrated that rapamycin may produce limited effects on cell viability, compared with AICAR, and this may be explained by a more marked inhibition of AICAR on mTOR and its downstream genes.

Previous studies have indicated that KLK12 splice variant KLK12sv3 may be used as a marker in predicting a desirable prognosis in breast cancer (19). Papachristopoulou et al (32) described the value of KLK12sv1/2 and KLK12sv3 in differentiating between benign and malignant breast tumors and their potential prognostic value. We hypothesized that KLK12 expression may also be associated with the survival and prognosis of patients with colorectal cancer, however, whether it may be used as a prognostic marker for colorectal cancer remains to be explored. However, by referring to other studies (33,34), the effects of KLK12 on cell proliferation and migration were verified. In the present study, it was identified that silencing KLK12 inhibited the cell viability and promoted apoptosis of human colorectal cancer HT-29 cells. However, colorectal cancer is a highly heterogeneous disease, and the role of KLK12 requires additional confirmation in multiple different colorectal cancer cells.

In conclusion, KLK12 down-regulation inhibited cell viability and metastasis via regulating AMPK/mTOR signaling pathway in HT-29 cells. The present results provided that KLK12 was an oncogene and could be used as a therapeutic target for treating colorectal cancer.

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Availability of data and materials
The analyzed data sets generated during the study are available from the corresponding author on reasonable request.

Authors' contributions
QL and HZ made substantial contributions to the conception and design of the study. XZ and ZF were responsible for data acquisition, data analysis and interpretation. HZ, XZ and QL were responsible for drafting the article and critically revising it for important intellectual content. All authors provided final approval of the version to be published. All authors are accountable for all aspects of the study in ensuring that questions related to the accuracy or integrity of the work are appropriately investigated and resolved.
Ethics approval and consent to participate

All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. The present study was approved by the Ethical Committee of The First Affiliated Hospital of Zhejiang Chinese Medical University and written informed consent was obtained from each patient.

Patient consent for publication

Written informed consent was obtained from each patient.

Competing interests

The authors declare that they have no competing interests.

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