TM4SF1 inhibits apoptosis and promotes proliferation, migration and invasion in human gastric cancer cells

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Abstract. Gastric cancer (GC) is associated with poor patient prognosis, and so it crucial to investigate the molecular mechanisms underlying the progression of GC. The aim of the present study was to investigate the role of transmembrane-4 L6 family member 1 (TM4SF1) in the progression of GC. TM4SF1 small interfering RNA (siRNA) and TM4SF1-expressing plasmids were employed to regulate TM4SF1 expression. *In vitro* experiments were performed to determine the effect of TM4SF1 on the expression of apoptosis-associated molecules and determine the role of TM4SF1 in apoptosis, proliferation, migration and invasion using human GC cell lines MGC803 and MKN45. The data of the present study demonstrated that TM4SF1 may regulate the expression of apoptosis-associated molecules at the mRNA and protein levels. TM4SF1 silencing reduced B-cell lymphoma 2 (Bcl2) expression, whilst caspase-3 and Bcl2-associated X expression increased, and upregulating TM4SF1 reversed these changes in GC cells. Furthermore, TM4SF1 knockdown promoted apoptosis while inhibiting the proliferation, migration and invasion of GC cells. Rescue experiments demonstrated that TM4SF1 upregulation reversed the changes induced by transfection with TM4SF1 siRNA. In summary, TM4SF1 is an anti-apoptosis protein associated with the progression of GC. Additional *in vivo* experiments and clinical trials are required to confirm the possible use of TM4SF1 in tumor therapy.

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Introduction

Gastric cancer (GC) is a disease that poses a serious threat to human health and quality of life globally (1). In 2015, GC is the third highest cause of cancer-associated moralities in males and the second highest in females in China (2). Currently, patients with GC generally have a poor prognosis, and early diagnosis is important to improve patient outcomes (3). Treatment primarily comprises of combined therapy of surgery with chemotherapy and radiotherapy; however, this treatment regime has limited effectiveness. Due to the majority of patients with GC being diagnosed at advanced stages, the resection rate for GC has been reported to be ~50% (4). Chemotherapy is an adjunctive treatment that is only able to alleviate the symptoms and prolong the survival of patients with GC, rather than treating the disease (5). Whether adjuvant radiotherapy can improve the overall survival of patients with GC remains controversial (6). Developing effective therapeutic strategies based on novel targets may help to improve the overall prognosis of patients with GC. Investigating the molecular mechanisms underlying GC progression is crucial in the identification of novel therapeutic targets.

Transmembrane-4 L6 family member 1 (TM4SF1), a low molecular weight protein consisting of 202 amino acids, is a member of the transmembrane-4 protein L6 superfamily. This family also includes TM4SF4, TM4SF5 and TM4SF18 (7). TM4SF1 is highly expressed in activated endothelial cells and can activate vascular endothelial growth factor (VEGF)-A or thrombin to stimulate angiogenesis (8). Additionally, TM4SF1 may also promote cell migration by increasing the formation of filopodia (9). TM4SF1 may therefore be a tumor promoter, and a number of studies have confirmed the important role of TM4SF1 in tumor progression. Cao *et al* (10) reported that TM4SF1 regulates pancreatic cancer migration and invasion *in vitro* and *in vivo*. Huang *et al* (11) reported that TM4SF1 promotes proliferation, invasion and metastasis in human liver cancer cells. Sun *et al* (12) confirmed that TM4SF1 regulates breast cancer cell migration and apoptosis; however, the effects of TM4SF1 on GC remain unclear.

A previous study reported that TM4SF1 was overexpressed in GC (13). Based on this, it was hypothesized in the present study that TM4SF1 may be an important regulator of GC development. The aim of the present study was to assess the effects of TM4SF1 on the proliferation, migration and invasion of GC cells. These data may improve the understanding of the

molecular mechanisms underlying GC progression, providing a basis for the development of effective therapeutic strategies.

Materials and methods

Cell culture and reagents. Human GC cell lines MGC803 and MKN45 were obtained from the Cell Bank of the Shanghai Institute of Biological Sciences, Chinese Academy of Sciences (Shanghai, China) and maintained in RPMI-1640 supplemented with 10% fetal bovine serum (FBS; Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA) and 1% penicillin/strep-tomycin at 37˚C in a humidified atmosphere containing 5% CO_2. Antibodies against B-cell lymphoma 2 (Bcl2; 1:500; ab692), caspase-3 (1:1,000; ab13847), Bcl2-associated X (Bax; 1:2,000; ab32503) and β-actin (1:2,000; ab8227) were purchased from Abcam (Cambridge, UK). Bound antibodies were detected with horseradish peroxidase-conjugated anti-body against mouse (1:10,000; ab6728; Abcam) or rabbit IgG (1:10,000; sc-2357; Santa Cruz Biotechnology, Inc., Dallas, TX, USA), followed by enhanced chemiluminescence detection (GE Healthcare Life Sciences, Little Chalfont, UK).

Gene transduction. The TM4SF1 small interfering RNA (siRNA), TM4SF1-expressing plasmid and their control vectors were obtained from Shanghai GenePharma Co., Ltd. (Shanghai, China). The sequence of TM4SF1 siRNA was as follows: Sense, 5'-GGCUCUUGGUCGGAAUUGAATT-3', and antisense, 5'-UUCAAUUCCACCAAGAGCCTT-3'. The TM4SF1-expressing plasmid was constructed by subcloning the human TM4SF1 cDNA by Shanghai GenePharma Co., Ltd.. MGC803 and MKN45 cells were seeded on to a 24-well plate at a concentration of $1x10^5$ cells/well for transfection. TM4SF1 siRNA (50 nM) and TM4SF1-expressing plasmid (1 µg/well) were transfected into MGC803 and MKN45 cells with Lipofectamine® 2000 (Invitrogen; Thermo Fisher Scientific, Inc.), according to the manufacturer's protocol. The control group represented MGC803 and MKN45 cells transfected with their control vectors. The siTM4SF1 group represented MGC803 and MKN45 cells transfected with TM4SF1 siRNA. The specificity of TM4SF1 siRNA was verified by a rescue experiment (14). In the rescue experiment, the TM4SF1 siRNA and TM4SF1-expressing plasmid were co-transfected into MGC803 and MKN45 cells, denoted as the siTM4SF1+TM4SF1 group. Changes in TM4SF1 expression levels were assessed using reverse transcription-quantitative polymerase chain reaction (RT-qPCR) at 48 h after transfection.

RT-qPCR. Total RNA was isolated from MGC803 and MKN45 cells using TRIzol® (Invitrogen; Thermo Fisher Scientific, Inc.). Purified RNA was reverse transcribed into cDNA with the M-MLV First Strand kit (Invitrogen; Thermo Fisher Scientific, Inc.), according to the manufacturer's protocol. RT-qPCR reactions were conducted using the SYBR® mix (Invitrogen; Thermo Fisher Scientific, Inc.) on an ABI PRISM 7500 Sequence Detection system (Applied Biosystems; Thermo Fisher Scientific, Inc.) (15). Briefly, after an initial denaturation step at 95˚C for 30 sec, amplifications were conducted with 40 cycles of 95˚C for 5 sec and 60˚C for 34 sec. Human β-actin was used as the housekeeping gene to normalize the expres-sion levels of the target genes (16,17). Primers used were as follows: TM4SF1, forward, 5'-CTGCTCTCACCAACAGCA AT-3', and reverse, 5'-TGCCAGTCTTTACAGGCGTT-3'; Bcl2, forward, 5'-CAGGAAACGGCCCGGAT-3', and reverse, 5'-CTGGGGCCTTTCATCCTCC-3'; Bax, forward, 5'-GGG TTGTCGCCCTTTTCTAC-3', and reverse, 5'-CTGGAGACA GGGACATCAGT-3'; caspase-3, forward, 5'-TGCTATTGT GAGGCGGTTGTAG-3', and reverse, 5'-GGCACACCCACC GAAAAC-3'; and β-actin, forward, 5'-CATTAAGGAGAA GCTGTGCT-3', and reverse, 5'-GTTGAAGGTAGTTTCGTG GA-3'. The relative expression levels were quantified with the $2^{-\Delta\Delta Cq}$ method (18).

Western blotting. The protein was extracted from cells using a mixture of Pierce RAPI Buffer (Thermo Fisher Scientific, Inc.) and Halt Protease Inhibitor Cocktail (Thermo Fisher Scientific, Inc.) at a ratio of 100:1. The protein concentration was determined using the bicinchoninic acid assay (Thermo Fisher Scientific, Inc.). Lysates of MGC803 and MKN45 cells were centrifuged at 4˚C for 10 min at 12,000 x g. Equal amounts of proteins (50 µg) were separated by 10% SDS-PAGE and transferred onto polyvinylidene fluoride membranes. Following blocking with 5% bovine serum albumin (Invitrogen; Thermo Fisher Scientific, Inc.)/PBS with Tween 20 buffer at 37˚C for 1 h, membranes were incubated with primary antibodies at 4˚C overnight. Subsequently, blots were incubated with horseradish peroxidase-linked secondary antibodies at 37˚C for 2-3 h. Immunoreactive proteins were detected using the SuperSignal West Pico Chemiluminescent substrate (Thermo Fisher Scientific, Inc.). Bands were quantified using ImageJ 1.50i (National Institutes of Health, Bethesda, MD, USA) (19).

Cell apoptosis assay. Apoptosis of MGC803 and MKN45 cells was evaluated using flow cytometry with an Annexin V-fluorescein isothiocyanate (FITC)/propidium iodide (PI) Apoptosis kit (BD Pharmingen; BD Biosciences, Franklin Lakes, NJ, USA), according to the manufacturer's protocol. Cells were washed with cold PBS twice and resus-pended in 100 µl 1X binding buffer, following which 5 µl Annexin V-FITC and 5 µl PI were added for 10 min at room temperature in the dark. Finally, 400 µl binding buffer was added to the cells, which were analyzed by a flow cytometer. All data were analyzed by FlowJo software version 7.6.1 (FlowJo LLC, Ashland, OR, USA).

Proliferation assay. An MTT assay was used to measure the viability of MGC803 and MKN45 cells. A 100 µl GC cell suspension containing 2,000 cells was seeded in each well of a 96-well plate and incubated for 4 h in an atmosphere containing 5% CO_2 at 37˚C. Subsequently, 100 µl MTT solution was added to each well and incubated for 4 h in an atmosphere containing 5% CO_2 at 37˚C. A total of 100 µl dimethyl sulfoxide was added to each well and incubated again for 4 h in an atmosphere containing 5% CO_2 at 37˚C, in order to dissolve the formazan crystals. Absorbance was measured at 570 nm using a Multiskan Plate Reader (Thermo Fisher Scientific, Inc.). Each experiment was repeated three times.

Transwell assay. Cell migration ability was measured using cell culture inserts (24-well type, 8-µm pore size;
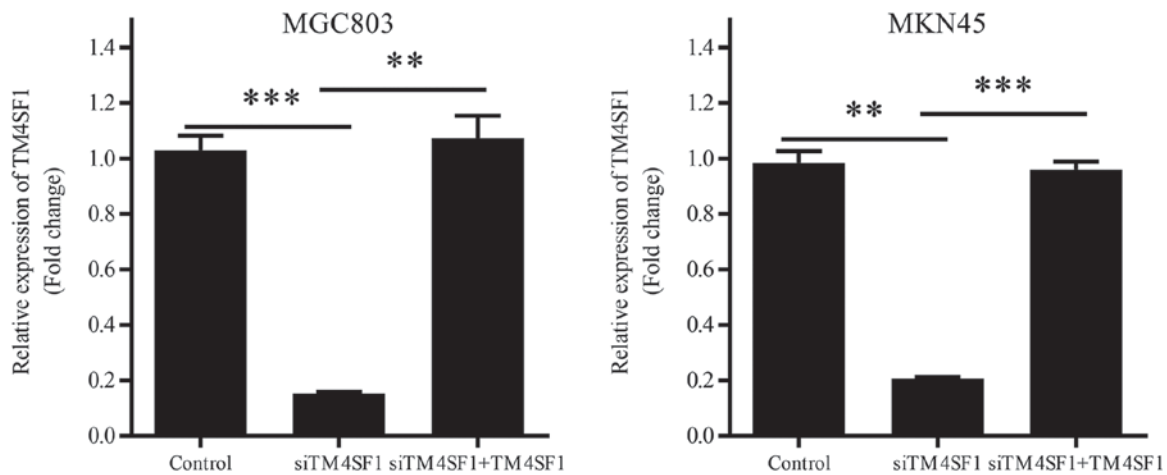
Figure 1. Efficiency of TM4SF1 siRNA and the TM4SF1-expressing plasmid. Cell lines MGC803 and MKN45 were transfected with blank vectors, TM4SF1 siRNA or TM4SF1 siRNA and the TM4SF1-expressing plasmid. TM4SF1 expression was measured using reverse transcription-quantitative polymerase chain reaction and expressed as fold-change. Each column is presented as the mean of three separate experiments. **P<0.01 and ***P<0.001. siRNA, small interfering RNA; TM4SF1, transmembrane-4 L6 family member 1; siTM4SF1, cells transfected with TM4SF1 siRNA; control, cells transfected with blank vectors; siTM4SF1+TM4SF1, cells transfected with TM4SF1 siRNA and the TM4SF1-expressing plasmid.

Corning Incorporated, Corning, NY, USA). Subsequently, $1x10^5$ MGC803 and MKN45 cells were added into the upper chambers with serum-free RPMI-1640 medium, while the lower chambers were filled with 500 μl RPMI-1640 supplemented with 10% FBS. Following 16 h of incubation at 37°C, the cells that had migrated to the lower chamber were fixed in 100% methanol at room temperature for 30 min and stained with 0.1% crystal violet at room temperature for 20 min. Cells were visualized and the number in 10 random fields were counted under a light microscope (Olympus Corporation, Tokyo, Japan; magnification, x400).

Wound-healing assay. MGC803 and MKN45 cells were seeded in 6-well plates ($5x10^5$ cells/well) with 2 ml RPMI-1640 medium supplemented with 10% FBS. Once 80% confluence was attained, scratches were produced using a 100-μl pipette tip. Wound healing was observed and images of the migration distance were captured at room temperature at 0 and 24 h under a light microscope (Olympus Corporation; magnification, x100).

Statistical analysis. Statistical analysis was performed using SPSS v21.0 (IBM Corp., Armonk, NY, USA) and GraphPad Prism v5.0 (GraphPad Software, Inc., La Jolla, CA, USA) software. Differences between ≥2 groups were assessed using one-way analysis of variance followed by Tukey's post-hoc test. Data are presented as the mean ± standard error of the mean. All experiments comprised three replicates and were performed at least twice independently. P<0.05 was considered to indicate a statistically significant difference, unless otherwise stated.

Results

Efficiency of TM4SF1 siRNA and TM4SF1-expressing plasmid. The efficiency of gene transduction was confirmed by RT-qPCR. The results for MGC803 and MKN45 were consistent with one another. TM4SF1 mRNA expression was

significantly downregulated in the siTM4SF1 group, compared with the control group (P<0.001 and P<0.01, respectively; Fig. 1). Transfection with the TM4SF1-expressing plasmid was demonstrated to significantly reversed the reduction of TM4SF1 mRNA expression, compared with the siTM4SF1 group (P<0.01 and P<0.001, respectively; Fig. 1).

TM4SF1 influences the expression of apoptotic molecules in MGC803 and MKN45 cells. The expression of apoptotic molecules was assessed using RT-qPCR. By comparing the siTM4SF1 group with the control group, it was demonstrated that TM4SF1 silencing significantly decreased Bcl2 mRNA expression, whilst caspase-3 and Bax expression levels were upregulated (P<0.001; Fig. 2A). By comparing the siTM4SF1 and siTM4SF1+TM4SF1 groups, it was demonstrated that TM4SF1 upregulation significantly reversed alterations in the mRNA expressions profile of MGC803 and MKN45 cells (P<0.001; Fig. 2A). Western blotting was used to assess these genes at the protein level. The results indicated that the effects of TM4SF1 on Bcl2, caspase-3 and Bax protein expression levels were consistent with the mRNA expression level results, and the comparison was statistically significant (P<0.05; Fig. 2B). Flow cytometry was used to verify the role of TM4SF1 in the apoptosis of MGC803 and MKN45 cells. The results demonstrated that the apoptosis of MGC803 and MKN45 cells in the siTM4SF1 group was significantly promoted, compared with the control group (P<0.01 and P<0.05, respectively; Fig. 3). Rescue experiments demonstrated that the apoptosis of MGC803 and MKN45 cells in the siTM4SF1+TM4SF1 group was significantly decreased, compared with the siTM4SF1 group (P<0.05 and P<0.01, respectively; Fig. 3). These results were consistent with changes in the expression of the aforementioned apoptotic molecules.

Roles of TM4SF1 in the proliferation, migration and invasion of MGC803 and MKN45 cells. To investigate the biological functions of TM4SF1 in GC, the proliferation, invasion and migration of MGC803 and MKN45 cells were assessed
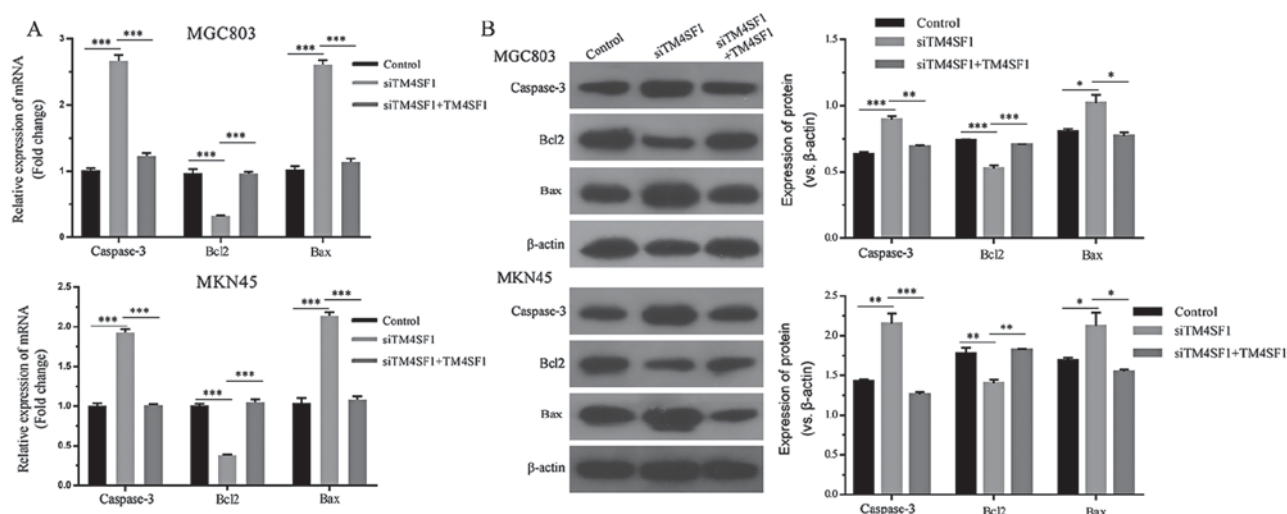
Figure 2. TM4SF1 influences the expression of apoptotic molecules in gastric cancer cells. (A) TM4SF1 silencing decreased the expression of Bcl2 mRNA and upregulated caspase-3 and Bax mRNA. Upregulating TM4SF1 reversed these effects. (B) Values represent fold-change compared with β-actin. Silencing the TM4SF1 gene decreased Bcl2 protein expression, and increased caspase-3 and Bax protein expression. Upregulating TM4SF1 reversed these changes. Each column is presented as the mean of three separate experiments. *P<0.05, **P<0.01 and ***P<0.001. Bcl2, B-cell lymphoma 2; Bax, Bcl2-associated X; TM4SF1, transmembrane-4 L6 family member 1; siTM4SF1, cells transfected with TM4SF1 siRNA; control, cells transfected with blank vectors; siTM4SF1+TM4SF1, cells transfected with TM4SF1 siRNA and the TM4SF1-expressing plasmid.

following the manipulation of TM4SF1 expression. The proliferation of MGC803 and MKN45 cells was measured using an MTT assay. Wound healing assays were used to measure the invasion of MGC803 and MKN45 cells. Transwell assays were used to measure the migration ability of MGC803 and MKN45 cells. The results demonstrated that TM4SF1 knockdown significantly reduced the proliferation of MGC803 and MKN45 cells, compared with the control group (P<0.05, and P<0.05, respectively; Fig. 4A), while TM4SF1 upregulation significantly increased the proliferation of MGC803 and MKN45 cells, compared with the siTM4SF1 group (P<0.05 and P<0.01, respectively; Fig. 4A). TM4SF1 knockdown significantly reduced the invasion of MGC803 and MKN45 cells, compared with the control group (P<0.05 and P<0.05, respectively; Fig. 4B), while TM4SF1 upregulation significantly increased the invasion of MGC803 and MKN45 cells, compared with the siTM4SF1 group (P<0.01 and P<0.001, respectively; Fig. 4B). TM4SF1 knockdown resulted in significantly reduced migration of MGC803 and MKN45 cells, compared with the control group (P<0.001 and P<0.05, respectively; Fig. 4C), while TM4SF1 upregulation significantly increased migration of MGC803 and MKN45 cells, compared with the siTM4SF1 group (P<0.001 and P<0.01, respectively; Fig. 4C).

Discussion

A number of studies have indicated that TM4SF1 may inhibit tumor apoptosis and promote tumor proliferation (10-12); however, the underlying mechanisms are notably complex. Sun *et al* (12) reported that the phosphoinositide 3-kinase (PI3K)/AKT/mechanistic target of rapamycin (mTOR) signaling pathway served a role in TM4SF1 regulation in breast cancer cell apoptosis. The PI3K/AKT/mTOR signaling pathway has been reported to regulate apoptosis in a number of tumor types, including breast cancer, glioma and nasopharyngeal cancer (20-22). The anti-apoptotic effects of the

PI3K/AKT/mTOR signaling pathway are dependent upon downstream apoptosis-associated proteins, including Bcl2, Bax and caspase-3 (23). Bcl2, as a member of the Bcl2 family, can prevent the release of cytochrome *c* from the mitochondria to the cytoplasm, thereby inhibiting apoptosis (24). The Bcl2 family is comprised of polarized groups of proteins containing pro-apoptotic proteins and anti-apoptotic proteins, and cell survival or apoptosis depend on the balance between these two types (25). Bax, a pro-apoptotic protein belonging to the Bcl2 family, can form a heterodimer with Bcl2 to inhibit its function (24). Caspase-3 is the most critical apoptotic protease involved in apoptosis. DNA-dependent protein kinase and poly adenosine diphosphate ribose polymerase are important DNA repair enzymes. Caspase-3 can hydrolysate these two enzymes to prevent DNA replication, transcription and injury repair (26). The results of the present study demonstrated that siTM4SF1 upregulates the expression of pro-apoptosis proteins Bax and caspase-3 and downregulates the expression of anti-apoptosis protein Bcl2. This indicates that TM4SF1 may serve as an anti-apoptotic protein in GC.

The results of the present study demonstrated the important role of TM4SF1 in tumor migration and invasion, and the complex mechanisms underlying has been verified in previous studies (27-29). Lekishvili *et al* (27) reported that TM4SF1 may be associated with tetraspanin-enriched microdomains, which are crucial for the pro-migratory activity of membrane proteins; furthermore, TM4SF1 overexpression downregulated the expression of tetraspanins cluster of differentiation (CD)63 and CD82, which are associated with the regulation of surface (28,29). CD82 can promote the proliferation of epidermal growth factor receptor to activate signaling cascades, including the FAK-Lyn-p130CAS-CrkII signaling pathway, which results in reduced cell motility (30). A number of additional molecules are associated with TM4SF1. A previous study demonstrated revealed an association between TM4SF1
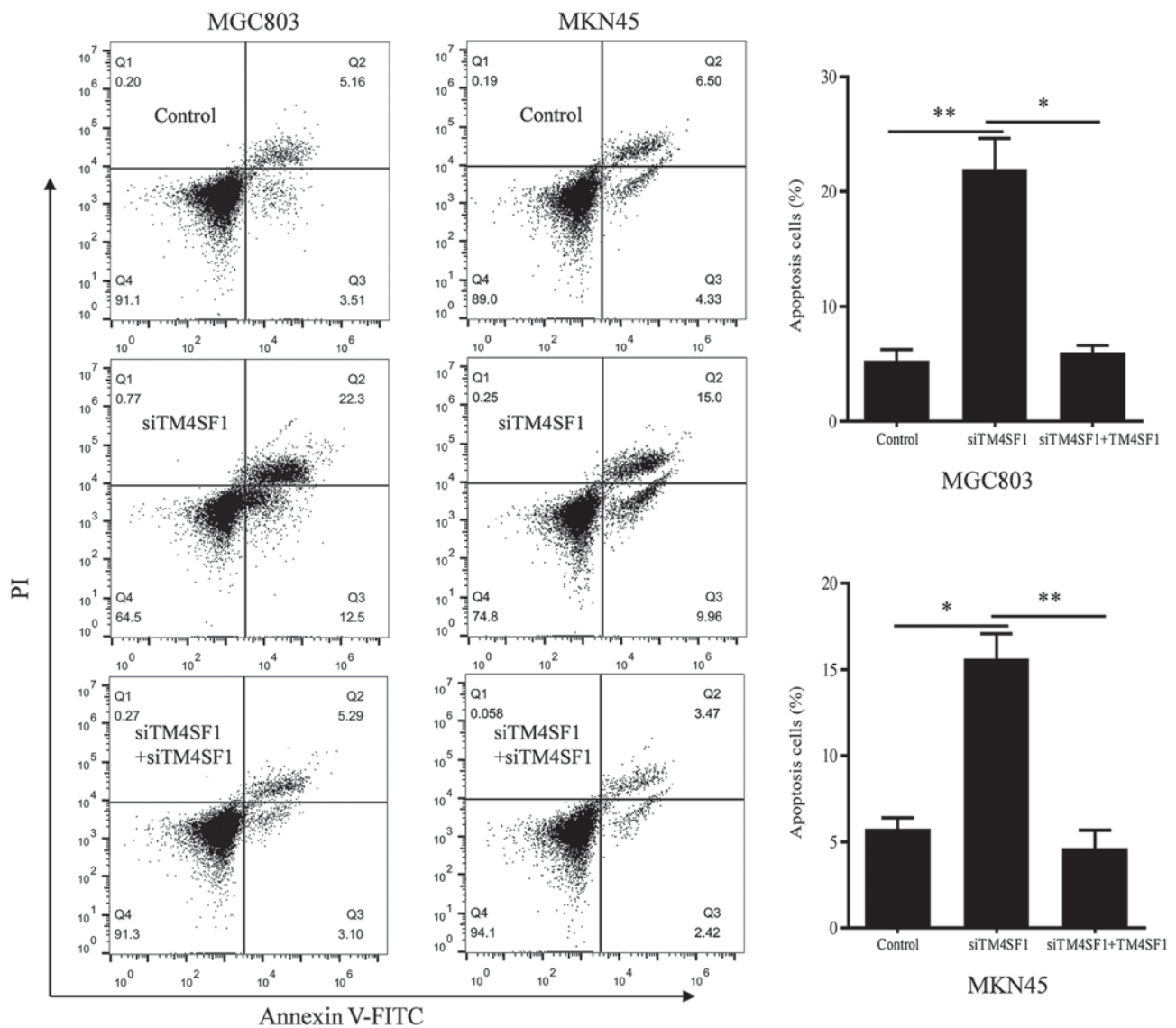
Figure 3. The role of TM4SF1 in gastric cancer cell apoptosis. Apoptosis was assessed using flow cytometry and representative images are depicted. Each column is presented as the means of three separate experiments. $^*P<0.05$, $^{**}P<0.01$. PI, propidium iodide; FITC, fluorescein isothiocyanate; TM4SF1, transmembrane-4 L6 family member 1; siTM4SF1, cells transfected with TM4SF1 siRNA; control, cells transfected with blank vectors; siTM4SF1+TM4SF1, cells transfected with TM4SF1 siRNA and the TM4SF1-expressing plasmid.

and CD13 using coimmunoprecipitation analysis, which can form and enhance cell migration in lung cancer cells (31). In prostate cancer, TM4SF1 was reported to directly target the androgen receptor to promote cell migration (32).

In the present study, functional experiments were performed to verify the effects of TM4SF1 knockdown and upregulation in GC cells. The results demonstrated that TM4SF1 may regulate the apoptosis, proliferation, migration and invasion of GC cells; however, tumor progression is a complicated process with numerous contributing factors, relying on not only the changes in tumor cells but also changes in the tumor microenvironment (33). It is therefore necessary to investigate the role of TM4SF1 in the tumor microenvironment.

Xue *et al* (34) demonstrated that matrix metallopeptidase (MMP)-2, MMP-9 and VEGF are the downstream proteins of TM4SF1, and TM4SF1 overexpression in turn upregulated MMP-2, MMP-9 and VEGF expression. VEGF is known to be a strong angiogenic factor, and the binding

of VEGF and its receptors may stimulate endothelial cell division, proliferation and migration, promote physiological and pathological neovascularization, increase microvascular permeability and promote embryonic hematopoiesis (35-37). VEGF overexpression promotes the secretion of MMP-2 and MMP-9, which may serve a crucial role in tumor invasion and metastasis (28,29). MMP-2, a zinc-dependent proteolytic enzyme that is secreted by tumor and stromal cells in the form of zymogens, has been reported to be closely associated with the development of tumors. Following hydrolysis, MMP-2 promotes the transformation of the extracellular matrix by degrading the main constituents of the basement membrane, including type IV, V, VI and X collagens and gelatin. This promotes tumor neovascularization, invasion and metastasis (38). MMP-9 is the enzyme with the largest molecular weight in the MMP family, and is also secreted in zymogen form. Following activation, MMP-9 can be transformed into type IV collagenase, which degrades and destroys
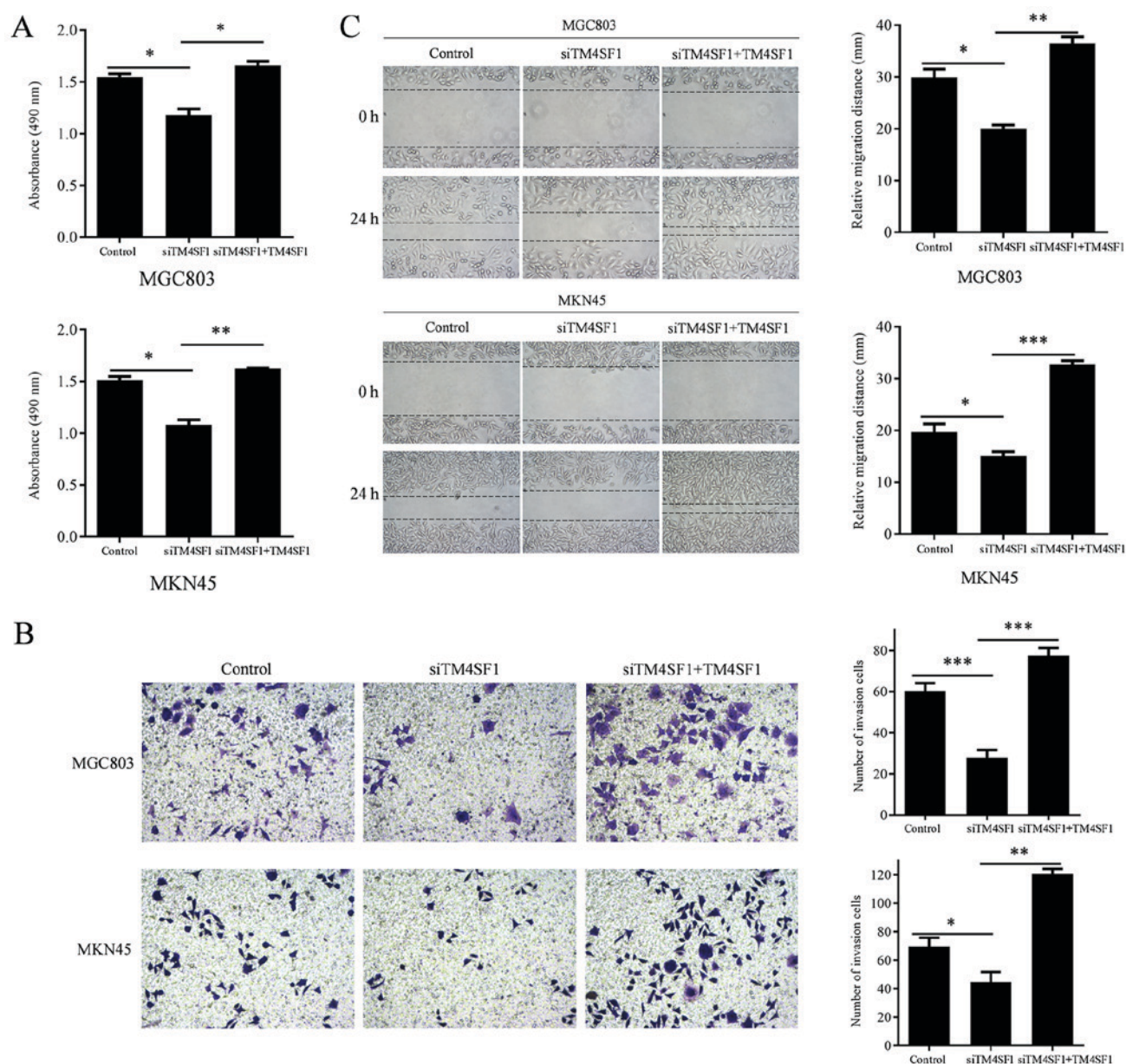
Figure 4. The role of TM4SF1 in the proliferation, migration and invasion of gastric cancer cells. (A) TM4SF1 knockdown reduced GC cell proliferation, while TM4SF1 upregulation increased GC cell proliferation. (B) TM4SF1 knockdown reduced GC cell invasion, while TM4SF1 upregulation increased GC cell invasion. (C) TM4SF1 knockdown resulted in reduced GC cell migration, while TM4SF1 upregulation increased GC cell migration. Each column is presented as the means of three separate experiments. $^*P<0.05$, $^{**}P<0.01$ and $^{***}P<0.001$. TM4SF1, transmembrane-4 L6 family member 1; siTM4SF1, cells transfected with TM4SF1 siRNA; control, cells transfected with blank vectors; siTM4SF1+TM4SF1, cells transfected with TM4SF1 siRNA and the TM4SF1-expressing plasmid.

type IV and V collagens as well as gelatin in the extracellular matrix around the tumor surface. Tumor cells are then able to infiltrate the surrounding tissue via the deficient basement membrane, which ultimately results in tumor invasion and metastasis (39).

The present study was not without limitations. Firstly, experiments were only conducted *in vitro*. Secondly, further research is required to identify the signaling pathways activated by TM4SF1 in GC. Despite these limitations, the results of the present study indicated that TM4SF1 is an anti-apoptotic protein that has the ability to promote the proliferation, migration and invasion of GC cells.

To conclude, the results of the present study indicated that TM4SF1 may serve an important role in the progression

of GC. In the future, TM4SF1 may be considered as a novel therapeutic target of GC; however, future *in vivo* experiments and clinical trials are required.

Availability of data and materials

All data that were generated or analyzed in this study are included in this manuscript.

Authors' contributions

YW conceived and designed the study. XS was a major contributor in writing the manuscript. XS and LL conducted the majority of the experiments. GC performed the statistical analysis. XC cultured the cells and drew the figures. YW conducted the data interpretation. HS performed the literature search and cell transfection. All authors read and approved the manuscript.

Ethics approval and consent to participate

Not applicable.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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