

Long non-coding RNA B3GALT5-AS1 contributes to the progression of gastric cancer via interacting with CSNK2A1

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Abstract. Gastric cancer is a type of cancer that is characterized by high morbidity and mortality rates. Long non-coding RNA (lncRNA) β -1,3-galactosyltransferase 5-AS1 (B3GALT5-AS1) was previously found to be highly expressed in the serum of patients with gastric cancer. However, the regulatory effects of B3GALT5-AS1 in gastric cancer remain poorly understood. The present study aimed to investigate the effects of B3GALT5-AS1 in gastric cancer cell lines. The expression levels of B3GALT5-AS1 were determined in different gastric cancer cell lines (AGS, HGC-27 and MKN-45) using reverse transcription-quantitative PCR. The potential interaction between B3GALT5-AS1 and casein kinase 2 α 1 (CSNK2A1) was evaluated using an RNA binding protein immunoprecipitation and RNA pull down assays. Western blot analysis was performed to measure protein expression levels. Cell Counting Kit-8 assay was utilized to determine cell viability, whilst cell invasion and migration were assessed using Transwell and wound healing assays, respectively. Apoptotic cells were evaluated using TUNEL assays. The results showed that B3GALT5-AS1 expression was upregulated in MKN-45 cells compared with the control group. In addition, B3GALT5-AS1 could bind to CSNK2A1 to regulate its expression. B3GALT5-AS1 knockdown attenuated cell viability, invasion and migration, whilst promoting cell apoptosis. These effects were partly reversed by CSNK2A1 overexpression. Overall, results of the present study revealed that interference with B3GALT5-AS1 impeded gastric cancer cell migration and invasion whilst promoting apoptosis by regulating CSNK2A1 expression. These findings suggested that B3GALT5-AS1 and CSNK2A1 may serve a tumorigenic role in the progression of gastric cancer and serve as therapeutic targets for this type of cancer.

Introduction

Gastric cancer is one of the most common and incurable types of cancer (1,2). Gastric cancer is associated with high rates

of incidence and mortality, which imposes a considerable economic burden on society with 1 million new cases annually in the United States (1,2). However, the overall survival rate of patients with early stages of gastric cancer is significantly higher compared with that in patients with advanced stages of gastric cancer (3). Therefore, diagnosis of gastric cancer in the early stages is critical for improving the survival rate. In addition to upper gastrointestinal endoscopy and the detection of common tumor biomarkers, including carcinoembryonic antigen (CEA), cancer antigen (CA)72-4 and CA19-9 in the serum (4-6), long non-coding RNAs (lncRNAs) is garnering interest for reported their role in the early diagnosis, therapy and prognosis of gastric cancer (7,8).

The effects of lncRNAs in gastric cancer are gradually becoming unraveled. Wei and Wang (9) showed that lncRNA maternally expressed 3 was highly expressed in gastric cancer tissues compared with that in adjacent normal tissues (9). In addition, other studies revealed that lncRNA small nucleolar host gene (SNHG)1 and SNHG7 overexpression could promote gastric cancer cell proliferation (10,11). A study demonstrated that B3GALT5-AS1 expression was markedly increased in patients with gastric cancer compared with those with normal colonic epithelia (12). Moreover, high serum B3GALT5-AS1 levels were found to be associated with TNM stage and lymph node metastasis (13), suggesting a role of B3GALT5-AS1 in gastric cancer occurrence and progression. Another study suggested that serum levels of B3GALT5-AS1 could also serve as a diagnostic biomarker of colorectal cancer (14). Indeed, B3GALT5-AS1 was demonstrated to be localized predominantly in the nucleus, where it could directly bind to the promoter of microRNA (miR)-203 to upregulate the expression of the target genes of miR-203, which in turn suppresses colon cancer metastasis to the liver (12). However, to date, the regulatory mechanism of B3GALT5-AS1 in gastric cancer remains elusive.

It has been previously demonstrated that casein kinase 2 α 1 (CSNK2A1) is associated with cell invasion and migration in several types of cancer, including lung (15) and breast cancer (16). Additionally, overexpression of CSNK2A1 in gastric cancer cells enhanced cell proliferation, invasion and migration (17), suggesting a regulatory role of CSNK2A1 in gastric cancer.

Therefore, in the present study, the expression profile of B3GALT5-AS1 was determined in gastric cancer cell lines. Additionally, whether B3GALT5-AS1 served a role in gastric

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cancer cell proliferation, migration, invasion and apoptosis and its potential relationship with CSNK2A1 was assessed.

Materials and methods

Cell culture. Gastric cancer cell lines, AGS, HGC-27 and MKN-45 were purchased from the China Infrastructure of Cell Line Resources, Institute of Basic Medical Sciences, Chinese Academy of Medical Sciences. The human gastric mucosa cell line GES-1 (cat. no. CL-0563) was obtained from Procell Life Science & Technology Co., Ltd.. All cells were cultured in RPMI-1640 medium (Gibco; Thermo Fisher Scientific, Inc.) supplemented with 10% FBS (Gibco; Thermo Fisher Scientific, Inc.) and maintained in an incubator with 5% CO₂ at 37°C.

Plasmid transfection. For B3GALT5-AS1 knockdown, two short hairpin RNAs (shRNA/sh), namely sh-B3GALT5-AS1-1 (5'-GCATAAGAGAGACCAACTTGG-3') and sh-B3GALT5-AS1-2 (5'-GCAAGACAGCGCATTGAT TGG-3'), were constructed using the pGPU6/Neo plasmid (Shanghai GenePharma Co., Ltd.). Scrambled shRNA [shRNA-negative control (NC)] served as the NC (5'-ACTTGC GCTTGCGAAAATCTATATAGC-3'). The pcDNA-MFHAS1 plasmid (Shanghai GenePharma Co., Ltd.) was used for CSNK2A1 overexpression (ov-CSNK2A1). The empty vector, ov-NC, served as the negative control for the overexpression experiments. Cells were seeded into six-well plates at a density of 5×10⁵ cells/well. When they reached 80% confluence, cells were transfected with 3 µg shRNA-encoding plasmids or 5 µg overexpression plasmids using Lipofectamine® 3000 (Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer's instructions. Following incubation for 48 h at 37°C, transfected cells were used for subsequent experiments.

RNA extraction and reverse transcription-quantitative PCR (RT-qPCR). Total RNA was extracted from the four cell lines (GES-1, AGS, HGC-27 and MKN-45) using TRIzol® reagent (Invitrogen; Thermo Fisher Scientific, Inc.). Following reverse transcription using AMV Reverse Transcription kit (cat. no. A3500; Promega Corporation), this reaction was performed at 25°C for 5 min, 50°C for 20 min, and then 75°C for 5 min. The FastStart Universal SYBR Green Master kit (Roche Diagnostics GmbH) was used for qPCR in an ABI PRISM 7900 HT system (Applied Biosystems; Thermo Fisher Scientific, Inc.). The primer sequences used were as follows: B3GALT5-AS1 forward, 5'-GATCCACGTCCA GGCTCACT-3' and reverse, 5'-GTGCTGGCTGTCAGGATG AG-3'; CSNK2A1 forward, 5'-GAACGCTTTGTCCACAGT GA-3' and reverse, 5'-TATCGCAGCAGTTTGTCCAG-3' and GAPDH forward, 5'-AACAGCGACACCCACTCCTC-3' and reverse, 5'-GGAGGGGAGATTTCAGTGTGGT-3'. The thermocycling conditions for PCR reaction was as follows: Initial denaturation for 5 min at 95°C, followed by 40 cycles of 30 sec at 95°C and 45 sec at 65°C. The relative mRNA expression was calculated using the 2^{-ΔΔC_q} method (18) and GAPDH served as the internal control.

Bioinformatics analysis. The starBase 2.0 database (<http://starbase.sysu.edu.cn/>) was used to predict the binding sites between B3GALT5-AS1 and CSNK2A1.

RNA immunoprecipitation (RIP) assay. RIP assay was performed using the Millipore Magna RIP™ RNA kit (EMD Millipore). Briefly, MKN-45 cells were resuspended in RIP lysis buffer (EMD Millipore) and incubated on ice for 5 min. Subsequently, the cell lysate (100 µl per antibody per RIP) was incubated with an anti-CSNK2A1 antibody (1:1,000; cat. no. ab70774; Abcam) or IgG antibody (0.2 µg/ml; cat. no. ab190475; Abcam) and 40 µl protein A/G magnetic beads (EMD Millipore) at 4°C overnight. The protein-RNA complexes were digested with 2 µl proteinase K buffer at 55°C for 30 min. The beads buffer was spun down at 2,000 x g for 30 sec, and the supernatant was transferred to a fresh tube. Finally, the extracted RNA was subjected to RT-qPCR.

RNA pull-down assay. RNA pull-down assay was performed using Pierce™ Magnetic RNA-Protein Pull-Down Kit (Thermo Fisher Scientific, Inc.). Briefly, MKN-45 cells were lysed by RIPA lysis buffer (Sigma-Aldrich; Merck KGaA). B3GALT5-AS1 primers were 5'-CCTTGAGAGACGAAG CAC-3' (sense) and 5'-ATTTACGGATGAGACGAC-3' (antisense). B3GALT5-AS1 was labeled with biotin using Pierce RNA 3' End Desthiobiotinylation Kit (Thermo Fisher Scientific, Inc.) and then bound to streptavidin magnetic beads (~10 µg), before this complex was incubated with 1 ml cell lysates at 4°C for 1 h. RNA-protein complexes were eluted by adding 30 µl SDS sample buffer to the beads and heating at 95°C for 5 min. The samples were cooled on ice for 1 min, 2 µl Benzonase added, and then incubated for 15 min at room temperature. The sample buffer was heated again at 95°C for 5 min. The magnetic beads were spun down at room temperature at 12,000 x g for 1 min. The presence of CSNK2A1 protein in the RNA-protein complexes was analyzed by western blotting.

Western blotting. Total proteins were isolated from cell lysates using RIPA lysis buffer (Sigma-Aldrich; Merck KGaA) and then quantified using a bicinchoninic acid protein assay kit (Thermo Fisher Scientific, Inc.). Subsequently, 20 µg protein extracts per lane were separated by 10% SDS-PAGE and transferred onto nitrocellulose membranes (EMD Millipore). Following blocking with 5% skim milk at room temperature for 2 h, membranes were incubated with antibodies against CSNK2A1 (1:1,000; cat. no. ab70774; Abcam), matrix metalloproteinase (MMP)2 (1:2,000; cat. no. AF0577; Affinity Biosciences), MMP9 (1:2,000; cat. no. AF5228; Affinity Biosciences), phosphorylated (p)-c-Met (1:1,000; cat. no. AF3128; Affinity Biosciences), c-Met (1:1,000; cat. no. AF6128; Affinity Biosciences), Bcl-2 (1:5,000; cat. no. AF6139; Affinity Biosciences), Bax (1:5,000; cat. no. AF0120; Affinity Biosciences), cleaved caspase-3 (1:2,000; cat. no. AF7022; Affinity Biosciences), caspase-3 (1:2,000; cat. no. AF6311; Affinity Biosciences) or GAPDH (1:5,000; cat. no. AF7021; Affinity Biosciences) at 4°C overnight. The next day, the membranes were incubated with corresponding HRP-conjugated secondary antibodies (1:5,000; cat. no. S0001; Affinity Biosciences) at room temperature for 2 h. The protein bands were visualized using the Immobilon Western Chemilum HRP substrate (EMD Millipore) and analyzed using Image Lab Software 3.0 (Bio-Rad Laboratories, Inc.).

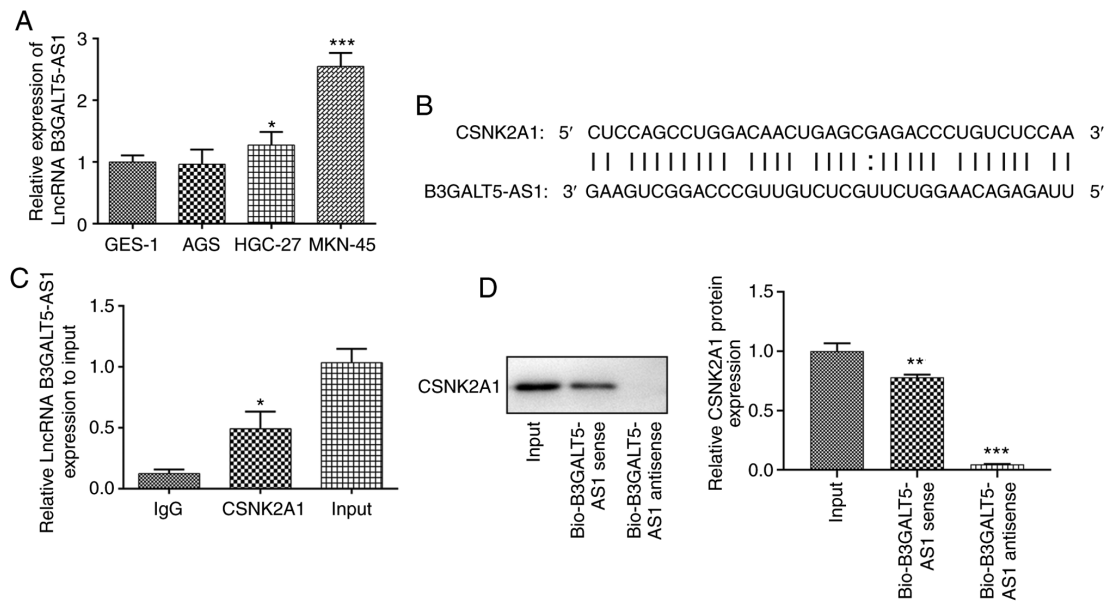


Figure 1. Interaction between B3GALT5-AS1 and CSNK2A1. (A) Expression levels of B3GALT5-AS1 in a range of gastric cell lines were determined by reverse transcription-quantitative PCR. * $P < 0.05$ and *** $P < 0.001$ vs. GES-1. (B) Potential binding sites between B3GALT5-AS1 and CSNK2A1 were predicted using the starBase 2.0 database. Interaction between B3GALT5-AS1 and CSNK2A1 was evaluated using (C) RNA immunoprecipitation. * $P < 0.05$ vs. IgG. (D) RNA pull-down assays. ** $P < 0.01$ and *** $P < 0.001$ vs. Input. B3GALT5-AS1, β -1,3-galactosyltransferase 5-AS1; CSNK2A1, casein kinase 2 α 1; LncRNA, long non-coding RNA.

Cell Counting Kit 8 (CCK-8) assay. For CCK-8 assays, MKN-45 cells were seeded into a 96-well plates at a density of 5,000 cells per well and incubated at 37°C overnight. The next day, 10 μ l CCK-8 reagent (Dojindo Molecular Technologies, Inc.) was added into each well and cells were incubated for 2 h at 37°C. Finally, the optical density (OD) in each well was measured using a microplate reader (Bio-Rad Laboratories, Inc.) at a wavelength of 450 nm.

Cell invasion assay. The upper chamber of the Transwell chamber (Corning, Inc.) was coated with 100 μ l Matrigel (1 mg/ml; BD Biosciences) and incubated at 37°C for 4 h. The cell density was then adjusted to 2×10^5 cells/ml and 100 μ l cell suspension in serum-free RPMI-1640 medium was added to the upper chamber and cultured at 37°C in the presence of 5% CO₂ for 24 h. The lower chamber was supplemented with 600 μ l RPMI-1640 medium containing 20% FBS. At 24 h post-incubation, invasive cells were fixed with 4% paraformaldehyde at room temperature for 0.5 h, stained with 0.1% crystal violet at 37°C for 10 min and photographed in randomly selected nine fields of view under a light microscope (magnification, $\times 100$).

Cell migration assay. MKN-45 cells were seeded into a six-well plate (5×10^5 cells/well) and when they reached 100% confluence, scratches were created using a 20- μ l pipette tip. The medium was then replaced with fresh medium containing 1% FBS and cells were cultured at 37°C for 24 h (19). Finally, images of the migrated cells were captured at 0 and 24 h in randomly selected fields of view under a light microscope (magnification, $\times 100$). The migration distance was calculated as the width of the scratch at 24 h minus the width of the scratch at 0 h. The relative migration rate was calculated by normalizing to the control group.

Cell apoptosis assay. Apoptotic cells were assessed using an One Step TUNEL Apoptosis Assay kit (Beyotime Institute of Biotechnology). Briefly, MKN-45 cells (3×10^5 cells/ml) were fixed with 4% paraformaldehyde at room temperature for 30 min and then incubated with 0.3% Triton X-100 at room temperature for 5 min. Subsequently, each well of the 24-well plate was supplemented with 50 μ l TUNEL detection reagent for 60 min at 37°C in the dark. DAPI staining solution (10 μ g/ml) was used to visualize all nuclei for 5 min at 37°C. The coverslips were then washed with PBS and mounted on slides with anti-fading solution. The apoptotic cells (green fluorescence) in three random fields were observed under a fluorescence microscope (magnification, $\times 100$).

Statistical analysis. All data in the study are expressed as the mean \pm SD and analyzed using an independent unpaired t-test or one-way ANOVA followed by a Tukey's post hoc test. Each experiment was repeated \geq three times. $P < 0.05$ was considered to indicate a statistically significant difference.

Results

Interaction between B3GALT5-AS1 and CSNK2A1. The expression levels of B3GALT5-AS1 were first determined in a range of gastric cell lines. The expression levels of B3GALT5-AS1 in HGC-27 cells were higher compared with that in the human gastric mucosa cell line GES-1 (Fig. 1A). Compared with that in GES-1 cells, B3GALT5-AS1 expression was significantly higher in MKN-45 cells (Fig. 1A). It has been previously suggested that B3GALT5-AS1 may originate from autocrine gastric tumor cells and is closely associated with gastric tumor metastasis (13). However, the specific regulatory mechanism underlying the effect of B3GALT5-AS1 on tumor metastasis remains unclear. Therefore, based on the expression

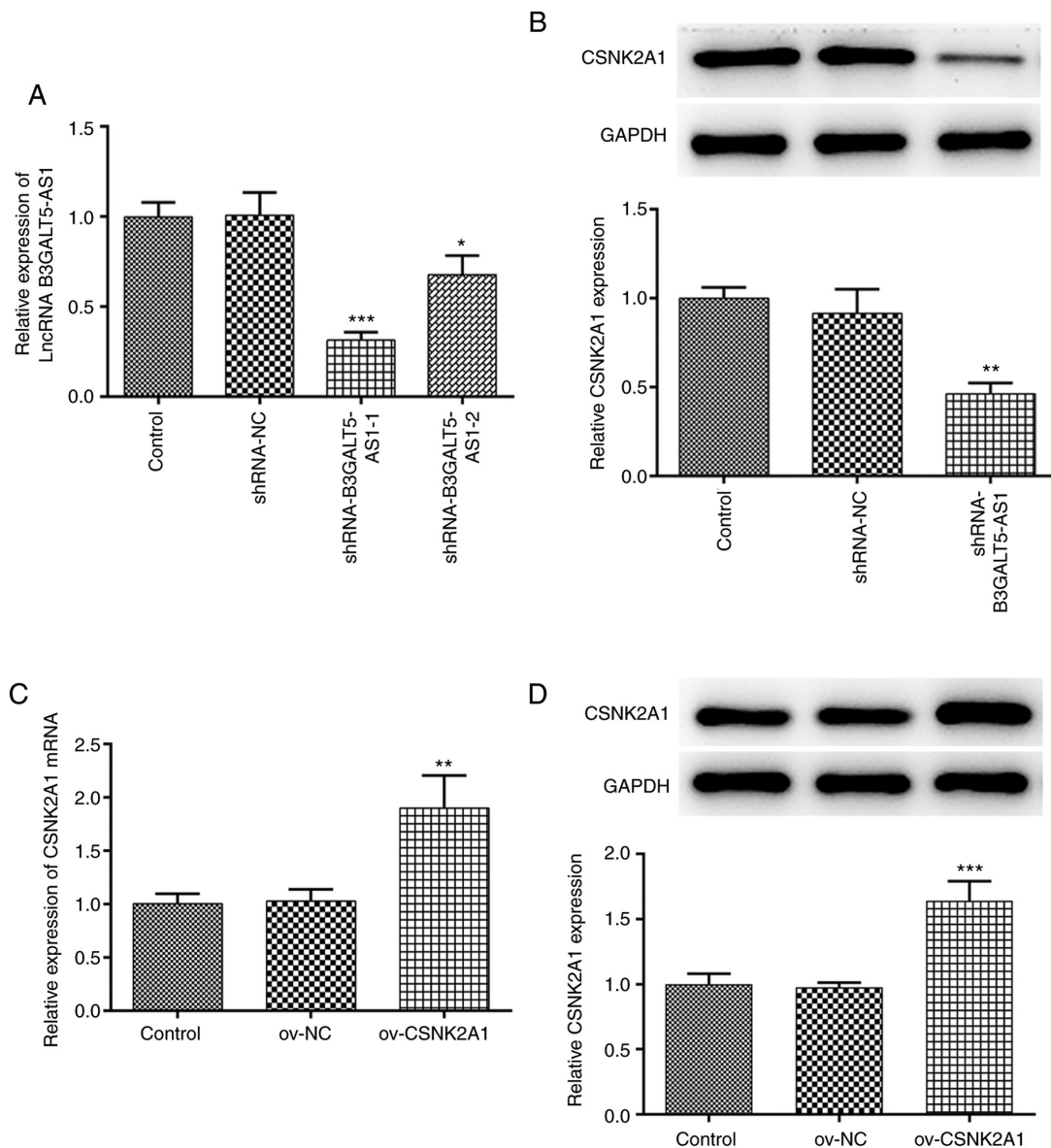


Figure 2. B3GALT5-AS1 silencing and CSNK2A1 overexpression on MKN-cells. (A) Expression levels of B3GALT5-AS1 in MKN-45 cells following transfection with shRNA-NC, shRNA-B3GALT5-AS1-1 or shRNA-B3GALT5-AS1-2 were determined by RT-qPCR. (B) Western blot analysis showing the protein expression levels of CSNK2A1. * $P < 0.05$, ** $P < 0.01$ and *** $P < 0.001$ vs. shRNA-NC. (C) mRNA expression levels of CSNK2A1 following transfection with ov-NC or ov-CSNK2A1 were assessed by RT-qPCR. (D) Western blot analysis showing the protein expression levels of CSNK2A1 following transfection of MKN-45 cells with ov-NC or ov-CSNK2A1. ** $P < 0.01$ and *** $P < 0.001$ vs. ov-NC. B3GALT5-AS1, β -1,3-galactosyltransferase 5-AS1; CSNK2A1, casein kinase 2 α 1; NC, negative control; RT-qPCR, reverse transcription-quantitative PCR; LncRNA, long non-coding RNA; Ov, expression; shRNA, short hairpin RNA.

of B3GALT5-AS1, the MKN-45 cell line was used for the subsequent experiments. mRNA of CSNK2A1 was located in Chr20:453618-453657. Bioinformatics analysis predicted that B3GALT5-AS1 could bind to the sequence of CSNK2A1 (Fig. 1B). Notably, bioinformatics analysis predicted binding sites between the sequences of B3GALT5-AS1 and CSNK2A1, suggesting that B3GALT5-AS1 may modulate CSNK2A1 expression by direct binding (12,20). As shown in Fig. 1C, the interaction between B3GALT5-AS1 and CSNK2A1 was evaluated using RIP assays. Further evidence of a CSNK2A1-complex interaction was found via a RNA pull down assay (Fig. 1D). Since B3GALT5-AS1 was found to be enriched in the CSNK2A1-complex, direct binding of CSNK2A1 to B3GALT5-AS1 was reproduced *in vitro* by a RNA pull-down assay.

Efficiency of B3GALT5-AS1 knockdown and CSNK2A1 overexpression. Since B3GALT5-AS1 expression was the highest in MKN-45 cells, these cells were transfected with plasmids encoding shRNAs to knockdown the expression B3GALT5-AS1. As shown in Fig. 2A, between the two shRNA clones, shRNA-B3GALT5-AS1-1 exhibited the more potent silencing effect compared with that mediated by shRNA-B3GALT5-AS1-2. The protein expression level of CSNK2A1 in MKN-45 cells transfected with shRNA-B3GALT5-AS1-1 was significantly lower compared with that in the shRNA-NC group. However, no significant differences were observed in the protein expression of CSNK2A1 between the control and shRNA-NC groups (Fig. 2B). Subsequently, MKN-45 cells were transfected with a CSNK2A1 overexpression plasmid. mRNA and protein

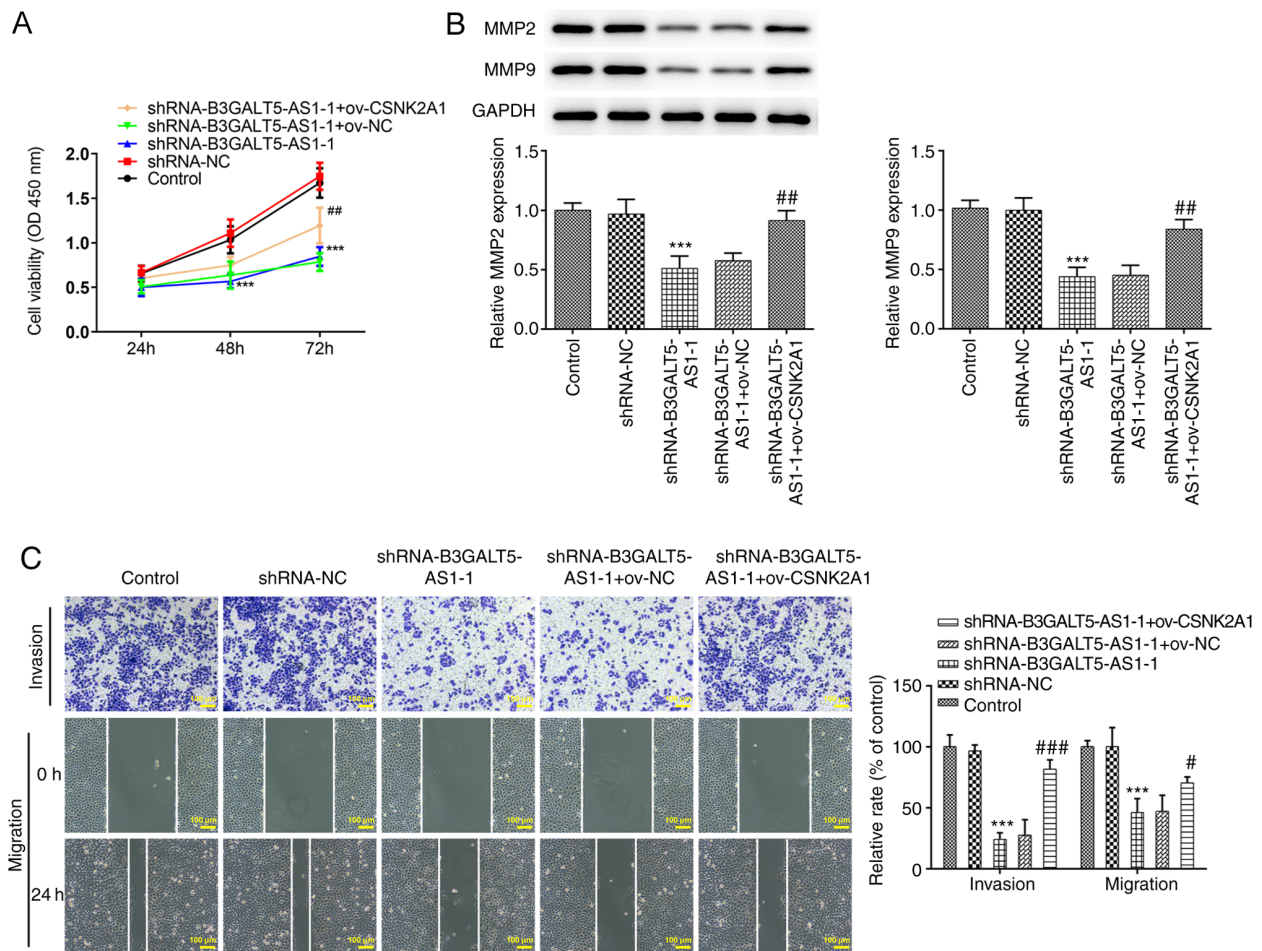


Figure 3. B3GALT5-AS1/CSNK2A1 axis regulates MKN-45 cell migration and invasion. (A) Cell viability in the different transfection groups was evaluated by Cell Counting Kit-8 assay. (B) Western blot analysis showing the protein expression levels of MMP2 and MMP9 in different groups. (C) Cell invasion and migration in different groups of cells after transfection. Scale bar, 100µm, *** $P < 0.001$ vs. shRNA-NC; # $P < 0.05$, ## $P < 0.01$ and ### $P < 0.001$ vs. shRNA-B3GALT5-AS1-1 + ov-NC. B3GALT5-AS1, β -1,3-galactosyltransferase 5-AS1; CSNK2A1, casein kinase 2 α 1; MMP, matrix metalloproteinase; LncRNA, long non-coding RNA; Ov, expression; shRNA, short hairpin RNA; OD, optical density.

expression of CSNK2A1 were both significantly upregulated following cell transfection with ov-CSNK2A1 compared with those transfected with ov-NC (Fig. 2C and D).

B3GALT5-AS1/CSNK2A1 axis controls cell migration and invasion. Subsequently, effects of gene knockdown or overexpression on the migratory and invasive behavior of MKN-45 cells were investigated. Cell viability was significantly decreased following cell transfection with shRNA-B3GALT5-AS1-1 compared with those transfected with shRNA-NC at 72 h. However, co-transfection with shRNA-B3GALT5-AS1-1 and ov-CSNK2A1 partially but significantly elevated cell viability compared with that in the shRNA-B3GALT5-AS1-1 + ov-NC group (Fig. 3A). MMPs are involved in the remodeling of the extracellular matrix and basement membrane, which is a key process in the invasion and metastasis of cancer cells (21,22). The protein expression levels of MMP2 and MMP9 were significantly reduced after cell transfection with shRNA-B3GALT5-AS1-1 compared with that in the shRNA-NC group, which was significantly reversed by CSNK2A1 plasmid co-transfection (Fig. 3B). Transwell assays showed that the invasive ability of MKN-45 cells was significantly decreased following B3GALT5-AS1

knockdown compared with that in the shRNA-NC group (Fig. 3C). B3GALT5-AS1 silencing also significantly attenuated the migratory ability of MKN-45 cells compared with that in cells transfected with shRNA-NC (Fig. 3C). For both migration and invasion, CSNK2A1 plasmid co-transfection significantly reversed the inhibitory effects of B3GALT5-AS1 knockdown (Fig. 3C).

B3GALT5-AS1/CSNK2A1 axis modulates cell apoptosis. Furthermore, a TUNEL assay was performed to measure cell apoptosis. The number of apoptotic cells in the shRNA-B3GALT5-AS1-1 group was markedly higher compared with that in the shRNA-NC or control group, which was reversed by CSNK2A1 plasmid co-transfection (Fig. 4A). The levels of the pro-apoptotic proteins Bax and cleaved caspase-3 were both significantly upregulated following B3GALT5-AS1 knockdown, but their protein levels were significantly restored after CSNK2A1 overexpression in addition to B3GALT5-AS1 knockdown (Fig. 4B). By contrast, the expression levels of the anti-apoptotic proteins Bcl-2 and p-c-Met were decreased in the shRNA-B3GALT5-AS1-1 group compared with those in the shRNA-NC group. However, co-transfection of MKN-45 cells with shRNA-B3GALT5-AS1-1 and ov-CSNK2A1 upregulated

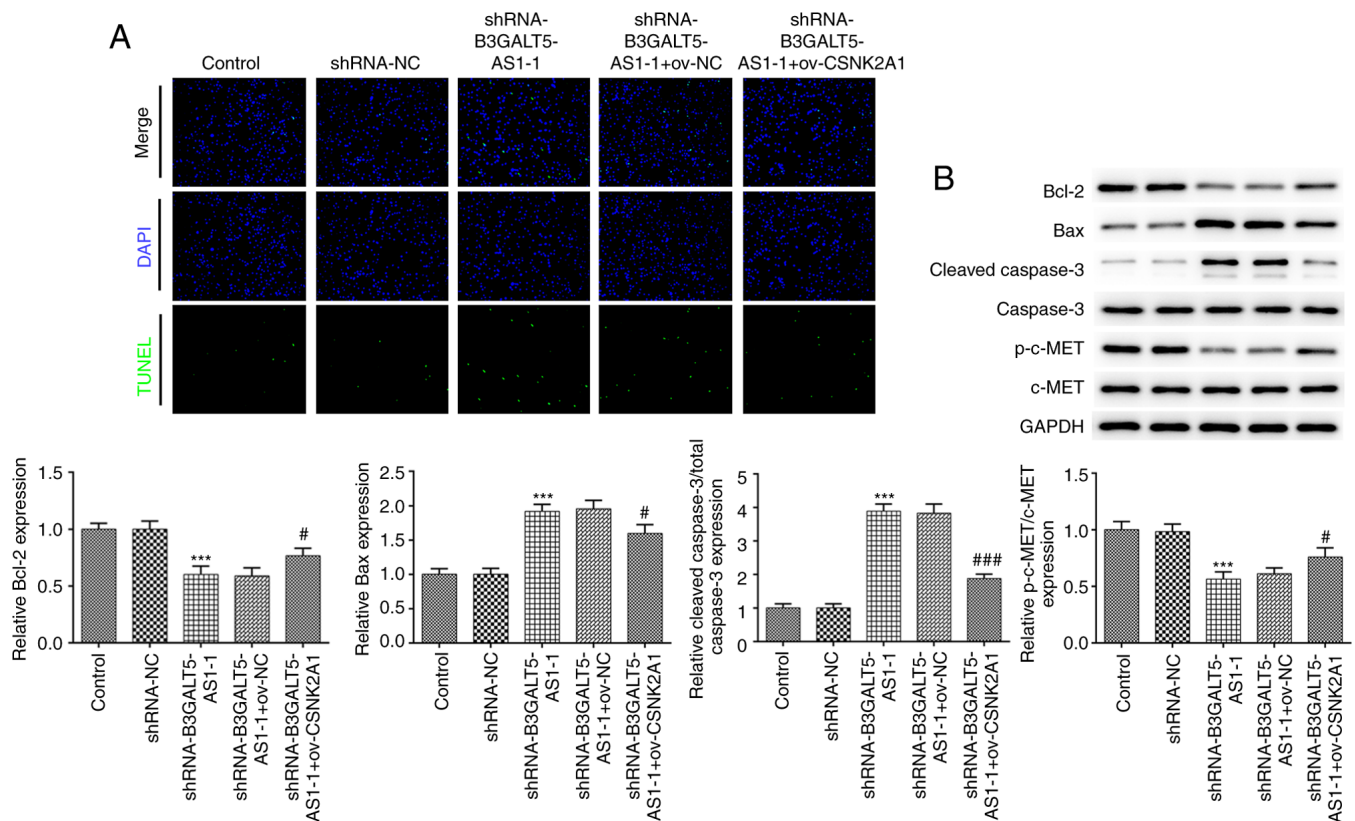


Figure 4. B3GALT5-AS1/CSNK2A1 axis regulates cell apoptosis. (A) Cell apoptosis in different transfection groups was assessed using a TUNEL assay. Magnification, x100. (B) Western blot analysis showing the protein levels of Bcl-2, Bax, cleaved caspase-3, caspase-3, p-c-Met and c-Met in different groups. *** $P < 0.001$ vs. shRNA-NC; # $P < 0.05$ and ### $P < 0.001$ vs. shRNA-B3GALT5-AS1-1 + ov-NC. B3GALT5-AS1, β -1,3-galactosyltransferase 5-AS1; CSNK2A1, casein kinase 2 α 1; p-, phosphorylated; Ov, expression; shRNA, short hairpin RNA; NC, negative control.

Bcl-2 and p-c-Met compared with cells transfected with shRNA-B3GALT5-AS1-1 and ov-NC (Fig. 4B). Dysregulation in c-Met has been implicated in the pathogenesis and development of gastric cancer, where its downregulation can induce apoptosis (23,24). These results aforementioned suggest that B3GALT5-AS1 knockdown promotes gastric cancer cell apoptosis by binding to CSNK2A1.

Discussion

The incidence rate of gastric cancer is particularly high in developing countries, such as China and India (25). Data suggests that gastric cancer ranks second in terms of incidence among all malignancies in China (26,27). Gastric cancer has long been considered to be a difficult challenge to treat clinically (28). Therapeutic interventions in patients with gastric cancer typically involve alterations in diet, psychological support and changes in daily life habits to optimize the therapeutic effects and improve the quality of life (29). The moderate diagnostic value of CEA and CA199 for the early detection of gastric cancer highlights the importance and urgency of developing novel biomarkers with high specificity and sensitivity (30). It has been reported that B3GALT5-AS1 can acts as a diagnostic and prognostic biomarker of gastric cancer (13).

The present study demonstrated that B3GALT5-AS1 was highly expressed in MKN-45 cells compared with that in normal gastric mucosa GES-1 cells. In addition, binding sites

between the sequences of B3GALT5-AS1 and CSNK2A1 were identified. B3GALT5-AS1 silencing repressed the protein expression of CSNK2A1. The catalytic subunit of CK2 α is encoded by CSNK2A1 (31) and is associated with cell proliferation and invasion in colorectal cancer (32). A previous study showed that the knockdown of CSNK2A1 expression attenuated the proliferative and invasive capabilities of breast carcinoma cells (16). In the present study, B3GALT5-AS1 knockdown in MKN-45 cells reduced cell viability, which was reversed by CSNK2A1 overexpression. The reduced MKN-45 cell migration and invasion, mediated by shRNA-B3GALT5-AS1, were also restored following CSNK2A1 overexpression. Furthermore, silencing of B3GALT5-AS1 expression induced cell apoptosis in a manner that was partially counteracted by CSNK2A1 overexpression. However, there are some limitations in the present study. These findings were based on *in vitro* cell model, which require further validation using *in vivo* animal models or human tissues in future studies. In addition, the underlying mechanism of the B3GALT5-AS1/CSNK2A1 axis regulating gastric cancer physiology has not been studied in depth in the present study.

To conclude, the present study demonstrated that knocking down B3GALT5-AS1 expression, which was upregulated in gastric cancer cells, could reduce cell viability and inhibit migration whilst inducing cell apoptosis. These effects could be restored by increasing the expression of CSNK2A1.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

PW and GBS acquired the data. GXD and BQW contributed to the study design and analysis of the data. PW drafted the manuscript and BQW revised it critically for important intellectual content. All authors read and approved the final manuscript. BQW and PW are responsible for confirming the authenticity if the raw data.

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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