Abstract. Wnt proteins have been reported to contribute to the progression of various types of cancer. Wnt6 is a member of the Wnt family and may promote tumorigenesis in gastrointestinal cancer and cervical cancer. In the present study, the expression of Wnt6 in human colon cancer cell lines was evaluated, in order to investigate the role of Wnt6 in the development of colon cancer. Additionally, the effects of Wnt6 upregulation or downregulation on proliferation, apoptosis, cell cycle and cell migration of colon cancer cells have been investigated. Furthermore, western blot analysis was employed to evaluate the expression of Wnt6, B-cell lymphoma 2-associated X protein (Bax), caspase-3 and matrix metalloproteinase (MMP)2. The results of the present study demonstrated that the expression of Wnt6 was increased in HCT116 and SW480 cells compared with the remaining colon cancer cell lines. Furthermore, overexpression Wnt6 resulting from transfection of pGPU6/GFP/Neo-Wnt6-Homo-1 plasmid promoted the proliferation, cell cycle and migration of HCT116 and SW480 cells, but inhibited cell apoptosis in vitro. The expression of caspase-3 and MMP2 was increased, whereas the expression of Bax was decreased in response to upregulation of Wnt6. These results suggested that Wnt6 may serve a vital function in the development of colon cancer.

Introduction

Colon cancer is a common malignant tumor of the gastrointestinal tract and is the fourth leading cause of cancer-associated mortality (1). The 5-year survival rate of patients with colon cancer is 50% (1,2). The onset of colon cancer is associated with genetic and environmental factors (e.g., exposure to carcinogens and smoking) (3). Inactivation of tumor suppressor genes and mutation of oncogenes lead to the development of malignant tumors (3). Previous studies investigated the effect of drugs on the proliferation, adhesion, invasion and migration of colon cancer cells, and the underlying molecular mechanisms of drug resistance in colon cancer (4-9).

The Wnt signaling pathway regulates diverse developmental processes, including cell adhesion, proliferation, differentiation, migration and apoptosis (10). Previous studies have demonstrated that numerous types of cancer, including melanoma, hepatocarcinoma, gastrointestinal, breast and ovarian cancer (11), are associated with abnormal Wnt signaling pathway. Abnormal activation of Wnt signaling pathway has been reported in colorectal cancer (12,13). Wnt family of proteins includes at least 19 secreted-type glycoproteins with conserved 22-24 cysteine residues, and serves vital functions in carcinogenesis and embryogenesis (14). Previous studies have reported that Wnt6 is upregulated in gastrointestinal cancer and cervical cancer, and overexpression of Wnt6 promotes physiological or pathological processes via activation of Wnt/β-catenin signaling pathway in various cancer cell lines (15-18). Wnt6 is highly expressed in colorectal adenoma and may be associated with increased risk of colorectal cancer (19). However, the role of Wnt6 in occurrence, progression and metastasis of colon cancer remains unclear.

In the present study, the expression of Wnt6 was evaluated in colon cancer cell lines (LoVo, SW480, HCT116, SW620 and HT29). The effects of overexpression and knockdown of Wnt6 on proliferation, cell cycle and apoptosis of colon cancer cells were investigated. The aim of the present study was to investigate the function of Wnt6 in tumorigenesis and progression of colon cancer and provide the basis for a novel therapeutic target in the treatment of colon cancer.

Materials and methods

Cell culture. Wnt6 high expression cell lines were selected from five human colon cancer cell lines (LoVo, SW480, HCT116, SW620 and HT29). Cells were maintained in Dulbecco's modified Eagle's medium (DMEM; Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA) supplemented with 10% fetal bovine serum (FBS, Gibco; Thermo Fisher Scientific, Inc.), 100 U/ml of penicillin and 100 µg/ml of streptomycin, and were cultured at 37°C in a humidified atmosphere containing 5% CO₂.
Reverse transcription-quantitative polymerase chain reaction (RT-qPCR). Total RNA was extracted from cells using TRIzol reagent (Ambion; Thermo Fisher Scientific, Inc.). RNA was reverse-transcribed into cDNA using PrimeScript™ 1st Strand cDNA Synthesis kit (Takara Biotechnology Co., Ltd., Dalian, China), and qPCR was performed using a KAPA SYBR Green qPCR kit (Kapa Biosystems, Inc., Wilmington, MA, USA). The primer sequences for Wnt6 were as follows: 5'-CGG AATGGTGTCAGAG-3' (forward) and 5'-CAGGATCGG TCAAAGG-3' (reverse). The primer sequences for β-actin were as follows: 5'-ACACTGTGGCATCTACG-3' (forward) and 5'-TGTCAGCAGATTTCC-3' (reverse). Each reaction (20 µl total volume) contained 10 µl SYBR, 0.40 µM of each primer and 0.2±0.02 µg cDNA template. The thermocycling conditions were as follows: Pre-denaturation at 95°C for 3 min, followed by 40 cycles of denaturation at 95°C for 5 sec, annealing at 60°C for 20 sec and elongation at 72°C for 20 sec. The threshold cycle (Ct) was determined for each reaction by using the 2^ΔΔCq method, which generated Ct values for each gene of interest normalized to the endogenous control gene (β-actin) (20). For each group, three replicates of each measurement were performed.

Cell transfection. Cells (HTC116 or SW480) in 1x10^5 cells/ml were seeded in 6-well plates and cultured until cells reached confluency (70-80%). Cells were then transfected with 800 ng/well of plasmids using Lipofectamine® 2000 (Invitrogen; Thermo Fisher Scientific, Inc.), according to the manufacturer's protocol. The primers for Wnt6 were as follows: 5'-CACCTGCGCCTTACCTCCC-3' (forward) and 5'-GATCCGGTGTCAGACGAGCC-3' (reverse). The corresponding cDNAs were inserted into the pGPU6/green fluorescent protein (GFP)/Neo vector to construct the recombinant pGPU6/GFP/Neo-Wnt6-1 plasmid, to overexpress Wnt6 (GenePharma Shanghai, China). An empty vector (EV) was used as a negative control. Additionally, specific short hairpin (sh)RNA-expressing vectors were employed to knockdown Wnt6. Negative control (NC) pGPU6/GFP/Neo-shNC (target sequence, GTTCTCCGACAGTGTCAGCT) and pGPU6/GFP/Neo-Wnt6-Homo-A (target sequence, AAG TGGTGGCAGAGCTAGCTC) vectors were obtained from Shanghai GenePharma Co., Ltd. (Shanghai, China). Untransfected HCT116 or SW480 cells consider as control group.

MTT assay. Cell proliferation was evaluated using an MTT assay. HCT116 and SW480 cells were seeded in 96-well plates and transfected with expression or empty vector with 1x10^5 cells in each well. At indicated timepoints (24, 48 and 72 h), 20 µl MTT (5 mg/ml; Bioswamp, Wuhan, China) was added to each culture prior to incubation at 37°C for an additional 4 h. Then, DMEM medium was removed and 150 µl dimethylsulfoxide was added to each well and mixed for 10 min. Absorbance was read at a wavelength of 490 nm.

Flow cytometric analysis of apoptosis. Apoptosis was detected by double staining with Annexin V-fluorescein isothiocyanate (FITC)/propidium iodide (PI) (BD Biosciences, Franklin Lakes, NJ, USA), according to the manufacturer's protocol. At 48 h post-transfection, HCT116 and SW480 cells (1x10^4 cells/ml) were washed three times with ice-cold PBS and incubated for 30 min at 4°C in the dark in 200 µl binding buffer containing 10 µl Annexin V-FITC and 10 µl PI. Apoptotic cells were analyzed using a flow cytometer (Beckman Coulter, Inc., Brea, CA, USA) and Cytomics FC 500 MCL with CXP software 5.0 network (Beckman Coulter, Inc.).

Analysis of cell cycle. Cell cycle was analyzed using flow cytometry. At 48 h post-transfection, HCT116 and SW480 cells in 1x10^5 cells/ml were washed with ice-cold PBS and then fixed with 70% ethanol at 4°C for 1 h. Following washing with PBS, cells were stained with PI [10 mM Tris (pH 7.0), 0.1% NP-40, 1 mM NaCl, 0.7 µg/ml ribonuclease A and 5 µg/ml PI] for 30 min in the dark. Cell cycle analysis was performed using a flow cytometer and Cytomics FC 500 MCL with CXP 5.0 software.

Cell migration assay. Cell migration was assessed using Transwell assays. HCT116 and SW480 cells (1x10^5 cells/ml) were starved in serum-free DMEM medium for 24 At 48 h, cells were fixed for 10 min using 4% paraformaldehyde (Bioswamp; Wuhan Beinlay Biological Technology Co., Wuhan, China) and incubated with 0.5% crystal violet (Bioswamp; Wuhan Beinlay Biological Technology Co.) for 30 min. A total of 100 µl cell suspension was seeded into transwell chambers with 8 µm pore polycarbonate membrane insert (Corning Incorporated, Corning, NY, USA). A total of 600 µl RPMI-1640 medium (Beijing Solarbio Science & Technology, Co., Ltd., Beijing, China) containing 20% FBS was added in lower chambers. Following incubation for 24 h in transwell chamber, cells remaining on the upper membrane were removed carefully with a cotton swab. Stained cells were counted in three fields using a light microscope (Nikon Corporation, Tokyo, Japan; magnification, x200).

Western blot analysis. Western blot analysis was performed as previously described (21). Antibodies against Wnt6 (cat. no. ab50030, 1:500 dilution; Abcam, Cambridge, UK), B-cell lymphoma 2 (Bcl-2)-associated X protein (Bax) (cat. no. ab32503, 1:2,000 dilution; Abcam), caspase-3 (cat. no. ab32351, 1:2,000 dilution; Abcam), matrix metalloproteinase (MMP)2 (cat. no. ab37150, 1:1,000 dilution; Cell Signaling Technology, Inc., Danvers, MA, USA) were used. HCT116 and SW480 cells were washed twice with PBS and homogenized in radioimmunoprecipitation assay lysis buffer (Beyotime Institute of Biotechnology, Haimen, China) containing protease inhibitor (cat. no. 58715; Cell Signaling Technology) and centrifuged at 12,000 g for 15 min at 4°C. The concentration of the proteins was measured using a bicinchoninic acid assay kit (cat. no. P0011; Beyotime Institute of Biotechnology). A total of 30 µg proteins were separated by 10% SDS-PAGE and transferred onto a polyvinylidene difluoride membrane (Merck KGaA, Darmstadt, Germany). The membranes were blocked with 5% skim milk for 2 h at room temperature in Tris-buffered saline. Then, the membranes were incubated with primary antibodies overnight at 4°C. Anti-β-actin antibody was selected as internal reference. Then, the membranes were washed with Tris-buffered saline and incubated in biotinylated goat IgG conjugated to horseradish...
peroxidase (HRP) secondary antibody (cat. no. ab7090, Abcam) for 2 h at room temperature. Immunoreactivity was visualized by colorimetric reaction using an enhanced chemiluminescence substrate buffer (Merck KgaA). Membranes were scanned with Gel Doz EZ imager (Bio-Rad Laboratories, Inc., Hercules, CA, USA). Bands were quantified using Quantity One 5.0 software (Bio-Rad Laboratories, Inc.).

Statistical analysis. Data were analyzed using SPSS software (version 18.0; SPSS, Inc., Chicago, IL, USA). The relevant data are expressed as the mean ± standard error of the mean. Statistical analysis was performed using one-way analysis of variance followed by Duncan’s multiple range test. P<0.05 was considered to indicate a statistically significant difference.
Results

Expression of Wnt6 in human colon cancer cell lines. RT-qPCR was employed to evaluate the expression levels of Wnt6 in human colon cancer cells. The results demonstrated that HCT116 and SW480 cells exhibited increased expression levels of Wnt6 (Fig. 1A). Therefore, we selected HCT116 and SW480 cells lines in the further studies, to explore the effect of Wnt6 expression on colon cancer. Additionally, pGPU6/GFP/Neo-Wnt6-Homo-1 plasmid was constructed to assess the effects of overexpression of Wnt6, whereas specific shRNA-expressing vectors were employed to knockdown Wnt6. Fig. 1B and C demonstrate the transfection efficiency of vectors using HCT116 and SW480 cells. As presented in Fig. 1D and E, shWnt6 (a plasmid carrying shRNA targeting Wnt6) significantly reduced the expression of Wnt6, whereas pWnt6 (Wnt6 overexpression plasmid) significantly increased the expression of Wnt6 in HCT116 and SW480 cells.

Overexpression of Wnt6 induced colon cancer cell proliferation. Cell proliferation was evaluated using an MTT assay. Cells were divided into the following groups: Control, shNC (transfection with negative control), shWnt6 (transfection with shWnt6), EV (transfection with empty vector) and shWnt6+pWnt6 (combined transfection with shWnt6 and pWnt6). The results demonstrated that cell proliferation was decreased in the shWnt6 group compared with that of the shNC group at 24, 48 and 72 h (Fig. 2A and B), indicating downregulation of Wnt6 inhibited the proliferation of HTC116 and SW480. Cell viability was significantly increased in the shWnt6+pWnt6 group in a time-dependent manner compared with the shWnt6 group (Fig. 2A and B). Therefore, knockdown of Wnt6 decreased cell proliferation and transfection with pWnt6 reversed this effect in HCT116 and SW480 cells.

Overexpression of Wnt6 inhibits the apoptosis of colon cancer cell. Cell apoptosis was evaluated using Annexin V-FITC/PI staining and flow cytometry. As presented in Fig. 3A and B, the percentage of apoptotic cells was significantly increased in the shWnt6 group compared with that of the shNC group. However, shWnt6+pWnt6 group exhibited decreased apoptosis compared with that of shWnt6 group. Knockdown of Wnt6 increased apoptosis and transfection with pWnt6 reversed this effect in HCT116 and SW480 cells.

Overexpression of Wnt6 promotes colon cancer cell cycle. The effect of Wnt6 on cell cycle was evaluated using flow cytometry. As presented in Fig. 4A and B, cells transfected with shWnt6 exhibited a significant G0-G1 cell cycle arrest accompanied with a reduction of cell numbers in S-phase, which was reversed in response to transfection with pWnt6. These results suggest that knockdown of Wnt6 may induce cell cycle arrest in G0-G1 phase in colon cancer cell lines and transfection with pWnt6 reversed this effect.

Overexpression of Wnt6 induces the migration of colon cancer cell. Cell migration was evaluated using Transwell assays. As presented in Fig. 5A and B, transfection with shWnt6 significantly suppressed the migration of cells compared with that of the shNC group, whereas shWnt6+pWnt6 group exhibited increased numbers of migrated cells. Knockdown of Wnt6 decreased the migratory ability of cells and transfection with pWnt6 reversed this effect in HCT116 and SW480 cells.

Overexpression of Wnt6 effects on the expression of apoptosis-associated proteins. The expression of Bax, caspase-3 and MMP2 was assessed using western blot analysis. The results confirmed that shWnt6-transfected cells exhibited decreased expression levels of Wnt6 compared with that of the shNC group, whereas the shWnt6+pWnt6 group exhibited increased expression levels of Wnt6 compared with that of the shNC group (Fig. 6A and B). Additionally, the expression levels of caspase-3 and MMP2 were decreased, whereas the expression levels of Bax were increased in response to shWnt6 (Fig. 6). The expression of caspase-3 and MMP2 was increased, whereas the expression of Bax was decreased following overexpression of Wnt6 (achieved by pWnt6) (Fig. 6A and B).

Discussion

The Wnt/β-catenin signaling pathway is triggered by a series of signaling cascade reactions, thus leading to transcription of target genes in the nucleus (10). The Wnt signaling pathway is involved in cell proliferation, apoptosis and epithelial-mesenchymal transition (EMT) (22). Additionally, previous studies demonstrated that aberrant Wnt signaling pathway promotes cell proliferation and tumorigenesis (23) in various types of cancer, including gastrointestinal (24), breast (25), kidney (26), pancreatic (27), prostate cancer (28), melanoma (29) and...
osteosarcoma (30). Wnt6 is a member of the Wnt protein family that has been reported to be involved various types of cancer (19,31). However, the function of Wnt6 in colon cancer remains unclear.

In the present study, the expression of Wnt6 was increased in HCT116 and SW480 cells compared with LoVo and HT29 cells. Therefore, HCT116 and SW480 cells were selected for subsequent experiments. The expression pattern of Wnt6 may
differ in colon cell lines due to differences in histological differentiation, tumor staging and biological characteristics. Kirikoshi et al (15) reported that Wnt6 is strongly expressed in SW480 cells. Wnt proteins have been reported to promote various types of cancer. Wnt1 regulates the progression of breast cancer by promoting cell proliferation and migration (32). Overexpression of Wnt2 contributes to tumorigenesis of colorectal and lung cancer (33,34). Wnt10b expression serves an important function in the development of endometrial cancer (35). In the present study, it was demonstrated that inhibition of Wnt6 may inhibit cell proliferation, cell cycle process and migration, and promote cell apoptosis, and overexpression...
of Wnt6 reversed this effect, thus upregulation of Wnt6 may contribute to tumorigenesis and development of malignant colon tumor.

The results demonstrated that the expression of caspase-3 and MMP2 was increased, whereas the expression of Bax was decreased following overexpression of Wnt6. Caspase-3 has been demonstrated to be cleaved in apoptotic cells and the expression of caspase-3 precursor decreased (36). The results of the present study demonstrated that overexpression of Wnt6 increased the expression of caspase-3 precursor, indicating that Wnt6 may inhibit cell apoptosis. MMP2 is involved in the breakdown of extracellular matrix. MMP2 is associated with the development of various malignant tumors and may promote EMT, a key process involved in cancer metastasis (37). Wnt6 increased the expression of MMP2 indicating that Wnt6 may promote cell migration. Bax is an apoptosis-promoting member of the Bcl-2 family (38). The results of the present study demonstrated that overexpression of Wnt6 decreased the expression of Bax, indicating that Wnt6 may inhibit cell apoptosis. Wnt1, Wnt3 and Wnt8 have been reported to activate Wnt/β-catenin signaling pathway (39,40). However, whether Wnt6 may activate the Wnt signaling pathway in colon cancer requires further investigation.

Figure 5. Wnt6 promotes the migration of HCT116 and SW480 cells. The migration of (A) HCT116 and (B) SW480 cells was evaluated using Transwell assays in CON, shNC, shWnt6, EV and shWnt6 groups. **P<0.01 vs. shNC; ***P<0.01 vs. shWnt6. sh, short hairpin; NC, negative control; EV, empty vector; p, plasmid; CON, control.
The present study demonstrated that HCT116 and SW480 cells exhibit increased expression levels of Wnt6. Downregulation of Wnt6 by shWnt6 inhibited cell proliferation and migration and induced cell apoptosis of HCT16 and SW480 cells. Additionally, overexpression of Wnt6 may promote the proliferation, cell cycle and...
migration of HCT116 and SW480 cells, but inhibit cell apoptosis by upregulation of the expression of caspase-3 and MMP2, and downregulation of the expression of Bax. These results indicated that Wnt6 may serve a vital function in the progression of colon cancer and may be utilized as a potential therapeutic target.

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Availability of data and materials

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

Authors' contributions

XZ was responsible for all the experiments and data analyses and editing of the manuscript; HY was responsible for the overall design of the study and responsible for providing the materials. All the authors approved the final submission.

Ethics approval and consent to participate

Not applicable.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

References


