Original Article

Low expression of Dapper1 induces malignancy via the Wnt signalling pathway and is associated with poor prognosis in gastric cancer

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Abstract: Gastric cancer is the second leading cause of cancer death and remains a major clinical challenge due to poor prognosis and limited treatment options. The Wnt signalling pathway is abnormally activated in gastric cancer as well as many other malignancies. The aim of this study was to explore the clinical significance and prognostic value of DAPP1 expression and its regulation of the Wnt signalling pathway in gastric cancer. Real-time PCR and immunohistochemistry were respectively performed to investigate the expression of DAPP1 at mRNA and protein level in both gastric cancer tissues and adjacent normal mucosa tissues. We found that DAPP1 mRNA and protein was commonly downregulated in gastric cancer tissues. The lower expression of DAPP1 was significantly correlated with tumor invasion, lymph-node metastasis, and TNM stage (P < 0.05). Survival analysis revealed that DAPP1 is a prognostic predictor of gastric cancer (P = 0.046). Overexpression of DAPP1 in SGC7901 cells significantly inhibited cell proliferation while increasing apoptosis. Overexpression of DAPP1 led to reduced expression of Dvl-2, β-catenin, survivin, and bcl-xl. Similarly, there was reduced expression of survivin, the canonical Wnt-signalling target gene. Finally, overexpressed DAPP1 inhibited tumour growth of SGC7901 cells transplanted into athymic nude mice. In conclusion, our results suggest that downregulation of Dapper1 gene expression in human gastric carcinoma promotes tumour development through regulating the canonical Wnt-signalling pathway.

Keywords: Gastric neoplasms, Dapper1, Wnt signalling pathway, prognosis

Introduction

Gastric cancer is a leading cause of cancer-related mortality worldwide [1]. Like other malignancies, the development of gastric cancer is a multi-step process involving aberrant signal transduction. In particular, the Wnt signalling pathway is abnormally activated in gastric cancer as well as many other malignancies [2]. Dishevelled (Dvl) is a central mediator of Wnt signalling in both the canonical and noncanonical pathways. It inhibits glycogen synthase kinase 3β-induced degradation of β-catenin. Dapper1 (Dpr1) is a Dvl-interacting protein that negatively regulates canonical Wnt signalling and is required for notochord formation [3, 4].

The human DAPP1 gene is located in the genome at nucleotide position 39,378,960-39,387,891 in the forward orientation (draft sequence NT_025892.9). DAPP1 mRNA is expressed in amnion, foetal brain, eye, heart, adult brain medulla, acute lymphoblastic leukaemia, germ cell tumour, chondrosarcoma, and parathyroid tumour [5]. Endogenous Dpr1 has been shown to enhance Wnt/β-catenin activity in zebrafish embryos that are hypomorphic for Wnt. Dpr1 synergizes with zebrafish Dvl2 and with the Dvl-interacting kinases, CK1e, Par1, and CK2, in activating target genes [6]. The inhibitory activity of Dpr1 on Wnt signalling is conserved from Xenopus to humans; Dpr1 antagonism of Wnt signalling via inducing Dvl degradation is similarly conserved [3]. Dpr1 regulates adipogenesis through its coordinated effects on gene expression that selectively alter intracellular and paracrine/autocrine compo-
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Components of the Wnt/β-catenin signalling pathway [7]. Human Dpr1 can inhibit Wnt signalling by promoting Dvl degradation in colorectal cancer [8, 9]. Dpr1 may play an important role in inhibiting the development of malignant tumours and may become a new target for cancer diagnosis and treatment. However, the expression profile of Dpr1 and its functional mechanism in gastric cancer have never been reported.

Materials and methods

Patients and tissues

Frozen specimens of gastric cancer and adjacent normal mucosa were obtained prospectively from 30 consecutive patients undergoing surgical resection for gastric cancer at Peking University People’s Hospital in 2010. The samples were immediately used for mRNA extraction to measure DAPPER1 expression.

An additional group of 84 gastric-cancer patients were confirmed by pathology and they underwent radical surgery at Peking University People’s Hospital from 2004 to 2006. No preoperative chemotherapy or radiotherapy was administered. From these patients, paraffin-embedded specimens were obtained and were analysed for Dpr1 protein expression by immunohistochemistry. All patients were followed by direct evaluation or phone interview until death or June 2012.

This study was approved by the Human Ethics Committee of Peking University People’s Hospital. All patients provided informed consent, and no children were enrolled.

Reverse transcription polymerase chain reaction (RT-PCR)

Total RNA was extracted from liquid-nitrogen-frozen human tissue using Trizol. After chloroform extraction, RNA was precipitated with isopropanol. DNase I was used to remove genomic DNA contamination. cDNA was synthesized from the extracted RNA using Superscript II reverse transcriptase (GIBCO) at 42°C for 60 min. PCR conditions were as follows: 32 cycles of 94°C for 45 sec, 60°C for 45 sec, and 72°C for 45 sec. After PCR products were electrophoresed at 100 V for 30 min, a gel image was taken and analysed semi-quantitatively to determine the grey value.

Western blot

Whole-cell extracts from gastric cancer cell lines, BGC823, MGC803, and SGC7901, were mixed with 2× sample buffer and heated at 100°C for 5 min. Proteins (50 μg/lane) were separated on a 10% SDS-PAGE separating gel and 4% stacking gel and were transferred onto a nitrocellulose membrane. Membranes were blocked for 2 h in 2.5% Blotto pre-hybridization solution. After blocking, membranes were incubated with the appropriate primary antibody (Dpr1, Dvl-2, β-catenin, Survivin, Bcl-2, or Bcl-xl rabbit anti-human antibody) in 2.5% Blotto pre-hybridization solution at 4°C overnight. After washing the membranes three times with 2.5% Blotto pre-hybridization solution for 15 min each, they were incubated with horseradish peroxidase-conjugated secondary antibody in 2.5% Blotto pre-hybridization solution for 45 min. Subsequently, membranes were washed twice with Tris-buffered saline with Tween (TBS-T) and once with TBS and were developed using the enhanced chemiluminescence (ECL) detection system. Quantitation of the signal for Dpr1, Dvl-2, β-catenin, Survivin, Bcl-2, and Bcl-xl was performed using a Density Scanner (ZEISS automatic image analyser, VIDAS data analysis).

Immunohistochemistry

Human tissues embedded in paraffin wax were sectioned to a thickness of 4 mm. Slides were stained with primary antibody (rabbit anti-human Dpr1) using 3% H2O2 solution as a blocking agent. Slides were incubated at 4°C overnight in a 1:100 dilution of the primary antibody and were washed three times with phosphate-buffered saline (PBS) for 5 min each. Slides were incubated with horseradish peroxidase-conjugated secondary antibody and were developed with 3,3’-diaminobenzidine (DAB) solution.

Flow cytometry

Efficiency of plasmid transfection and apoptosis of gastric cancer cell lines were determined by fluorescence microscopy (Olympus IX70) and FAC Sort flow cytometer (Becton Dickinson). Cultured cells were collected by centrifuging for 5 min, washed twice in PBS, and resuspended in binding buffer. Cells were stained with ANNEXIN V-FITC (1 μg/ml) and PI (2.5 μg/
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Figure 1. Relationship between Dpr1 expression and survival of 84 gastric cancer patients. The overall five-year survival rate was 41.7%, while the survival rate of patients with Dpr1-negative expression was 29.6% and the positive was 47.9%.

Statistical analysis

All data were analysed using SPSS 16.0 statistical software, and differences were considered significant when \( P < 0.05 \). The Chi-square test was used to analyse the relationship between Dpr1 expression variation in tumour tissue and the clinicopathological parameters of 84 gastric-cancer patients. The Kaplan-Meier method was used to draw the survival curve of 84 gastric-cancer patients, and the Log-rank method was used to analyse the relationship between Dpr1 expression variation in tumour tissue and patient outcomes.

Results

DAPP1 expression in gastric cancer tissues

DAPP1 mRNA expression was significantly lower in tumour tissue than in adjacent normal mucosa from 17 of 30 gastric-cancer patients (Supplementary Figure 1). Dpr1 protein expression was lower in tumour tissue than in adjacent normal mucosa from 59 of 84 (70.2%) gastric-cancer patients. Immunohistochemistry results showed that Dpr1 protein was mainly localized in the cytoplasm, while the nucleus was free of expression (Supplementary Figure 2).
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Dpr1 protein expression was absent from gastric tumour tissues of 36 patients (44%) and present in gastric tumour tissues of 48 patients (56%). The level of Dpr1 protein expression correlated with depth of tumour invasion \( (P = 0.046) \), lymph-node metastasis \( (P = 0.016) \), and TNM stage \( (P = 0.048) \) but not with the patient’s gender or age or degree of tumour differentiation (Borrmann type).

Kaplan-Meier survival curves showed that survival of patients whose tumours were positive for Dpr1 protein expression was significantly higher than patients whose tumours were negative for Dpr1 protein expression \( (Figure 1, P = 0.046) \).

Effect of DAPPER1 overexpression on gastric cancer cell lines

Dpr1 protein expression level was lower in SGC7901 cells than in MGC803 cells; β-catenin protein expression level was higher in SGC7901 than in BGC823 cells or MGC803 cells; Dvl-2 protein expression level was similar among the three gastric cancer cell lines \( (Supplementary Figure 3) \). Based on these results, SGC7901 and BGC823 cell lines were selected for further study.

In SGC7901 cells, DAPPER1 mRNA expression was significantly higher in the experimental group than in the control group and empty-vector group \( (Figure 2) \). Proliferation of SGC7901 cells was reduced by 19% in the experimental group as compared to the control group and empty-vector group at both 48 and 72 h after transfection \( (P < 0.05) \) \( (Supplementary Figure 4) \).

In SGC7901 cells, the apoptosis rate was significantly higher in the experimental group than in the control group and empty-vector group \( (Figure 3) \). The apoptosis rates were 13.96%, 2.89%, and 3.67%, respectively. Similarly, in BGC823 cells, the apoptosis rate was significantly higher in the experimental group than in the control group and empty-vector group. The apoptosis rates were 4.89%, 2.01%, and 2.08%, respectively.

Effect of DAPPER1 overexpression on Wnt pathway proteins and their target genes

In SGC7901 cells, levels of Dvl-2 and β-catenin proteins were significantly lower in the experimental group than in the control group and empty-vector group. In SGC7901 cells, Survivin protein levels were significantly lower in the experimental group than in the control group and empty-vector group. Similarly, Survivin mRNA levels in SGC7901 cells were significantly lower in the experimental group than in the control group and empty-vector group \( (Supplementary Figure 4) \). In BGC823 cells, there were no differences in Survivin protein or mRNA levels among the experimental group, control group, and empty-vector group 48 h after transfection (data not shown).

Effect of DAPPER1 gene overexpression on apoptosis-related protein Bcl-2 family members

There were no significant differences in Bcl-2 protein expression among the experimental group, control group, or empty-vector group in either cell line (data not show). Bcl-xl protein levels in both cell lines were significantly lower in the experimental group than in the control group and empty-vector group, while Bcl-xl mRNA levels did not significantly differ among the three groups \( (Supplementary Figure 5) \).
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Figure 3. Apoptosis in gastric cancer cells overexpressing DAPPER1. Upper: The lower right quadrant is early apoptotic cells; upper right quadrant is late apoptotic cells. *apoptotic rate. Lower: Apoptosis of SGC7901 cells assessed by fluorescence microscopy at 100× magnification. Annexin V has green fluorescence and is located in the membrane; PI has red fluorescence and is located in the nucleus.
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In the present study, DAPPER1 mRNA levels were significantly lower in gastric tumour tissue than in the adjacent normal mucosa in 57% of cases (17/30), which is consistent with Yau’s finding from hepatocellular carcinoma [13]. Accordingly, Dpr1 protein levels were significantly lower in gastric tumour tissue than in the adjacent normal mucosa in 70% of cases. There was no significant difference in tumour size between the control group and empty-vector group (Figure 5).

Discussion

In 2002, Cheyette isolated a novel Dvl-binding protein from the Japanese toad body using two-dimensional electrophoresis technology [4]. Unlike previously characterized Dvl-binding proteins, this protein inhibited the function of Dvl when bound to it; therefore, Cheyette named it the Dapper (Dpr). Subsequently, Jushua divided Dpr proteins into Dpr1 and Dpr2 according to their functions [6]. Dpr1’s main function is in regulating the classic Wnt/β-catenin pathway, while Dpr2’s main function is in regulating non-classic Wnt pathways. Thereafter, Jushua, Barbara, Gloy, and Hikasa discovered that Dpr1 regulated cell proliferation and differentiation and played an important role in embryogenesis and organogenesis [10-12]. Therefore, they speculated that Dpr1 might also influence tumour development.

Xenograft growth rate was significantly lower in the experimental group than that in control group and empty-vector group. From the second week, tumour volume in the experimental group was significantly lower than that in the control group and empty-vector group. There was no significant difference in tumour size between the control group and empty-vector group (Figure 5).

Effect of DAPPER1 overexpression on tumorigenicity of SGC7901 cells in nude mice

There was no swelling, ulceration, or other inflammatory response at the site of cell-culture injection. Tumours of approximately 1-mm diameter could be seen at the injection site subcutaneously one week after the inoculation.
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(59/84). Thus, we speculate that reduction of Dpr1 protein expression is associated with the development of gastric cancer. Furthermore, Dpr1 expression level was correlated with depth of tumour invasion, lymph-node metastasis, and TNM stage. Thus, we deduce that Dpr1 expression level is related to the development of gastric cancer.

Our immunohistochemistry staining results suggested that Dpr1 was mainly expressed in the cytoplasm, which is consistent with previous reports [4]. In addition, high expression of Dpr1 was identified in the cytoplasm of normal stromal cells within the tumour tissue, and Dpr1 expression level was similar to that in the adjacent normal mucosa, suggesting that Dpr1 is generally expressed at a high level in normal human gastric tissue. Results from in-vitro experiments confirmed that DAPPER1 overexpression inhibited cell proliferation, increased apoptosis, and reduced tumourigenicity of gastric cancer cells. Based on these results, we concluded that decreased DAPPER1 expression is a feature of malignant tissue and is related to the development of gastric cancer.

Dpr1’s functional mechanism in inducing gastric-cancer development remains unclear. Cheyette and others have suggested that Dpr1 inhibits the classic Wnt pathway [3, 4, 13]. Waxman has suggested that Dpr1 activates the classic Wnt pathway [6]. Cheyette introduced Dpr1 antisense oligonucleotides into HEK293 cells, which resulted in increased β-catenin expression [4]. Our study demonstrated that DAPPER1 overexpression in the gastric-cancer cell line SGC7901 inhibits β-catenin, the core protein of the classic Wnt pathway, as well as mRNA expression of Survivin, the classic Wnt-pathway target gene. These results are consistent with those of Cheyette’s study. In addition, we found relatively low β-catenin expression in the gastric-cancer cell line BGC823. DAPPER1 overexpression in those cells did not increase the expression of Survivin mRNA or protein, suggesting that Dpr1 did not activate the classic Wnt pathway. As such, we conclude that Dpr1 inhibits the classic Wnt pathway and thus promotes the development of gastric cancer.

Interestingly, we obtained a contradictory finding. Overexpression of DAPPER1 significantly reduced Survivin protein expression in the gastric-cancer cell line SGC7901 but not in the gastric-cancer cell line BGC823. Nevertheless, overexpression of DAPPER1 significantly increased the apoptosis rate in both cell lines, indicating that Dpr1 may promote apoptosis through a Wnt-independent pathway. Therefore, we measured the expression of other apoptotic proteins (Bcl-2 and Bcl-xL) after overexpression of Dpr1. Overexpression of DAPPER1 decreased Bcl-xL protein expression in both SGC7901 and BGC823 cells. Catlett-Falcone reported that Stat3 bound to the Bcl-x gene at the APRE sequence of Stat3-binding sites, a fragment of about 600 bp, and initiated its transcription [14]. Jing and others have shown that the Wnt/β-catenin signalling pathway upregulates Stat3 gene expression in mouse embryonic carcinoma cells [15]. Based on these findings, we speculate that Dpr1’s effect on apoptosis is achieved through simultaneous interaction with the Wnt and Stat3 signal transduction pathways.

Most importantly, DAPPER1 expression was closely correlated with prognosis among gastric-cancer patients. Previous studies have indicated that lower levels of N-cadherin expression are associated with reduced patient survival. However, Dpr1 has never been used as a prognostic factor. Our results indicate that patients with Dpr1-positive tumours have better prognoses than those with Dpr1-negative tumours. Therefore, Dpr1 may be a valuable prognostic biomarker for resectable gastric cancer.

Conclusions

In conclusion, downregulation of DAPPER1 gene expression in human gastric carcinoma induced tumour development chiefly through regulating the Wnt signalling pathway. Dpr1 may be a valuable prognostic factor and a novel therapeutic target for human gastric cancer.

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Disclosure of conflict of interest

None.
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References

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Supplementary Figure 1. DAPPER1 mRNA level in gastric carcinoma tissue and adjacent normal mucosa: Tx represents tumor tissue, Nx represents adjacent normal mucosa.

Supplementary Figure 2. Dpr1 protein expression in gastric carcinoma tissue and adjacent normal mucosa: A: Adjacent normal mucosa. B: Tumor tissue (immunohistochemistry staining using the SP method at 100× magnification). A: The cytoplasm of adjacent normal mucosal cells was stained dark brown, while the nucleus was not stained. B: The nuclei and cytoplasm of cancer cells were not stained. The cytoplasm of stromal cells was stained dark brown.

Supplementary Figure 3. Dpr1, β-catenin, and Dvl-2 protein expression in gastric cancer cell lines.

Supplementary Figure 4. Cell proliferation of SGC7901 cell after DAPPER1 overexpression.
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**Supplementary Figure 5.** Bcl-xl protein and mRNA expression in SGC7901 and BGC823 cells with DAPPER1 over-expression.