Association between endogenous gene expression and growth regulation induced by TGF-β1 in human gastric cancer cells

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Abstract
AIM: To investigate the association between endogenous gene expression and growth regulation including proliferation and apoptosis induced by transforming growth factor-β1 (TGF-β1) in human gastric cancer (GC) cells.

METHODS: Reverse transcription polymerase chain reaction (RT-PCR) was performed to detect the main components of the TGF-β1/Smads signal pathway in human poorly differentiated GC cell line BGC-823. Localization of Smad proteins was also determined using immunofluorescence. Then, the BGC-823 cells were cultured in the presence or absence of TGF-β1 (10 ng/mL) by semi-quantitative RT-PCR and Western blot, and the effects of TGF-β1 on proliferation and apoptosis were measured by cell growth curve and flow cytometry (FCM) analysis. The ultrastructural features of BGC-823 cells with or without TGF-β1 treatment were observed under transmission electron microscope. The apoptotic cells were visualized by means of the terminal deoxynucleotidyl transferase (TdT)-mediated dUTP in situ nick end-labeling (TUNEL) method. Meanwhile, the expression levels of endogenous p15, p21 and Smad7 mRNA and the corresponding proteins in the cells were detected at 1, 2 and 3 h after culture in the presence or absence of TGF-β1 (10 ng/mL) by semi-quantitative RT-PCR and Western blot, respectively.

RESULTS: The TGF-β1/Smad signaling was found to be intact and functional in BGC-823 cells. The growth curve revealed the most evident inhibition of cell proliferation by TGF-β1 at 48 h, and FCM assay showed G1 arrest accompanied with apoptosis induced by TGF-β1. The typical morphological changes of apoptosis were observed in cells exposed to TGF-β1. The apoptosis index (AI) in TGF-β1-treated cells was significantly higher than that in the untreated controls (10.7±1.3% vs 0.32±0.06%, P<0.01). The levels of p15, p21 and Smad7 mRNA and corresponding proteins in cells were significantly up-regulated at 1 h, but gradually returned to basal levels at 3 h following TGF-β1 (10 ng/mL) treatment.

CONCLUSION: TGF-β1 affects both proliferation and apoptosis of GC cells through the regulation of p15 and p21, and induces transient expression of Smad 7 as a negative feedback modulation of TGF-β1 signal. Our results suggest a novel functional role of p21 as an accelerant of TGF-β1-mediated apoptosis in GC cells.

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Key words: Gastric cancer; Transforming growth factor-β1; Apoptosis; Gene expression

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INTRODUCTION
Gastric cancer is one of the most common malignant tumors in the world[1]. Gastric carcinogenesis refers to accumulation of genetic alterations of multiple genes such as oncogenes, tumor suppressor genes and mismatch repair genes. However, the exact molecular mechanism remains to be fully elucidated.

Numerous data have shown that TGF-β1, as a prototypic member of TGF-β superfamily of signaling molecules, has a diverse range of functions, including cell growth control (both negative and positive), differentiation, apoptosis (induction or inhibition) and synthesis of extracellular matrix proteins[2]. The TGF-β1 signal is transduced across the membrane by the type I and II receptors (TßRI and II), which contain intrinsic serine/threonine kinase activities. Identification and isolation of the Smads as TGF-ß1’s intracellular signaling mediators have enabled the TGF-ß1 pathway to be traced from the cell surface to the nucleus[3]. Smad proteins can be classified into three categories[4]; receptor-specific Smads (R-Smads: Smad2 and Smad3) that interact with and are directly phosphorylated by a particular TßRI; common partner Smad (Co-Smad: Smad4) that acts as a common mediator of TGF-ß1 signaling by interacting with all R-Smads; inhibitory Smad (I-Smad: Smad7) that competively inhibits R-Smads’ association with activated TßRI and thereby prevents ongoing signaling[5]. TGF-ß1 signal transduction relies upon the rapid TßRI-mediated phosphorylation of R-Smads followed by R-Smad/Co-Smad dimerization and translocation to the nuclei, where the dimmer can directly or indirectly regulate specific gene transcription, through interactions with other transcription factors. I-Smad functions to antagonize this signal by a negative feedback modulation of TGF-ß1[5]. It has been noted that TGF-ß1 shows biphasic effects on contributing to tumorigenesis[6]. In the initial stage of tumorigenesis, TGF-ß1 may play a negative or tumor suppression role in tumor development by inhibiting cell growth. However,
a requirement for TGF-β1 in progression of late stage tumor, which allows the tumor to invade and metastasize through its effects on angiogenesis, immunosuppression and synthesis of extracellular matrix proteins, suggests a second role of TGF-β1 in cancer progression [6-9].

Previous studies have suggested that TGF-β1 not only inhibits cell proliferation [10], but also induces cell apoptosis [11] by regulating the activities of cyclin, cyclin-dependent kinases (CDKs) and CDK inhibitors (CDKIs), which lead to the arrest of the cell cycle at G1 phase. Nevertheless, the possible role of TGF-β1 signal in the simultaneous modulation of GC cells’ proliferation and apoptosis remains unclear. Therefore, elucidation of the mechanism by which TGF-β1 inhibits cell growth in GC cells not only provides insights into the molecular events involved in tumor progression, but also allows us to better understand how cells regulate those events contributing to tumor progression through the cell cycle. In this study, we investigated the effects of TGF-β1 on cell kinetics and the expression of p15, p21 and Smad7 genes and the potential role of these genes in regulation of cell kinetics in human GC cells, in order to reveal the role of this signaling in gastric carcinogenesis.

**MATERIALS AND METHODS**

**Cell culture**

Human poorly-differentiated gastric adenocarcinoma cell line, BGC-823, was obtained from the Shanghai Institute of Cell Biology, Chinese Academy of Sciences. Cells were grown in RPMI-1640 medium supplemented with 10% fetal bovine serum (FBS), L-glutamine and penicillin/streptomycin (100 U/mL), and maintained at 37°C with 95% air.

**Reverse transcription polymerase chain reaction (RT-PCR)**

Total RNA was extracted from BGC-823 cells by TriZol RNA isolation (GIBCO-BRL, USA) and purified with DNase I (Invitrogen, USA). The integrity of RNA was determined by electrophoresis on 1% agarose gel containing ethidium bromide (Invitrogen, USA). RNA (1 μg) was reverse-transcribed to the first strand of cDNA using superscript reverse transcriptase system (Invitrogen, USA) according to the manufacturer’s instructions. cDNA amplification was used to detect expression of the main components of TGF-β1/Smad signaling, including TGF-β1, TβRI, TβRII, Smads 2, 3, 4 and 7.

**Immunofluorescence assays**

BGC-823 cells were seeded onto coverslips, and allowed to grow 20-30% confluency. After washing with cold 1×PBS, cells were fixed in 4% paraformaldehyde for 15 min, and then treated with 0.1% Triton X-100 for 15 min. Subsequently, cells were incubated with Smads 2, 3, 4 or 7 primary antibodies (Santa Cruz, USA) overnight at 4°C. Primary antibodies were diluted at 1:25 with PBS containing 1% BSA. After washing with cold PBS, cells were incubated with Fluorescein (FITC)-conjugated affinity-purified anti-goat IgG secondary antibodies (1:50 with 1% BSA in PBS, Zhongshan, Beijing) for 2 h at room temperature. After washing with cold PBS, slides were immersed for 30 min in 10 ng/mL Hoechst dye 33258 to stain cellular nuclei, and then subjected to an additional wash in cold PBS. All slides were observed under a microscope equipped with epifluorescence optics (Zeiss Axioskop, Germany).

**Cell growth curve**

Cells were seeded in 24-well plates at a concentration of 5×10^4/well, and allowed to attach overnight, followed by a 24-h serum starvation to synchronize and subsequently treated with 1640 culture containing 1% PBS in the presence or absence of TGF-β1 (10 ng/mL, R&D systems, USA). Duplicate wells were trypsinized, stained with trypan blue and counted manually per 12 h for 3 d. The cell growth curve was produced from the mean data of three independent experiments using Microsoft Excel 2000.

**Flow cytometry (FCM) analysis**

Cells were plated in 25 cm² culture flasks at a density of 5×10⁶ cells, followed by synchronization, and then incubated in 1640 medium containing 1% PBS in the presence or absence of TGF-β1 (10 ng/mL) for 24 and 48 h. All floated and adherent cells were harvested, centrifuged at 1 000 r/min for 10 min. Cell pellet was washed with 1×PBS, centrifuged at 1 000 r/min for

| Table 1 Oligonucleotide primers for RT-PCR (Sangon, Shanghai, China)  |
|-------------------------------------------------|
| Gene   | cDNA primer sequence | Annealing temperature (℃) | Product size (bp) |
| TGF-β1 | Forward: 5'CGACTGCGACGATGTTAT-3'  | 61 | 284 |
|       | Reverse: 5'ATAGTGGTGCACGGGCTCG-3'  | 380 |  |
| TβRI   | Forward: 5'AGCCGTTACAGTGTCTCTG-3'  | 358 |  |
|       | Reverse: 5'CCACCGCGTTATACTCGG-3'  | 688 |  |
| TβRII  | Forward: 5'ACGCAACTGACGATCCCTC-3'  | 489 |  |
|       | Reverse: 5'ACTACAGACTTCTACAGG-3'  | 57 |  |
| Smad2  | Forward: 5'ATCCCTACAGAATCTCCCGC-3'  | 57 |  |
|       | Reverse: 5'GTGCTTTTTCAGGCGGTG-3'  | 284 |  |
| Smad3  | Forward: 5'GGAGGGCAAGCCTTGGGAAAAATG-3'  | 57 |  |
|       | Reverse: 5'CCCCTCCCCCCGGCCACCCACATT-3'  | 284 |  |
| Smad4  | Forward: 5'AAAGTGAAGGTATGTTTG-3'  | 56 | 264 |
|       | Reverse: 5'CTGATAAGCTGGAATGACT-3'  | 294 |  |
| Smad7  | Forward: 5'GGGGGCGGTCTACTGTC-3'  | 57 | 436 |
|       | Reverse: 5'ACTTCTGGATGCTCTTCCC-3'  | 58 |  |
| p15    | Forward: 5'GGCGGGAAGCTGCGCTG-3'  | 60 |  |
|       | Reverse: 5'GACCGAGCTAGTTAATCTCA-3'  | 284 |  |
| p21    | Forward: 5'AGTGGACAGCGACCGAGA-3'  | 57 |  |
|       | Reverse: 5'ATCTTTAGACATGACACCAGA-3'  | 284 |  |
10 min, fixed with 70% ethanol, made into a single cell suspension and stored at 4 °C overnight. Then the cells were stained with propidium iodide (PI) solution (with the final concentration of 100 μg/mL) on ice for 30 min and assayed using a FCM (Bio-Rad, USA).

Transmission electron microscopy (TEM)
Cells were plated in 25 cm² culture flasks at a density of 5 × 10⁵ cells, and then incubated in 1600 medium containing 1% FBS in the presence or absence of TGF-β1 (10 ng/mL) for 48 h. All floated and adherent cells were collected, centrifuged at 1 000 r/min for 15 min, fixed in 2.5% cold glutaraldehyde and stored at 4 °C overnight. Then the cells were dehydrated, embedded, cut into ultrathin sections and stained routinely. The ultrastructural features of cells were observed under JEM-1220 transmission electron microscope (JEOL, Japan).

Apoptosis assay
BGC-823 cells were seeded on coverslips, and then incubated in 1600 medium containing 1% FBS in the presence or absence of TGF-β1 (10 ng/mL) for 48 h. Terminal deoxynucleotidyl transferase (TdT)-mediated dUTP in situ nick end-labeling (TUNEL) assay was performed using an in situ Apoptosis Detection kit (Boehringer Mannheim, Germany) according to the manufacturer’s instructions. After washing with cold PBS, the cells were fixed in 4% paraformaldehyde for 30 min, then immersed for 30 min in 0.3% H₂O₂ to block endogenous peroxidase at room temperature. Subsequently, cells were treated with 0.1% TritonX-100 (containing 0.1% sodium citrate) for 15 min, fixed in 4% paraformaldehyde for 30 min at 37 °C. After washing with cold PBS, the cells were fixed in 4% paraformaldehyde for 30 min at 37 °C, then subjected to an additional wash in cold PBS. Cells were colorized with dianinobenzidine tetrahydrochloride (DAB, Zhongshan, Beijing), and the reaction was terminated with a wash in cold PBS after the appearance of positive cells with brown nuclei under a light microscope. Slides were reconstitute with hematoxylin slightly and sealed up with neutral resin. Five high power microscopic views (×200) were selected on each slide, and 1 000 cells were counted. The apoptotic index (AI) was calculated as follows: AI (%) = (number of apoptotic cells/total number counted) × 100%. The cells with 50 μL dUTP solution were used as negative controls, and cells treated with DNase I for 20 min were used as positive controls. The experiments were repeated three times independently. All data were summarized as mean±SD and analyzed using χ²-test. P<0.01 was considered statistically significant.

Semi-quantitative RT-PCR
Cells were plated in 25 cm² culture flasks at a density of 10⁶ cells, followed by synchronization and then incubated in 1600 medium containing 1% FBS in the presence or absence of TGF-β1 (10 ng/mL) for 1, 2 and 3 h. Total RNA was extracted and reverse-transcribed to the first strand of cDNA. p15, p21 and Smad7 cDNA were amplified by PCR under the conditions in Table 1. β-actin cDNA fragment was used as an internal control to eliminate the systematic and quantitative errors. The experiments were repeated three times independently.

Western blot analysis
Cells were plated in 25 cm² culture flasks at a density of 10⁶ cells, followed by synchronization. After incubated in 1600 medium containing 1% FBS in the presence or absence of TGF-β1 (10 ng/mL) for 1, 2 and 3 h. Total proteins were extracted and quantified using DU 640 protein analyzer (BECKMAN, USA). A precasted low-molecular-weight rainbow marker (Invitrogen, USA) and the protein sample (50 μg) were separated on 12% or 15% SDS gels by SDS-PAGE. Separated proteins were transferred onto a 0.45 μm/L polyvinylidene difluoride membrane (Millipore, USA) that was blocked at room temperature for 2 h in Tris-buffered saline with 0.1% Tween-20 (TBS-T) containing 5% skim milk and probed with primary antibodies. The primary antibodies against P15, P21 (Zhongshan, Beijing) and Smad7 (Santa Cruz, USA) were diluted at 1:200 in TBS-T, and the β-actin primary antibody (Santa Cruz, USA) was diluted at 1:500 in TBS-T. Secondary antibodies included horseradish peroxidase (HRP)-labeled anti-rabbit IgG against P15, anti-mouse IgG against P21 and anti-goat IgG against Smad 7 and β-actin. All secondary antibodies were diluted at 1:1 000 with TBS-T. Protein bands were visualized by ECL (Amersham, UK) according to the manufacturer’s instructions and developed on a film (Kodak, USA). Western blot of β-actin was performed for an internal control.

RESULTS

Localization and expression of main TGF-β1/Smad signaling pathway components
The results of RT-PCR demonstrated that the BGC-823 cell line expressed all main TGF-β1/Smad signaling pathway components, including TGF-β1, TβRI, TβRII, Smads 2, -3, -4, and -7 (Figure 1). In addition, all Smad proteins were located in the cytoplasm as demonstrated by immunofluorescence (Figure 2).

Semi-quantitative RT-PCR

Table 2

| Time of TGF-β1 treatment (h) | Cell cycle (%) |
|-----------------------------|---------------|
|                            | G0-G1 | S  | G2-M | Apoptosis |
| 0                          | 52.72 | 32.39 | 14.89 | 0.34 |
| 24                         | 57.07 | 33.82 | 9.11  | 6.61 |
| 48                         | 60.20 | 29.54 | 10.26 | 9.86 |

Effect of TGF-β1 on proliferation and apoptosis of BGC-823 cells

BGC-823 cell growth was inhibited by TGF-β1 treatment, and the phase of significant inhibition was at 48 h. Thenceforth, this inhibitory effect attenuated gradually, and cells resumed normal growth speed (Figure 3). FCM analysis showed that cells at G1 phase increased while cells at S phase decreased, and the hypodiploid peak, called apoptotic peak indicating reduced DNA content in apoptotic cells, appeared before the G1 phase after exposure to TGF-β1 (Figure 4). Table 2 shows that TGF-β1 treatment induced G1 arrest and apoptosis of tumor cells in a time-dependent manner and apoptotic rate reached 6.61% and 9.86%, respectively, at 24 h and 48 h after TGF-β1 stimulation.
Figure 2 Localization and expression of Smads in BGC-823 cells by immunofluorescence.

Figure 3 Effect of TGF-β1 on proliferation of BGC-823 cells.

Figure 4 FCM analysis of BGC-823 cells treated with TGF-β1. A: Controls; B: Treated with TGF-β1 for 24 h; C: Treated with TGF-β1 for 48 h.

Figure 5 TGF-β1-induced ultrastructural changes in BGC-823 cells under TEM. A: Controls, ×5 000; B: Treated with TGF-β1 for 48 h, ×8 000.
Under transmission electron microscope, control cells were big and round, with normal organelles, intact nuclear membrane, obvious nucleoli and low density in nuclear chromatin (Figure 5A). However, the cells treated with TGF-β1 exhibited typical characteristics of apoptosis including cell membrane shrinkage, vacuole formation and mitochondria swelling in cytoplasm, morphologic alteration of nuclei, condensation and fragmentation of nuclear chromatin adjacent to nuclear membrane, although the cell membrane and a few organelles were relatively intact. Cells in late phase of apoptosis with pyknotic nuclei were occasionally observed (Figure 5B). The TUNEL assay revealed that TGF-β1 induced evident apoptosis of BGC-823 cells, and AI (10.7±1.3%) at 48 h after TGF-β1 treatment was significantly higher than that of control group (0.32±0.06%, P<0.01) (Figure 6).

Up-regulation of endogenous genes, p15, p21 and Smad 7 by TGF-β1 treatment
Transcript levels of p15, p21 and Smad 7 in BGC-823 cells were significantly elevated, and gradually recovered after 1 to 3 h following TGF-β1 (10 ng/mL) treatment using semi-quantitative RT-PCR (Figure 7). Subsequently, the accordant changes of p15, p21 and Smad7 proteins were confirmed by Western blot analysis. Although the basal levels of those proteins were relatively low or almost undetectable, they were rapidly increased in response to TGF-β1 treatment (Figure 8). Up-regulation of p15 protein peaked at 1 h after TGF-β1 stimulation, whereas the induction of p21 and Smad7 proteins was most significant at 2 h. The alteration of p15 was ahead of p21 appreciably. Thereafter those protein levels decreased but were still paranormal 3 h after TGF-β1 administration. These observations suggested that p15, p21 and Smad7 gene were rapidly but transiently induced by TGF-β1 treatment.

DISCUSSION
The dynamic balance between cell proliferation and apoptosis is very important to maintain the homeostasis in human body.
and gastric carcinogenesis is related to this imbalance\textsuperscript{[13]}. Many cancers develop resistance to the growth-inhibitory effects of TGF-\(\beta\)-1, and mutations in signaling pathway components may underlie tumor progression. Several researches on human GC cell lines have revealed that transcriptional repression of the \(T\betaRI\) gene by 5'Cpg island hypermethylation and decreased or truncated expression of \(T\betaRI\) gene result in the development of resistance to antiproliferation of TGF-\(\beta\)-1\textsuperscript{[14,15]}. Apoptosis of GC cells induced by TGF-\(\beta\)-1 is also mediated by functional \(T\betaR\)\textsuperscript{[16]}. Both the \(T\betaRI\) expression and AI show negative correlations with the degree of gastric mucosal lesions such as chronic superficial gastritis, intestinal metaphases, atypical hyperplasia to gastric adenocarcinoma. The loss of \(T\betaRI\) expression may play an important role in the inhibition of apoptosis and is involved in the development of GC\textsuperscript{[17]}. No mutation of Smad2 gene is detected in human sporadic gastric carcinomas\textsuperscript{[18]}. In digestive tumors, mutations of Smad4 gene are mainly found in terminal cancer and correlate with tumorigenesis and metastasis\textsuperscript{[19,20]}. In this study, BGC-823 cells expressed all main components of TGF-\(\beta\)-1/Smad signal pathway, including TGF-\(\beta\)-1, \(T\betaRI\), \(T\betaRII\), Smad2, -3, -4 and -7, and Smads were located in the cytoplasm. However, it has been shown that the subcellular localization of Smad7 has specificity in various cell types\textsuperscript{[21]}. TGF-\(\beta\)-1 either simultaneously regulates cell proliferation and apoptosis or just inhibits proliferation without induction of apoptosis\textsuperscript{[22,23]}. In this study, growth of BGC-823 cells was inhibited in response to TGF-\(\beta\)-1 treatment, and the phase with significant inhibition was at 48 h after exposure to TGF-\(\beta\)-1. Thereafter, this effect attenuated gradually, and cells resumed normal growth speed. FCN analysis revealed the existence of G1 arrest and apoptosis in tumor cells treated with TGF-\(\beta\)-1, and the decreased proportion of cells at S phase. TEM and TUNEL also indicated that TGF-\(\beta\)-1 induced apoptosis in BGC-823 cells. Our results suggest an intact and functional TGF-\(\beta\)-1/Smad signal pathway in this cell line, which supports the responsiveness of this cell line to TGF-\(\beta\)-1. The negative growth regulation includes both proliferation inhibition and apoptosis enhancement. In diverse cell types, the mechanism of TGF-\(\beta\)-1 in modulating the activity of CDKIs and consequent antiproliferation involve the direct induction of p15 and p21 gene transcription\textsuperscript{[22,23]} and the regulation of p27 binding shift from CDK4 to cyclin-CDK2\textsuperscript{[26]}. p15 and p21 are known to regulate the activity of cyclin-CDK complexes, which is essential to the cell cycle. As for the key process of cell division, the transcriptional induction of p15 and p21 genes has been postulated at least as partially responsible for the anti-proliferative action of TGF-\(\beta\)-1\textsuperscript{[24]}. TGF-\(\beta\)-1 has been shown to regulate p15 gene expression at two levels, i.e., mRNA accumulation and protein stability\textsuperscript{[26]}. Furthermore, Smads 2, -3 and -4 cooperate with Sp1 to induce p15 transcription in response to TGF-\(\beta\)-1\textsuperscript{[20]}. The increased transcripts of p15 gene result in combination of p15 protein with cyclinD-CDK6/CDK4 in G1 phase and subsequent repression of kinase activity, suggesting that the role of p15 is to prevent the phosphorylation of intracellular targets including retinoblastoma protein (Rb) by cyclin-CDK complexes during TGF-\(\beta\)-1-mediated arrest. On the other hand, accumulation and overexpression of p21 mRNA inhibits the activities of multiple cyclins-CDKs such cyclinE-CDK2 and cyclinD-CDK4, and prevents cells from entering into S phase. The mechanism of cell apoptosis stimulated by TGF-\(\beta\)-1 has not been fully elucidated. TGF-\(\beta\)-1 may induce apoptosis of multiple tumor cells and consequent inhibition of carcinogenesis\textsuperscript{[31,32]} by down-regulating the expression of Bcl-2 and Bcl-xl and promoting the transcription of the death-associated protein kinase and activating caspases\textsuperscript{[13,38]}. It has been found that TGF-\(\beta\)-1 induces apoptosis in some GC cell lines in vitro, which is related to the activation of caspase-3\textsuperscript{[33,34,39]}. Recently, the effect of CDKIs, especially p21, on cell apoptosis has attracted more and more attentions\textsuperscript{[40]}. Patients with multiple myeloma have a delayed cell apoptosis if p15 gene is methylated\textsuperscript{[41]}. A number of studies have shown that endogenous or exogenous p21 may assume both pro- and anti-apoptotic functions in response to different anti-tumor agents depending on cell type and cellular context\textsuperscript{[42]}. The uncertain function of p21 in the apoptosis of GC cells is bewildering\textsuperscript{[47,50]}. The precise mechanism of p21 in regulating apoptosis is still poorly understood, but some researchers believe that p21 activates some pro- or anti-apoptotic signal pathways. There is a possible relation between the enhancement of apoptosis by p21 and activation of caspase-3 and caspase-9\textsuperscript{[49,51]}. p21 promotes apoptosis by increasing the expression of Bax, thus modulating the molecular ratio of Bcl-2: Bax in human hepatocarcinoma cells\textsuperscript{[45]}. On the other hand, p21 inhibits apoptosis by down-regulating the Bax/Bcl-2 ratio in human colon cancer cells\textsuperscript{[52]}, suggesting that the role of p21 in apoptosis remains uncertain\textsuperscript{[45]} and the association between p21 and cell apoptosis induced by TGF-\(\beta\)-1 is still unclear. TGF-\(\beta\)-1 induces apoptosis of cultured bronchiolar epithelial cells and GC cell line SNU-16 via caspase-3 activation and p21 repression\textsuperscript{[56,50]}. However, TGF-\(\beta\)-1-induced apoptosis of head and neck squamous cell carcinoma cells and hepatoma cells is associated with increased expression of p21 and p15 and reduced ratio of phosphorylated Rb (pRb), implicating that TGF-\(\beta\)-1-induced apoptosis occurs downstream of the pRb/E2F pathway\textsuperscript{[54,55]}. In our experiments, the expression levels of p15, p21 and Smad7 mRNA were rapidly and significantly up-regulated after exposure to TGF-\(\beta\)-1, which is in agreement with previous findings\textsuperscript{[56,57]}. The accordant changes in p15, p21 and Smad7 proteins were confirmed by Western blot analysis. Cells showed modest reduction at S phase, and the obvious cell apoptosis may be associated with the up-regulation of p21 by TGF-\(\beta\)-1 and consequent activation of some pro-apoptotic pathways. Based on our data, we believe that p15 and p21 synergistically inhibit the cellular transition from G1 to S phase\textsuperscript{[45]} in response to TGF-\(\beta\)-1 by acting on different cyclin-CDK complexes in BGC-823 cells. The net effects of this responsiveness are to repress the proliferation and enhance the apoptosis, in which p21 may serve as a critical regulator. Therefore, our results suggest a novel functional role of p21 as an accelerant of TGF-\(\beta\)-1-mediated apoptosis in GC cells. The idea that Smad7 antagonizes TGF-\(\beta\)-1-mediated antiproliferation has been widely accepted. The rapid and transient induction of Smad7 gene in this study suggests that Smad7, as an antagonist of TGF-\(\beta\)-1 antiproliferation, is also a TGF-\(\beta\)-1-inducible early target gene, and functions as a negative feedback loop in GC cells\textsuperscript{[5]}. Recent studies have shown that Smad7 may modulate the apoptosis induced by TGF-\(\beta\)-1 as either an accelerant\textsuperscript{[58-60]} or a depressor\textsuperscript{[59,61]} depending on various cell type and cellular context. The molecular mechanism of apoptosis regulated by Smad7 is largely undefined, and may be related to inhibition of the activity of NFXB and activation of p38 and caspases\textsuperscript{[59,60,62,63]}. It is postulated that Smad7 modulates TGF-\(\beta\)-1-mediated apoptosis independent of its role in antagonizing the antiproliferative function of TGF-\(\beta\)-1. Further studies should focus on the exact mechanisms of how endogenous p21 and Smad7 regulate apoptosis induced by TGF-\(\beta\)-1 and their interactions with known apoptosis-associated proteins.
pathological factors such as other members of the TGF-β superfamily, tumor suppressor genes, other growth factors, cytokines, and hormones may enhance or antagonize the function of p15, p21 and Smad7 genes, and regulate their expression by known or unknown mechanisms. The full scope of the interplay of those ligands and TGF-β1 is unclear, and requires further studies.

In conclusion, all observations in this study provide strong evidence that the TGF-B1/Smad signal pathway is intact and capable of regulating expression of endogenous genes, consequently inhibiting cell growth in BGC-823 cells. Thus manipulation of p15, p21 and Smad7 may provide a novel strategy to cancer therapy.

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