C-terminal binding protein-2 mediates cisplatin chemoresistance in esophageal cancer cells via the inhibition of apoptosis

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Abstract. C-terminal binding protein-2 (CtBP2) is a transcriptional co-repressor that is associated with tumorigenesis and tumor progression. It has been reported to predict a poor prognosis in several human cancers, including esophageal squamous cell carcinoma (ESCC). The present study aimed to investigate the involvement of CtBP2 in the cisplatin (DDP) resistance of the ECA109 ESCC cell line and its effect on the expression of apoptosis-associated proteins. Constructed recombinant lentiviruses were used for the knockdown or overexpression of CtBP2 in ECA109 cells, and the expression of CtBP2 was measured using reverse transcription-quantitative polymerase chain reaction and western blotting following transfection. MTT assays, Hoechst 33342 staining and flow cytometry (FCM) were applied to detect the influence of CtBP2 on the DDP-induced viability and apoptosis of the transfected ECA109 cells. In addition, the levels of apoptosis-associated proteins, including p53, B-cell lymphoma 2 (Bcl-2), Bcl-2-associated X protein (Bax) and activated caspase-3 were investigated in the transfected ECA109 cells. Stable ECA109 cells with CtBP2 overexpression or knockdown were successfully established. The results of the MTT, Hoechst 33342 and FCM assays demonstrated that overexpression of CtBP2 attenuated the reduction of cell viability and inhibited the cell apoptosis induced by DDP. Furthermore, the western blotting results indicated that CtBP2 overexpression inhibited the DDP-induced apoptosis of ECA109 cells via the reduction of p53, activated caspase-3 and Bax expression, and promotion of Bcl-2 expression. Therefore, the present study indicated that CtBP2 reduced the susceptibility of ECA109 cells to DDP by regulating the expression of apoptosis-related proteins, suggesting that it may be a promising therapeutic target in ESCC in the future.

Introduction

Esophageal cancer has a high morbidity rate and is the sixth most common cause of cancer-related mortality worldwide (1-3). A survey published in 2013 reported that there were 450,000 new cases of esophageal cancer annually (4), and China accounts for more than half of all cases of this type of cancer (5). Each year, it is estimated that 150,000 individuals succumb to esophageal cancer in China, and the 5-year survival rate of patients is usually <30% (6-8). Esophageal cancer has two main subtypes, namely esophageal squamous cell carcinoma (ESCC) and esophageal adenocarcinoma (9,10). Currently, the most commonly used treatments for ESCC are surgery, chemotherapy, radiation therapy and comprehensive treatment (11,12). The preferred treatment for ESCC is surgical resection, but postoperative recurrence and distant metastasis are clinical problems for which effective treatments remain to be identified (13). Cisplatin (DDP) is a first-line drug in the treatment for ESCC, and the development of DDP resistance in ESCC cells is the main cause of chemotherapy failure (14,15). The effectiveness of chemotherapy depends on the sensitivity of the tumor cells to chemotherapy drugs (16), and ESCC usually exhibits a high resistance to chemotherapy (17,18). Therefore, the identification of oncogenes that may be targeted to combat resistance is likely to be a great benefit to clinical practice.

C-terminal binding protein 2 (CtBP2) acts as a transcriptional co-repressor, and modulates certain essential cellular processes, such as proliferation, migration and (19). It has been reported that CtBP2 has a critical function in tumorigenesis and tumor progression (20,21). CtBP2 is overexpressed in a number of different tumor types, including hepatocellular carcinoma (22), prostate cancer (23), breast cancer (24,25) and ovarian cancer (26). Furthermore, preliminary studies conducted by the present research team revealed that the
expression of CtBP2 was upregulated in ESCC tissues (27,28). In addition, CtBP2 predicts a poor prognosis in human cancers, including ESCC (27). Therefore, it appears that CtBP2 serves an oncogene-like role in tumorigenesis and tumor progression. It has been reported that CtBP2 represses the sensitivity of breast cancer cells by p53-dependent and -independent mechanisms (29). However, the involvement of CtBP2 in the drug resistance of ESCC remains unknown.

In order to further understand the DDP resistance mechanisms of ESCC, the present study investigated the effect of CtBP2 on DDP resistance in ECA109 cells. To the best of our knowledge, the present study is the first to investigate this. The effect of knocking down CtBP2 on the susceptibility of ESCC cells to DDP was evaluated, and the underlying mechanism, such as the regulation of the expression of apoptosis-related proteins, was investigated. The results may indicate the potential of CtBP2 as a therapeutic target for ESCC.

Materials and methods

Cell culture. The human ESCC cell line ECA109 was supplied by the Cell Resource Center of Shanghai Institute for Biological Sciences, Chinese Academy of Sciences (Shanghai, China). Cell culture was performed in Dulbecco's modified Eagle's medium (DMEM; Thermo Fisher Scientific, Inc., Waltham, MA, USA) containing 10% fetal calf serum (FCS; Hyclone; GE Healthcare Life Sciences, Logan, UT, USA), and 10 kU/ml penicillin and 10 mg/ml streptomycin (Beyotime Institute of Biotechnology, Haimen, China).

Knockdown or overexpression of CtBP2. Firstly, recombinant lentiviral vectors were constructed to knockdown CtBP2 (LV-CtBP2-RNAi) or overexpress CtBP2 (LV-CtBP2). The sequences were designed using the software EPTiRNA (http://optirna.unl.edu/) and synthesized by GeneChem Co., Ltd. (Shanghai, China). The vector construction and virus packaging were conducted by GeneChem Co., Ltd. For the former, small hairpin RNA (shRNA) of CtBP2, whose target sequence was 5'-GCGCCTTGGTCAGTAATAG-3', was cloned into a GV248 vector (GeneChem Co., Ltd.) via EcoRI and AgeI restriction endonuclease sites. For the latter, the coding sequence of CtBP2 was cloned into a GV492 vector (GeneChem Co., Ltd.) via BamHI and AgeI restriction endonuclease sites. The primers used for overexpression were as follows: 5'-AGG TCG ACT CTA GAG GAT CCC GCC ACC ATG GCC CTT GTG TCA ACC ATG GCC TTC TTA GAT AAG CAC-3' (forward) and 5'-TCC TTG TAG TCC ATA CTA TGC TCG TTA GGG TGC TCT CGA TTG-3' (reverse). Schematic diagrams of the recombinant lentiviral vector constructs are presented in Fig. 1A.

ECA109 cells were transfected with LV-CtBP2-RNAi or LV-CtBP2 to knockdown or overexpress CtBP2, respectively. ECA109 cells transfected with empty vector GV248 and empty vector GV492 served as the negative controls, LV-CtBP2-RNAi and LV-CtBP2, respectively. The following formula was used to calculate to volume of virus to be added: Virus volume = multiplicity of infection x cell number/virus titer. The cells (1x10^6 cells/well) were seeded into 6-well plates prior to transfection. In order to improve the transfection rate, 5 µg/ml polybrene and enhanced infection solution (without FCS) were incubated overnight with the cells at 37°C with 5% CO₂. Puromycin (2 µg/ml; Gibco; Thermo Fisher Scientific, Inc.) was added to the culture medium (DMEM containing 10% FCS) on the third day and refreshed every 2 days for 1 week to select the transfected cells. The knockdown or overexpression of CtBP2 in the ECA109 cells was confirmed using the observation of green fluorescence, reverse transcription-quantitative polymerase chain reaction (RT-qPCR) and western blotting.

RNA extraction and RT-qPCR. Total RNA was extracted from the CtBP2 knockdown or overexpressing cells using TRIzol reagent (Thermo Fisher Scientific, Inc.) in accordance with the manufacturer's protocol. First-strand cDNA synthesis was conducted using a RevertAid First Strand cDNA Synthesis kit (Thermo Fisher Scientific, Inc.). The temperature protocol was 60 min at 42°C, with termination of the reaction by heating at 70°C for 5 min. qPCR was performed in triplicate using SYBR Green Master mix (Roche Diagnostics, Basel, Switzerland) in a 7500 Real-Time PCR instrument (Applied Biosystems; Thermo Fisher Scientific, Inc.) to test the mRNA expression level of CtBP2. The thermocycling conditions were as follows: Firstly 95°C for 10 min, followed by 95°C for 10 sec, 60°C for 15 sec and 72°C for 20 sec, for 40 cycles. The relative expression of mRNA was calculated using the 2^{-ΔΔCq} method with GAPDH as an internal reference (30). The following primer sequences were used: GAPDH, 5'-GAC CTG ACC TGC CGT CTA-3' (sense) and 5'-AGG AGT GGG TGT CGC TGT-3' (antisense); CtBP2, 5'-CTG AGT TCC GGG TCC TTC TG-3' (sense) and 5'-GAC TTG ATA TCC GCG TCC TC-3' (anti-sense).

Western blot analysis. Briefly, cells were homogenized in lysis buffer containing 1 mM phenylmethylsulfonyl fluoride and complete protein inhibitor mixture (Beyotime Institute of Biotechnology) for 15 min on ice, and then centrifuged at 13,400 x g for 10 min to collect the supernatant. The supernatant was diluted in 5X sodium dodecyl sulfate (SDS) loading buffer (Beyotime Institute of Biotechnology), boiled for 5 min and then cooled on ice. The protein concentration was measured at 280 nm using a One Drop Spectrophotometer (Wuyi Technology Co., Ltd., Nanjing, China) prior to loading the protein onto a gel (50 µg protein/lane). The proteins were separated by 10% SDS-PAGE and then blotted to polyvinylidene difluoride membranes (EMD Millipore, Billerica, MA, USA). The membrane was blocked with Tris-buffered saline and 0.1% Tween-20 (TBST) supplemented with 5% non-fat milk for 2 h at room temperature, and then reacted with the following primary antibodies overnight at 4°C: Anti-CtBP2 (sc-17759; 1:200; Santa Cruz Biotechnology, Inc., Dallas, TX, USA), anti-activated caspase-3 (cleaved) (AB3623; 1:200; EMD Millipore), anti-B-cell lymphoma 2 (anti-Bcl-2; Ab-1; 1:1,000; EMD Millipore), anti-Bcl-2-associated X protein (anti-Bax; ab53154; 1:500; Abcam, Cambridge, MA, USA) and anti-β-actin (ab8227; 1:1,000; Abcam). After washing with TBST three times, the membrane was then reacted with horseradish peroxidase (HRP)-conjugated goat anti-rabbit IgG (A8919; 1:1,000) or HRP-conjugated rabbit anti-mouse IgG (A9044; 1,100; both from Sigma-Aldrich; Merck KGaA, Darmstadt, Germany) for 2 h at 37°C. The protein bands were
detected using ECL chemiluminescence reagent (Thermo Fisher Scientific, Inc.) and imaged using a chemiluminescence detection system (Tanon Science and Technology Co., Ltd., Shanghai, China). ImageJ software (National Institutes of Health, Bethesda, MD, USA) was used to analyze the density of the bands and β-actin was used as a reference for normalization. All experiments were repeated three times.

**DDP treatment and cell viability assay.** ECA109 cells were seeded in triplicate in a 96-well plate and cultured in 100 µl DMEM medium containing 10% FCS. When the cells had become attached to the bottom of well, the cells were treated with DDP (Sigma-Aldrich; Merck KGaA) in serial dilutions (final concentration 1.5x10⁻³, 1.5x10⁻⁴, 1.5x10⁻⁵, 1.5x10⁻⁶, 1.5x10⁻⁷ or 1.5x10⁻⁸ M) in DMEM. Following incubation for 48 h, methythiazolyl tetrazolium solution (MTT, Beyotime Institute of Biotechnology; 10 µl; 5 mg/ml) was added to each well and the plate was incubated for a further 4 h. The medium was eliminated and 100 µl dimethylsulfoxide (Merck KGaA) was added to each well. The plate was shaken to dissolve the MTT-formazan crystals and the absorbance at a wavelength of 570 nm was read using an ELX800 microplate reader (BioTek Instruments, Inc., Winooski, VT, USA).

On the basis of the MTT results for the various concentrations of DDP, the half maximal inhibitory concentration (IC₅₀) of DPP was determined. This concentration of DDP was subjected to further testing of cell viability with different treatment times (0, 12, 24 and 48 h) in order to determine the appropriate treatment duration. For subsequent analysis of the role of CtBP2 in the cytotoxicity of DDP, transfected and untransfected ECA109 cells were exposed to the concentration of DPP closest to the IC₅₀ using the treatment duration identified to be appropriate in this assay.

**Hoechst 33342 staining.** ECA109 cells on glass coverslips in 24-well plates (5x10⁴ cells/well) were fixed with 4% paraformaldehyde for 30 min at room temperature and then washed with 0.01% PBS three times, for 10 min each time at room temperature. The cells were then stained with Hoechst 33342 (Shanghai Yeasen Biotechnology Co., Ltd., Shanghai, China) for 10 min at room temperature. The slides were mounted with anti-fade solution (Beyotime Institute of Biotechnology). The apoptotic cells were identified by detecting the condensation and fragmentation of the cell nuclei under a fluorescence microscope (Zeiss AG, Oberkochen, Germany). Apoptotic cell numbers were counted in three randomly selected fields to calculate the apoptosis rate in triplicate.

**Flow cytometry (FCM) assay.** Following the aforementioned treatments, the cells were digested with trypsin and washed with 0.01% PBS twice. The cells were then stained for 10-20 min with Annexin V-fluorescein isothiocyanate and propidium iodide (Abcam) solution at room temperature for 1 h in the dark, and then subjected to FCM (BD Biosciences, San Jose, CA, USA). A total of 10,000 fluorescence signals of each group were collected. The data were analyzed using FACSuite software (BD Biosciences).

**Statistical analysis.** All data were analyzed using SPSS 17.0 software (SPSS, Inc., Chicago, IL, USA). One-way analysis of variance was used to analyze the data and Tukey’s post hoc test was used to analyze the differences between specific groups. P<0.05 was considered to indicate a statistically significant difference.

**Results**

**Successful establishment of stable cell lines.** In order to detect the impact of CtBP2 on DDP resistance in ESCC, the ECA109 cells were transfected with lentivirus to knockdown or overexpress CtBP2. As the recombinant lentiviral vector contained the enhanced green fluorescent protein (EGFP) gene, the rate of transfection could be determined by direct observation under a fluorescence microscope. The percentage...
of cells positive for EGFP was >90% in the ECA109 cells transfected with recombinant LV-CtBP2-RNAi or LV-CtBP2. Fluorescence images of the transfected cells are presented in Fig. 1B.

The expression levels of CtBP2 in the ECA109 cells transfected with recombinant LV-CtBP2-RNAi or LV-CtBP2 (the LV-CtBP2-RNAi$^+$ and LV-CtBP2$^+$ groups, respectively) were further confirmed by RT-qPCR and western blot analysis. The expression of CtBP2 mRNA was revealed to be significantly changed by RT-qPCR analysis. Compared with the blank and negative control (LV-CtBP2-RNAi$^-$) groups ($P<0.01$), the relative expression of CtBP2 mRNA in the LV-CtBP2$^+$ transfection group was increased ~6-fold ($P<0.01$). By contrast, the relative expression of CtBP2 mRNA in the LV-CtBP2-RNAi$^+$ transfection group was decreased by more than half compared with that in the blank and negative control (LV-CtBP2-RNAi$^-$) groups ($P<0.01$). No significant difference in CtBP2 mRNA levels was detected between the blank group and the LV-CtBP2, or LV-CtBP2-RNAi$^-$ group ($P>0.05$; Fig. 2A and B).

The protein levels of CtBP2 detected by western blotting exhibited similar trends to the expression levels of CtBP2 mRNA in the ECA109 cells subjected to CtBP2 knockdown or overexpression. Compared with the blank and respective negative control groups, the expression of CtBP2 was increased ~2-fold in the LV-CtBP2$^+$ transfection group ($P<0.05$) and decreased by two-thirds in the LV-CtBP2-RNAi$^+$ transfection...
group (P<0.01; Fig. 2C and D). Furthermore, no significant difference was detected between the blank group and the LV-CtBP2- or LV-CtBP2-RNAi- group (P>0.05).

These results suggest that stable cell lines with the knockdown or overexpression of CtBP2 were successfully obtained following recombinant lentiviral transfection for use in the following experiments.

**CtBP2 overexpression promotes ECA109 cell viability following DDP treatment.** In order to determine the optimized concentration and treatment time for DDP, ECA109 cells were treated with DDP solution in serial dilutions for different time periods. Firstly, an MTT assay was conducted to test the changes in cell viability following treatment with serial dilutions of DDP. The cell viability was markedly decreased following treatment with increasing concentrations of DDP for 24 h. Compared with the blank group, the cell viability in the 1.5x10^{-3}, 1.5x10^{-4} and 1.5x10^{-5} M DDP treated groups was significantly reduced (P<0.01). Notably, the cell viability of the 1.5x10^{-5} M DDP treatment group was reduced by 52.15% (Fig. 3A).

Secondly, the effects on cell viability of incubation with 1.5x10^{-5} M DDP for different treatment times were examined via MTT assay (Fig. 3B) and by observation under an inverted phase contrast microscope (Fig. 3C). When observed under the microscope, the number of dead cells appeared to increase gradually as the treatment time with DDP was prolonged. The MTT assay demonstrated that compared with the blank group, the viability of the cells treated with DDP for 24 and 48 h was significantly decreased (P<0.01), but the reduction in viability of the cells treated with DDP for 12 h was not significant (P>0.05; Fig. 3B).

These results indicate that the effect of DDP on cell viability was dependent on concentration and reaction time. The optimized concentration and treatment time for DDP in ECA109 cells were 1.5x10^{-5} M and 24 h, respectively, and were used in the following experiments.

The impact of CtBP2 on the viability of the DDP-treated ECA109 cells was examined by microscopy and MTT assay (Fig. 4). The microscopy images revealed that cell shrinkage occurred following treatment with DDP, and the number of dead cells was increased. The viability of the ECA109 cells treated with DDP for 24 h was significantly reduced compared with that of the control cells (P<0.01). The cell viability of the LV-CtBP2-RNAi+ + DDP group was significantly reduced compared with that of the LV-CtBP2-RNAi+ + DDP group (P<0.05). Furthermore, the cell viability of the LV-CtBP2* + DDP group was significantly increased compared with that of the LV-CtBP2* + DDP group (P<0.01). No statistically significant difference was detected between the LV-CtBP2* + DDP and LV-CtBP2-RNAi+ + DDP groups and the blank + DDP group (P>0.05). These results indicate that the overexpression of CtBP2 attenuated the reduction of cell viability induced by DDP, and the knockdown of CtBP2 augmented the DDP-induced reduction of cell viability.
CtBP2 overexpression reduces the DDP-induced apoptosis of ECA109 cells. Hoechst 33342 staining and FCM were used to investigate the effect of CtBP2 on the cell apoptosis induced by DDP. The results of Hoechst 33342 staining demonstrated that the numbers of apoptotic bodies were significantly increased (P<0.01) in the DPP-treated cells compared with the untreated control group. The number of apoptotic bodies was increased significantly in the LV-CtBP2-RNAi + DDP group compared with the LV-CtBP2-RNAi + DDP group, and decreased significantly in the LV-CtBP2 + DDP group compared with the LV-CtBP2 + DDP group (P<0.05; Fig. 5A). On the basis of these results, it appears that CtBP2 overexpression attenuates the increase of apoptotic bodies induced by DDP.

FCM was used to further verify the impact of CtBP2 on the DDP-induced apoptosis of ECA109 cells. The results are consistent with those of Hoechst 33342 staining. The percentages of apoptotic cells in the untreated control, blank + DDP, LV-CtBP2-RNAi + DDP and LV-CtBP2-RNAi + DDP groups were 5.34, 13.7, 13.98 and 21.59% respectively. Compared with the blank + DDP group and the LV-CtBP2-RNAi + DDP group, the percentage of apoptotic cells in the CtBP2 knockdown (LV-CtBP2-RNAi + DDP) group was significantly increased (P<0.05). The percentages of apoptotic cells in the blank + DDP, LV-CtBP2 + DDP and LV-CtBP2 + DDP groups were 13.7, 14.75 and 5.86% respectively. The percentage of apoptotic cells in the CtBP2 overexpression (LV-CtBP2+ DDP) group was significantly lower than those of the LV-CtBP2 + DDP and blank + DDP groups (P<0.05; Fig. 5B). The FCM results further indicate that CtBP2 overexpression inhibited the DDP-induced apoptosis of ECA109 cells.

Mechanisms underlying the effect of CtBP2 on DDP chemoresistance. Caspase serves essential roles in cell apoptosis, which is a cellular event considered as programmed cell death (31,32). Caspase-3 is one of the crucial downstream effectors of apoptosis (33). The effects of CtBP2 on the protein levels of p53, Bcl-2, Bax and activated caspase-3 were analyzed by western blotting (Fig. 6). The results shown in Fig. 6A and B demonstrate that the expression of p53 was significantly increased (P<0.01) in the ECA109 cells treated with 1.5x10^{-5} M DDP for 24 h compared with the untreated control cells. Notably, the expression of p53 in the CtBP2 overexpression (LV-CtBP2+ DDP) group was significantly decreased compared with that of the LV-CtBP2 + DDP group (P<0.05). The changes in cleaved caspase-3 levels (Fig. 6D) exhibited a similar pattern to those of p53. These results demonstrate...
that the overexpression of CtBP2 inhibited the DDP-induced increase in the protein levels of p53 and activated caspase-3 in ECA109 cells.

The protein levels of Bax and Bcl-2 were also detected using western blotting. The results clearly demonstrate that Bcl-2 expression was downregulated while Bax expression was upregulated in the CtBP2 knockdown (LV-CtBP2-RNAi+ + DDP) group. Therefore, the Bcl-2/Bax ratio in the LV-CtBP2-RNAi+ + DDP group was significantly decreased compared with those in the untreated control, blank + DDP and LV-CtBP2-RNAi + DDP groups (P<0.05). By contrast, the ratio of Bcl-2/Bax in the CtBP2 overexpression (LV-CtBP2+ + DDP) group was significantly increased compared with those in the blank + DDP and LV-CtBP2+ + DDP groups (P<0.05; Fig. 6C). These results indicate that CtBP2 overexpression alleviated the DDP-induced apoptosis of ECA109 cells by reducing the protein levels of p53, cleaved caspase-3 and Bax, and increasing Bcl-2 expression.

Discussion

Understanding of the biology and molecular mechanisms underlying ESCC development and progression is required for advances in the treatment of ESCC (34). Chemotherapy is one of the main therapeutic strategies for the treatment of ESCC (35,36). However, chemoresistance to anticancer drugs greatly reduces the effectiveness of these drugs, and is a huge obstacle to the discovery of a successful therapy for ESCC (37). Therefore, it is urgently necessary to identify an effective potential therapeutic target for combating drug resistance in ESCC. DDP is a first-line drug in the treatment for ESCC, and DDP resistance remains a serious challenge.
A previous study conducted by the present research team indicated that the expression of CtBP2 was upregulated in ESCC tissues compared with adjacent non-tumorous tissues (17). However, the effect of CtBP2 on the susceptibility of ESCC cells to DDP was unclear. Therefore, the present study established stable ECA109 cells with the overexpression or knockdown of CtBP2 via recombinant lentiviral transfection in order to investigate the effect of CtBP2 on these cells when treated with DDP.

In the present study, the optimized concentration and treatment time of DDP for use in vitro were determined by MTT assay; 1.5x10^{-5} M DDP treatment for 24 h was selected for further investigation. The optimized concentration (1.5x10^{-5} M) in the study is similar to the clinically used dose of DDP (20 mg/m^2/day) (38,39). The effect of CtBP2 on the cell apoptosis induced by DDP was investigated by Hoechst 33342 staining and FCM. The overexpression of CtBP2 attenuated the reduction of cell viability and inhibited the cell apoptosis induced by DDP. By comparison, the knockdown of CtBP2 increased the susceptibility of ECA109 cells to DDP, as it increased the number of apoptotic cells following treatment with DDP. These findings indicate that CtBP2 is a potential target for the chemotherapy of ESCC, via which drug resistance may be reversed in patients with ESCC.

There are various mechanisms by which tumor cells develop resistance to chemotherapy. CtBP2 has been demonstrated to reduce the chemosensitivity of breast cancer cells to various chemotherapeutic drugs via p53-dependent and p53-independent effects (29). The present study attempted to further investigate the effect of CtBP2 on the chemoresistance of ECA109 cells. The protein levels of the apoptosis-related proteins p53, Bcl-2, Bax and cleaved caspase-3 in the DDP-treated ECA109 cells with CtBP2 overexpression or knockdown were determined using western blotting. The overexpression of CtBP2 reduced the protein levels of p53, cleaved caspase-3 and Bax, and increased the protein levels of Bcl-2 in DDP-treated ECA109 cells, and the knockdown of CtBP2 exhibited opposing effects. These results indicated that CtBP2 reduced the chemosensitivity of ECA109 cells to DPP via the inhibition of p53, caspase-3 and Bax. This information supplements the findings of a previous study by the present research team, which demonstrated that CtBP2 promotes the progression of ESCC via the negative transcriptional regulation of p16^{INK4A} (27). Nuclear p16 has been reported to be important to the chemosensitivity of multiple cancers (40,41), and in ESCC, CtBP2 promotes chemoresistance through the negative transcriptional regulation of p16^{INK4A}. However, a limitation of the present study is that only a single cell line was investigated, and the further investigation of other cell types of esophageal cancer is required to confirm the findings of the study.

In conclusion, the results of the present study indicate that CtBP2 attenuated the susceptibility of ESCC cells to DDP by...
regulating the expression of apoptosis-related proteins and thereby inhibiting cell apoptosis. This knowledge may provide a new strategy for decreasing the chemoresistance to DDP in the treatment of ESCC.

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Availability of data and materials
The analyzed data sets generated during the study are available from the corresponding author on reasonable request.

Authors' contributions
MJ, YZ, HS, YM and QJ designed and did the research. QJ, MY, HS, YM and QJ designed and did the research. QJ, WY, WB and PW analyzed the data. All authors have read and approved the final manuscript.

Ethics approval and consent to participate
Not applicable.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests with respect to the data appearing in the manuscript.

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