Effects of poly (ADP-ribosyl) polymerase (PARP) inhibitor on cisplatin resistance & proliferation of the ovarian cancer C13* cells

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Background & objectives: Drug resistance is the primary cause of failure in the treatment of cancers. It has been suggested that the enhancement of DNA repair capability may be responsible for the drug resistance of the tumour cells, and poly(ADP-ribosyl)ation plays an important role in DNA repair. This study investigated the effect of PARP inhibitor 3-aminobenzamide (3-AB) on the cisplatin resistance and proliferation of the cisplatin-resistant ovarian cancer C13* cells in vitro.

Methods: C13* cells were treated with various concentrations of 3-AB in vitro. MTT assay was used to determine the effect of 3-AB on the cisplatin sensitivity and proliferation of cells. The expression levels of PARP-1 mRNA and protein in the C13* cells were examined using reverse transcription-polymerase chain reaction (RT-PCR) and Western blot, and changes caused by 3-AB treatment were investigated. Immunofluorescence microscopy was used to detect the localization and expression of the PARP-1 proteins before and after treatment with 5 mmol/l 3-AB.

Results: The inhibitory ratio and the cisplatin sensitivity of C13* cells significantly increased with the increase of the concentration of 3-AB (P<0.05). The RT-PCR analysis revealed that the expression of PARP-1 mRNA was decreased when platinum (Pt) and 3-AB were combined. The expression levels of PARP-1 protein were decreased by 23.15 ± 2.53, 59.11 ± 2.23 and 73.24 ± 3.88 per cent, respectively, in C13* cells with the increase of the concentration of 3-AB (P<0.05). The immunofluorescence microscopy results indicated that the expression level of PARP-1 protein was significantly decreased after treatment with 3-AB (P<0.05).

Interpretation & conclusions: 3-AB inhibited the proliferation activity of C13* cells, and increased the cellular sensitivity to cisplatin. Our findings show that the PARP inhibitor 3-AB can downregulate the expression of PARP-1 at transcriptional and translational levels in C13* cells.

Key words 3-aminobenzamide - cisplatin - drug resistance - ovarian cancer - poly(ADP-ribosyl)ation
Ovarian cancer is the second most common gynaecological malignancy and the leading cause of death of patients with gynaecological cancers. Most of these cases lack early symptoms and are diagnosed at an advanced stage (approximately 70%), when the tumour has metastasized. The traditional treatment of advanced ovarian cancer is based on cytoreductive surgery, followed by platinum-based chemotherapy. Unfortunately, despite the high response rate to the initial chemotherapy, there remains a significant risk for recurrence and resistance to the chemotherapy, and the long-term survival is relatively poor with only 10-15 per cent of advanced-disease patients surviving at 10 years. Consequently, seeking promising novel agents for chemotherapy and elucidating drug resistance mechanisms are crucial for individualizing the treatment and prognosis of ovarian cancer patients.

Chemotherapy resistance is a significant problem and is the major obstacle in ovarian cancer treatment. The enhancement of DNA damage repair capability could be a critical mechanism for chemotherapy. Poly(ADP-ribose)lation is a post-translational modification found in mammalian cells. In humans, the conserved PARP [poly(ADP-ribose) polymerase] signature motif has been identified in 17 homologous genes, including parp-1, parp-2, parp-3, vault-parp, and tankyrases. PARP-1 is an essential member of the family of PARPs, which plays an important role in DNA repair, genomic stability, energy metabolism, transcriptional regulation, inflammation and cell death. DNA repair is a complex and multifaceted process, which is critical to cell survival.

In our earlier study, the expression levels of PARP-1 protein in cisplatin-resistant ovarian cancer C13* cells were found to be higher than cisplatin-sensitive ovarian cancer OV2008 cells and PARP-1 RNA interference can significantly downregulated PARP-1 expression and effectively reversed resistance of C13* cells to cisplatin. However, the RNA interference still has some limitations for clinical application. Therefore, in the present study PARP inhibitor 3-aminobenzamide (3-AB) was used to reduce the expression of PARP. The cisplatin-resistant ovarian cancer C13* cells were treated with various concentrations of 3-AB, and the effect of decreased expression of the PARP-1 protein in C13* cells was studied on proliferation activity and cisplatin sensitivity of cells.

**Material & Methods**

**Cell culture:** Cisplatin-resistant ovarian cancer C13* cells, (kindly provided by Dr Du Meirong, Department of Gynecology, Fudan University), were cultured in RPMI1640 medium (Hyclone, Logan, UT, USA) supplemented with 10 per cent foetal bovine serum (FBS; Hyclone, Logan, UT, USA) and 1 per cent penicillin/streptomycin (Hyclone, Logan, UT, USA) at 37°C in a 5 per cent CO₂ atmosphere.

**Effect of 3-AB on the proliferation of C13* cells:** C13* cells (4 × 10⁴ /ml) were plated in each well of a 96-well plate containing 100 µl growth medium. On the following day, cells were treated with increasing concentration of 3-AB (0, 0.5, 1, 2.5, 5, 7.5 and 10 mmol/l) and cultured for 24 h, each group contained three samples. After 24 h, cells were added with 20 µl 3-[4,5-demethylthiazol -2y] -2,5 diphenyltetrazolium broide MTT (5 mg/ml; Sigma, St Louis, MO, USA) and incubated for 4 h. Cells were collected and centrifuged at 3000 g for 2 min, and the supernatant was discarded. The formazan crystals were dissolved in 150 µl Dimethyl sulfoxide (DMSO), and shaken for 15 min. The absorbance value (A) at 490 nm was read using an automatic multiwell spectrophotometer (Bio-Tek, USA). The inhibitory ratio was calculated as follows: Inhibitory ratio (%) = 1-(mean experimental absorbance mean control absorbance)× 100%.

**Effect of 3-AB on the cell survival of C13* cells:** C13* cells (3 × 10⁴ /ml) were plated in each well of a 96-well plate containing 100 µl growth medium, and treated with 5 mmol/l 3-AB, each group contained six samples and plated six board. One board was taken out every day, the absorbency value was determined using the MTT assay and cell growth curve was drawn.

**Cellular sensitivity to cisplatin after treatment with 3-AB:** C13* cells (4 × 10⁴ /ml) were plated in each well of a 96-well plate containing 100 µl growth medium. Cells were incubated for 16 h to allow adhesion, and then treated with combination of 3-AB (0, 2.5, 5 or 7.5 mmol/l) plus cisplatin (0, 1.25, 2.5, 5, or 10 µl /ml) (Qilu Pharmaceutical Co. Ltd, Jinan, China). Each group contained three samples. After cells were cultured for 24 h, the inhibitory ratio was determined using the MTT assay.

**Reverse transcription-polymerase chain reaction (RT-PCR):** Total RNA was extracted using Trizol (Takara, Japan) according to the manufacturer’s protocol. The first strand cDNA was generated by reverse transcription reagent kit (Takara, Japan). After a sufficient amount of cDNA was obtained, DNA was amplified by performing PCR with primer of PARP-1;
forward, 5'-AAGGCGAATGCCAGCGTTAC-3'; reverse, 5'-GGCACTCTTGGAGACCAGTGCA-3' (Takara, Japan). Amplification was carried out for 30 cycles with 30 sec of incubation at 94°C, 30 sec at 55°C and 1 min at 72°C. The products were analyzed on a 2 per cent agarose gel and examined by staining with ethidium bromide. Their amount was normalized to the value of β-actin as the standard value.

Western blot: Control and treated cells were lysed, and protein concentration was measured by using the BCA protein assay kit (Source?). Cell lysates (60 μg/lane) were separated by SDS-PAGE (10% polyacrylamide gel), transferred to an nitrocellulose (NC) membrane. After blocking with 5 per cent non-fat milk in Tris-buffered saline, the blots were incubated with human anti-rabbit parp-1 antibody (1:1000; Cell Signaling Technology, Beverly, MA, USA) at 4°C overnight. Mouse anti-beta-actin monoclonal antibody (1:1000; Cell Signaling Technology, Beverly, MA, USA) served as a control for protein loading in each lane. Membranes were incubated with horse radish peroxidase (HRP)-conjugated secondary antibodies (1:8000; Santa Cruz, CA, USA), for 1 h at room temperature. The protein bands were visualized by ECL detection system, and the density of the bands was quantified by Alpha Imager 2200 (Alpha Innotech, San Leandro CA, USA).

Immunofluorescence: Control and treated cells were washed with PBS, fixed in 4 per cent paraformaldehyde, and permeabilized with 0.2 per cent Triton-X 100 for 20 min in a humidified tissue culture incubator at 37°C, then processed using standard immunofluorescence staining procedures8. Cells were incubated with a human anti-rabbit parp-1 antibody (1:200; Cell Signaling Technology, Beverly, MA, USA) at 4°C overnight. One well was incubated with PBS instead of the primary antibody and thus was used as the negative control. Fluorescein isothiocyanate (FITC)-conjugated secondary antibodies (1:300; Santa Cruz, CA, USA), for 30 min at 37°C. The nuclei were stained with 4′,6-diamidino-2-phenylindole (DAPI) solution for 2 min. The cells were observed using fluorescence microscopy (Thermo Hybaid, USA) and photographed with a CCD camera (Olympus, Japan).

Statistical analysis: Each experiment was repeated three times. All statistical analyses were performed using SPSS 18.0 software (SPSS, Inc., Chicago, IL, USA). Differences among various treatment groups were determined by analysis of variance followed by Dunnett’s t test.

Results

The effects of 3-AB on the proliferation of C13* cells: The inhibitory ratios of the C13* cells treated with 3-AB at 0.5, 1, 2.5, 5, 7.5 or 10 mmol/l were 5 ± 1.4, 13 ± 1.2, 21 ± 1.3, 30 ± 1.5, 41 ± 1.8, and 59 ± 1.0 per cent, respectively. Hence, the inhibitory ratio was associated with the concentration of 3-AB (Fig. 1; P<0.05).

Effect of 3-AB on the cell survival of C13* cells: After treated with 5 mmol/l 3-AB, the proliferation capacity of C13* cells was inhibited, and cell growth slowed down. The difference was significant at 2, 3, 4, 5, and 6 days compared to C13* group (Fig. 2, P<0.05).

Cellular sensitivity to cisplatin after treatment with 3-AB: When the Pt concentration was fixed (2.5 µg/ml), the inhibitory ratio of C13* cells exhibited a significant increase in response to higher 3-AB (5 mmol/l) concentrations (P<0.05). Similarly, when the 3-AB concentration was fixed, the inhibitory ratio of C13* cells was also increased by higher concentrations of Pt (P<0.05). These results indicated that the PARP inhibitor 3-AB inhibited the proliferation activity of C13* cells, and increased the cellular sensitivity to cisplatin (Table).

RT-PCR analysis: The RT-PCR analysis revealed that after cells were treated with 2.5 µg/ml Pt, 5 mmol/l 3-AB, or 2.5 µg/ml Pt plus 5 mmol/l 3-AB, the expression levels of PARP-1 mRNA in the C13* cells were 0.6188 ± 0.03, 0.4348 ± 0.01, and 0.2219 ± 0.02, respectively, compared to the blank control (0.6844 ± 0.06). Our data indicated that the expression

![Fig. 1. Influence of 3-AB on the cellular proliferation (inhibitory ratio) detected by MTT. The inhibitory ratios of the C13* cells were associated with the concentration of 3-AB (0, 0.5, 1, 2.5, 5, 7.5 and 10 mmol/l).](image-url)
of PARP-1 mRNA was unaffected by Pt treatment, but was significantly decreased when Pt and 3-AB were combined ($P<0.05$).

**Western blot analysis**: The Western blot analysis revealed that after C13* cells were treated with 3-AB (0, 2.5, 5 or 7.5 mmol/l) for 24 h, the expression levels of PARP-1 in C13* cells were 1.69 ± 0.08, 1.30 ± 0.04, 0.69 ± 0.04 and 0.45 ± 0.07, respectively. Compared to the blank control (0 mmol/l 3-AB), the PARP-1 expression levels of 3-AB treated cells were significantly decreased by 23.15 ± 2.53 per cent for the 2.5 mmol/l 3-AB group, 59.11 ± 2.23 per cent for the 5 mmol/l 3-AB group, and 73.24 ± 3.88 per cent for the 7.5 mmol/l 3-AB group ($P<0.05$). These results suggested that 3-AB significantly downregulated the PARP-1 expression at the translational level, and further implicated that the overexpression of PARP-1 protein in the ovarian cancer C13* cells was related to the cellular sensitivity to cisplatin.

**Immunofluorescence**: PARP-1 proteins were largely located in the nuclei, and a small fraction of the proteins appeared in the cytoplasm (Fig. 3). After 3-AB treatment, the expression level of PARP-1 in the C13* cells was decreased. These data demonstrated that 3-AB significantly downregulated the expression of PARP-1.

**Discussion**

Platinum-based chemotherapy has been used as primary treatment for recurrent ovarian epithelial cancer. However, the median time to recurrence is not more than 2 years, with a 5-years survival rate of 20-25 per cent\(^{10}\). Acquired resistance to chemotherapy is the primary cause of failure in the treatment of cancers. Therefore, identification of the potential molecular pathways involved in cisplatin-resistant chemotherapy has become one of the significant areas in the research of tumour treatment.

Platinum is a non-specific anti-tumour medicine, and its cytotoxicity against tumour cells has been thought to be mediated through several mechanisms\(^{11}\), including formatting and enhancing the platinum-DNA adduct repair capacity, decreasing platinum accumulation, and increasing drug inactivation and its ability to tolerate platinum-DNA damage\(^{12}\). The DNA repair capability of cells may be a rate-limiting factor for cisplatin resistance and sensitivity in cisplatin resistant ovarian cancer cells\(^{13}\).

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<th>Table. Influence of 3-AB on the cellular sensitivity to cisplatin (inhibitory ratio) detected by MTT</th>
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Fig. 2. Effect of 3-AB on the cell survival of C13* cells detected by MTT. The difference of cell growth was significant at 3, 4, 5 and 6 days compared to control group.

Fig. 3. Detection of the PARP-1 protein expression in C13* cells by Immunofluorescent microscopy. (A) PARP-1 protein expression and localization in C13* cells (red). (B) PARP-1 protein expression and localization in C13* cells after treated with 5 mmol/l 3-AB (red) for 24 h. Nuclei stained with DAPI (blue). Original magnification, × 400.
Poly(ADP-ribo)ylation is post-translational type of protein modification involved in the regulation of cell cycle, transcription, initiation of the DNA repair and apoptosis. The poly(ADP-ribo)ylation process can regulate a series of molecular events through regulating enzymatic activities and macromolecular interactions between proteins, DNA and RNA. PARP-1 is responsible for sensing the signals of DNA damage. DNA strand breaks can be caused by exposures to ionizing radiation, UV light, chemotherapy, or products of cellular metabolism. Activation of PARP-1 can lead to the addition of poly-(ADP-ribose) branched chains onto the damaged DNA, as well as the assembling of proteins involved in DNA repair. In addition to its role in DNA repair, poly(ADP-ribo)ylation can also protect the naked DNA from being degraded by nuclease.

It has been suggested that PARP-1 facilitates diverse inflammatory responses through upregulating the expression of inflammation-relevant genes that encode oxidation-reduction-related enzymes, cytokines, and adhesion molecules. Further, excessive activation of PARP-1 can induce arteriosclerosis, myocardial infarction, hypertension and cardiovascular complications of diabetes. However, the relationship between PARP and tumours has been rarely reported. The overexpression of PARP in breast cancer, glioblastoma, liver cancer, prostatic carcinoma and pancreatic cancer can facilitate the proliferation of tumour cells. Gurung et al. suggested that the inhibition of PARP-1 and telomerase in MEFs rendered cells is more susceptible to DNA-damaging agents. Other studies demonstrated that inhibiting the activation of PARP could enhance glioblastoma and lung cancer sensitivity to radiation. The PARP-1 inhibitor ANI and the suppression of PARP-1 expression by siRNA can significantly sensitize human liver cancer cells to doxorubicin treatment.

The inhibitory effects of 3-AB on PARP-1 expression have been well established. Valenzuela et al. showed that a concentration of 5mM of 3-AB after 2 h-exposure efficiently blocks PARP activity avoiding PARP-polymer formation in response to DNA damage in primary mouse embryonic fibroblast. Wang et al. reported that 3-AB was able to reduce PARP-1 activity in the millimole range in the repair proficient human glioma M059K cells. Zheng et al. indicated that 3-AB suppressed cell growth, cell invasion and enhanced the suppressive effects of cisplatin in vitro in U2OS cells. They suggested that 3-AB may be developed into an effective agent for the treatment of human osteosarcoma.

In summary, our study indicated that the PARP inhibitor 3-AB inhibited the proliferation activity of C13 cells, and increased the cellular sensitivity to cisplatin. We thus proposed that combined treatment of PARP inhibitor with cisplatin can reverse the drug resistance of ovarian cancer cells, reduce the dosage and side effects of cisplatin chemotherapy, and improve the prognosis of patients suffering from ovarian cancer. However, the PARP protein family includes 17 members, and chemical inhibitors may have nonspecific suppression effects. Systemic long-term administration of PARP inhibitors may harm the DNA repair and genomic stability in normal cells, which can lead to secondary tumours years after. Therefore, novel PARP-1 inhibitor with low toxicity and high specificity need to be tested in future. With the progress in the PARP research and combination therapy, targeted therapy using PARP inhibitors may play a vital role in the prevention of tumours.

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Conflicts of interest: There is no conflicts of interest to declare.

References


27. Gurung RL, Balakrishnan L, Bhattacharjee RN. Inhibition of poly(ADP-ribose) polymerase-1 in telomerase deficient mouse embryonic fibroblasts increases arsenite-induced genome instability. Genome Integr 2010; 1 : 5.


