Abstract. Ovarian cancer is a common tumor of the reproductive system, and primarily responds to cytoreductive surgery and cisplatin (DDP)-based chemotherapy. However, chemoresistance results in high ovarian cancer mortality. Therefore, the aim of the present study was to investigate the effects of metformin on the apoptosis and autophagy of ovarian cancer drug-resistant SKOV3/DDP cells. To do so, MTT assay, flow cytometry, electron microscopy and western blotting were used in the present study. Metformin could inhibit the growth of SKOV3 and SKOV3/DDP cells in a concentration- and time-dependent manner (P<0.05). The half-inhibitory concentration (IC50) values of DDP and methotrexate (MTX) were 14.35 and 4.21 µg/ml for SKOV3 cells, and 70.26 and 15.27 µg/ml for SKOV3/DDP cells, respectively. In addition, the resistance index of SKOV3/DDP for DDP and MTX was 4.89 and 3.62, respectively. After combining metformin with DDP and MTX, the IC50 values for SKOV3 cells were 11.20 and 2.80 µg/ml, and 6.21 and 2.74 µg/ml for SKOV3/DDP cells, respectively. Metformin decreased the IC50 of DDP and MTX in drug-resistant cancer cells SKOV3/DDP by 11.31- and 6.18-fold. This indicated that cell proliferation was inhibited when treated with the combination of metformin and chemotherapeutic agents, compared with chemotherapeutic agents alone. In addition, autophagy was not observed in SKOV3 and SKOV3/DDP cells; however, it was observed in SKOV3/DDP cells following incubation with 10 mmol/l metformin for 48 h. Furthermore, the expression levels of microtubule-associated protein 1 light chain 3-II protein in SKOV3/DDP cells were upregulated compared with in SKOV3 cells (P<0.05). These results demonstrated that metformin can sensitize drug-resistant ovarian cancer cells to chemotherapeutic agents, and that it may be associated with the induction of autophagy.

Introduction

Ovarian cancer has one of the highest rates of mortality in the USA, and is the leading cause of cancer-associated mortality in women (1). Diagnosis of ovarian cancer is a problem, as early stages have no noticeable signs or symptoms (2). The majority of patients are diagnosed at an advanced stage of the disease, which is mainly treated with cytoreductive surgery and platinum-based chemotherapy. However, 60-65% of patients will subsequently relapse, and a low five-year survival rate is observed following the initial diagnosis (3). The high mortality rate for ovarian cancer is primarily caused by chemoresistance, which remains an important obstacle to successful chemotherapy (4). Therefore, it is necessary to find a novel therapeutic approach to improve the effects of ovarian cancer treatment.

Cell death mainly includes apoptosis, autophagy and necrosis. Disruption in the balance between cell proliferation and apoptosis leads to uncontrollable cell growth and tumorigenesis; therefore, apoptosis can inhibit tumor growth (5,6). The majority of chemotherapy drugs kill cancer cells by inducing apoptosis (5). It has previously been reported that chemoresistance of tumor cells is not only associated with inhibiting apoptosis and reducing apoptotic susceptibility, but is also associated with autophagy (7,8). Autophagy is a self-stabilizing mechanism in eukaryotic cells that allows the orderly degradation and recycling of cellular components (9). This self-degradation process not only serves a crucial role in cell protection but can also be involved in cell killing (10). Hence, autophagy and apoptosis can work individually, and can affect the sensitivity of tumor cells to chemotherapy in a reciprocal manner (11,12). However, to the best of our knowledge, whether autophagy protects or kills cells remains unclear.

Metformin as an insulin sensitization agent was formally approved as a first-line treatment for diabetes by the American Diabetes Association and European Association for the Study of Diabetes (13). Previous studies have revealed that metformin can not only effectively reduce blood sugar levels and regulate lipid metabolism, endocrine function, hypercoagulability and hyperthrombocytopenia, but may
also possess antitumor effects (13-15). It has been reported that metformin can inhibit breast, prostate, renal, pancreatic and colon cancer in animal and cell experiments (16-19). The antitumor effects of metformin are induced by the inhibition of tumor cell proliferation, promotion of apoptosis in tumor cells, activation of tumor sensitivity to chemotherapeutic agents and inhibition of tumor angiogenesis (19). However, research into the effects of metformin on ovarian cancer is currently limited.

The present study investigated the effects of metformin on cisplatin (DDP) and methotrexate (MTX)-induced apoptotic cell death in epithelial ovarian cancer parental SKOV3 and drug-resistant SKOV3/DDP cells. Metformin inhibited the growth of SKOV3/DDP cells and enhanced the sensitivity of SKOV3/DDP cells to chemotherapeutic agents.

Materials and methods

Chemical reagents. Metformin, MTT, DDP, MTX, ribonuclease A (RNase A), DAPI, fluorescein isothiocyanate isomer (FITC), propidium iodide (PI), dimethyl sulfoxide (DMSO), ethanol, formaldehyde, glutaraldehyde, epoxy resin, acetone and Nonidet P-40 were purchased from Sigma-Aldrich (Merck KGaA). RPMI 1640 medium, fetal bovine serum (FBS), penicillin and streptomycin were purchased from Thermo Fisher Scientific, Inc. Polyclonal antibodies against microtubule-associated protein 1 light chain 3 (LC3), goat anti-rabbit IgG conjugated to horseradish peroxidase (HRP) and anti-β-actin antibody were purchased from Santa Cruz Biotechnology, Inc. An Annexin V-FITC apoptosis detection kit was purchased from Santa Cruz Biotechnology, Inc. An Annexin V-FITC apoptosis detection kit was purchased from Xi’an Jiaoda Bao Sai Bio-technology Co., Ltd. A Pierce™ Bicinchoninic Acid (BCA) Protein Assay kit was supplied by Thermo Fisher Scientific, Inc. Total cell protein extraction kit was purchased from Shanghai BestBio Biotechnology Co., Ltd. All other chemicals and solvents were purchased from Sigma-Aldrich (Merck KGaA).

Cell culture. The ovarian cancer cell lines SKOV3 and SKOV3/DDP were provided by the Institute of Basic Medicine, Chinese Academy of Medical Sciences & Peking Union Medical College. Both parental and drug-resistant cells were cultured in RPMI 1640 medium, supplemented with 10% FBS, 100 U/ml penicillin and 100 µg/ml streptomycin at 37°C with 5% CO₂.

MTT assay. An MTT assay was used to evaluate the resistance index (RI) of SKOV3 and SKOV3/DDP cells. Cells were seeded in 96-well microtiter plates (1x10³ per well). After 12 h, different concentrations of DDP (12.5, 25, 50 and 100 µg/ml) or MTX (2, 4, 8 and 16 µg/ml) were added to the cells for 72 h at 37°C. Cells were washed with PBS and incubated with MTT (50 µl; 0.5 mg/ml). After 4 h incubation at 37°C, the formazan precipitates were dissolved in DMSO (150 µl/well). The optical density (OD) was measured at a wavelength of 570 nm using a Benchmark microplate reader (Bio-Rad Laboratories, Inc.). The OD values were used to calculate the inhibition rates and half-inhibitory concentration (IC₅₀) of chemotherapeutic agents. The ratio of IC₅₀ for drug-resistant and parental cells (RI), was calculated to evaluate the drug-resistance of SKOV3/DDP cells. The MTT assay was performed in triplicate.

The MTT assay was also used to evaluate the effect of the tested compounds on the viability of SKOV3/DDP cells. After the cells were seeded in a 96-well microtiter plates for 12 h, different concentrations of metformin (2.5, 5 and 10 mmol/l) were added. The cells were incubated for 24, 48 and 72 h at 37°C prior to the addition of MTT. To determine the susceptibility to metformin as a chemotherapeutic agent, the cells were incubated with 10 mmol/l metformin for 12 h at 37°C, and different concentrations of DDP (12.5, 25, 50 and 100 µg/ml) or MTX (2, 4, 8 and 16 µg/ml) were subsequently added to the wells. The cells and agents were co-cultured for another 48 h at 37°C prior to the MTT assay.

Detection of apoptosis by flow cytometry. Following incubation of SKOV3 and SKOV3/DDP cells with different concentrations of metformin for 48 h, attached and floating cells were harvested and washed twice with PBS. The cells were fixed with 70% ethanol at 4°C overnight, and then incubated in RNase A (100 µg/ml in PBS) at 37°C for 30 min. Detection of apoptotic cells was performed using an Annexin V-FITC apoptosis detection kit including recombinant human Annexin V labelled by FITC, according to the manufacturer’s protocol. After incubation with RNase A, cells (1x10⁶) were washed with cold PBS prior to incubation with FITC (10 µl) and PI (5 µl) for 15 min in the dark at 4°C. The apoptotic rate was analyzed using flow cytometer (Beckman Coulter, Inc.) and quantified with CellQuest Pro V5.1 software (BD Biosciences).

Detection of apoptosis by DAPI staining. Following incubation of SKOV3 and SKOV3/DDP cells with indicated drugs (10 mmol/l DDP or 10 mmol/l metformin or 10 mmol/l DDP + 10 mmol/l metformin) for 48 h, attached and floating cells were both harvested and fixed at 4°C for 2 h with PBS solution containing 3.7% formaldehyde, 0.5% Nonidet P-40 and 0.1 mg/l DAPI. Apoptosis was assessed by fluorescence microscope (Olympus Corporation; magnification, ×200) of condensed chromatin and micronucleation. Cells with a condensed and/or fragmented nucleus were considered as apoptotic cells. At least two independent experiments were carried out for each treatment condition and a minimum of 300 cells were counted by ImageJ V1.8.0 software (National Institutes of Health) in each experiment.

Detection of autophagy by electron microscopy. Metformin (10 mmol/l) was added to SKOV3 and SKOV3/DDP cells for 48 h. Following centrifugation at room temperature for 10 min at 200 x g and cleaning, cells were fixed with 2.5% glutaraldehyde at 4°C overnight. Fixed cells were stained with 1% uranium acetate at room temperature for 10 min, washed with double distilled water for 20 sec, subsequently stained with 0.2% lead citrate at room temperature for 10 min and washed with double distilled water for 20 sec. Then, the specimens were rinsed with PBS, dehydrated in a graded acetone series (50, 70, 80, 90 and 100%) and embedded in an epoxy resin/acetone (1:1) mixture at room temperature for 1 h. Ultra-thin sections (50-60 nm) were examined by the FEI TecnaiG2 F20 transmission electron microscope (TEM; magnification, x15,000; FEI Company).
Western blot analysis for the detection of protein-associated apoptosis and autophagy. Total proteins were extracted from cells following the aforementioned treatments with the total cell protein extraction kit (cat. no. BB-3101; Shanghai BestBio Biotechnology Co., Ltd.) and proteins were quantified using a BCA Protein Assay kit (Thermo Fisher Scientific), according to the manufacturer's protocol. Equal amounts of proteins (80 µg/lane) were separated via SDS-PAGE (12% gels), and transferred to nitrocellulose membranes (Bio-Rad Laboratories, Inc.). The membrane was probed with polyclonal antibodies against LC3 (cat. no. sc398822; 1:500) overnight at 4˚C, followed by incubation with HRP-conjugated secondary antibody (cat. no. sc2030; 1:3,000) at room temperature for 2 h. Visualization was achieved using Super Signal enhanced chemiluminescence (Applygen Technologies, Inc.). The data were analyzed via densitometry using Quantity One V4.6.2 software (Bio-Rad Laboratories, Inc.). To determine equal lane loading, the membrane was stripped and re-probed with anti-β-actin antibody (cat. no. sc47778; 1:5,000).

Statistical analysis. Statistical analysis was performed using SPSS software (version 17.0; SPSS, Inc.) and SAS 9.4 (SAS Institute Inc.). All experiments were repeated as least in triplicate, and the values were presented as the mean ± standard deviation. A two-way analysis of variance was applied to compare the main and interactive effects of variables, followed by Bonferroni post hoc test. P<0.05 was considered to indicate a statistically significant difference.

Results

Metformin reduces viability and induces apoptosis in ovarian cancer cells. Metformin could inhibit the growth of SKOV3 and SKOV3/DDP cells, whereas inhibition rates were increased in a drug concentration- and time-dependent manner (Table I), suggesting that both cells were sensitive to metformin. In particular, when the concentration of metformin was 10 mmol/l, the inhibitory effect was increased in SKOV3/DDP cells compared with SKOV3 cells (P<0.05). For the detection of metformin-induced apoptosis, flow cytometric analysis was performed. As presented in Fig. 1, when 2.5, 5 and 10 mmol/l metformin were incubated for 48 h, the apoptotic rates of SKOV3/DDP cells were 11.45±2.78, 33.28±1.30 and 40.36±2.58%, respectively, whereas the apoptotic rates of SKOV3 cells were 10.56±1.08, 31.31±2.21 and 35.60±3.24, respectively (Tables II and III).

Effects of metformin on the cytotoxicity of SKOV3/DDP ovarian cancer cells. As presented in Table IV, the inhibition rates of DDP and MTX on SKOV3 and SKOV3/DDP cells were increased as the concentration of chemotherapeutic agents increased. The IC_{50} values of MTX and DDP were 4.21 and 14.35 µg/ml for SKOV3 cells and 15.27 and 70.26 µg/ml for SKOV3/DDP cells, respectively (Table V). Therefore, the inhibition rates of DDP and MTX on SKOV3 cells were significantly increased compared with SKOV3/DDP cells (P<0.05). The SKOV3/DDP cell was resistant to DDP and MTX, and the RI values were 4.89 and 3.62, respectively.

To evaluate the effects of metformin on the cytotoxicity of chemotherapeutic agents, an MTT assay was performed. As indicated in Table V, the IC_{50} values of metformin combined with MTX and DDP were 2.80 and 11.20 µg/ml for SKOV3 cells, and 2.47 and 6.21 µg/ml for SKOV3/DDP cells, respectively. Compared with the IC_{50} values for when chemotherapeutic agents were used alone, metformin decreased the IC_{50} of MTX and DDP in drug-resistant cancer cells, SKOV3/DDP, by

| Table I. Inhibition rates of metformin with indicated concentrations and incubation times in SKOV3 and SKOV3/DDP cells. |
|-----------------------------|-----------------------------|-----------------------------|-----------------------------|
| Concentration (mmol/l)     | Time (h)        | SKOV3 (%±SD) | SKOV3/DDP (%±SD) | t-statistic | P-value |
| 2.5                        | 24             | 10.80±1.78   | 13.14±2.94       | 0.290       | 0.779   |
|                            | 48             | 18.41±1.94   | 16.84±1.90       | -0.470      | 0.651   |
|                            | 72             | 22.20±1.19   | 24.89±2.22       | 3.275       | 0.061   |
| 5                          | 24             | 16.25±1.60   | 18.82±1.92       | 3.301       | 0.050   |
|                            | 48             | 23.01±2.76   | 22.46±2.56       | -0.330      | 0.750   |
|                            | 72             | 27.62±1.62   | 31.84±2.25       | 3.398       | 0.009   |
| 10                         | 24             | 20.30±1.69   | 27.97±1.82       | 6.904       | <0.001  |
|                            | 48             | 27.95±2.69   | 34.35±2.39       | 3.984       | 0.004   |
|                            | 72             | 33.30±3.56   | 38.04±2.90       | 2.309       | 0.041   |

N=3/group. DDP, cisplatin; SD, standard deviation.

| Table II. Comparison by two way ANOVA of the effects of metformin concentration and cell type, alone and combined, on apoptotic rates. |
|-------------------------------------------------|-----------------|-----------------|-----------------|
| Factor                                          | F-value | P-value |
| Metformin concentrationa                          | 707.130  | 0.000   |
| Cell typeb                                       | 11.070   | 0.004   |
| Metformin concentration * Cell typec              | 4.212    | 0.022   |

aMain effect of metformin concentration on apoptotic rates; bMain effect of cell type on apoptotic rates; cInteraction effect between metformin concentration and cell type on apoptotic rates.
Figure 1. Metformin-induced apoptosis in SKOV3 and SKOV3/DDP cells. SKOV3 and SKOV3/DDP cells were incubated with different concentrations of metformin for 48 h. FITC, fluorescein isothiocyanate isomer; PI, propidium iodide; DDP, cisplatin.
6.18- and 11.31-fold, respectively. These results indicated that SKOV3/DDP cell viability was significantly inhibited when treated with the combination of metformin and chemotherapeutic agents compared with chemotherapeutic agents alone.

The effects of metformin on apoptosis were also evaluated using DAPI staining. Compared with chemotherapeutic agents alone, the combination of metformin and DDP significantly increased apoptotic rates. As indicated in Table VI, when DDP was replaced by the combination of metformin and DDP, the apoptotic rates for SKOV3/DDP significantly increased from 7.02 to 75.22%; similarly, the apoptotic rates for SKOV3 cells increased from 24.50 to 74.12%.

Metformin upregulates the expression of LC3-II in ovarian cancer cells. LC3 possesses two forms, including an 18-kDa cytosolic protein (LC3-I) and a processed 16-kDa form (LC3-II). The expression levels of LC3-II and the ratio of LC3-II/LC3-I represent autophagic activity (11). Therefore, LC3-I and LC3-II were detected in SKOV3 and SKOV3/DDP cells incubated with different concentrations of metformin for 48 h using western blot analysis. As indicated in Fig. 3, the expression levels of LC3-II in SKOV3/DDP cells were higher than those in SKOV3 cells,
indicating that the autophagic activity of drug-resistant cells was increased compared with that of the parental cells. Metformin may induce autophagy in ovarian cancer cells. Following incubation with 10 mmol/l of metformin for 48 h, the expression levels of LC3-II protein were significantly increased in SKOV3 and SKOV3/DDP cells, and the expression levels in SKOV3/DDP cells were higher compared with SKOV3 cells (Fig. 4). These results demonstrated that metformin could promote the autophagy of ovarian cancer cells, and served a crucial role in promoting autophagy in drug-resistant ovarian cancer cells.

Discussion

Ovarian cancer had the highest mortality rate among gynecologic malignant tumors in the USA in 2008 (20). At present, the leading therapeutic method for ovarian cancer is operative treatment combined with platinum-based chemotherapy; however, chemoresistance severely influences the prognosis of the patient, and results in high mortality rates. Therefore, overcoming chemoresistance has become the key to treating ovarian cancer (4). Metformin is a member of the biguanides group of drugs, and is a first-line drug for the treatment of type 2 diabetes (14). Metformin not only reduces blood sugar levels, but also possesses other pharmacological effects, including cardiovascular protective effects (21) and delaying Alzheimer's disease (22). Those patients with type 2 diabetes who received metformin had a decreased rate of cancer-associated mortality compared with those receiving sulfonylureas (3.5 vs. 4.9%) (23). Metformin has been reported to enhance the antiproliferative effects of paclitaxel and cisplatin in breast cancer, lung cancer and other cells (24-27).

In the present study, metformin not only suppressed the growth of ovarian cancer parental SKOV3 cells and drug-resistant SKOV3/DDP cells, but it also sensitized drug-resistant SKOV3/DDP cells to chemotherapeutic agents. The inhibition rates of metformin increased in SKOV3 and SKOV3/DDP cells following drug treatment with increasing concentration and incubation time. In particular, when the concentration of metformin was 10 mmol/l, significantly increased inhibition was observed in SKOV3/DDP cells compared with SKOV3 cells. In addition, the apoptotic rates of the two cell types incubated with metformin for 48 h increased with increasing drug concentration. These results indicated that the two cell types were sensitive to metformin.

Metformin-induced chemosensitivity of ovarian cancer cells was examined via MTT assays and DAPI staining. Cell viability was markedly inhibited in both types of cells when treated with a combination of metformin and a chemotherapeutic agent (DDP or MTX) compared with those treated with chemotherapeutic agents alone. The IC50 values for the combination of metformin with DDT (or MTX) were reduced by 11.31- (or 6.18-) fold in SKOV3/DDP cells compared with DDT (or MTX) alone, but only 1.28- (or 1.50-) fold in SKOV3 cells. Similarly, following incubation with DDP
alone for 48 h, the apoptotic rates were 24.50 and 7.02% for SKOV3 and SKOV3/DDP cells, respectively, and the former were notably higher compared with the latter. With the combination of metformin and DDP, instead of DDP only, the apoptotic rates for SKOV3 and SKOV3/DDP cells were not significantly different; 74.12 and 75.22%, respectively. These results demonstrated that metformin can enhance the cytotoxicity of chemotherapeutic agents on drug-resistant cancer cells.

Autophagy, as a type of cell death, degrades cellular components, and damaged organelles and macromolecular material are wrapped by cytomembrane to form a complete autophagosome; then, the autophagosome is combined with a lysosome and digested to complete the autophagy process (9,10). Autophagy is considered a double-edged sword in the process of tumor development (10). The effects that autophagy has on tumor cells are complicated and controversial. Autophagy has been demonstrated to suppress carcinogenesis via the elimination of oncogenic molecules and damaged organelles (9). However, once an invasive cancer is established, autophagy promotes tumor growth via the intracellular recycling of degraded metabolites, which further stimulates the metabolism of cancer cells (28). In addition, autophagy is enhanced by chemotherapy and radiation therapy, and functions as an adaptive response that mediates resistance to these treatments (29). In particular, autophagy is associated with the occurrence and development of ovarian cancer, and also with DDP-resistance in ovarian cancer cells (30). In the present study, autophagy was significantly increased in SKOV3/DDP cells following incubation with metformin for 48 h, resulting in cell death. The aforementioned, however, was not observed in SKOV3 cells. LC3-II is a marker of autophagosome formation. Incubation with metformin induced a significant increase in the expression levels of LC3-II, and the expression levels of LC3-II in drug-resistant cells were upregulated compared with in parental cells. Metformin enhanced the autophagy activity of ovarian cancer cells, and the effect was more prominent in SKOV3/DDP cells compared with that in SKOV3. The apoptotic rates of drug-resistant cells, with the incubation of combined metformin and DDP, were roughly equivalent to the apoptotic rates of the parental cells. These results indicated that metformin may sensitize drug-resistant SKOV3/DDP cells to chemotherapeutic agents through the induction of autophagy.

In conclusion, metformin not only induced apoptosis of ovarian cancer parental SKOV3 and drug-resistant SKOV3/DDP cells, but also enhanced autophagic activity in SKOV3/DDP cells. Therefore, the fact that metformin can sensitize drug-resistant ovarian cancer cells to chemotherapeutic agents may indicate an association with inducing autophagy. Although the specific underlying molecular mechanism remains unclear in the present study, metformin is a potential anticancer drug, particularly for the treatment of drug-resistant ovarian cancer. The results of the present study will be verified in future studies that include additional ovarian cancer cell lines and animal models.

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Availability of data and materials

The datasets used and/or analyzed during the present study are available from the corresponding author on reasonable request.

Authors' contributions

CY and NZ performed the experiments. CY and YC drafted the manuscript. CY, DL and GZ analyzed the data. YC designed the study. All authors read and approved the final manuscript.

Ethics approval and consent to participate

Not applicable.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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