

# PI3K Inhibition Sensitize the Cisplatin-resistant Human Ovarian Cancer Cell OVCAR3 by Induction of Oxidative Stress

Sahar Baghal-Sadriforush<sup>1</sup>, Morteza Bagheri\*<sup>2</sup>, Isa Abdi Rad<sup>2</sup>,  
Fattah Sotoodeh nejadnematalahi<sup>1</sup>

## Abstract

**Background:** This study evaluates the effect of simultaneous AKT inhibition and cisplatin therapy in changes of *Reactive Oxygen Species* (ROS) production, apoptosis induction, and cell survival in cisplatin-resistant OVCAR3 cell.

**Methods:** OVCAR3 cancer cells were treated with cisplatin, Ly 294002 (LY), and cisplatin+Ly to investigate the cytotoxicity effect of the mentioned groups via MTT assay. Then, DCFH-DA (2', 7'-dichlorodihydro fluorescein diacetate) assay kit is used to assess the potential of treated groups in intracellular ROS generation. Protein expression levels of caspase-3, cleaved caspase 3, PI3K, Akt, p-Akt, XIAP, and Survivin are estimated through immunoblotting assay in all three experimental groups.

**Results:** The results showed that all three treated groups, including cisplatin and Ly alone and co-administration of cisplatin+Ly, could reduce the cell vitality of OVCAR3 cancer cells, induced intracellular production of ROS and increased the expression level of activated caspase 3 and Akt protein, whereas down-regulated the phosphorylation of Akt protein. However, the effect of combination therapy was more tangible compared to single therapy and control groups. In contrast, the expression amount of XIAP, Survivin, and PI3K did not show detectable changes in comparison with the control group.

**Conclusions:** The results showed that the AKT inhibition by Ly could sensitize the OVCAR3 cancer cells to the cisplatin and lower the effective dose of cisplatin through hyperactivation of oxidative stress.

**Keywords:** Caspase-3, Cisplatin, Ovarian cancer, PI3K/Akt signaling.

## Introduction

Ovarian cancer with the fourth most common cause of cancer-related death can be considered as an aggressive and under-recognized gynecological malignancy in women across the world (1-3). Lack of effective screening methods and little or no specific symptoms resulted in ovarian cancer diagnosis only at an advanced stage (stage III or IV) and, subsequently, a high mortality rate (2, 4). Surgical debulking followed by the combination of platinum-based and taxane-based drugs (cisplatin and carboplatin) are

first-line treatments used to treat advanced ovarian cancer in clinical practice (5, 6). However, the development of chemoresistance and tumor relapse is the primary clinical obstacle that poses a significant impediment, and restriction in optimal cancer therapy led to a considerable reduction in survival rates (7-9). Therefore, a high concentration of platinum-based treatments and secondary chemotherapy drugs are needed to overcome drug resistance resulted in adverse side effects and severe toxicity such as gastrointestinal disorder,

1: Department of biology, Science and Research branch, Islamic Azad University, Tehran, Iran.

2: Cellular and Molecular Research Center, Cellular and Molecular Medicine Institute, Urmia University of Medical Sciences, Urmia, Iran

\*Corresponding author: Morteza Bagheri; Tel: +98 4433457277; E-mail: mortezabagheri@umsu.ac.ir.

Received: 9 Oct, 2021; Accepted: 17 Oct, 2021

asthenia, renal, and neurological consequences (1, 10). The resistance of ovarian cancer cells to chemotherapy drugs is created firstly due to the complex interaction of multiple factors and multi-gene. For instance, the deregulated apoptosis pathway and also PI3K/Akt pathway are major contributors factors to drug resistance and cancer development (11).

For this reason, it is currently essential to identify a novel approach to improve the efficiency of therapeutic agents and curb their side effects and also molecules to target multiple drug-resistant mechanisms and consequently tackle chemotherapy resistance and increase the sensitivity of cancer cells to chemotherapeutic agents (12, 13). Cisplatin is one of the most effective chemotherapy drugs, which is implied in the form of either monotherapy or combination to treat ovarian cancer, induce apoptosis via cysteine proteases called caspases cascades in ovarian cancer cells (10, 11). In ovarian cancer, activation of phosphatidylinositol 3-kinase/protein kinase B (PI3K/Akt) pathway as a result of down-regulation of phosphatase and tensin homolog (PTEN), RET/PTC oncogenes and activated p21 plays a vital role in biological processes of ovarian tumor cells including tumorigenesis, proliferation, growth regulation, loss of apoptosis, metabolism as well as resistance to apoptosis (14-17). Strategies aimed at the down-regulation of this signaling pathway in combination with conventional anti-cancer drugs might bring about suppression of cell proliferation and cell death and take into account as a suitable approach to overcome chemoresistance in clinical. Hence, LY294002 as an inhibitor of PI3K class I catalytic subunit cause inactivation of this protein led to apoptosis induction and inhibition of cancer cell growth *in vitro* and *in vivo* (2, 4).

In this study, we aimed to investigate the effects of Ly and cisplatin either alone or simultaneously on ROS generation, apoptosis-related proteins, caspase 3 and xXIAP activation, and cell survival-related proteins PI3K, AKT and surviving regulation.

## Materials and Methods

### Materials

MTT [3-(4, 5-dimethylthiazol-2-yl)-2, 5-diphenyltetrazolium bromide] powder and cisplatin as a conventional anti-cancer drug were purchased from Sigma Chemical Co. (St. Louis, MO). Human ovarian cancer cells line (OVCAR3 cell line NCBI code: C209), were obtained from the National Cell Bank of Iran (NCBI), Pasteur Institute. All cell culture reagents were provided from an appropriate, commercially available supplier. All antibodies were from Santa Cruz Biotechnology Company.

### Cell culture

Cisplatin-resistant ovarian cancer cell line OVCAR3 were incubated at 37 °C in a humidified atmosphere with 5% CO<sub>2</sub>. RPMI 1640 as a standard media (Sigma-Aldrich, USA) containing 10% heat-inactivated fetal bovine serum (FBS) (Gibco, USA), 100 U/ml penicillin and 100 µg/ml streptomycin (Gibco, USA) were used routinely to culture cells.

### Cell viability assay

Cancer growth inhibitory effect was demonstrated using (3-(4, 5-Dimethylthiazol-2-yl)-2, 5-Diphenyltetrazolium Bromide (MTT) (Sigma-Aldrich, USA) assay. Briefly,  $1 \times 10^4$  cells/well were seeded into a 96-well plate cultured in RPMI-1640 medium supplemented with 10% FBS and allowed to grow overnight. Later on, the cells were treated with different concentrations of either cisplatin or Ly in the course of 24, 48, and 72 h cultured in the same media complemented with 2% FBS. Next, 20 µl of MTT solution (5 mg/ml) was added to each well and incubated in the dark for another 4h at 37 °C. After the aspiration of culture media, 200µl of dimethyl sulfoxide (DMSO) used to solubilize MTT formazan crystals. Optical density (OD) was determined at a wavelength of 630 nm via a microplate reader (Biotek Instruments, USA). The concentration of cisplatin or Ly caused 50% inhibition of OVCAR3 cell activity named IC<sub>50</sub>, which is expressed as (OD

control-OD experimental)/OD control × 100% and calculated via dose-response curve of cell viability percentage against various concentrations of cisplatin or Ly. Regarding either additive or synergistic effect of simultaneous administration of cisplatin and Ly, the IC<sub>50</sub> value of Ly after 24h combined with various concentrations of cisplatin for 24 h to determine the impact of this combination on IC<sub>50</sub> of Cisplatin. Each evaluation was displayed in triplicate.

### **Measurement of ROS**

The intracellular ROS production was carried out using a DCFH-DA (2', 7'-dichlorodihydro fluorescein diacetate) assay kit. DCFH is oxidized to a highly fluorescent agent such as dichlorofluorescein (DCF) in the existence of cellular oxidizing agents. OVCAR3 cells were seeded in a 96-well plate and then incubated at 37 °C in 5% CO<sub>2</sub> and 95% air. At the end of incubation time, treated cells were harvested and washed with PBS PH 7.4 for two times. Followed by resuspending cells in 100 µl of DCFH-D dissolved in a fresh media in the dark for 30 min at 37 °C and in a humidified atmosphere containing 5 % CO<sub>2</sub> and allowed activation of dye. After removing the free dye by washing with PBS, the cells were subjected to photography by a fluorescence microscope (Olympus IX70, Tokyo, Japan). The results of ROS production are expressed as the Mean Fluorescence Intensity (RIF) of DCF at excitation and emission wavelengths of 530 nm and 485 nm, respectively.

### **Protein extraction and Western blot analysis**

Immunoblotting assay (RIPA) used for protein expression analyses. Firstly, treated cells were suspended in 100 µl RIPA ice-cold lysis buffer (25 mm HEPES, 1% Triton X-100, 2 mm EDTA, 0.1 m NaCl, 25 mm NaF, 1 mm Sodium Orthovanadate) compromised protease inhibitors during 30 min. Afterward, cell lysates were centrifuged at 12000 g for 20 min at 4 °C, and the supernatant was collected to obtain total proteins. Equal amounts of each protein sample (20 µg), which is quantified via Bradford protein assay were separated with

12% Sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE). They subsequently shifted to polyvinylidene difluoride (PVDF) membrane (Roche, UK). After blocking the membrane at room temperature for 1 h with bovine serum albumin (BSA) 3% in Tris-buffered saline (pH 7.5) to remove Non-specific binding reactions, the membrane was maintained with diluted primary antibodies, X-linked inhibitor of apoptosis protein (sc-55550), phospho-Akt (sc-52940), Akt (sc-5298), β-actin (sc-47778), Survivin (sc-17779) and caspase 3 (sc-136219) at 4 °C overnight. After that, the membrane was washed with PBS and then probed with HRP-conjugated secondary antibody for 2 h at room temperature. Eventually, Protein expression was visualized through the enhanced chemiluminescence (ECL) detection kit (Pierce, Rockford, IL). The protein band intensities were pictured and analyzed by Image J software (version 1.62, National Institutes of Health, Bethesda, MD, USA) and normalized to β-actin as an internal loading control (18).

### **Statistical analysis**

All the experiments were carried out in triplicate and repeated three times. Statistical analysis was performed using the GraphPad Prism 6 Scientific software (GraphPad Software, Inc., La Jolla, CA). The results were presented as the mean±SEM. One-way analysis of variance (ANOVA) followed by Tukey post-hoc test to determine the differences between the experimental groups. A p value of < 0.05 was considered to indicate statistically significant.

## **Results**

### **MTT assay**

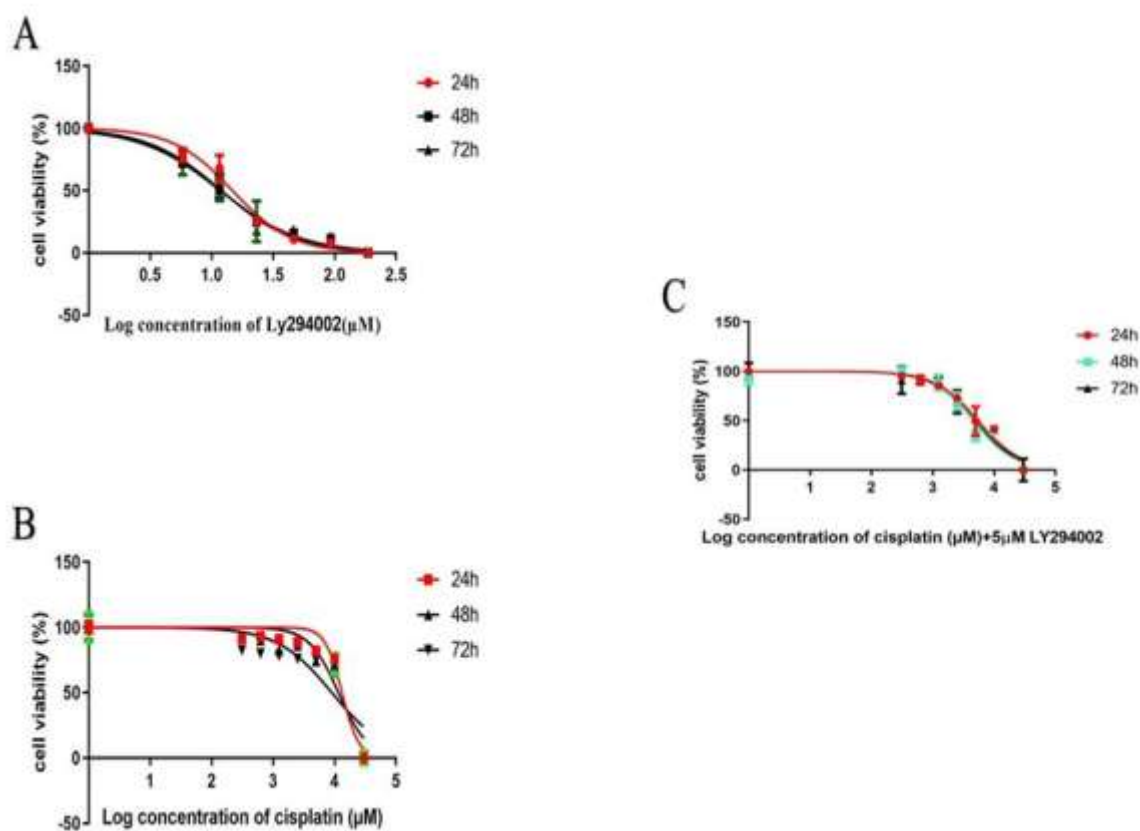
The effect of Ly, cisplatin, and the combination of cisplatin and Ly on cell viability was examined via MTT assay on OVCAR3 cancer cells, and the results showed in Figures 1A, 1B, and 1C. Based on obtained results IC<sub>50</sub> value of cisplatin alone became 13.18 µM, 11.55 µM, and 9.02 µM for 24, 48, and 72 h, respectively. Besides, Cis+Ly

reduced cell viability more than single therapy with an IC50 amount of 5.46  $\mu$ M, 4.60  $\mu$ M, and 4.82  $\mu$ M during 24, 48, and 72 h,

respectively, which proved the increased sensitivity of OVCAR3 cancer cells to cisplatin in the presence of Ly (5  $\mu$ M).

**Table 1.** Comparison of IC50 amount of cisplatin, cisplatin in combination with Ly. \*p<0.05, \*\* p<0.01 and \*\*\* p<0.001 versus control.

Time	Cisplatin IC50	Cisplatin+Ly IC50	p-value
24	13.18	5.46	p<0.001
48	11.55	4.60	p<0.001
72	9.02	4.82	p<0.01

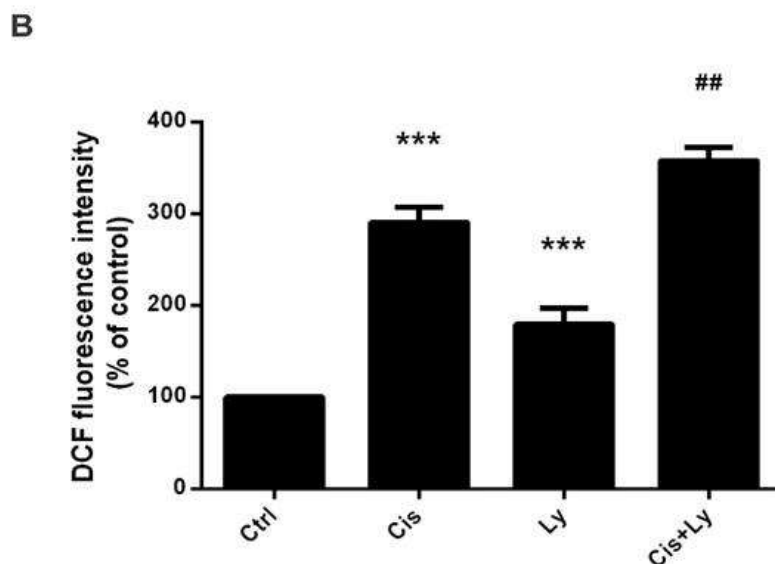
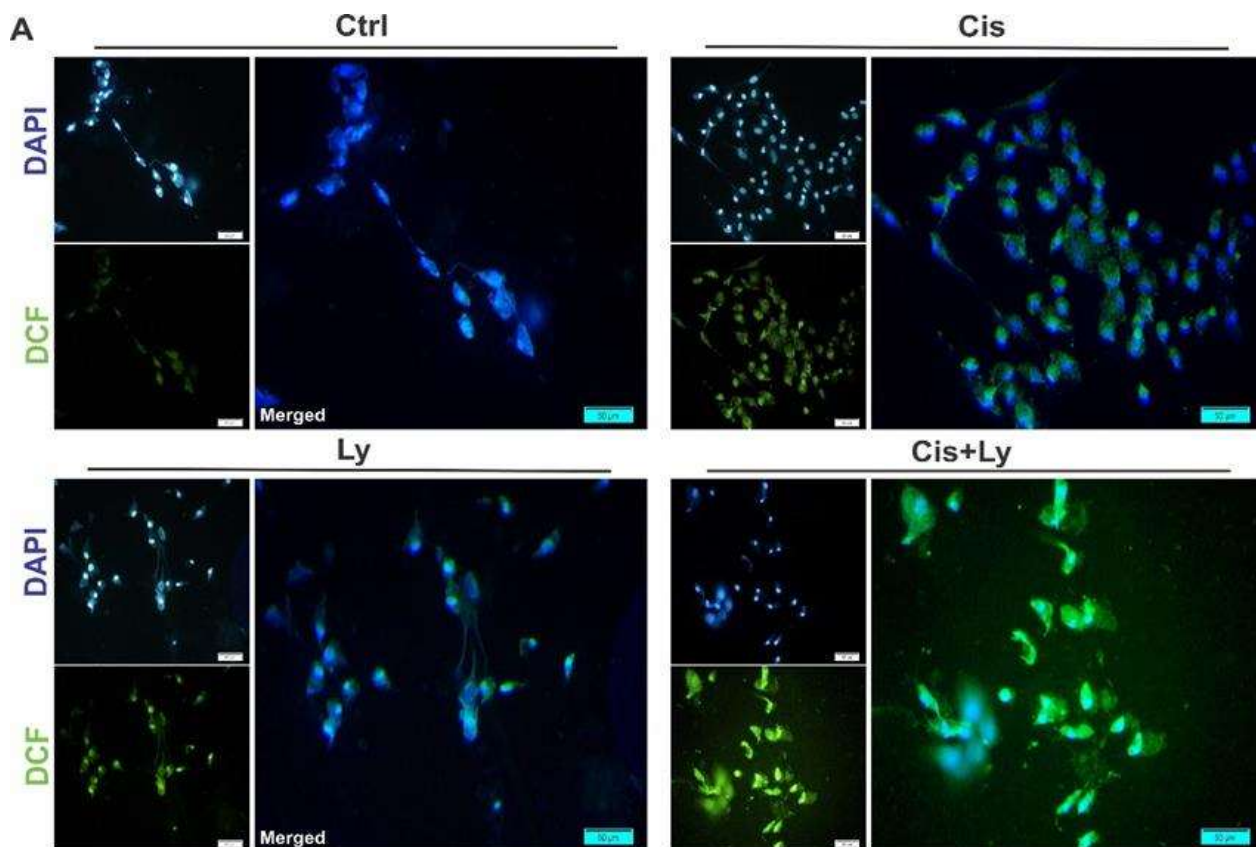


**Fig. 1.** effect of Ly, Cisplatin, and Cisplatin+Ly on the proliferation of ovarian cancer cells alone or in combination. OVCAR3 cells were treated with a fixed concentration of 5  $\mu$ M of Ly and different concentrations of cisplatin and cisplatin+ Ly during 24, 48, and 72 h. Obtained triplicate data were expressed as mean $\pm$ SEM.

**Intracellular ROS levels**

H2DCF-DA was used to examine ROS generation in OVCAR3 cells. The results in Fig. 2 revealed that Ly, cisplatin, and co-administration of cisplatin and Ly significantly induced intracellular ROS generation versus

control. Among them, combined treatment with cisplatin and Ly displayed considerable augmentation in the ROS level versus to each of them alone (p<0.01). Hence, the combination of cisplatin with Ly is the better choice for ROS production in OVCAR3 ovarian cancer cells.



**Fig. 2.** Intracellular detection of reactive oxygen species (ROS) generation after treatment with cisplatin, Ly, and cisplatin+ LY in OVCAR3 cells. Results are presented as a percentage of the control and express the mean±SEM of three independent experiments. \* $p < 0.05$ , \*\*  $p < 0.01$  and \*\*\*  $p < 0.001$  versus control. ##  $p < 0.01$  versus cis (Ctrl: control; Cis: cisplatin, Ly, Cis+Ly: Cisplatin+Ly).

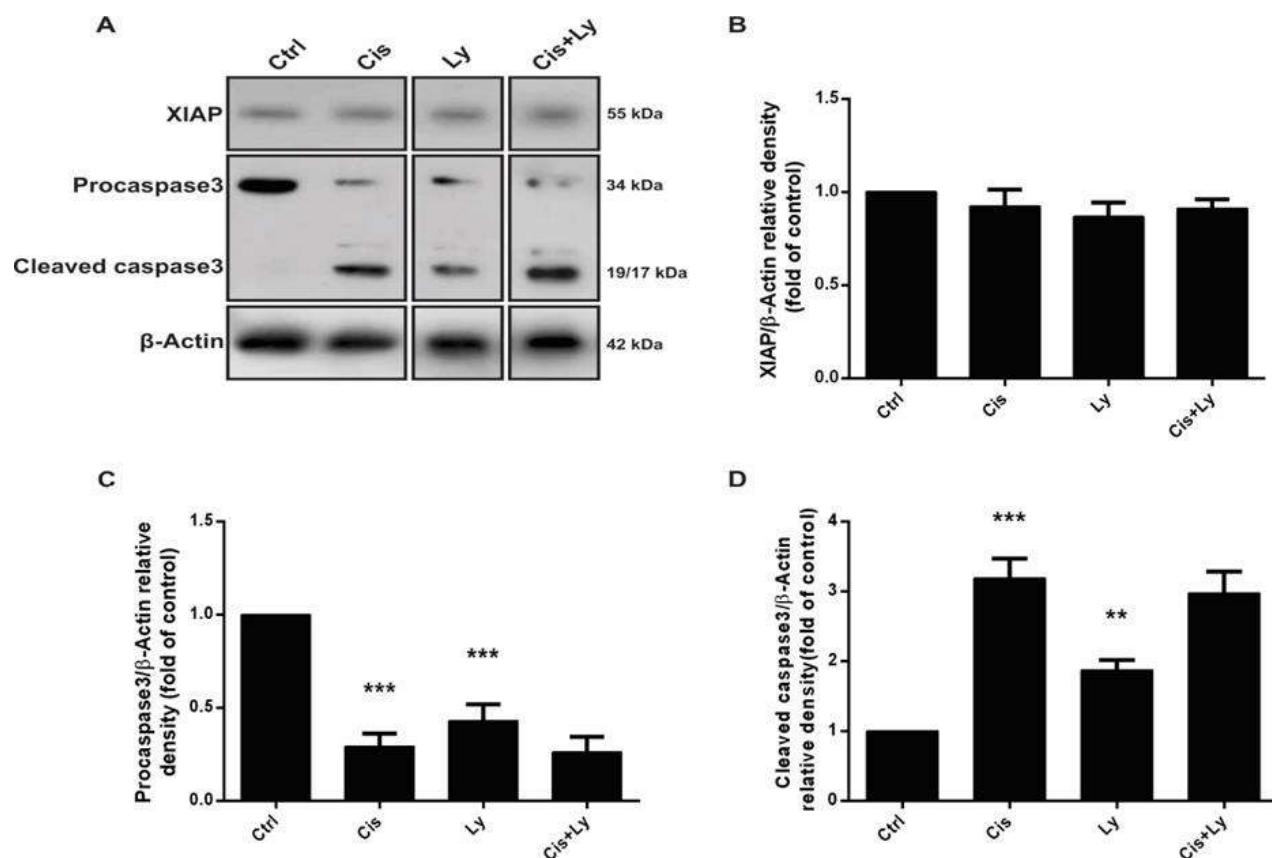
**Apoptosis markers**

To determine the apoptosis induction in experimental conditions, the expression of

apoptosis-related genes was analyzed in treated cells. Based on the obtained results in the western blotting test, treatment with all three

mentioned groups had no detectable effect on XIAP expression, whereas treatment with cisplatin and Ly alleviated caspase 3 activations ( $p < 0.001$ ). However, cisplatin plus Ly showed a considerable decrease in total caspase 3 expressions (cleaved+procaspase3) versus control chemoresistant cells (Fig. 3A). When it comes to changes in activation of caspase 3, treatment with cisplatin+Ly increased the cleaved caspase 3

versus Ly ( $p < 0.05$ ); however, no significant change was detected versus cisplatin ( $p = 128$ ). It seems that the up-regulation of cleavage of caspase 3 protein did not have a remarkable difference in the cisplatin + Ly as compared with cisplatin alone (Fig. 3D). Nevertheless, it is noticeable that approximately half of the concentration of cisplatin has been used in combination conditions to achieve these levels of apoptosis.



**Fig. 3.** The protein expression level of XIAP, caspase 3, and cleaved caspase 3 when they were treated with cisplatin, Ly alone, and their combination in the OVCAR3 cells and examined with western blotting assay. A) Western blotting image of XIAP, procaspase 3, cleaved caspase 3, and  $\beta$ -actin proteins determined via Western blot. Quantification of band densities, B), C), and D) protein expression fold of control of XIAP, pro-caspase 3, and cleaved caspase 3, respectively. The data represent the means $\pm$ SEM of three separate experiments. \* $p < 0.05$ , \*\*  $p < 0.01$  and \*\*\*  $p < 0.001$  versus Ctrl group.

### Akt activity

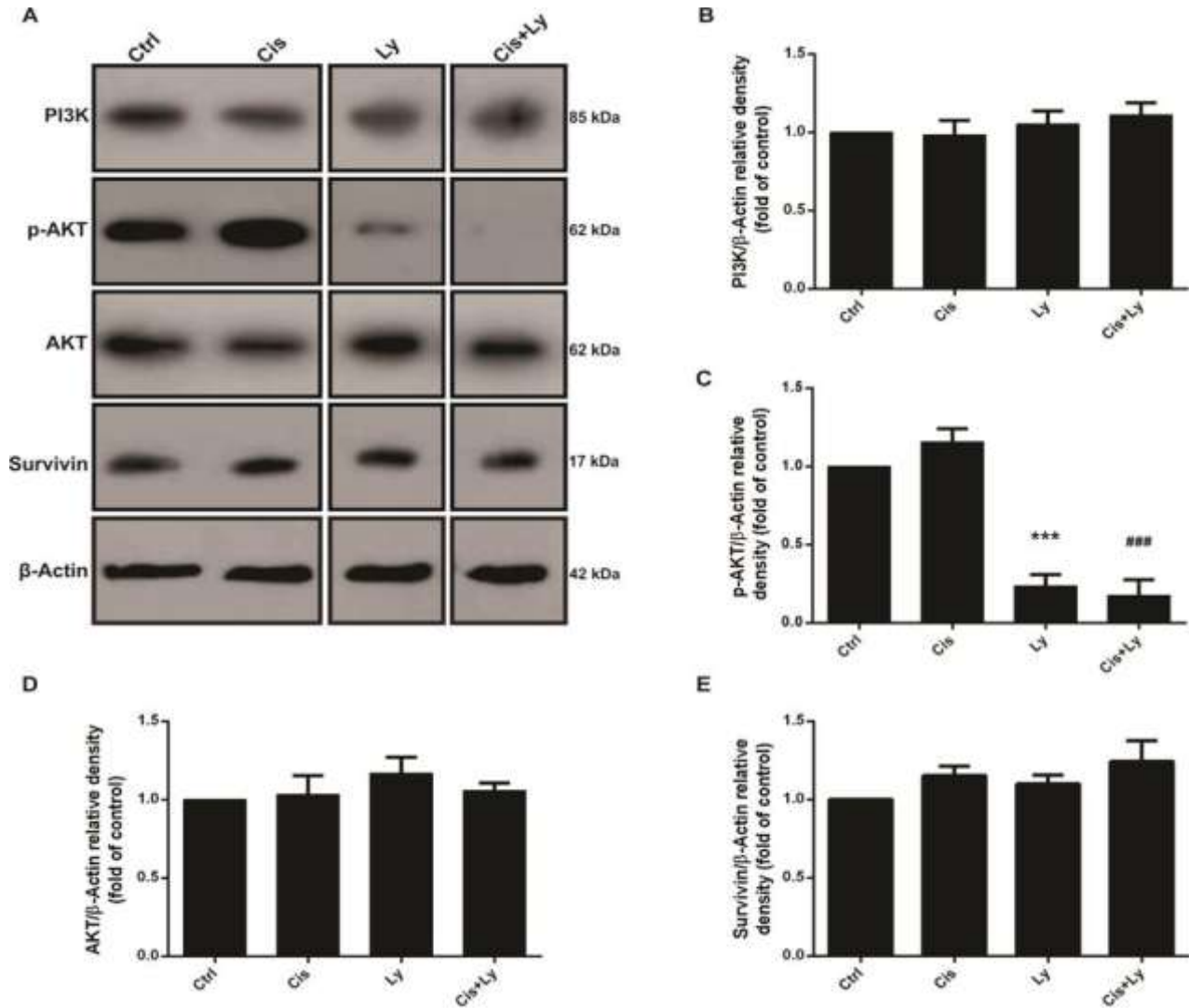
To investigate the effect of cisplatin, Ly and co-treatment of them (cisplatin + Ly) on the PI3K/Akt signaling pathway due to involvement in cisplatin-based chemotherapy resistance, immunoblotting tests were performed to determine the protein expression level of PI3K, p-Akt, Akt, and Survivin in OVCAR3 cancer cells (Fig. 4). As indicated by the obtained results, the expression of PI3K remained

unchanged based on treatment with all the mentioned groups (Fig. 4B). Regarding P-Akt protein content, treatment with cisplatin not only did not reduce the protein expression ratio but also surprisingly, the significant increase was detected in comparison to the control cell ( $p < 0.05$ ). In contrast, Ly ( $p < 0.001$ ) decreased the P-Akt protein level sharply, confirming the strong inhibition of AKT



phosphorylation. A significant decrease in P-AKT levels was detected in cisplatin + Ly versus cisplatin alone ( $p < 0.001$ ). However, the difference between Ly and cisplatin + Ly was insignificant compared with Ly alone

(Fig. 4C). Moreover, treatment with all studied groups neither did not alter the expression level of Akt nor survivin protein significantly in comparison with untreated OVCAR3 cells (Figs. 4D and 4E).



**Fig. 4.** The effect of cisplatin, Ly, and cisplatin plus Ly treatment on the expression level of proteins involved in the PI3K/Akt signaling pathway in the OVCAR3 cells. A) Immunoblotting image of all proteins, including PI3K, p-Akt, Akt, and Survivin. Besides, B), C), D), and E) the expression percentage of PI3K, p-Akt, Akt, and Survivin, respectively. The proteins were analyzed through western blotting assay, and the intensity of proteins band estimated based on the expression of  $\beta$ -actin as a control for protein loading. The experiments were repeated thrice as means $\pm$ SEM (n= 3): \*\*  $p < 0.001$  versus the control group. ###  $p < 0.001$  versus Cis group. (Ctrl: control; Cis: cisplatin, Cis+Ly: Cisplatin + Ly).

### Discussion

AKT/PI3K is a critical biological pathway involved in resistance to standard chemotherapeutic agents (19-21). Therefore, AKT/PI3K inactivation is of crucial importance in tumor prevention based on cytotoxic drugs

and can be considered as one of the primary approaches for estimation of the clinical effectiveness of cancer cell sensitizing agents (22, 23).

Cisplatin is a well-known anti-cancer agent named as cisplatinum or cis-diamminedichloroplatinum (II). Various types of human cancer, including head and neck, ovarian, bladder, lung, and testicular cancers, are being treated with cisplatin. The mechanism of action is related to the crosslinking ability of cisplatin with the purine bases on DNA molecules that led to intervention in DNA repair mechanism and, consequently, DNA damage and apoptosis induction in cancer cells. However, drug resistance, allergic reactions, decrease immunity to infections, gastrointestinal disorders, and hearing loss are serious problems limiting single therapy-based cisplatin. Therefore, co-administration of cisplatin, along with other types of anti-cancer drugs, can be considered as a novel therapy method to overcome drug resistance and reduce adverse side effects (24). LY294002 can reversibly inhibit 3 out of 4 isoforms of the catalytic subunit of class I in PI3K enzyme, p110 $\alpha$ , p110 $\beta$ , p110 $\gamma$ , and p110 $\delta$ . Ly acts on the ATP binding site of the catalytic subunit. The IC50s for p110 $\alpha$ ,  $\beta$ , and  $\delta$  are 500 nM, 973 nM, and 570 nM (2, 25, 26) LY294002 is also a potent autophagy inhibitor by blocking autophagosome formation (5).

In this study, we used combination therapy of Ly as a synthetic inhibitor of PI3K to block PI3K/Akt pathway along with cisplatin as a conventional chemotherapy agent to target multiple molecules in the OVCAR3 cells.

In other words, the main aim of combination is down-regulation of cell survival pathway PI3K/Akt, up-regulation of apoptosis-related genes (caspases 3), and hyperactivation of oxidative stress.

Firstly, we compared the anti-proliferative effect of the combination of cisplatin and Ly with a single administration of cisplatin by MTT assay on the OVCAR3 cancer cell. The cell cytotoxicity results showed that administration of the 5.46 $\mu$ M cisplatin + 5 $\mu$ M Ly (IC50 in OVCAR3) inhibits the proliferation of ovcAR3 cells as much as 13.18 $\mu$ M cisplatin alone. This result suggests that Akt inhibition by Ly, in combination with cisplatin, could reduce the effective dose of cisplatin and

therefore lower the side effects and drug resistance. However, the side effect of the Ly administration should also be investigated in comprehensive examines.

Phosphatidylinositol 3-kinase (PI3K) is an upstream protein of AKT, Phosphorylating Akt in response to various stimuli including protein phosphatase inhibitors, growth factor, stimulations, stress such as oxidation and Ras activation and extracellular matrix molecules (27, 28). PI3K performs a balance between cell survival and apoptosis, along with regulating various essential cellular responses (29, 30). After confirming of approximately complete inhibition of AKT phosphorylation in LY treated cells, Consistent with previous studies, we also found a remarkable decrease in AKT phosphorylation in cancer cells treated with Ly+cisplatin in proportion to cisplatin alone. This data is in line with the results of the MTT assay, which is mentioned above.

As aforementioned, the PI3K/Akt pathway contributes to developing ovarian cancer and induces chemoresistance to cisplatin and inhibits cisplatin-induced apoptosis of ovarian cancer cells (30-32). This study can be evidence of a slight increase in Akt protein level and phosphorylation of AKT in a single treatment of OVCAR3 cancer cells with cisplatin in our experiment. In a study, Lin et al. implied MK-2206 as an inhibitor of the PI3K/Akt pathway in administrated with Taxol or cisplatin to assess the combined effect of mentioned groups. The results depicted that inhibiting PI3K/Akt pathway enhanced the effects of cisplatin and Taxol, highlighting the critical roles of the Akt pathway in reversing chemoresistance in ovarian cancer cells (15).

ROS is a metabolic product of cellular aerobic metabolism that increases in its generation brings about mitochondrial damage, which consequently leads to cell death via apoptotic cascade reaction.

Previous studies have been shown that ROS production leads to apoptosis induction in ovarian cancer cells. Furthermore, ROS, as a crucial regulator of apoptosis, is contributed to a diverse array of pathological and physiological processes, including the



mitochondria-mediated apoptosis pathway. The decrease in P-Akt protein expression in this study could increase the accumulation of intracellular ROS generation. Hence, our results prompt us to speculate that knockdown of PI3K/Akt signaling pathway via the combination treatment of cisplatin with Ly act as a leading factor of apoptosis induction via generation of ROS (17). In line with our obtained results YAMADA et al. depicted that down-regulation of PI3K/Akt signaling pathway via a combination of LY294002 as a synthetic inhibitor of PI3K and novel histone deacetylase (HDAC) inhibitor-induced ROS generation in RCC cancer cells (28).

Caspases are a group of intracellular cysteine proteases that are responsible for the activation of apoptosis signaling cascade. It has been reported that caspases are activated throughout apoptosis in various tumor cells. Cleaved caspase 3 as one of the main molecules is necessary for the onset of apoptosis irreversibly via both extrinsic and intrinsic pathways. Based on the intrinsic apoptotic pathway, cytochrome c release into cytosol and then binds to Apaf-1 and activates caspase-9, followed by caspase-3 activation (33). We showed that treatment with Cisplatin, Ly, and co-administration of them reduced remarkably the content of pro-caspase 3 compared to control groups. In contrast, cleaved caspase 3 significantly increased in all treated cells in comparison to untreated cells, among which the effect of combination therapy was considerable compared to the cisplatin-treated cells. This result shows that the hyper induction of apoptosis coincides with down-regulation of P-Akt and hyperactivation of oxidative stress.

In line with our study, Lou et al. showed that treatment of cancer cells with Quercetin induces apoptosis via increasing the expression level of LC3 and ERK, cytoplasm p53, cleaved Caspase-3, and PARP and down-regulation protein expression of activated PI3K/AKT and Bcl-2 (34).

Previous studies have been shown that ROS production arises out of the down-regulation of Survivin and XIAP, which subsequently lead to apoptosis induction in ovarian cancer cells.

Besides, elevated X-linked IAP (XIAP) expression as a critical member of the inhibitor of apoptosis (IAP) family is of considerable significance in proliferation, cell invasion, and chemoresistance in ovarian cancer (32, 33). Overexpression of XIAP as a most potent caspases (including caspases-3, -7, and -9) inhibitory protein protects cancer cells bearing apoptosis induced with various anti-cancer agents (22, 35) in other hands, Survivin, as the other member of (IAP) family, display multiple functions, including regulation of apoptosis and mitosis. It is reported that the up-regulation of Survivin resulted in resistance of cancer cells toward conventional chemotherapeutic agents such as paclitaxel and cisplatin and, subsequently, prevention of tumor cells from apoptosis (28). Hence, suppression of survivin expression led to drug-induced apoptosis and also is one of the main indexes in opting novel anti-cancer drug. Based on such data we investigated the change in protein expression of XIAP and Survivin (34). Surprisingly we did not detect any significant decrease in the protein expression level of XIAP in three treated groups, including cisplatin, Ly, and LY+cisplatin, compared to control groups. Therefore, the activation of caspase 3 and decrease in cell proliferation and not change in XIAP and surviving expression promote us to assume that these molecules (XIAP and survivin) have been inactivated via an unknown way which deserves to investigate in future.

Overall, our obtained results displayed that co-treatment with cisplatin and Ly could sensitize the drug-resistant ovcAR3 cancer cell line to cisplatin that is partially mediated by hyperactivation of oxidative stress.

Taken together, the results of this study depicted that the Ly could decrease the cisplatin-resistance in OVCAR3 cancer cells through the induction of apoptosis. It is noteworthy that the induction of apoptosis in combination therapy may be the result of an increase in ROS generation in OVCAR3 cancer cells.

### Acknowledgements

We would like to give a special thanks to all individuals that have helped us in present study.

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