Nedaplatin sensitization of cisplatin-resistant human non-small cell lung cancer cells

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Received January 5, 2015; Accepted February 4, 2016

DOI: 10.3892/ol.2016.4276

Abstract. Cisplatin (DDP) has been one of the most widely used chemotherapy drugs for advanced non-small cell lung cancer. However, the increase in the number of DDP-resistant cancer cells has become a major impediment in the clinical management of cancer. In the present study, for the first time, the 3-(4,5-dimethylthiazol-2-y1)-2,5-diphenyltetrazolium bromide assay was used to demonstrate that nedaplatin (NDP) could have a stronger inhibitory effect than DDP alone in DDP-resistant A549 (A549DDP) cells and that it could attenuate the resistance of these cells. Additionally, flow cytometry analysis showed that the apoptosis rate of these resistant cells when exposed to NDP was markedly increased and the number of cells in the G2 stage of the cell cycle was significantly increased. Furthermore, western blot analysis indicated that NDP decreased the protein expression of P-glycoprotein, tumor protein p53 and B-cell lymphoma 2, and increased the expression of Bcl-2-associated X protein, all of which could possibly improve the NDP intracellular drug concentration and promote cell apoptosis. These observations suggested that NDP could have higher efficacy in DDP-resistant lung cancer cells, and further studies applying more detailed analyses are warranted to elucidate the mechanism(s) behind this effect.

Introduction

Lung cancer is the leading cause of cancer-associated mortality worldwide, and ~85% of lung cancer diagnoses are of non-small cell lung cancer (NSCLC) (1,2). Although great progress has been made in small molecular-targeted drugs for treating NSCLC, particularly epidermal growth factor receptor (EGFR) tyrosine kinase inhibitors such as gefitinib and erlotinib, EGFR mutations are detected in only 10% of NSCLC patients in the United States and in 35% of NSCLC patients in East Asia. Thus, platinum-based combination chemotherapies remain the mainstay of advanced NSCLC treatment, and cisplatin (DDP) is widely used in clinical therapy (3). However, the overall 5-year survival rate for lung cancer is ~15%, and this rate has improved only slightly over the last 30 years despite the advancement of modern chemotherapy, a problem which is mainly caused by drug resistance to platinum (4).

The problem of resistance to DDP-based chemotherapy remains one of the major obstacles to the treatment of lung cancer. A number of mechanisms have been proposed to explain cancer cell resistance to chemotherapy (5,6). These mechanisms generally involve an increase in the level of multidrug resistance-1/P-glycoprotein (P-gp) (7,8), and the regulation of apoptosis-related genes and proteins such as tumor protein p53 (p53) and B-cell lymphoma 2 (Bcl-2) family members (9-11). However the underlying mechanisms are not yet fully understood. Thus, there is an urgent requirement to learn how to improve the efficacy of platinum or to identify a novel generation of platinum agents.

Nedaplatin (NDP), which is a second-generation DDP analog, was developed by the Shionogi Pharmaceutical Company (Japan) in 1983, in order to provide a treatment with a level of effectiveness similar to that of DDP, but with decreased gastrointestinal and renal toxicities (12). A number of previous clinical studies have demonstrated the efficacy of NDP to be higher than that of DDP in patients with DDP-resistant lung cancer (13,14). Conversely, there are comparatively few in vitro studies to support the consensus. Moreover, it is unclear why NDP is not completely cross-resistant with DDP.

The purpose of the present study was to demonstrate the efficacy of NDP in DDP-resistant A549 (A549DDP) cells in vitro directly. Moreover, the study aimed to detect the expression of DDP resistance-related proteins, such as P-gp, p53, Bcl-2-associated X protein (Bax) and Bcl-2, to investigate the possible mechanisms behind NDP efficacy in the A549DDP cells.

Materials and methods

Cell culture. The human NSCLC A549 cell line and the human DDP-resistant cell strain, A549DDP, were used in this study. The cells were obtained from Shanghai Cell Biology, an Institute of the Chinese Academy of Sciences (Shanghai, China).
The A549 cell line was cultured in RPMI-1640 medium supplemented with 100 U/ml penicillin, 100 mg/ml streptomycin and 10% fetal bovine serum (Invitrogen; Thermo Fisher Scientific, Inc., Waltham, MA, USA). The A549DDP cells were cultured in high glucose Dulbecco's modified Eagle's medium supplemented with 100 U/ml penicillin, 100 mg/ml streptomycin and 10% fetal bovine serum; 2 µg/ml DDP (Jiangsu Haosen Pharmaceutical Co., Ltd., Lianyungang, China) was dissolved into this solution in order to maintain drug resistance. However, the A549DDP cells were grown in the absence of DDP 2 days prior to treatment. These cells were incubated in a standard cell culture incubator (Series 8000 Water-Jacketed CO2 Incubator; Thermo Fisher Scientific, Inc.) at 37°C with 5% CO2, and passaged once or twice a week. Cells in the algorithm growth phase were used in the following experiments.

Cell proliferation and the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay. The half maximal inhibitory concentration (IC50) of the A549DDP and A549 cells was determined by the MTT assay (Beyotime, Shanghai, China). The A549 and A549DDP cells were seeded into 96-well plates (1x103-1x105 cells per well), and treated with DDP and NDP (Jiangsu Aosaikang, Nanjing, China) at different concentrations (A549 cells: 2, 4, 6, 8 and 10 µg/ml; A549DDP cells: 10, 15, 20, 25 30 µg/ml) for 48 h. Following incubation, 5 µl MTT (20 µl/well) was added to the media and the cells were further incubated in an atmosphere of 5% CO2, at 37°C for 4 h. Dimethylsulfoxide (150 µl; Sigma-Aldrich, St. Louis, MO, USA) was added to the cells in each of the wells after the media was removed, and the cells were further incubated for 10 min. The optical density (OD) of each well was measured using a microplate reader (Multiskan™ GO Microplate Spectrophotometer; Thermo Fisher Scientific, Inc.) at 560 nm. All experiments were performed in triplicate according to the following formula: Cell inhibitory rate (%) = (1 - OD of test group / OD of control group) x 100.

Apoptosis detection and cell cycle analysis. The rate of apoptosis induced by the anticancer regimens was analyzed by flow cytometry using an annexin V-fluorescein isothiocyanate/propidium iodide kit (Kaijibio, Nanjing, China). Adherent and floating cells were harvested and gently disaggregated to a single-cell suspension. Staining was performed according to the manufacturer's protocols. The data were analyzed immediately by flow cytometry using CXP software (Beckman Coulter, Inc., Brea, CA, USA).

Protein isolation and western blot analysis. Subsequent to exposure to DDP and NDP for 48 h, cell protein extracts were determined using 500 µl radioimmunoprecipitation assay lysis buffer with 5 µl phenylmethylsulfonyl fluoride and protease inhibitor (Abcam, Cambridge, UK). Total proteins were quantified using the bicinchoninic acid assay (Beyotime, Shanghai, China) according to the manufacturer's protocols. Total protein (20 µg) was loaded onto an 8% sodium dodecyl sulfate-polyacrylamide gel and transferred to a polyvinylidene difluoride membrane (EMD Millipore, Billerica, MA, USA). The membrane was incubated for 1 h at 25°C in 5% (w/v) skimmed dried milk and then washed three times for 5 min at 25°C using blocking buffer [Tris-buffered saline with Tween 20 (TBST) buffer: 0.1% Tween 20, 13.7 mM NaCl, 0.27 mM KCl and 2.4 mM Tris)]. Next, the membrane was incubated overnight at 4°C with monoclonal mouse anti-human primary antibodies for P-gp (clone, JSB-1; catalog no., ab3366), p53 (clone, PAb 1801; catalog no., ab28), Bax (clone, 2D2; catalog no., ab77566) and Bcl-2 (clone, Bcl2/100; catalog no., ab117115). All antibodies were diluted to 1:1,000 and purchased from Abcam. Subsequent to being washed three times with TBST for 5 min each, the membranes were incubated for 1 h with horseradish peroxidase-conjugated secondary antibodies (goat anti-rabbit IgG; 1:5,000; catalog no., sc-2004; Santa Cruz Biotechnology Inc., Dallas, TX, USA). In order to evaluate of protein expression accurately, β-actin (mouse monoclonal; clone, AC-15; catalog no., ab6276; Abcam) and histone H3 protein (mouse monoclonal; clone, mAbcam 1220; catalog no., ab1220; Abcam) were used as an internal standard. Band intensity was analyzed with an imaging and analysis system (Peijing JS-780; Hai Pei Qing Technology Co., Ltd., Shanghai, China), and protein expression was presented as the ratio of the protein band intensity to β-actin or Histone H3 in the same blot.

Statistical analysis. The values presented represent the mean ± standard deviation calculated from the data. All analyses were performed using the Statistical Package for Social Sciences, version 13 (SPSS Inc., Chicago, IL, USA). Differences were evaluated using Student's t-test or an analysis of variance.

Results

Cell inhibitory measurement by MTT assay. An MTT assay was used to determine the sensitivity of A549DDP cells to DDP, to investigate whether these cells are resistant to DDP...
and to determine whether NDP has a stronger inhibitory effect than DDP in A549DDP cells.

The inhibition rate is shown in Figs. 1 and 2, and Tables I and II. For the first part of the MTT assay, the IC\textsubscript{50} values of the A549 and A549DDP cells treated with DDP were 2.53±0.12 and 23.36±1.41 µg/ml, respectively, and the difference between them was significant (P<0.001), which verified that A549DDP cells exhibit resistance to DDP.

In the second part, of the MTT assay, the IC\textsubscript{50} of the A549 cells treated with DDP and NDP was 2.53±0.12 and 2.49±0.78 µg/ml (Fig. 1; Table I), and the difference was not significant (P=0.834). However, the IC\textsubscript{50} values of the A549DDP cells treated with DDP and NDP were 23.36±1.41 and 19.97±0.88 µg/ml (Fig. 2; Table II); this difference was significant, suggesting that NDP had a better effect than DDP on the A549DDP cells.

Cell apoptosis and the cell cycle, as shown by flow cytometry. Flow cytometry was used to investigate the differences in cell apoptosis and the cell cycle between the A549DDP cells treated with DDP and NDP. After 48 h of intervention with DDP (20 µg/ml) and NDP (20 µg/ml), the levels of cell apoptosis in the NDP and DDP groups increased and were significant when compared with the control group (each P<0.01). In comparison with the DDP group, the degree of early and late apoptosis in the NDP group increased, and this difference was significant (P<0.05). DDP, cisplatin; NDP, nedaplatin.

Table I. Inhibition A549 cells after 48 h of intervention with DDP and NDP at different concentrations.

<table>
<thead>
<tr>
<th>Drug concentration, µg/ml</th>
<th>Group</th>
<th>2</th>
<th>4</th>
<th>6</th>
<th>8</th>
<th>10</th>
<th>IC\textsubscript{50}</th>
</tr>
</thead>
<tbody>
<tr>
<td>DDP</td>
<td></td>
<td>42.78±2.50</td>
<td>63.21±2.73</td>
<td>66.79±1.76</td>
<td>72.06±2.83</td>
<td>73.68±2.51</td>
<td>2.53±0.12</td>
</tr>
<tr>
<td>NDP</td>
<td></td>
<td>44.84±2.32</td>
<td>62.34±1.97</td>
<td>67.28±1.96</td>
<td>73.56±2.63</td>
<td>76.88±3.04</td>
<td>2.49±0.78</td>
</tr>
<tr>
<td>P-value</td>
<td></td>
<td>0.354</td>
<td>0.677</td>
<td>0.765</td>
<td>0.537</td>
<td>0.232</td>
<td>0.834</td>
</tr>
</tbody>
</table>

Table II. Inhibition of A549DDP cells after 48 h of intervention with DDP and NDP at different concentrations.

<table>
<thead>
<tr>
<th>Drug concentration, µg/ml</th>
<th>Group</th>
<th>10</th>
<th>15</th>
<th>20</th>
<th>25</th>
<th>30</th>
<th>IC\textsubscript{50}</th>
</tr>
</thead>
<tbody>
<tr>
<td>DDP</td>
<td></td>
<td>24.79±1.53</td>
<td>36.28±2.85</td>
<td>43.25±2.04</td>
<td>50.38±1.95</td>
<td>60.54±1.66</td>
<td>23.36±1.41</td>
</tr>
<tr>
<td>NDP</td>
<td></td>
<td>27.74±1.48</td>
<td>39.20±2.91</td>
<td>49.93±3.22</td>
<td>59.05±2.56</td>
<td>63.21±1.93</td>
<td>19.97±0.88</td>
</tr>
<tr>
<td>P-value</td>
<td></td>
<td>0.074</td>
<td>0.283</td>
<td>0.038\textsuperscript{a}</td>
<td>0.009\textsuperscript{a}</td>
<td>0.144</td>
<td>0.024\textsuperscript{a}</td>
</tr>
</tbody>
</table>

Table III. Cell apoptosis rate induced by DDP and NDP, as detected by flow cytometry.

<table>
<thead>
<tr>
<th>Group</th>
<th>Q1</th>
<th>Q2</th>
<th>Q3</th>
<th>Q4</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control, %</td>
<td>3.49±1.74</td>
<td>2.60±0.47</td>
<td>92.78±1.69</td>
<td>1.11±0.17</td>
</tr>
<tr>
<td>DDP, %</td>
<td>4.09±3.35</td>
<td>3.63±2.06</td>
<td>83.81±5.55</td>
<td>8.47±1.54</td>
</tr>
<tr>
<td>NDP, %</td>
<td>9.64±3.75</td>
<td>11.80±1.50</td>
<td>60.97±6.70</td>
<td>17.59±2.81</td>
</tr>
</tbody>
</table>

Q1 represents necrotic cells, Q2 represents late apoptotic cells, Q3 represents living cells and Q4 represents early apoptotic cells. DDP, cisplatin; NDP, nedaplatin.

Table IV. Detection of DDP- and NDP-induced cell cycle arrest by flow cytometry.

<table>
<thead>
<tr>
<th>Group</th>
<th>G1</th>
<th>S</th>
<th>G2</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control, %</td>
<td>68.04±3.50</td>
<td>15.81±3.04</td>
<td>14.44±2.59</td>
</tr>
<tr>
<td>DDP, %</td>
<td>54.53±5.84</td>
<td>19.73±6.93</td>
<td>25.73±5.84</td>
</tr>
<tr>
<td>NDP, %</td>
<td>51.52±8.59</td>
<td>17.34±5.39</td>
<td>31.47±4.76</td>
</tr>
</tbody>
</table>

DDP, cisplatin; NDP, nedaplatin.
the proportion in the G1 stage decreased (P=0.023 and P=0.031, respectively) following intervention with DDP and NDP, and the difference was significant (Fig. 4; Table IV). Furthermore the difference in the percentage of cells in the G2 phase between the DDP and NDP groups was not significant (P=0.228).

Protein expression and western blot analysis. Western blotting results revealed that the expression levels of P-gp, p53, Bax and Bcl-2 in the NDP group were different from the levels of expression in the control and DDP groups.

P-gp relative expression in the NDP group was 0.50±0.03, which was significantly higher than that of the control (0.80±0.12; P=0.015) and DDP (0.63±0.05; P=0.014) groups (Fig. 5). However, compared with the control group, the expression of P-gp in the DDP group was not significantly different (P=0.094).
The relative expression levels of p53 in the control, DDP and NDP groups were 0.77±0.05, 0.62±0.06 and 0.45±0.05, respectively. p53 expression in the NDP and DDP groups was significantly less than in the control group (P=0.002 and P=0.020, respectively) (Fig. 5). Compared with the DDP group, the NDP group exhibited a lower p53 expression level, and the difference was significant (P=0.020).

The relative expression of Bax in the NDP group was 0.57±0.04, which was significantly higher than that of the control (0.39±0.06; P=0.012) and DDP (0.46±0.03; P=0.024) groups (Fig. 5). By contrast, the Bax expression in the DDP group, when compared to the control group, did not exhibit a significant difference (P=0.112).

However, the western blot analysis revealed that the relative expression of Bcl-2 in the control, DDP and groups was 0.80±0.10, 0.60±0.04 and 0.45±0.04, respectively. Bcl-2 expression in the NDP and DDP groups was significantly lower than that in the control group (P=0.005 and P=0.005, respectively) (Fig. 5). Compared with the DDP group, the NDP group exhibited lower Bcl-2 expression, and the difference was also significant (P=0.008).

Discussion

The present study results showed that at the same concentration, NDP had a higher cell inhibition rate than DDP in A549DDP cells, particularly for concentrations of 20 and 25 µg/ml (P=0.038 and P=0.009, respectively). It was also found that the blockage of the cell cycle at the G2 phase in the A549DDP cells increased significantly following intervention with DDP and NDP, but that the difference between these two groups was not significant. However, compared with the DDP group, the NDP group exhibited significantly greater early and late apoptosis ratios. Therefore, it was concluded that the use of NDP was more advantageous than the use of DDP in A549DDP cells. In this study, the results also showed that the expression levels of P-gp, p53 and Bcl-2 in the NDP group were significantly lower than those of the other two groups, and that the expression of Bax in the NDP group was significantly higher. Moreover, it was found that the difference in Bcl-2 expression between the NDP and DDP groups was more pronounced (P=0.008).

P-gp is one of the major drug efflux transporters; it increases the efflux of drugs out of cells against the concentration gradient, thereby reducing the intracellular concentration of the drug below the effective level, which finally leads to drug resistance (8,15). It has been verified that a number of anticancer drugs, including DDP, etoposide and vinblastine, are P-gp substrates (16). The results of the present study showed that the P-gp expression level of cells exposed to NDP was significantly lower than that of cells exposed to DDP. Thus, it was concluded that NDP could possibly inhibit the P-gp expression level, thereby decreasing the efflux of drugs out of the A549DDP cells. This would improve the NDP intracellular drug concentration, which would suppress A549DDP cell proliferation.

Lung cancer cells have been shown to possess a higher p53 mutation rate (70%); the mutation of the gene could result in abnormal expression of the p53 protein (17). The wild-type p53 protein is able to exert a range of anti-proliferative effects, including the induction of apoptosis and causing a marked increase in the sensitivity of these cells to DDP (18,19). However, malignancies with mutated p53 genes and aberrant p53 proteins in laboratory studies and one clinical study have been observed to be less responsive to chemotherapy agents that induce DNA damage, such as DDP (20,21). A number of studies have suggested that the overexpression of the mutant p53 protein may directly enhance tumor cell resistance to
anticancer agents in a way that is dependent on the particular mutation and the mechanism of action of the drug (22-24). In a situation of cellular stress, such as DNA damage, the mutated p53 genes and aberrant p53 proteins participate in the process of inducing cell-cycle arrest, and can enhance DNA repair or cell death and upregulate the expression of P-gp (25,26).

In the present study, compared with DDP intervention, NDP intervention led to a significant downregulation of p53 protein. Combined with the greater downregulation of P-gp protein following NDP intervention, this results indicates that the mutant p53 protein was likely detected in the A549DDP cells, and that NDP could inhibit the expression of the mutant p53 protein, thereby decreasing the upregulation of P-gp expression in order to withstand DDP resistance.

There are numerous members in the Bcl-2 family, and while certain members, such as Bcl-2, are anti-apoptotic, other, such as Bax, are pro-apoptotic. The ratio between pro- and anti-apoptotic Bcl-2 family members is a significant determinant of cell survival and cell death. A number of cancer chemotherapeutic agents ultimately act on these factors causing cells to undergo apoptosis (27). The Bcl-2 family plays a significant role in the cellular response to chemotherapy. The overexpression of Bcl-2 increases the resistance to drug-induced apoptosis, and the survival of Bcl-2-negative tumors is less than that of Bcl-2-positive tumors (28,29). In the present study, it was found that the Bcl-2 expression level of NDP-exposed cells was significantly lower than that in cells exposed to DDP, while the Bax expression level in the NDP-exposed cells was significantly greater. It was concluded that NDP could regulate Bcl-2 and Bax expression, thereby promoting the apoptosis of A549DDP cells by allowing NDP to withstand DDP resistance.

Clinically, an association has been observed between NDP and an improved response in DDP-resistant cancer (13,30,31). In summary, the present study suggested that NDP could improve the apoptotic function of A549DDP cells, and an improved response in DDP-resistant cancer (13,30,31).

**References**


