Detection of Five Differentially Expressed Proteins of Capsulation-resistant and Sensitive Osteosarcoma Tissues by Western Blot Technique in Vitro

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Abstract. The tissue samples of cisplatin-resistant and sensitive patients with human osteosarcoma were obtained through chemosensitivity testing before. Five typical differentially expressed proteins of cisplatin-resistant and sensitive osteosarcoma tissue samples were analyzed by two-dimensional electrophoresis and the relevant differentially expressed proteins were obtained before. By Western Blot technique in vitro, Two three-representative-tissues on cisplatin-resistant and sensitive osteosarcoma were chosen to test which is the differently expressed of 5 proteins. It is concluded that ALDOA and PGK1 may be the right markers in cisplatin chems in trating human osteosarcoma.

Introduction

The treatment of tumors resistant to cytotoxic drugs is not only an important reason for treatment failure, but also an important factor to limit the chemotherapy \cite{1-4}. The mechanism is complex, it is determined by the character of tumor, such as the proportion of viable cells, the adequacy of the blood supply, the specific cellular mechanisms and MDR phenotype \cite{2-4}. Among them, multi drug resistance (MDR) is one of the main obstacles in clinical cancer chemotherapy \cite{5,6}. It is a particular phenomenon produced by tumor cells resistant. Its characteristics are that once the cells resistant to certain drugs generated, the drugs with other different structures and mechanisms of action produce cross-resistance phenomenon. In the past 10 years, many scholars dedicated to research of multidrug resistance in tumor \cite{7-9}.

Cisplatin can attain its effect of chemotherapy by inducting osteosarcoma cell autophagy activation \cite{10-12}. Because of this, it is one of the three main drugs osteosarcoma neoadjuvant chemotherapy Rosen T Series program \cite{13}. From the experiments completed for the first \cite{14} and second step \cite{15}, it has been verified that cisplatin is a rational drug for ongoing study of multidrug resistance. Cisplatin-resistant and sensitive osteosarcoma tissue in the first step experiment is selected, and a typical one of cisplatin-resistant and sensitive osteosarcoma tissue samples were analyzed by two-dimensional electrophoresis and the relevant differentially expressed proteins were obtained. Five differentially expressed proteins were obtained through the analysis of mass spectrometry technology and identified comparing with protein databases. This experiment is the third step of the multi-drug resistance study.

Experimental methods

General Material. Choose two three-representative-tissue on cisplatin-resistant and sensitive osteosarcoma.

Reagents. Stock solution and working solution preparation.

Tissue protein extraction

1. Tissue blocks were weighed. 2. Using liquid nitrogen, pulverized tissue blocks in a mortar. 3 Add RIPA buffer (per gram of tissue 3 ml RIPA), PMSF (per gram of tissue 30μl, 10 mg / ml PMSF), further homogenized using Polytron (15,000 rpm / min 1 min) and maintained 4°C.4 Add PMSF (per gram of tissue 30μl, 10 mg / ml PMSF), incubated on ice for 30 minutes. 5. Ingraft into
centrifuge tubes 4°C approximately 20,000 g (15,000 rpm) for 15 minutes. 6. Supernatant of cell lysates, divided into equipment stored at -20 °C. 7. Bradford colorimetric assay for protein concentration.

**SDS-polyacrylamide gel electrophoresis**

1. Installed shelves. 2. Separation gel prepared according to the following formula (Total: 8ml) Adding a layer of distilled water in the gel above to promote better glue agglutination. 3. Preparation of concentrated gel after separating gel set. (Total: 3.5ml) Insert a pre-prepared comb then. 4. After gelling be set, sample, electrophoresis. Upper gum set with 60-80V voltage, when the sample to separation gel, set 100-120V. General electrophoresis was 1.5 hours.

Adding a layer of distilled water in the gel above to promote better glue agglutination.

**Electrotransfer (semidrying process)**

1. experiment condition selection. Currently 1mA-2mA/cm2 was used. We usually use 100mA / film, to choose transfer time according to the size of the protein molecule and gum concentration, and adjustments could be taken specifically based on the actual appropriate. 2. Experimental operation (1) The filter paper and membrane preparation (2) transfusion

**Block** Before the end of the transferation prepared 5% Milk (TBST solution). Transferred into the milk in the film after the end of block (be sure to put a clean container to avoid contamination and should be sufficient to cover the film), tidy and clean filter paper used for the next use. Block 4 °C O / N, or RT 1 hr.

**No 1 antibody incubation** The diluted antibody and membrane were incubated. Generally used RT 1 hr. According to the volume and membrane antigen antibody appropriate to extend or shorten the time. Ablution Quick first washed three times with TBST, wash out the milk as soon as possible. Then 5mins * 5. Washing is to wash away the first antibody and antigen non-specific binding. The effect of washing directly affect the results of the background shades. **No 2 antibody incubation** RT incubation 1 hours. Generally used the HRP-conjugated secondary antibody. 1:5000 dilution. Ablution Quick first washed three times with TBST, wash out the milk as soon as possible. Then 5mins for 5 times. Color (HRP enzyme) by enhanced chemiluminescence (ECL).

**Results**

**2-D electrophoresis** We cisplatin-sensitive three osteosarcoma cells as well as three resistant cells described above five related proteins detected. The results show ALDOA and PGK1 kinases expression of the two groups was significant difference, while the other three proteins there was no significant difference. The results shown in Figure 1.

![Fig.1 Five related proteins of osteosarcoma cell from western blot results.](image)
1,2,3: cisplatin-sensitive osteosarcoma cells; 4,5,6: cisplatin-resistant osteosarcoma cells.

Table 1. Determination of protein expression results gray value (gray value detecting protein / protein β-ACTIN internal reference grayscale value) *p<0.05,n=3

<table>
<thead>
<tr>
<th></th>
<th>HMGB1</th>
<th>ALDOA</th>
<th>PGK1</th>
<th>ENO1</th>
<th>GAPDH</th>
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</thead>
<tbody>
<tr>
<td>cisplatin-sensitive</td>
<td>0.91±0.02</td>
<td>0.56±0.12</td>
<td>0.78±0.26</td>
<td>0.48±0.05</td>
<td>2.56±0.23</td>
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<tr>
<td>osteosarcoma cells</td>
<td></td>
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<tr>
<td>cisplatin-resistant</td>
<td>1.02±0.023</td>
<td>1.45±0.15*</td>
<td>1.36±0.37*</td>
<td>0.52±0.09</td>
<td>2.79±0.33</td>
</tr>
<tr>
<td>osteosarcoma cells</td>
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Conclusion

Expression of ALDOA and PGK1 was significant different in cisplatin’s treating human osteosarcoma with statistical significance. The two proteins may be the right marker in cisplatin chems in treating osteosarcoma.

References


