Proteomic identification of differentially expressed proteins associated with the multiple drug resistance in methotrexate-resistant human breast cancer cells

SIYING CHEN1, JIANGXIA CAI1, WEIPENG ZHANG1, XIAOWEI ZHENG1, SASA HU1, JUN LU1, JIANFENG XING2 and YALIN DONG1

1Department of Pharmacy, The First Affiliated Hospital of Medical College, 2Department of Pharmacy, College of Medicine, Xi’an Jiaotong University, Xi’an, Shaanxi 710061, P.R. China

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Abstract. Methotrexate (MTX), as a chemotherapeutic drug, is widely used in the therapy of several cancer types. The efficiency of drug treatment is compromised by the appearance of multidrug resistance (MDR), and the underlying molecular mechanisms remain incompletely understood. We investigated the mechanism of MDR in the MTX-induced breast cancer MCF-7 cells (MCF-7/MTX) using proteomic analysis. MCF-7 drug-sensitive cells (MCF-7/S) were exposed in progressively increasing concentrations of MTX to establish the drug-resistant cell line MCF-7/MTX. The biological characteristics of the cells were analyzed by MTT, flow cytometry, quantitative PCR, western blotting and the global protein profiles of MCF-7/MTX and MCF-7/S were compared using a proteomic approach. The resistance factor of MCF-7/MTX cells was 64, and it possessed significant MDR. Seventeen differentially expressed proteins between MCF-7/MTX and MCF-7/S cells were identified, seven proteins were upregulated and 10 proteins were downregulated in MCF-7/MTX cells. We verified that the protein levels of nucleophosmin (NPM), α-enolase (ENO1) and vimentin (VIM) were upregulated, and heterogeneous nuclear ribonucleoprotein (hnRNP C1/C2), phosphoglycerate mutase 1 (PGAM1) and proteasome subunit α type-2 (PSMA2) were downregulated in MCF-7/MTX cells. The mRNA levels of NPM, VIM, hnRNP C1/C2, PGAM1 and PSMA2 were consistent with the protein expressions, but the gene expression of ENO1 was slightly downregulated. Surprisingly, knockdown of NPM by siRNA sensitized MCF-7/MTX cells to MTX and attenuated the multidrug resistance. The proteins identified, particularly NPM provides new insights into the mechanism of MDR and is expected to become a crucial molecular target for breast cancer treatment.

Introduction

Breast carcinoma is a common malignant tumor among women, the incidence and mortality rates continue to rise rapidly worldwide along with lifestyle changes in diet and environmental factors (1). In Asian countries, the incidence of breast cancer increase doubled or even tripled in the past few decades, similarly, in China ~20-30% of the new cases of breast cancer occurred in the past decade (2). Currently, most breast cancer patients receive chemotherapy, but the long-term stimulation of cytotoxic agents accelerate the occurrence of severe acquired resistance, especially multidrug resistance (MDR) has become an important cause of tumor recurrence after surgery and chemotherapy has failed (3). However, the molecular mechanisms of MDR are rather complicated, it involves in increased drug efflux, drug metabolic biotransformation and alteration of repair ability for DNA damage (4). Therefore, investigating the other potential mechanisms and screening the molecular targets of drug resistance have become key factors in the study of MDR of breast cancer.

Methotrexate (MTX), an antifolate antineoplastic, mainly prevents tumor cells growth and proliferation through potent inhibition of dihydrofolate reductase (DHFR) for purine and pyrimidine nucleotide biosynthesis and blockage of DNA replication. As a classical anticancer drug, it possesses strong efficacy and low price, which is widely used for the treatment of human leukemia, lymphoma, breast cancer, head and neck cancers and other solid tumors, both alone and in combination with other chemotherapeutics (5). Although antifolates are important and effective components of different chemotherapeutic regimens currently used for various cancer therapies, drug resistance frequently arises from the long-term use for treatment in clinic, and the molecular mechanism of antifolate-resistance in cancer cells is believed to be a multifactorial process, it relates to the loss of folate carrier (RFC) function, overexpression and mutation of DHFR and thymidylate synthase (TS), increased ATP-driven MDR efflux
transports, decreased polyglutamate synthase, the high expression of γ-glutamyl hydroxide and tetrahydrofolic acid (THF) (6). However, the MTX resistance is still an extremely complex process, and the mechanism of action may involve changes in a variety of pathways and expressions of different resistant proteins.

With the aim of gaining further insight into the molecular mechanisms of MDR and to discover new biological targets of MTX, the functional proteomic analysis using two-dimensional gel electrophoresis (2-DE) and matrix-assisted laser desorption ionization time of flight mass spectrometry (MALDI-TOF-MS) were performed to identify the differential protein profiles between the MTX-resistant breast cancer cell line MCF-7/MTX and drug sensitive cell line MCF-7/S.

Materials and methods

Establishment of MCF-7/MTX cell line. The parental human breast cancer cell line MCF-7 was obtained from the Chinese Academy of Science (Shanghai, China). These cells were cultured in RPMI-1640 medium containing 10% fetal bovine serum (Gibco), 1% penicillin and streptomycin, at 37°C in a humidified atmosphere of 5% CO₂. The MTX-resistant cell line MCF-7/MTX was established from its sensitive cell line MCF-7/S by progressively increasing methotrexate (Shanxi Pude Pharmaceutical, China) over a period of 12 months. The culture condition was the same as the MCF-7/S cell line. The drug concentration was increased by 1.5-fold at each step of resistance, from 4 to 220 nM. The cells were cultured in MTX of each step for at least 2 weeks before each experiment.

Drug sensitivity assay. The MTT assay (Sigma) was used to measure the drug sensitivity. Cells were seeded at a density of 4x10⁴ cells per well in 96-well plate with each well containing 100 µl medium. The cells were grown for 24 h, then the culture medium was replaced with 200 µl fresh medium containing MTX at different concentrations for 72 h. The medium was removed and added with 180 µl of RPMI-1640 and 20 µl MTT (5 mg MTT/ml) and incubated for 4 h. Following medium removal, 150 µl of DMSO was added to solubilize the formazan. The absorbance was read at 490 nm on a microplate reader (BioTek ELx808, USA). The 50% growth inhibitory concentration (IC₅₀) of drug was determined to evaluate the drug sensitivity. Resistance factor (RF) was calculated by the ratio of the IC₅₀ values of MCF-7/MTX to MCF-7/S cells.

MDR detection. The sensitivity of MCF-7/S and MCF-7/MTX cells to doxorubicin (Zhejiang Haizheng Pharmaceutical, China), paclitaxel (Nanjing Sike Pharmaceutical, China), 5-fluorouracil (Shanghai Xudonghaipu Pharmaceutical, China), mitoxantrone (Shandong Luxin Pharmaceutical, China), vinorelbine (Hangzhou Minsheng Pharmaceutical, China), gemcitabine (Jiangsu Haosen Pharmaceutical, China), cisplatin, docetaxel and pemetrexed (Shandong Qilu Pharmaceutical, China), were determined by MTT assay as described above, the IC₅₀ value and RF for each drug were calculated.

Analysis of cell cycle distribution. MCF-7/S and MCF-7/MTX cells were seeded at a density of 4x10⁴ cells per well into 6-well plates and incubated for 24 h. Cells were trypsinized, washed twice with cold PBS and fixed by 70% ethanol overnight at 4°C, then washed and re-suspended with PBS containing RNase and propidium iodide. After 30 min in the dark, the cell cycle was analyzed using a flow cytometer (FACSCanto™II, BD Biosciences, USA).

Preparation of proteins. MCF-7/S and MCF-7/MTX cells were harvested, washed with PBS and then lysed in the lysis buffer (8 M urea, 2 M thiourea, 4% CHAPS, 40 mM Tris, 50 mM DTT, 1 mM PMSF, 0.5% (w/v) Bio-lyte and 0.5% IGP buffer) at 4°C for 30 min. After centrifugation at 14,000 rpm for 40 min at 4°C, the supernatants were purified by ReadyPrep2-D Cleanup kit (Bio-Rad, Hercules, CA, USA) according to the manufacturer’s instructions and stored at -80°C for use. The protein concentration was measured with Bradford method.

2-DE and image analysis. Each protein sample (800 µg) was applied to pH 3.0-10.0 (non-linear 17 cm) IPG strip (Bio-Rad) using a passive rehydration method. After rehydration for 16 h, IEF was performed successively for 1 h at 250 V, 1 h at 1,000 V, and 5 h at 10,000 V to give a total of 60 kVh on an IPGphor. Then, the gel strip was equilibrated for 15 min with DTT and iodoacetamide, respectively. The second dimension was performed using 12% SDS-PAGE until bromphenol blue reached the bottom of the gel. The gel was stained with sensitive colloidal Coomassie blue G-250 and scanned with UMax Powerlook 2110XL (Umax). Three independent experiments were made for each cell line to ensure the accuracy of the analyses. The images were analyzed by Imagemaster 2D Platinum (GE Amershams). The gel spot pattern of each gel was normalized and matched. The differentially expressed proteins between MCF-7/S and MCF-7/MTX in a given spot were calculated as the ratio of all the normalized spot values. Only spots that showed significant differences (± >1.5-fold, P<0.05) were selected for further analysis.

In-gel digestion. Differential protein spots were excised from the stained gels, destained in 30 mM potassium ferri cyanide/100 mM sodium thiocyanate (1:1 v/v) for 20 min and washed with water. Then the spots were incubated in 0.2 M NH₄HCO₃ for 20 min and freeze-dried. Each spot was digested with 12.5 ng/ml trypsin in 25 mM NH₄HCO₃ at 37°C overnight. The peptides were extracted three times with 60% ACN/0.1% TFA, and the extracts were pooled and dried in a vacuum centrifuge to complete desiccation at room temperature.

MALDI-TOF-MS analysis. The mass spectra were analyzed using MALDI-TOF-TOF instrument (4800 proteomics analyzer; Applied Biosystems). For protein identification, combined peptide mass fingerprinting (PMF) and the MS/MS queries were performed automatically by searching the EBI database using the MASCOT search engine 2.2 (Matrix Science, UK). The database search was carried out with the following parameters: enzyme, trypsin; the allowance of one missed cleavage; carbamidomethylation was set as fixed modification; oxidation of methionine was allowed as variable modification; the peptide and fragment mass tolerance were set at 100 ppm 0.4 Da, respectively. A GPS Explorer protein confidence index ≥95% was used for further manual
validation. Proteins with probability based MOWSE scores exceeding their threshold (P<0.05) were considered to be positively identified.

Western blot analysis. Western blot assay was performed after 10% SDS-PAGE separation, transfer to a PVDF membrane (Millipore) and saturation with 5% non-fat dry milk for 2 h at room temperature. Followed by incubation with primary rabbit polyclonal antibodies including anti-P-gp (1:500 dilution; GeneTex), anti-MRP1 (1:600 dilution; GeneTex), anti-RFC (1:1,500 dilution; Abcam), anti-hnRNP C1/C2 (1:600 dilution; GeneTex), rabbit monoclonal antibodies including anti-BCRP (1:800 dilution; Epitomics), anti-DHFR (1:20,000 dilution; Epitomics), anti-NPM (1:1,000 dilution; Epitomics), anti-ENO1 (1:2,500 dilution; Epitomics), anti-VIM (1:600 dilution; Epitomics), anti-PGAM1 (1:1,000 dilution; Epitomics), anti-PSMA2 (1:2,000 dilution; Epitomics), or anti-β-actin (1:800 dilution; Beijing Biosynthesis Biotechnology, China) as a loading control overnight at 4˚C, then the membranes were incubated with horseradish peroxidase-conjugated goat anti-rabbit IgG (1:20,000 dilution; CWbiotech, China) for 2 h at room temperature. Protein bands were detected by ECL detection system. Images were analyzed using a quantitative analysis system (Image-Pro Plus).

Quantitative PCR analysis. Total RNA was extracted using RNAfast2000 kit (Fastagen) according to the manufacturer's instructions. Quantitative PCR was performed with PrimeScript RT Master Mix Perfect Real Time Kit (Takara DRR036A) and SYBR Premix Ex Taq II (Takara). The primer sequences and product length are listed in Table I. The experiments were run on the Bio-Rad CFX96™ Real-time system (Bio-Rad); pre-degeneration for 95˚C, 30 sec, 1 cycle, and PCR reaction, 95˚C 5 sec followed by 60˚C, 30 sec, 40 cycles, and 95˚C for 15 sec, 60˚C for 30 sec, 95˚C for 15 sec for dissociation. β-actin was used as an internal control gene to normalize expression levels.

Small interfering RNA (siRNA) transfection. MCF-7/MTX cells were seeded in a 6-well plate at a density of 5x10^5 cells per well in RPMI-1640 without antibiotics. After 24 h, the NPM or non-specific control siRNA (Shanghai GenePharma, China) was transfected with Lipofectamine 2000 (Invitrogen) according to the manufacturer's instructions. The concentration of NPM siRNA was 50 nM. The efficiency of RNA interference was checked by quantitative PCR and western blot analysis, respectively. The transfected cells were used for proliferation assay, other genes and proteins detection.

Results

Methotrexate-resistant cell line. After a continuous induction with low concentration of MTX in a gradually increasing manner for 12 months, the MCF-7/MTX-resistant cell line grew steadily in the culture medium with 220  nM MTX. The IC_{50} values of MTX for MCF-7/S cells and MCF-7/MTX cell lines were 44±4.25 nM and 2818±97.9  nM, respectively (Fig. 1). The resistance of MCF-7/MTX cells to MTX was 64-fold higher than that of MCF-7/S cells, which implied the MTX-resistant cell line was successfully established.

MDR phenotype of MCF-7/MTX cells. Both MCF-7/S and MCF-7/MTX cell lines were treated with different concentrations of chemotherapeutic drugs, and the IC_{50} and RF values are summarized in Table II. The cross-resistance of MCF-7/MTX cells to doxorubicin, paclitaxel, 5-fluorouracil, mitoxantrone, vinorelbine, cisplatin, docetaxel, pemetrexed and gemcitabine significantly increased compared with that of MCF-7/S cells, which indicated the typical MDR phenotype of MCF-7/MTX cells.
mRNA and protein levels of MDR1, MRP1, BCRP, RFC and DHFR by quantitative PCR and western blot analyses. The mRNA levels of MDR1, MRP1, BCRP and DHFR in MCF-7/MTX cells were higher than those in MCF-7/S cells, while the mRNA expression of RFC was significantly decreased in MCF-7/MTX cell line (Fig. 2A). The protein levels of these molecules were consistent with gene expressions in both cell lines (Fig. 2B). These results demonstrated that the established MCF-7/MTX cell line is a valuable model to study MDR mechanism.

**Cell cycle distribution.** In order to further explore the effect of MTX on cells, the cell cycle profiles of MCF-7/S and MCF-7/MTX was performed using flow cytometry analysis. We found that the proportion of MCF-7/MTX cells in G0/G1 and G2/M phases showed a gradual increasing trend, from 63.37 to 65.69% and from 11.78 to 19.23%, respectively, while the proportion of S phase dropped from 24.85 to 15.07% (Fig. 3). These results suggested that MTX could block the cell cycle at the G0/G1 phase of progression.

**The proteome comparison of MCF-7/MTX and MCF-7/S cells.** To compare the proteome between MCF-7/MTX and MCF-7/S cells, we performed 2-DE to separate the global protein lysates of these two cell lines. The analysis was carried out in six gels made from three independent protein samples of each cell line. The images of protein spots were assessed using PDQuest6.0 analysis software. We selected 25 differentially expressed protein spots (±1.5-fold) between these two cell lines for further identification using MALDI-TOF-MS, and 17 proteins that were successively identified were marked with arrows in Fig. 4A, among which 7 proteins were upregulated and 10 proteins were downregulated in the MCF-7/MTX cell line (Table III).

**Validation of protein expression.** To further confirm the expression of identified proteins, western blotting was performed to validate six differentially expressed proteins (Fig. 4B). Consistent with the results of 2-DE analysis, three proteins nucleophosmin (NPM), α-enolase (ENO1), and vimentin (VIM) were significantly upregulated, while other three proteins including heterogeneous nuclear ribonucleoprotein (hnRNP C1/C2), phosphoglycerate mutase 1 (PGAM1), and proteasome subunit α-type-2 (PSMA2) were obviously downregulated in MCF-7/MTX cells (Fig. 5A).

**Validation of mRNA expression.** To further validate whether the mRNA levels of these six differentially expressed proteins had variations, quantitative PCR assay was applied to detect the mRNA expression of NPM, ENO1, VIM, hnRNP C1/C2, PGAM1 and PSMA2 (Fig. 5B). Compared with MCF-7/S cells, the mRNA levels of NPM and VIM were increased while hnRNP C1/C2, PGAM1 and PSMA2 genes were decreased in MCF-7/MTX cell line, which was consistent with the alteration of the protein levels. Unexpectedly, the mRNA expression of

<table>
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<th>Drugs</th>
<th>MCF-7/S</th>
<th>MCF-7/MTX</th>
<th>MCF-7/MTX RNAi</th>
<th>IC_{50} MCF-7/MTX/IC_{50} MCF-7/S</th>
<th>RF</th>
<th>IC_{50} MCF-7/MTX RNAi/IC_{50} MCF-7/S</th>
<th>RF</th>
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<td>Methotrexate</td>
<td>44±4.25</td>
<td>2,818±97.9</td>
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<td>64.0</td>
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<td>Doxorubicin</td>
<td>71±2.57</td>
<td>245±7.39</td>
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<td>Paclitaxel</td>
<td>20±0.45</td>
<td>48±1.62</td>
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<td>5-fluorouracil</td>
<td>3,890±178</td>
<td>7,943±55.3</td>
<td>6,892±57.9</td>
<td>2.04</td>
<td>1.77</td>
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<td>Mitoxantrone</td>
<td>186±16.0</td>
<td>295±24.8</td>
<td>285±21.0</td>
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<td>1.53</td>
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<td>Vinorelbine</td>
<td>51±7.18</td>
<td>1,023±91.6</td>
<td>256±10.4</td>
<td>20.1</td>
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<td>Cisplatin</td>
<td>2,042±74.2</td>
<td>5,495±66.2</td>
<td>3,987±187</td>
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<td>1.95</td>
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<tr>
<td>Docetaxel</td>
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<td>22±5.42</td>
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<tr>
<td>Pemetrexed</td>
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<td>9,550±111</td>
<td>6,424±245</td>
<td>2.63</td>
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<tr>
<td>Gemcitabine</td>
<td>955±47.0</td>
<td>6,026±568</td>
<td>2,698±109</td>
<td>6.31</td>
<td>2.83</td>
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</table>

MCF-7/MTX versus MCF-7/S; *P<0.01. MCF-7/MTX RNAi versus MCF-7/MTX; **P<0.01.
Figure 2. MCF-7/MTX cells have classic MDR and MTX resistance. (A) The mRNA expression of MDR1, MRP1, BCRP, RFC and DHFR in MCF-7/S and MCF-7/MTX cells was compared by quantitative PCR. The results are shown by fold change (± SEM) relative to the control. (B) The protein levels of P-gp, MRP1, BCRP, RFC and DHFR in MCF-7/S and MCF-7/MTX cells were detected by western blot analyses (left panel). Relative expression is shown as mean ± SD (right panel). All experiments were performed three times (*P<0.05; **P<0.01).

Figure 3. Cell cycle distribution of MCF-7/S and MCF-7/MTX cells. The DNA content was evaluated by flow cytometry after ethanol fixation and PI staining. Data are represented as proportion of cells in G0/G1, S and G2/M phases of the cell cycle from three independent experiments.
ENO1 was slightly downregulated in the MCF-7 MTX cells, which was different from the protein level.

Reversal of MDR in MCF-7/MTX cells by knockdown of NPM.
We found the protein and mRNA expressions of NPM were both significantly upregulated in the MCF-7/MTX cells. To study the functions of the increased expression of NPM in the MDR of MCF-7/MTX cells, MTT assay was performed to determine the sensitivity of MCF-7/MTX cells to MTX and other antitumor drugs following knockdown of NPM by siRNA. Our results showed that siRNA significantly downregulated the gene and protein of NPM in the MCF-7/MTX cells (Fig. 6A). Knockdown of NPM by siRNA in MCF-7/MTX cells obviously enhanced the inhibitory effect of MTX on proliferation. In addition, the IC$_{50}$ value and RF of MCF7/MTX cells exposed to doxorubicin, paclitaxel, 5-fluorouracil, vinorelbine, cisplatin, docetaxel, pemetrexed and gemcitabine were significantly decreased after downregulating the expression of NPM by siRNA (Table II). Surprisingly, the protein and mRNA levels of drug resistance-related molecules, MDR1, MRP1, BCRP and DHFR were also significantly decreased while the protein and gene expressions of RFC were increased after knockdown of NPM in MCF-7/MTX cells (Fig. 6B).

These observations demonstrate that NPM has a prominent role in the multidrug resistance of MCF-7/MTX cells.

Discussion
The antifolate MTX is applied to the treatment of a wide variety of cancers, it is usually given in combination with cyclophosphamide and 5-fluorouracil to cure breast cancer. However, drug resistance is frequently observed upon treat-
ment with MTX for cancer patients, thus compromising the drug effectiveness. Although the alteration of gene expressions also contribute to MTX resistance, such as increase in S100A4, UGT1A6, caveolin-1, enolase-2, PRKCA and the decrease of miR-224 or E-cadherin (7-10), the resistance mechanism of MTX, especially the MDR caused by MTX is not fully understood. In the present study, we have successfully established the MTX-resistant human breast cancer cell line MCF-7/MTX, which displays cross-resistance to a variety of commonly used chemotherapy drugs and multidrug resistance phenotype. Using the proteomic assay, we integrally analyzed the protein expression differences between resistant cells and parental cells to identify potential molecular targets for MDR of breast cancer.

In this study, 17 differentially expressed proteins were identified, seven proteins were upregulated and 10 proteins were downregulated in MCF-7/MTX cells. Moreover, six significantly expressed proteins with distinct functions were validated by western blot and quantitative PCR assays at protein and mRNA levels, respectively. These proteins are involved in tumorigenesis, metabolism, glycolysis, cell proliferation, apoptosis and the invasion process. Their roles in the formation of drug resistance and molecular mechanism are discussed below.

α-enolase (ENO1), known as 2-phosphate-D-glycerate hydrolase, catalyzes the formation of phosphoenolpyruvate from 2-phosphoglycerate in the process of glycolysis, which was a key role in cellular energy metabolism (11).
ENO1 has been recognized as a conserved and single function protein, but recent studies found that the effect of this enzyme is involved in transcription, regulation of apoptosis and cell differentiation processes. Research has shown that the expression of ENO1 significantly increased in certain tumor cells, including liver cancer, lung cancer, head and neck cancer, which could be used as prognostic indicators for clinical treatment (12-14). It was reported that the elevated expression of ENO1 was closely related to tamoxifen and adriamycin resistance in breast cancer (15,16). The expression of ENO1 was regulated by ERK1/2 appearing to be mediated by c-Myc, which changed the level of extracellular ATP, thus affecting cell survival (17). In our study, the protein level of ENO1 in MCF-7/MTX cells are significantly increased, but its gene level was slightly lower in the resistant cells, which indicates the mRNA level is not fully present in the protein level, due to the mRNA possessed storage, transport, degradation, translational regulation and post-translational processing of the product, which affects the quality and quantity of the protein (18). We have found the correlation between ENO1 and MDR in breast cancer, which may be regulated by activating ERK1/2 pathway, further affecting tumor cell proliferation.

Vimentin (VIM) is a member of intermediate filament protein family, which constitutes the cytoskeleton with microtubules and actin filaments, it is involved in cell adhesion, migration, apoptosis and cell signal transduction (19). In prostate cancer and triple-negative breast cancer cells, the high expression of VIM was significantly associated with cell invasiveness, moreover, the activation of ERK signaling pathway promotes the overexpression of VIM and increased cell migration and invasion in head and neck cancer (20-22). In recent research, VIM was intimately associated with the resistance to chemotherapeutic drugs, it was significantly higher expression in resistant cancer cells including paclitaxel-resistant prostate cancer, temozolomide-resistant malignant glioma and tamoxifen-resistant breast cancer (23-25). Our results show that the protein and mRNA levels of VIM are significantly increased in the MTX-resistant cells, which may be related to the variation of cell adhesion, thereby possibly influencing cell metastasis and invasion, and its resistance mechanisms need to be further explored.

Heterogeneous nuclear ribonucleoprotein C1/C2 (hnRNP C1/C2) is a member of hnRNPs family, as an RNA-binding protein, it participates in the regulation of pre-mRNA splicing and post-translational modification (26). In addition, studies

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### Table III. MS identification of differentially expressed proteins between MCF-7/MTX and MCF-7/S cells.

<table>
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<th>Spot no.</th>
<th>Accession no.</th>
<th>Protein name</th>
<th>MW (kDa)/ PI</th>
<th>Peptide matched</th>
<th>Score</th>
<th>Sequence coverage (%)</th>
<th>Expression in MCF-7/MTX/MCF-7/S</th>
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have shown that hnRNPs were associated with tumor development and prognosis (27). Previously, hnRNP C1/C2 has been considered as an apoptosis-related protein, and the protein level of hnRNP C1/C2 was p53-dependently upregulated, but its gene transcription did not depend on the p53 role in colon cancer cells after mitomycin C treatment (28,29). Furthermore, in A549 lung cancer cells of wild-type p53, the expression of hnRNP C1/C2 was significantly reduced in the process of apoptosis, it regulated the synthesis of p53 isoforms with p53 IRES interaction and affected tumor cell death (30,31). On the contrary, hnRNP C1/C2 was associated with DNA repair function, knockdown of hnRNP C1/C2 by RNA interference could increase the sensitivity of HeLa cells to chemotherapeutic drugs (32). In the present study, both protein and mRNA expression of hnRNP C1/C2 are downregulated in MCF-7/MTX cells, which may be connected with mutations of apoptosis regulatory factors and affects the normal process of cell apoptosis, while the exact mechanism needs further investigation.

Phosphoglycerate mutase 1 (PGAM1), as a key enzyme of glycolysis and gluconeogenesis process, catalyzes the conversion of 3-phosphoglycerate (3-PG) to 2-phosphoglycerate (2-PG). It was found that PGAM1 was overexpressed in certain tumor cells including liver cancer, breast cancer, and glioma, its upregulation may facilitate the proliferation and metabolism of cancer cells, which could be used as a potential therapeutic target for tumors (33-36). Moreover, PGAM1 was also negatively correlated with the tumor suppressor p53, which further illustrates the interaction between PGAM1 and cell apoptosis (37). However, we discovered that the protein and mRNA levels of PGAM1 are significantly reduced in MTX-resistant cells, abnormal glucose metabolism may be involved in mediating the role of MDR in breast cancer, and the resistance mechanism is still unclear and requires further study.

Proteasomes are macromolecular complexes with a variety of proteolytic functions, acting as catalysts in the ubiquitin-proteasome pathway of intracellular protein degradation. However, the ubiquitin-proteasome pathway may control numerous important
physiological functions, such as cell cycle, signal transduction, DNA damage repair and cell apoptosis, additionally, abnormal alteration of this pathway is closely bound with prognosis and development of malignant tumor. Increased expression of proteasome subunit protein can lead to formation of tumors, especially to protect anti-apoptotic effects of cancer cells, therefore, the proteasome subunit proteins could be recognized as potential antitumor targets. Some studies demonstrated that PSMA7 could inhibit proliferation and invasion in A549 cells (38), it regulated cell transcription, cell cycle and tumor development with other important proteins interaction (39). It was found that PSMA5 was highly expressed in 5-fluorouracil-resistant colorectal cancer cells, which showed certain anti-apoptotic activity (40). Whereas, in our established MTX-resistant cell line, we first discover that the protein and gene levels of PSMA2 are significantly down-regulated, which reflects a correlation between PSMA2 and MDR in breast cancer, this provides the basis for future investigation of resistance mechanisms.

Nucleolar phosphoprotein (NPM) is abundant and highly conserved phosphoprotein which shuttles rapidly between nucleus and cytoplasm, it participates in the assembly and synthesis of ribosome, replication of chromosomes and centrosomes and intracellular signal transduction, which plays an important role in cell growth, proliferation and transformation. An increasing number of studies manifested that there may exist a connection between NPM and the occurrence of tumors, high expression of the proto-oncogene c-Myc could lead to upregulation of NPM transcriptional level; its overexpression could inactivate the function of p53 and enhance the anti-apoptosis effect in cancer cells; NPM was able to regulate tumor progression by activating the phosphorylation of MAPK/ERK and c-Myc signaling pathway (41). The overexpression of NPM defended the p53-mediated cellular senescence and growth arrest in colorectal cancer cells, which showed its role in promoting tumor growth (42), conversely, reducing the expression of NPM by shRNA sensitized the resistant leukemia cells to chemotherapy (43). There is also evidence indicating that NPM has a tumor suppressor property maintaining the gene stability and regulating the ARF tumor suppressor (44). Similarly, in invasive breast cancer MDA-MB-231 cells, the overexpression of NPM inhibited cell growth, which could be used as a tumor suppressor factor and predictor of poor prognosis (45). Although the above demonstrated functions of NPM are controversial, in our study, the protein and gene levels of NPM are both significantly increased in MCF-7/MTX cells. We discovered the connection between NPM and MTX resistance mechanism, which may induce MDR through acting on MAPK/ERK and c-Myc signaling pathway in breast cancer. Subsequently, we reduced the expression of NPM by siRNA interference, which attenuates the resistance of MCF-7/MTX cells to various drugs and the expressions of cell factors related to MDR and MTX resistance mechanism. These results indicate that NPM contributes to MTX resistance in breast cancer cells and suggest that NPM may play a significant role in the occurrence of MDR of cancer cells.

In conclusion, we have successfully established the MTX-resistant human breast cancer cell line and identified 17 differentially expressed proteins. NPM, involving in tumorigenesis and signal transduction, plays a prominent role in the MDR of MCF-7/MTX cells. This study investigated the resistance mechanisms of breast cancer and provided a theoretical basis for the clinical diagnosis of MDR.

Acknowledgements

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References


