Abstract. The present study was designed to evaluate the gene polymorphisms of organic anion transporting polypeptide 1B1 (OATP1B1) in predicting the therapeutic efficacy of tamoxifen (TAM) for MCF-7. Established plasmids OATP1B1 wild-type 388GG and 521CC were transfected into MCF-7 cells and used to determine whether the gene polymorphisms affected the therapeutic efficacy of TAM for MCF-7. The established plasmids pcDNA3.1(-)-OATP1B1 wild-type 388GG and 521CC were digested by restriction enzymes and analyzed by gene sequencing. The gene polymorphisms of OATP1B1 in MCF-7 breast cancer cells were examined by RT-PCR and western blot analysis. The results showed that the mutations of OATP1B1 388GG and 521CC led to a decrease of the inhibition and apoptotic rates of MCF-7 cells, albeit not significantly compared to the OATP1B1 group. The G0/G1 phase length ratio was reduced, and the S and G2M phases were increased in the OATP1B1 388GG and 521CC groups, although not significantly compared to the OATP1B1 group. The mutations of OATP1B1 388GG and 521CC inhibited the activity of OATP1B1 protein, restrained the turnover capacity of OATP1B1 and reduced the entrance of TAM into MCF-7 cells, resulting in weakened efficacy of TAM in the treatment of breast cancer.

Introduction

Breast cancer continues to be the most common type of cancer diagnosed among women in China, and it was estimated that there were ~232,340 new cases of invasive breast cancer in 2013 (1). The American Cancer Society projected ~39,620 breast cancer-associated mortalities in 2013 (2). In the past three decades, breast cancer has been the second leading cause of cancer death after lung cancer (3). However, breast cancer is a complex disease entity with different biological characteristics and clinical behavior. Many studies have shown that there are no treatment guidelines for triple-negative breast cancer.

Adjuvant endocrine therapy is an important method used for the comprehensive treatment of breast cancer (4). Chemotherapy for breast cancer is closely associated with hormone levels. Tamoxifen (TAM) is non-synthetic cholesterol antiestrogen, which is the basic drug utilized in adjuvant endocrine therapy for breast cancer. TAM is widely used to treat breast cancer patients (5). However, the differences in the gene polymorphism of patients who are administered the drug may lead to individual differences regarding efficacy and toxicity (6). Genetic factors play an important role in these differences.

Organic anion transporting polypeptide 1B1 (OATP1B1) is a type of intake body, distributed in multiple human organs and particularly distributed in specific organs (7). It belongs to the superfamily of solute uptake, of which the encoding genes are known as the \textit{SLCO1B1} gene collectively. In the Chinese population, mutation frequencies of OATP1B1 388A>G and OATP1B1 521T>C are 73.4 and 14.0%, respectively, which are similar to those of the Japanese population (8,9). The incidence rate of OATP1B1 521T>C is 16% in Asians and 18% in Caucasians, thus this incidence rate is varied based on ethnicity (10). Studies have shown that in African-Americans, the mutation rate of OATP1B1 388G is 2.5-fold that of Europeans and Americans, with the mutation rate of OATP1B1 521C being higher among the Finnish population. The mutations of OATP1B1*15 (388G and 521C) is 43.4% in the Japanese population simultaneously. However, the incidence rate of OATP1B1*15 is 14% in Caucasians (11). The majority of current studies concerning the genetic polymorphism of OATP1B1 have focused on statin drugs, which enter the liver cells to exert an inhibitory effect on HMG-CoA reductase. Therefore, OATP1B1 is important in most internalization processes of statin drugs into hepatocytes (12).
Recent findings have shown that the genetic polymorphism of OATP1B1 affects the pharmacokinetic process of irinotecan in cancer patients significantly, including OATP1B1*1a, OATP1B1*ib, OATP1B1*5 and OATP1B1*15 (13,14).

There were no significant effects on simvastatin pharmacokinetics by SLCO1B1 388A>G, SLCO1B1 521T>C, SLCO1B1 11187G>A, SLCO1B1 571T>C and SLCO1B1 597C>T. Ritonavir intracellular concentrations were associated with OATP1B1 521T>C polymorphism (15). Previous results suggest that the gene mutation of OATP1B1 521T>C influences the transport of rosuvastatin (16). However, no association between the OATP1B1 521T>C polymorphism and the cholesterol synthesis response and rosuvastatin plasma concentration has been identified (17). OATP1B1 388A>G and 521T>C have been shown to lead to altered pharmacokinetics of pravastatin. The pharmacokinetics of irinotecan, estrone-3-sulfate and estradiol-17β-glucuronide also affect OATP1B1 388A>G and 521T>C, although without statistical significance (18).

In the present study, we found that the specific mutation of OATP1B1 significantly reduced the uptake activity of OATP1B1 protein, which may affect the internalization of many endogenous substances and drug uptake into hepatocytes and significantly influence the metabolism and efficacy of many drugs. OATP1B1 also influences the adverse reactions of some drugs and the interaction between drugs. A gene polymorphism expression platform of OATP1B1 388GG and 521CC was also established and utilized to determine whether OATP1B1 388GG and OATP1B1 521CC gene polymorphisms can affect TAM treatment of breast cancer in vitro.

Materials and methods

Total RNA extraction. Approximately 100 mg tissue was ground into a powder in liquid nitrogen, and moderate RNAiso Plus (Qiagen, Hilden, Germany) was added for 5-10 min at room temperature. The supernatant was collected by centrifugation at 12,000 rpm for 5 min at 4˚C followed by the addition of chloroform. Miscible liquids were agitated and placed for 5-10 min at room temperature. The liquid was centrifuged at 12,000 rpm for 10 min at 4˚C, and the supernatant was added to the centrifuge tube. The isopropanol, which was the same volume as that of the supernatant, was added to the centrifuge tube. The liquid was centrifuged at 12,000 rpm for 5 min to obtain sediment again. The sediment obtained was dried at room temperature, and then treated in 20-100 µl DEPC-H2O. Total RNA was maintained at -80˚C.

cDNA synthesis. Total RNA (1 µl), 1 µl 0.5 µg/µl oligo(dT)18 and 10 µl DEPC-H2O was mixed, subjected to PCR at 70˚C for 5 min and cooled rapidly on ice. Four microliters 5X reaction buffer, 1 µl 20 U/µl RNA enzyme inhibitors and 2 µl 10 mmol/l dNTP were dissolved in miscible liquids and subjected to PCR at 37˚C for 5 min. One microliter 200 U/µl reverse transcriptase was added and the mixture was subjected to PCR. Cycling conditions used were: 42˚C for 60 min and 70˚C for 10 min.

cDNA synthesis was maintained at -80˚C.

DNA extraction. The primers were designed according to gene coding sequence (CDS) of OATP1B1 in GenBank. KpnI and NotI restriction sites were subsequently inserted. The primers used were: forward: 5'-TTACGCGCCGCATGGACCCAA ATCAAACATT-3' and reverse, 5'-GCCGGTACCTTAACACAG ATGGTTTTCATCTCG-3'.

cDNA (2-4 µl), 2 µl sense primer (10 pM), 2 µl reverse primer (10 pM), 4 µl dNTP (2 mM), 5 µl 10X PCR buffer, 1 µl Taq DNA Polymerase (Takara, Otsu, Japan) and moderate DEPC-H2O were mixed in the centrifuge tube, and the total volume was 50 µl. Cycling conditions used were: 94˚C for 10 min followed by 40 cycles at 95˚C for 30 sec, 58˚C for 90 sec and 72˚C for 30 sec, and then 72˚C for 7 min. DNA was kept at -80˚C. DNA OATP1B1 was analyzed by electrophoresis of 0.5 or 1% agarose gel followed by gene sequencing.

Site-directed mutagenesis of OATP1B1 388GG and 521CC. The site-directed mutagenesis primers of OATP1B1 388GG and OATP1B1 521CC were designed according to OATP1B1 gene CDS in GenBank (Table I). The plasmids were constructed and transfected into MCF-7 cells using Lipofectamine 2000 (Invitrogen) according to the manufacturer’s instructions in a
humidified atmosphere at 37°C with 5% CO₂. After 6 h, the transfusion medium was replaced with Dulbecco's modified Eagle's medium (DMEM) containing 10% fetal bovine serum (FBS) without antibiotic in a humidified atmosphere at 37°C with 5% CO₂. The OATP1B1 gene polymorphisms were examined in MCF-7 cell lines by RT-PCR and western blot assay.

**Cell culture.** MCF-7 human breast cancer cells were obtained from the Shanghai Institute of Cell Biology in the Chinese Academy of Sciences for the present study. MCF-7 cells were maintained in DMEM containing 10% FBS supplemented with 100 U penicillin/streptomycin in a humidified atmosphere at 37°C with 5% CO₂.

**Experiments.** The experiment was divided into six groups: A group was MCF-7; B group was MCF-7 with TAM (10 µM); C group was MCF-7 transfected with pcDNA3.1(-) plasmid and provided with TAM (10 µM); D group was MCF-7 transfected with pcDNA3.1(-)-OATP1B1 plasmid and provided with TAM (10 µM); E group was MCF-7 transfected with pcDNA3.1(-)-OATP1B1 388GG plasmids and provided with TAM (10 µM); and F group was MCF-7 transfected with pcDNA3.1(-)-OATP1B1 521CC plasmids and provided with TAM (10 µM).

**MCF-7 cell proliferation assay.** MCF-7 cell proliferation was examined using the colorimetric assay using 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assays. Cells were plated at 2.5×10⁵ cells/well in a 96-well plate and incubated for 24 h. The plasmids were transfected into MCF-7 cells. The cells were treated with TAM solutions at concentrations of 10 µM for 24 and 48 h. PBS (pH 7.4) alone was used as the vehicle-group. TAM was dissolved in PBS and was used as treat-groups. For the cell cycle analysis, the cells of the vehicle- and treat-groups were trypsinized without EDTA and centrifuged at 2,000 x g for 5 min. The cells were harvested, washed in PBS (pH 7.4) and fixed in 70% with ice-cold ethanol at 0-4°C overnight. The cells were collected and centrifuged at 2,000 x g for 5 min and the supernatant was removed. The cell pellet was resuspended and washed in PBS (pH 7.4). The cell solution was transferred to a flow tube, and incubated with 10 ml of RNase A (10 mg/ml) for 1 h at 37°C. The cell samples were incubated with 50 ml of PI (250 mg/ml) for 30 min in the dark at 0-4°C. The cell cycle proliferation index was subsequently calculated using the formula: PI = (G₂/M + S)/(G₀/G₁ + S + G₂/M) x 100%. The cell cycle distribution was analyzed by flow cytometry (BD Biosciences).

**Quantification of apoptotic cells by flow cytometry.** Apoptotic cells were measured by flow cytometry (BD Biosciences, San Diego, CA, USA). The cells were stained using an Annexin V-FITC Apoptosis Detection kit according to the manufacturer's instructions. Briefly, MCF-7 cells (5.0×10⁵ cells/well) were plated in 6-well plates and incubated for 24 h. The plasmids were subsequently transfected into MCF-7 cells. The cells were treated with TAM solutions at concentrations of 10 µM for 24 and 48 h. PBS (pH 7.4) alone was used as the vehicle-group. TAM was dissolved in PBS and was used as treat-groups. For the apoptotic cell analysis, the cells of the vehicle- and treat-groups were trypsinized without ethylenediaminetetraacetic acid (EDTA) and centrifuged at 2,000 x g for 5 min. The cells were collected, washed in PBS (pH 7.4), and centrifuged at 2,000 x g for 5 min. The cell samples were suspended with 500 µl mixed buffer solution. Then, 5 µl fluorescein isothiocyanate (FITC)-Annexin V and 10 µl propidium iodide (PI) were added into the cell samples and incubated for 5-15 min at room temperature in the dark. Apoptotic cells were analyzed by flow cytometry (BD Biosciences).

**Statistical analysis.** The experiments were repeated at least three times. Data are presented as the mean ± SEM. The differences between means were analyzed using the two-tailed Student's t-test. Statistical analyses were performed using SPSS 17.0 software. Differences with P<0.05 were considered to indicate a statistically significant result.

**Results**

**DNA fragmentation of OATP1B1.** DNA fragmentation of OATP1B1 was analyzed by electrophoresis of 1% agarose gel, which showed that the amplified gene fragment was 2,096 bp (Fig. 1A). The DNA fragmentation of OATP1B1 was analyzed by gene sequencing, which showed that the amplified gene fragment identified the 388 site as A, and the 521 site as T (Fig. 1B and C).

**Restriction enzyme and gene sequence of pcDNA3.1(-)-OATP1B1.** The established plasmids of pcDNA3.1(-)-OATP1B1 were digested by restriction enzymes (KpnI and NotI). The products were analyzed by electrophoresis of 0.5% agarose gel, which showed that pcDNA3.1(-) was 5.4 kb, while the OATP1B1 fragment was 2,096 bp (Fig. 2A). Gel extraction of the OATP1B1 fragment was performed and the fragment was analyzed by gene sequencing, which showed that the amplified gene fragment identified the 388 site as A, and the 521 site as T (Fig. 2B and C).

**Restriction enzymes and gene sequencing of pcDNA3.1(-)-OATP1B1 388GG and 521 CC.** The established plasmids of pcDNA3.1(-)-OATP1B1 388GG and 521 CC were digested by restriction enzymes, KpnI and NotI. These products were analyzed by electrophoresis of 0.5% agarose gel, which showed that pcDNA3.1(-) was 5.4 kb, and the OATP1B1 fragment was 2,096 bp (Fig. 3A). The OATP1B1 fragment was extracted using gel extraction and analyzed by gene sequencing. The amplified gene fragment identified at the 388 site was G (Fig. 3B).
The established plasmids of pcDNA3.1(-)-OATP1B1 were digested by restriction enzymes, KpnI and NotI. The products were analyzed by electrophoresis of 0.5% agarose gel, which showed that pcDNA3.1(-) was 5.4 kb, and the OATP1B1 fragment was 2,096 bp (Fig. 3C). The OATP1B1 fragment was extracted using gel extraction and was analyzed by gene sequencing, which showed that the amplified gene fragment identified at the 521 site was T (Fig. 3D).

**Figure 2.** Restriction enzyme and gene sequence of pcDNA3.1(-)-OATP1B1. The established plasmids of pcDNA3.1(-)-OATP1B1 were digested by restriction enzymes: (A) left of image is pcDNA3.1(-) and right of image is DNA marker. The established plasmids of pcDNA3.1(-)-OATP1B1 were analyzed by gene sequencing: (B) OATP1B1 388 site was A and (C) OATP1B1 521 site was T.

The established plasmids of pcDNA3.1(-)-OATP1B1 521CC were digested by restriction enzymes, KpnI and NorI. The products were analyzed by electrophoresis of 0.5% agarose gel, which showed that pcDNA3.1(-) was 5.4 kb, and the OATP1B1 fragment was 2,096 bp (Fig. 3C). The OATP1B1 fragment was extracted using gel extraction and was analyzed by gene sequencing, which showed that the amplified gene fragment identified at the 521 site was C (Fig. 3D).

**Cataphoresis of plasmid transfection.** RNA was extracted from the transfection of MCF-7 cells of each group, and PCR was used to amplify double-stranded DNA. The electrophoretic analysis was performed on the product using 1% agarose gel, which showed that the plasmid was successfully expressed (Fig. 4).

**Western blotting of plasmid transfection.** The OATP1B1 gene polymorphisms in MCF-7 cell lines were examined by western blot assay. A was MCF-7 cell, B was pcDNA3.1(-) was transfected into MCF-7 cell, C was pcDNA3.1(-)-OATP1B1 was transfected into MCF-7 cell, D was pcDNA3.1(-)-OATP1B1 388GG was transfected into MCF-7 cell, E was
pcDNA3.1(-)-OATP1B1 521CC was transfected into MCF-7 cell. A and B did not express OATP1B1 protein, whereas C-E expressed OATP1B1 protein (Fig. 5).

TAM effect on the cell inhibition rate of MCF-7 by plasmid transfection. Effect of TAM (10 µM) on the inhibition rate of MCF-7 cells at (A) 24 h and (B) 48 h. *P<0.01 compared with MCF-7 group, P>0.05 compared with pcDNA3.1(-)-OATP1B1 transfected in the MCF-7 cell group. TAM, tamoxifen.
groups was slightly decreased (P>0.05, n=3) compared to the D group, although the difference was not statistically significant (Fig. 6A).

After the effect of TAM (10 µM) on the proliferation of MCF-7 cells at 48 h, the inhibition rate of D group was markedly increased (P<0.05, n=3), compared to the B group (Fig. 6B). However, the inhibition rate of MCF-7 cells of the E and F groups were slightly decreased (P>0.05, n=3) compared to the D group, although the difference was not statistically significant (Fig. 6B).

**Figure 7. TAM effect on the cell apoptotic rate of MCF-7 by plasmid transfection.** Effect of TAM (10 µM) on the cell apoptotic rate of MCF-7 cells at (A) 24 h and (B) 48 h. *P<0.01 compared with MCF-7 group, †P>0.05 compared with pcDNA3.1(-)-OATP1B1 transfected into MCF-7 cell group. TAM, tamoxifen.

**TAM effect on the cell apoptotic rate of MCF-7 by plasmid transfection.** We established expression platforms of OATP1B1 genetic polymorphisms to assess the role of OATP1B1 genetic polymorphisms on TAM (10 µM) with regard to the apoptosis of MCF-7 cells. The results showed that at 24 h, apoptosis in the D group was markedly increased (P<0.05, n=3), compared to the B group (Fig. 7A). However, apoptosis of MCF-7 cells of the E and F groups was slightly decreased (P>0.05, n=3) compared to the D group, although the difference was not statistically significant (Fig. 7A and B).
After the effect of TAM (10 µM) on the apoptosis of MCF-7 cells at 48 h, apoptosis of the D group was markedly increased (P<0.05, n=3), compared to the B group (Fig. 7C and D). However, the apoptosis of MCF-7 cells of the E and F groups was slightly decreased (P>0.05, n=3) compared to the D group, although the differences were not statistically significant (Fig. 7C and D).

**TAM effect on the cell cycle of MCF-7 by plasmid transfection.** We established expression platforms of OATP1B1 genetic polymorphisms to assess the role of OATP1B1 genetic polymorphisms concerning the effect of TAM (10 µM) on the cell cycle of MCF-7 cells. After the effect of TAM (10 µM) on the cell cycle of MCF-7 cells at 24 h, the G0/G1 phase in D group was markedly increased (P<0.05, n=3), the S and G2M phases were markedly decreased (P<0.05, n=3), compared to the B group (Table II, Fig. 8A). In E and F groups, the G0/G1 phase was slightly decreased (P>0.05, n=3), while the S and G2M phases were slightly increased (P>0.05, n=3), respectively, compared to the B group.
Table II. TAM effect on the cell cycle of MCF-7 by plasmid transfection.

<table>
<thead>
<tr>
<th>Time</th>
<th>Group</th>
<th>G0/G1 (%)</th>
<th>S (%)</th>
<th>G2/M (%)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>24 h</td>
<td>A</td>
<td>68.7±3.12</td>
<td>22.2±4.21</td>
<td>5.7±1.02</td>
<td>28.97</td>
</tr>
<tr>
<td></td>
<td>B</td>
<td>74.3±2.24</td>
<td>12.7±1.06</td>
<td>12.8±1.01</td>
<td>25.58</td>
</tr>
<tr>
<td></td>
<td>C</td>
<td>74.7±2.31</td>
<td>14.8±1.11</td>
<td>9.9±1.03</td>
<td>24.85</td>
</tr>
<tr>
<td></td>
<td>D</td>
<td>79.5±1.69</td>
<td>10.1±0.77</td>
<td>9.8±1.09</td>
<td>20.08a</td>
</tr>
<tr>
<td>48 h</td>
<td>A</td>
<td>64.3±2.53</td>
<td>21.4±2.31</td>
<td>13.9±2.13</td>
<td>35.49</td>
</tr>
<tr>
<td></td>
<td>B</td>
<td>75.9±2.72</td>
<td>12.3±1.58</td>
<td>11.0±1.35</td>
<td>23.53</td>
</tr>
<tr>
<td></td>
<td>C</td>
<td>76.6±2.69</td>
<td>13.0±1.62</td>
<td>10.3±1.40</td>
<td>23.34</td>
</tr>
<tr>
<td></td>
<td>D</td>
<td>81.2±3.20</td>
<td>9.1±1.18</td>
<td>9.6±1.21</td>
<td>18.78a</td>
</tr>
<tr>
<td></td>
<td>E</td>
<td>80.1±2.11</td>
<td>10.6±1.35</td>
<td>8.8±1.25</td>
<td>19.54a,b</td>
</tr>
<tr>
<td></td>
<td>F</td>
<td>81.0±2.10</td>
<td>9.3±1.20</td>
<td>9.3±1.17</td>
<td>18.76a</td>
</tr>
</tbody>
</table>

A group was MCF-7; B group was MCF-7 with TAM (10 µM); C group was MCF-7 transfected with pcDNA3.1(-) plasmid with TAM (10 µM); D group was MCF-7 transfected with pcDNA3.1(-)-OATP1B1 plasmid with TAM (10 µM); E group was TAM-transfected with pcDNA3.1(-)-OATP1B1 388GG plasmids with TAM (10 µM); F group was MCF-7 transfected with pcDNA3.1(-)-OATP1B1 521CC plasmids with TAM (10 µM). *P<0.01 compared with the TAM group, **P<0.05 compared with the MCF-7 group.

After the effect of TAM (10 µM) on the cell cycle of MCF-7 cells at 48 h, the G0/G1 phase in the D group was markedly increased (P<0.05, n=3), while the S and G2/M phases were markedly decreased (P<0.05, n=3), compared to the B group (Table II, Fig. 8B). In the E and F groups, the G0/G1 phase was slightly decreased (P>0.05, n=3), while the S and G2/M phases were slightly increased (P>0.05, n=3), respectively, compared to the D group, although the differences were not statistically significant (Table II, Fig. 8B).

Discussion

The functions of OATP1B1 are numerous, of which the specificity is distributed in the basolateral membrane of liver cells of the human. A variety of endogenous substances and drugs enter the liver cells via the portal system, including thyroid hormones, prostaglandin, bile acid, and statins. The OATP1B1 gene is located on chromosome 12, with the full-length 10.86 kb (7). The encoding gene of SLCO1B1 includes 2,076 bases, encoding 691 amino acids, including 15 exons and 14 introns, as well as 20 function-genetic polymorphisms. The OATP1B1 gene has multiple single-nucleotide polymorphisms that are associated with abnormal uptake in in vitro or in vivo experiments, of which the OATP1B1 gene 388A>G and 521T>C are the most common mutations (19). In a previous study, it was found that the SLCO1B1 gene 388A>G polymorphism was not statistically significantly associated with gallstone formation in a north Indian population (20). SLCO1B1 521 T>C plays a significant role in susceptibility to colorectal cancer risk in the Turkish population (14). SLCO1B1 388A>G and 521T>C polymorphisms were also not associated with response to atorvastatin or simvastatin (21).

In the present study, a molecular biology technique was applied to construct the pcDNA3.1(-)-OATP1B1 plasmid. The site-directed mutagenesis method was then used to establish pcDNA3.1(-)-OATP1B1 388GG and 521CC plasmids successfully. The pcDNA3.1(-)-OATP1B1-MCF-7 gene polypeptide expression platform demonstrated that OATP1B1 388GG and 521CC mutations lead to a decrease of the inhibition and apoptotic rate of breast cancer cells, compared to the OATP1B1 group, although the differences were not statistically significant.

Breast cancer cells in the G0/G1 phase were reduced in the OATP1B1 388GG and 521CC groups, compared to the OATP1B1 group. Breast cancer cells in the S and G2/M phases were increased in the OATP1B1 388GG and 521CC groups, compared to the OATP1B1 group, although the differences were not statistically significant. Therefore, pre-synthesis DNA of breast cancer cells in the OATP1B1 388GG and 521CC group were reduced, while DNA increased in the synthesis and post-synthesis phase, causing the decreased inhibition rate of MCF-7 cells, thereby reducing the effect of TAM on MCF-7 cells. This conclusion is consistent with the inhibition rate and apoptosis results.

In summary, our present investigation has shown that OATP1B1 388GG and 521CC mutations may inhibit the activity of the OATP1B1 protein. OATP1B1 388GG and 521CC mutations may lead to restraining of the turn-over capacity of OATP1B1 and a reduction of TAM that is internalized into MCF-7 cells, resulting in weakened TAM treatment for breast cancer.

Nozawa et al (22) and Iwai et al (23) employed HEK293 cells and used estrone sulfate and estradiol-17β-glucuronide as a substrate, finding OATP1B1 388GG and 521CC had no significant effect on OATP1B1 absorbing ability, consistent with the results of the present study. OATP1B1 388GG and 521CC mutations cause decreased capacity of OATP1B1 in absorbing TAM, without statistical significance compared with the gene groups that have not mutated. This occurs due to OATP1B1 388GG and 521CC mutations being single mutation sites, and these changes of the DNA sequence result in such limited change of the turn-over capacity that effective change for the turn-over capacity cannot be markedly inhibited. This phenomenon is extremely common in gene mutations. The results of that study found that CYP2D6*4 or *10 mutations have a limited impact on the efficacy of TAM, and the difference was not statistically significant (24). Previous findings have shown that CYP2C9*2 and *3 mutations have limited impact on the efficacy of TAM, and the difference was not statistically significant (25).

The ability of SLCO1B1*15 (388G/521C) in absorbing estrone sulfate and estradiol-17β-glucuronide was found to be significantly reduced (26), indicating that the double mutation (SLCO1B1*15) may have greater significance on the transport of TAM. In summary, 388GG and 521CC of OATP1B1 gene polymorphism expression platforms were established in this experiment. The results show that...
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OATP1B1 388GG and 521CC mutations lead to the decreased turn-over capacity of OATP1B1 and a reduction of TAM in the treatment of breast cancer, although the difference was not statistically significant. Thus construction of a OATP1B1*15 double mutation plasmid to study the changes in drug absorption should be carried out in future investigations.

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