Therapy-relevant aberrant expression of MRP3 and BCRP mRNA in TCC-/SCC-bladder cancer tissue of untreated patients

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Abstract. Multidrug resistance (MDR) is a critical factor, which results in suboptimal outcomes in cancer chemotherapy. One principal mechanism of MDR is the increased expression of ATP-binding cassette (ABC) transporters. Of these, multidrug resistance-associated protein 3 (MRP3) and breast cancer resistance protein (BCRP) confer MDR when overexpressed in cancer cell lines. We measured the mRNA expression of MRP3 and BCRP in primary untreated bladder cancer specimens using reverse transcription-quantitative PCR (RT-qPCR) in comparison to normal bladder tissue. The MRP3 and BCRP expression in the two major histotypes of bladder cancer; transitional cell carcinoma (TCC; urothelial type of bladder cancer) and squamous cell carcinoma (SCC; ‘Schistosoma-induced’ bladder cancer) were compared. Furthermore, the association between MRP3 and BCRP expression and tumor grade and stage were investigated. MRP3 mRNA expression in bladder cancer specimens was increased ~13-fold on average compared to normal bladder tissue (n=36, P<0.0001). BCRP mRNA expression was decreased in bladder cancer specimens to ~1/5 on average, compared to normal bladder tissue (n=38, P<0.0001). TCC showed significantly increased MRP3 mRNA expression compared to SCC of the bladder (P<0.0001). BCRP mRNA expression was similar in TCC and SCC of the bladder (P=0.1072). The increased MRP3 mRNA expression was not related to bladder tumor grade (P=0.3465) but was, however, significantly higher in superficial than in invasive bladder tumors (P=0.0173). The decreased expression of BCRP was not related to bladder tumor grade (P=0.1808) or stage (P=0.8016). The current data show that bladder cancer is associated with perturbed expression of MRP3 and BCRP. Representing drug resistance factors, determining the expression of these transporters in native tumors may be predictive of the outcome of chemotherapy based-treatment of bladder cancer.

Introduction

Worldwide, bladder cancer is the ninth most common cause of cancer for both sexes combined. The highest recorded annual incidence and mortality rates of bladder cancer are that found in Egypt (1-3). TCC of the urinary bladder is the major histological cell type of non-Bilharzial bladder cancer, hence is the major histological cell type in Western countries. SCC, the major histological cell type of bilharzial bladder cancer represents less than 5% of the cases of bladder cancer in Western countries (2). In Egypt, both SCC and TCC are predominant (4) where schistosomiasis/Bilharziasis as well as increased smoking and exposure to environmental carcinogens prevail.

ABC transporters constitute a superfamily of 49 transport proteins, organized into seven subfamilies, ABCA - ABCG, on the basis of their sequence homology and domain organization (5,6). ABC transporters play an important role in limiting cellular exposure to toxic xenobiotics and carcinogens by actively pumping their substrates out of the cells. In cancer cells, overexpression of ABC transporters leads to decreased cellular drug accumulation below cytotoxic levels (7). Thus, ABC transporters contribute to MDR in cancer cells. Historically, the first of the human ABC transporters to be identified as a multidrug resistance protein was the P-glycoprotein (MDR1/ABCB1). In addition, the multidrug resistance-associated protein (MRP/ABCC) family of ABC transporters is included, which consists of nine proteins, eight of which have been determined to function as cellular resistance factors for various anticancer agents (8). The MRP1/ABCC1 is the founding member of the
MRP family of ABC transporters and is nearly ubiquitously expressed, while the MRP2/ABCC2 is present mainly in the canalicular membrane of hepatocytes. The third member of the MRP membrane transporter family is the MRP3/ABCC3. The MRP3 shares a high degree of amino acid homology with MRP1 (58%) (9-11) and is a 170 kDa protein with 18 transmembrane domains encoded by the ABCC3 gene (chromosome 17q21.3). While in the liver (basolateral membrane of cholangiocytes and hepatocytes), pancreas, and the intestinal tract (basolateral membrane of enterocytes; small, intestine, and colon) MRP3 is expressed at high levels (9-11), the urinary bladder shows lower, yet considerable levels of MRP3 mRNA (12). With respect to its physiological function, MRP3 is involved in the transport and, hence, regulation of bile salts, transports leukotrienes as well as glucuronic acid conjugates and glucosides. In MRP3-transfected cancer cell lines it was demonstrated that MRP3 has the ability to transport certain classes of cytotoxic anticancer agents (21-23), mainly drug compounds with a molecular weight above 0.45-0.5 kDa as well as drug or metabolite conjugates. Therefore, increased MRP3 expression during carcinogenesis implies possible involvement of MRP3 in the acquisition of an ab initio drug-resistant phenotype in some tumors.

A second unique ABC transporter is the breast cancer resistance protein (BCRP/ABCG2). The BCRP transporter is a 72 kDa ‘half transporter’ encoded by the ABCG2 gene, consisting of six transmembrane domains and functions as a homodimer or homotetramer. It belongs to the G subfamily of ABC transporters of the National Cancer Institute (NCI), Cairo University and the German University in Cairo, and informed consents were obtained in accordance with the Declaration of Helsinki. A total of 44 biopsies from patients with primary untreated bladder tumors were collected from the endoscopy unit, National Cancer Institute-Cairo, Egypt after obtaining informed consent. Immediately after transurethral resection, specimens were submerged in approximately 10 volumes of RNAlater RNA Stabilization Reagent (Qiagen). All specimens were less than 0.5 cm thick for effective RNA stabilization. Specimens were incubated overnight at 4°C before storage at -20°C until RNA extraction process started. Before RNA isolation, each specimen was cut longitudinally into two equal fragments. One fragment was used for RNA isolation and subsequent gene expression analysis. The other fragment was used for preparation of hematoxylin and eosin stained paraffin embedded tissue sections for histopathological examination. Specimens were washed two times for 15 min in phosphate buffered saline (PBS) before preparation of hematoxylin and eosin stained paraffin embedded tissue sections. In addition, routine pathology examination reports for those patients were obtained from the National Cancer Institute for further confirmation of diagnosis. The 1997 TNM stage and the 1973 WHO grade classifications were used.

Total RNA isolation and cDNA synthesis. Total RNA was isolated from 30 mg RNA-stabilized bladder cancer specimens using RNeasy Plus Mini kit (Qiagen) according to the manufacturer’s directions. Tumor sample total RNA was eluted in 40 µl nuclease-free water and stored at -80°C. Total RNA integrity and purity were electrophoretically verified by ethidium bromide staining and by A260/A280 nm absorption ratio >1.85. Total RNA (5 µg) was reverse transcribed using Sprint™ PowerScript™ PrePrimed Single Shots kit (Clontech Laboratories, Inc.) according to the manufacturer's directions in a final reaction volume of 20 µl.

RT-qPCR. The mRNA levels of BCRP and MRP3 were measured by RT-qPCR based on SYBR Green I chemistry and quantified using MX3005P™ quantitative real-time PCR system (Stratagene). In addition, the mRNA levels of a reference gene, aminoacyl tRNA synthetase complex-interacting multifunctional protein 1 (AIMP1/EMAP) was measured and used to normalize the mRNA levels of the BCRP and MRP3. The primer pairs for the BCRP, MRP3, and AIMP1 (Table I) were as previously described (17-19), and purchased from Metabion GmbH. The forward and reverse primers were run in a BLASTn for aligning their sequences against human transcript database to ensure that the primers do not coamplify homologous transcripts. Primers amplify across an intron/exon boundary or span large introns, thereby preventing amplification of residual genomic DNA. RT-qPCR was performed in triplicate reactions using Power SYBR Green PCR Master Mix (Applied Biosystems) for BCRP expression analysis and QuantiFast SYBR Green PCR Master Mix (Qiagen) for MRP3. Each reaction mixture contained 12.5 µl Master Mix, 1 µl (~50 ng) tumor sample cDNA as a template, 300 nM forward and reverse primers for BCRP and MRP3 and 200 nM forward and reverse primers for AIMP1 in a final reaction volume.

Materials and methods

Tumor specimen acquisition and preservation. The use of all human tissue was approved by the local ethics commit-
of 25 µl. Reactions were pipetted in semi-skirted 96-well plates (Eppendorf AG) and sealed with optical adhesive PCR film (Eppendorf AG). For BCRP expression analysis, denaturation was performed at 95°C for 10 min, followed by 40 PCR cycles with the following specifications: 95°C for 30 sec, 60°C for 1 min. For MRP3 expression analysis, denaturation was performed at 95°C for 10 min, followed by 40 PCR cycles with the following specifications: 95°C for 15 sec, 60°C for 20 sec, 72°C for 20 sec. No template controls (NTCs) were included in all the assays. At the end of each RT-qPCR experiment, a dissociation curve was generated using the following thermal profile: 95°C for 1 min, 55°C for 15 sec, 60°C for 20 sec. The specificity of each assay was further verified on 1% agarose gels (Fig. 2B). The amplification behavior of the AIMP1 was comparable to that of the BCRP and MRP3. The AIMP1 was tested in 15 specimens representing the different histotypes, grades and stages of bladder cancer against β-actin, a commonly used reference gene in various studies on bladder cancer and its expression was found to be stable.

<table>
<thead>
<tr>
<th>Gene name</th>
<th>GenBank accession number</th>
<th>5' to 3' Primer sequence (forward/reverse)</th>
<th>BLASTn score - E-value (forward/reverse)</th>
<th>Size of amplicon (forward/reverse)</th>
<th>Location of the primers (forward/reverse)</th>
</tr>
</thead>
<tbody>
<tr>
<td>BCRP</td>
<td>NM_004827</td>
<td>TGGCTGTCATGGCTTCAGTA/GCCACGTGATTCTCCTACAA</td>
<td>40.1-0.015/40.1-0.015</td>
<td>206 bp</td>
<td>1690-1709/1895-1876</td>
</tr>
<tr>
<td>MRP3</td>
<td>NM_003786</td>
<td>CCAAGGCACATTTGAGCCAG/CCTCTGCACCTTCCAAACGC</td>
<td>42.1-0.005/38.2-0.038</td>
<td>263 bp</td>
<td>2357-2377/2617-2599</td>
</tr>
<tr>
<td>AIMP1</td>
<td>NM_004757</td>
<td>TGCTTTCCCCAGGAGGCC/CACCCCAAGGGAACTCCCTTTG</td>
<td>36.1-0.15/44.1-0.001</td>
<td>124 bp</td>
<td>762-779/885-865</td>
</tr>
</tbody>
</table>

The efficiency-adjusted ∆∆Cq method for the calculation of relative expression ratios (R) takes into consideration the difference between PCR amplification efficiency (E) of the target gene and the reference gene. Expression of the drug transporters in the bladder cancer specimen were compared to control/calibrator samples. The controls/calibrators were two normal bladder cDNA samples reverse transcribed from human bladder total RNA purchased from two suppliers; Ambion, Inc. (1 subject) and Clontech Laboratories, Inc. (20 subjects). Relative expression ratios against these two calibrators were averaged according to:

$$R = \frac{(E_{target})^{\Delta Ct_{target(control - unknown)}}}{(E_{reference})^{\Delta Ct_{reference(control - unknown)}}}$$

The log transform of relative expression ratios were used in the statistical analysis. Data were tested for normality using the Kolmogorov and Smirnov normality test. Extreme outliers were tested using the Grubb's test at P=0.01, as described in the International Standard Organization document ISO 5725-2 (23). The two-tail one-sample t-test was used to test for significant difference in BCRP and MRP3 mRNA expression in cancer versus normal bladder tissue. The relation between the relative BCRP and MRP3 mRNA expression and tumor histological type, grade, and stage were investigated using the parametric two-tail unpaired t-test, or one-way ANOVA.
when appropriate. The interrelationship between BCRP and MRP3 was tested using Pearson correlation ($r_p$). All the statistical tests were performed using the GraphPad Prism version 5.00 for Windows, GraphPad Software, San Diego, CA, USA (www.graphpad.com). A difference was considered statistically significant at $P<0.05$.

**Results**

**Malignancy characterization.** The characteristics of the bladder cancer patients with respect to age and sex are summarized in Table II. Moreover, the overview summarizes patient statistics during a 2 months period at the National Cancer Institute-Cairo University, showing that the majority of incoming patients were male. Data given in Table II also include the pathophysiological characteristics for resected tissues: histological type, grade, and invasiveness of their bladder neoplasms. In 6 out of 44 patients undergoing transurethral resection of bladder tumors, no evidence for malignancy was detected through histopathology. In 2 tissue samples with SCC, indications for bilharzia infections were detected, i.e., Schistosoma eggs were present.

**BCRP and MRP3 mRNA expression.** mRNA expression of BCRP was quantified in 44 bladder tumor specimens relative to normal bladder tissue represented as two normal bladder cDNA samples reverse transcribed from human bladder total RNA purchased from two suppliers; Ambion, Inc. (1 subject) and Clontech Laboratories, Inc. (20 subjects). Histopathological examination revealed that 6 of the 44 specimens collected had no evidence of malignancy. In 95% of the bladder cancer specimens (36 of 38 specimens), the BCRP mRNA expression was less than in normal bladder tissue (log relative expression ratio <0). The decreased expression of BCRP was statistically significant; mean: -0.6832±0.07950, 95% CI: -0.8443 to -0.5220, $P<0.0001$. This means that BCRP mRNA expression was decreased to ~1/5 on average, relative to normal bladder tissue. The MRP3 expression was measured in 36 of the 38 bladder cancer specimens. In 97% of the bladder cancer specimens (35 of 36 specimens), the MRP3 expression was higher than in normal bladder tissue (log relative expression ratio >0). The increased expression of MRP3 was also statistically significant; mean: 1.117±0.09456, 95% CI: 0.9245 to 1.309, $P<0.0001$. This means that MRP3 mRNA expression was increased to ~13-fold on average compared to normal bladder tissue. Fig. 3 summarizes the pattern of expression of BCRP and MRP3 mRNA in bladder cancer specimens as well as specimens with no evidence of malignancy. Analysis with regard to the potential interrelationship between BCRP and MRP3 revealed that the BCRP mRNA levels were not related to those of MRP3; $r_p$=0.1245, $P=0.4693$ (Fig. 4).

**BCRP and MRP3 mRNA expression versus histopathological parameters of bladder cancer.** The various pathological processes, as characterized via the given histological parameters, that could affect the decreased or increased expression of BCRP or MRP3, respectively, were biometrically investigated by employing multivariate analysis. The first parameter tested is the bladder tumor histotype; TCC or SCC of the bladder. Only specimens with pure TCC or SCC phenotype were included in this analysis. The relative expression of BCRP mRNA was similar in TCC and SCC of the bladder cancer specimens as determined by two-tail unpaired t-test; $P=0.1072$. The relative expression of MRP3 mRNA was, however, significantly...
higher in TCC than in SCC of the bladder cancer specimens; P=0.0101. Fig. 5 summarizes the pattern of expression of BCRP and MRP3 mRNA in TCC vs. SCC of the bladder.

The second parameter tested is the degree of bladder tumor differentiation or bladder tumor grade; G1, G2, G3. The decreased expression of BCRP mRNA was not related to the degree of bladder tumor differentiation (G1, G2, and G3); P=0.1808, nor was the increased expression of MRP3 mRNA to the degree of bladder tumor differentiation; P=0.3465. Fig. 6 summarizes the comparison of BCRP and MRP3 mRNA expression to bladder tumor grade; G1, G2, and G3.

The third parameter tested is the invasiveness of bladder tumor; superficial (Ta/Tis/T1) vs. muscle-invasive (>T1) bladder tumors. The expression of BCRP mRNA was similar in superficial (Ta/Tis/T1) and muscle-invasive (>T1) bladder tumors; P=0.8016. The expression of MRP3 mRNA was, however, significantly higher in superficial than in invasive bladder tumors; P=0.0173. Fig. 7 summarizes the comparison of BCRP and MRP3 mRNA expression to the invasiveness of bladder tumor.

**Discussion**

Being expressed at the epithelial barriers of several tissues, ABC transporters are believed to be a component of the lines of defense of tissues by actively effluxing toxic endobiotics and xenobiotics out of cells (7,9,14,24). On the other hand, overex-
expression of ABC transporters in cancer cell lines is associated with resistance to a variety of commonly used chemotherapeutic agents (9,25,26), a phenomenon which directed several studies to investigate the relevance of ABC transporters in clinical MDR. However, to ascertain their relevance in clinical MDR, studies investigating the expression of ABC transporters in specific tumor types are still needed (27-30). In addition, there are very few studies examining the expression of ABC transporters in native (untreated) cancer tissues making the potential role that ABC transporters play in the pathophysiology of cancer and/or clinical MDR still unknown (16).

Of interest, BCRP and MRP3 are expressed at the transitional epithelium of normal bladder. However, their expression data in primary TCC and SCC bladder tumors are still lacking. In the present study we measured the mRNA levels of MRP3 and BCRP in untreated primary bladder tumors compared to normal bladder tissue using RT-qPCR. Furthermore, the relative mRNA expression in the two major histotypes of bladder cancer; TCC and SCC were compared. In addition, the association between the mRNA expression and tumor characteristics of grade (degree of tumor differentiation) and stage (invasiveness of bladder tumor) were investigated. RT-qPCR is considered the prime technique and the method of choice to measure differential gene expression (20,31,32). Its high sensitivity enables the detection of small differences in mRNA expression that other commonly used methods such as immunohistochemistry or western blotting would fail to, as in cases of measuring gene expression of ABC transporters (33-35). The levels of BCRP mRNA were previously reported to be consistent with the protein expression as determined by western blot analysis in head and neck squamous cell carcinoma cell lines (36). Similar observation was shown when MRP3 protein levels were analyzed in breast cancer by immunohistochemistry (37).

Downregulation of BCRP mRNA. Significant downregulation of BCRP was detected in bladder cancer compared to normal bladder tissue in the current study. Although BCRP mRNA expression in SCC of the bladder was lower than that in TCC of the bladder, the difference between cell types was not statistically significant. This clearly indicates that downregulation of BCRP in bladder cancer occurs in general, irrespective of the histological type of the tumor. Previous studies investigating the expression of BCRP in bladder cancer documented no or very weak expression of the BCRP protein (38,39). In another study on the expression of BCRP in locally advanced bladder cancer, positive immunohistochemical staining of BCRP protein was
found in only 23 (28%) of 82 patients and expression was not associated with the therapeutic outcome (40). Doyle and Ross detected mRNA expression of BCRP in normal bladder tissue using a commercially available dot blot that contained RNA from 50 human tissues (7). In addition, Fetsch et al confirmed the localization of BCRP protein at the transitional epithelium of normal bladder (14). To our best knowledge, this is the first study to report decreased expression of BCRP mRNA in the different histotypes of untreated bladder tumors relative to normal bladder tissue.

Previously, low mRNA levels of BCRP were documented in breast cancer (41,42). In addition, immunohistochemical staining of breast cancer tissues revealed positive staining in normal duct cells versus no staining in carcinoma cells (41). In addition, BCRP mRNA expression was significantly lower (approximately 100-fold) in tumoral colon tissue compared to the normal colon (43). Using immunohistochemistry, strong BCRP staining was observed in only 11 of 41 gallbladder carcinoma specimens, while the majority of the normal and cholelithiasis samples showed intense staining (44). In addition, BCRP mRNA and protein were decreased several-fold in human colorectal cancer tissues as well as in cervical cancer tissues in comparison to their matched normal tissues (16); a result that drove the investigators to examine BCRP expression using a commercial array of cDNA from 19 different cancer tissues, including bladder cancer, paired with their matched cDNA from normal tissue from the same patient. Significant downregulation was observed in breast, ovarian, lung, kidney, liver, prostate, uterine, rectal, thyroid, testicular, as well as colon and cervical cancer (16); a result that led the investigators to hypothesize that downregulation of BCRP is a widespread phenomenon in human cancers. In contrast to our finding that BCRP is downregulated in bladder cancer, the study of Gupta et al showed no significant difference in BCRP mRNA expression between cancer and normal bladder tissue (16). However, the number of bladder cancer samples in the cDNA array hybridization used in their study was only 5 samples compared to 38 bladder cancer samples included in our study, besides the high sensitivity of quantitative real-time PCR compared to array hybridization (45). Taken together and in addition to our finding that BCRP is downregulated in the different histotypes of untreated bladder tumors relative to normal bladder tissue.

Localization of BCRP protein at the transitional epithelium of normal bladder (14) would decrease cellular accumulation of its substrates of chemical carcinogens, hence, protect the bladder epithelium against chemical-induced carcinogenesis. A variety of chemical carcinogens are known to be transported by BCRP. These include; 2-amino-1-methyl-6-phenylimidazo[4,5-
bpyridine (PhIP), 2-amino-3-methylimidazo[4,5-f]quinoline (IQ), 3-amino-1,4-dimethyl-5H-pyrido[4,3-b]indole (Trp-P1), aflatoxin B1, 2-amino-3,8-dimethylimidazo[4,5-f]quinoline (MeIQx), and the sulfate and glucuronide conjugates of benzo[a]pyrene (BP) (46-49). Decreased expression of BCRP in bladder cancer would lead to increased cellular accumulation of such chemical carcinogens increasing DNA damage, mutations, and subsequently increasing malignant transformations.

Given that protoporphyrin IX is a substrate of BCRP (50-52) and is the precursor for heme, downregulation of BCRP would lead to the accumulation of protoporphyrin IX and heme in cancer cells. Increased cellular levels of protoporphyrin IX and heme generates potentially DNA damaging reactive oxygen species (53). Downregulation of BCRP by RNA interference technology and the FTC (a BCRP inhibitor) treatment of embryonic stem cells resulted in increased cellular levels of protoporphyrin IX and reactive oxygen species (53). Moreover, DNA damage signals were triggered suggesting that enforced downregulation of BCRP in embryonic stem cells resulted in DNA damage as a result of the increased cellular levels of reactive oxygen species (53). In addition, heme is a cofactor for the product of the iNOS gene; the inducible nitric oxide synthase. Inducible nitric oxide synthase utilizes arginine to synthesize nitric oxide (NO). Previously, it has been demonstrated that malignancy is associated with overexpression of the iNOS gene inducing oxidative stress through the overproduction of nitric oxide (54,55). Therefore, downregulation of BCRP may provide an explanation for the increased availability of heme required for the increased activity of inducible nitric oxide synthase in cancer cells. NO can damage DNA and itself induces angiogenesis therefore has the potential to promote tumor development, growth, as well as invasiveness and metastatic ability (55).

In our study, decreased BCRP mRNA expression was not associated with high grade or invasive bladder tumors suggesting that decreased BCRP mRNA expression can rather be an early event in bladder tumorigenesis. The most intensively investigated solid tumor type was breast cancer (33,41,42). Similar to our results Burger et al documented no association between BCRP mRNA expression and tumor grade or stage of breast cancer (33). Similarly, Kanzaki et al reported that BCRP mRNA expression was independent of tumor stage (42). In contrast, Faneyte et al reported that high tumor malignancy grade was associated with decreased BCRP mRNA expression in breast cancer (41). In pancreatic carcinoma, there was no correlation of the mRNA expression level to carcinoma stages or grades (56).

**Potential impact of expression profiles on cytotoxic therapies.** Previously, it has been reported that BCRP plays a role in the maintenance of cellular folate homeostasis (61) by its unique capability to export mono-, di-, and triglutamates of folates out of cells (62,63). In contrast, other ABC transporters (MRP1 through MRP4) have restricted ability to export only monoglutamate forms of folates. Cellular retention of folates is aided by the addition of glutamate residues by the folylpoly-γ-glutamate synthase (FPGS) enzyme (64). Neoplastic dividing cells have an absolute requirement for reduced folates, in order to properly initiate and complete DNA replication and mitosis. This is due to the fact that reduced folate cofactors play a key role in one-carbon transfer reactions in the de novo biosynthesis of purines and thymidylate. Ifergan et al showed that downregulation of BCRP expression and efflux activity is an essential component of cellular survival under conditions of folate deficiency (61). Thus, in the context of malignancy, downregulation of BCRP is consistent with the need to increase cellular retention of folates to support the requirements of the rapidly dividing cancer cells for DNA synthesis.

Of interest, methotrexate which is commonly used in drug combination regimens for treatment of bladder cancer has a susceptibility to polyglutamation. The ability of MRP3 to export polyglutamated forms of methotrexate is limited (65). This suggests that bladder cancer would respond optimally to anticancer regimens including methotrexate. The fact that polyglutamation of methotrexate results in massive enhancement of cytotoxicity is consistent with our finding that BCRP is downregulated while MRP3 is upregulated in bladder cancer. Therefore, polyglutamated methotrexate will be retained into the cells and the increased MRP3 expression will not compensate for the decreased BCRP expression in effluxing polyglutamated methotrexate.
In conclusion, bladder cancer is associated with perturbed expression of BCRP and MRP3. BCRP expression is decreased in TCC and SCC of the bladder with no significant difference between the two histotypes of bladder tumors. MRP3 expression is, however, increased in TCC and SCC of the bladder with TCC expressing MRP3 at significantly higher levels than SCC. The manner in which the two transporters are aberrantly regulated in bladder cancer needs further investigation. Whether this aberrant expression has the potential to affect therapeutic outcome of cancer chemotherapy needs further investigation as well. Moreover, the aberrant expression of BCRP could be linked to the pathogenesis of bladder cancer.

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