EXPRESSION, LOCALIZATION, AND FUNCTIONAL CHARACTERISTICS OF BREAST CANCER RESISTANCE PROTEIN IN CACO-2 CELLS

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ABSTRACT:

The function of breast cancer resistance protein (BCRP) and its role in drug absorption, distribution, and elimination has recently been evaluated. The objective of the present study was to examine the expression, localization, and functional characteristics of BCRP in Caco-2 cells, a widely used human intestinal epithelial cell model for investigating intestinal drug absorption. The expression of BCRP in Caco-2 cells was measured by Western blotting using the antibody BXP-21. Localization of BCRP was determined by an immunofluorescence technique using both antibodies BXP-21 and BXP-34. The drug efflux function of BCRP was evaluated via the epithelial transport of methotrexate (MTX) and estrone-3-sulfate (E3S) across Caco-2 cell monolayers in the presence or absence of the BCRP inhibitors Ko143 or GF120918. The drug efflux function of BCRP was evaluated via the epithelial transport of methotrexate (MTX) and estrone-3-sulfate (E3S) across Caco-2 cell monolayers in the presence or absence of the BCRP inhibitors Ko143 or GF120918. The drug efflux function of BCRP was evaluated via the epithelial transport of methotrexate (MTX) and estrone-3-sulfate (E3S) across Caco-2 cell monolayers in the presence or absence of the BCRP inhibitors Ko143 or GF120918.

Breast cancer resistance protein (BCRP) was originally cloned and sequenced from genomic DNA, from highly mitoxantrone-resistant S1-M1-80 human colon carcinoma cells, and from MCF7 AdVp human breast cancer cells selected in doxorubicin (Doyle et al., 1998; Miyake et al., 1999). BCRP is a member of the ATP-binding cassette transporter G family and is also known as ABCG2 or ABCP or MXR (Ejendal and Hrycyna, 2002; Doyle and Ross, 2003). It is a 655-amino acid polypeptide (72 kDa), containing six putative transmembrane domains and four potential N-glycosylation sites. BCRP is similar to half the duplicated P-glycoprotein (P-gp) or multidrug resistance protein 1 (MRP1) molecules and functions as a homodimer bridged by disulfide bonds (Doyle et al., 1998; Kage et al., 2002).

BCRP is endogenously expressed at a high level in human placenta and to a lesser extent in liver, small intestine and colon, ovary, veins, capillaries, kidney, adrenal, and lung, with little to no expression in brain, heart, stomach, prostate, spleen, and cervix (Doyle et al., 1998; Litman et al., 2001; Maliepaard et al., 2001; Scheffer and Schepfer, 2002). Importantly, BCRP is expressed in the human jejunum at levels considerably higher than those of many other ABC transporters (Taipalensuu et al., 2001). BCRP has been demonstrated to exist on the apical membrane of intestinal epithelium and has limited the oral absorption of topotecan in mice and humans (Jonker et al., 2000; Kruijtzer et al., 2002a). Given the liver and intestinal localization pattern, BCRP, similar to P-gp, may act as a barrier to uptake and absorption and limit the oral bioavailability of drugs as well as mediating hepatobiliary excretion of drugs (Jonker et al., 2000; Jorritsma et al., 2002; Kruijtzer et al., 2002b).

Caco-2 cells are derived from human colonic adenocarcinoma cell line, and exhibit morphological and functional similarities to intestinal enterocytes. It has been widely used as a model of human intestinal epithelium for studies of intestinal drug absorption and metabolism. Many active transport systems such as P-gp (encoded by MDR-1 gene, also named ABCB1) and multidrug resistance-associated protein 2 (MRP2, or ABCC2) have been characterized in Caco-2 cells (Makhey et al., 1998; Doppenschmitt et al., 1999; Gutmann et al., 2000). BCRP mRNA has been detected in Caco-2 cells, although its level is 100-fold lower than that in human jejunum (Taipalensuu et al., 2001).

The present study was aimed at characterization of the protein expression, localization, and efflux function of BCRP in Caco-2 cells. We have used BXP-21 monoclonal antibody (mAb) to determine the

ABBREVIATIONS: BCRP, breast cancer resistance protein; ABC, ATP-binding cassette transporter; A-to-B, apical-to-basolateral; B-to-A, basolateral-to-apical; E3S, estrone-3-sulfate; FBS, fetal bovine serum; MDR-1, human multidrug resistance gene 1; mAb, monoclonal antibody; MRP, multidrug resistance-associated protein; MTX, methotrexate; Papp, apparent permeability coefficient; P-gp, P-glycoprotein; TEER, trans-epithelial electrical resistance; GF120918, N-(4-[2-(1,2,3,4-tetrahydro-6,7-dimethoxy-2-isouquinolinyl]ethyl]-phenyl)-9,10-dihydro-5-methoxy-9-oxo-4-acridine carboxamide; LY335979, zosuquidar trihydrochloride; MK571, 3-[3-[(7-chloroquinolin-2-yl)vinyl]phenyl]-(2-dimethylcarbamoyl-ethylsulfanyl)methylsulfanyl]propionic acid; 2,4-DNP, 2,4-dinitrophenol; FITC, fluorescein isothiocyanate.
BCRP protein expression, and BXP-21 and BXP-34 mAbs to characterize the subcellular distribution of BCRP in Caco-2 cells. In addition, the efflux function of BCRP in Caco-2 cells was determined using estrone-3-sulfate (E3S) and methotrexate (MTX) as substrates. Knowledge of the properties of BCRP in Caco-2 cells is valuable to investigate the absorption mechanism of drug molecules using this in vitro model system.

Materials and Methods

Materials. [3H]Estrone-3-Sulfate (specific activity, 45 Ci/mmol) and [3H]methotrexate (specific activity, 33.5 Ci/mmol) were purchased from Moravek Biochemicals (Brea, CA). GF120918 and LY335979 were synthesized at Millenium Pharmaceuticals, Inc. Ko143 was obtained from the Netherlands Cancer Institute (Amsterdam, Netherlands). MK571 was purchased from Alexis Biochemicals (San Diego, CA). Unlabeled E3S, MTX, prazocin, and 2,4-dinitrophenol (2,4-DNP) were purchased from Sigma-Aldrich (St. Louis, MO). Cell culture media and supplies were obtained from Invitrogen (Carlsbad, CA). BXP-21 and BXP-34 murine monoclonal antibodies were purchased from Signet Laboratories (Bedford, MA). Alexa Fluor isothiocyanate (FITC)-conjugated goat anti-mouse IgG was obtained from Abcam Inc. (Cambridge, MA). Horseradish peroxidase-conjugated goat anti-mouse IgG2a, mouse IgG2a, and mouse polyclonal antibody 11B2 for β-actin were purchased from Santa Cruz Biotechnology, Inc. (Santa Cruz, CA). All supplies for Western blot studies were obtained from Invitrogen.

Caco-2 Cell Culture. The Caco-2 cells (passage 18) were obtained from American Type Culture Collection (Manassas, VA). The stock cells were cultured in 150-cm² tissue culture T-flasks for subsequent plating onto 24-Transwell plates (0.33 cm²/well, 0.4-μm pore size; Costar, Cambridge, MA). Briefly, 1 × 10⁶ cells were suspended in 0.2 ml of culture medium [Dulbecco’s modified Eagle’s medium with 0.1 mM nonessential amino acids, 2 mM l-glutamine, 4.5 g/l glucose, and 10% fetal bovine serum (FBS)]. Cell cultures were cultured in 150-cm² tissue culture T-flasks for subsequent plating onto 24-Transwell plates (0.33 cm²/well, 0.4-μm pore size; Costar, Cambridge, MA). The confluent cell monolayers were obtained within 5 to 7 days after plating. The transepithelial electrical resistance (TER), measured as an epithelial volt-ohm-meter (World Precision Instruments, Inc., Sarasota, FL), progressively increased and reached a plateau after day 5, indicating the formation of tight junctions. Cell monolayers were obtained within 5 to 7 days after plating. The transepithelial electrical resistance (TEER), as measured by an epithelial volt-ohm-meter (World Precision Instruments, Inc., Sarasota, FL), gradually increased and reached a plateau after day 5, indicating the formation of tight junctions.

Western Blotting. Cells were scraped and subsequently lysed in hypotonic lysis buffer, consisting of 150 mM NaCl, 50 mM Tris-HCl (pH 7.4), 1% SDS, 1% Triton X-100 supplemented with protease inhibitors (aprotinin, leupeptin, and phenylmethylsulfonyl fluoride) (Yu and Sinko, 1997). Cell lysates were sonicated and centrifuged at 10,000 rpm for 20 min. The supernatant was aliquoted and stored at −80°C. Protein levels were determined using the BCA protein assay (Pierce Endogen, Rockford, IL). Cells lysates, with and without treatment of reducing agent (7–13% dithiothreitol), were separated on a 4 to 12% gradient polyacrylamide gel and subsequently transferred electrophoretically to a polyvinylidene difluoride membrane. Proteins were hybridized using BXP-21 (1:150 optimal dilution) and horseradish peroxidase-conjugated goat anti-mouse IgG2a (1:1000), and further visualized by enhanced chemiluminescence (ECL) (Amersham Biosciences Inc., Piscataway, NJ). The protein bands were semi-quantified by NIH Image Software.

Immunohistochemistry. Cell monolayers cultured on the Transwells were washed with phosphate-buffered saline, pH 7.4; similar phosphate-buffered saline washes were included between each subsequent step. Cell monolayers were fixed in 3.7% (v/v) formaldehyde solution for 30 min at room temperature. Cell membranes were permeabilized by saponin (2%, 2 min at room temperature). Nonspecific binding sites were blocked by incubation in 10% FBS for 45 min. Cells were incubated with anti-human BCRP mAb (BXP-21 or BXP-34 in 1% FBS) for 2 h. BCRP staining was revealed by incubation with FITC-conjugated goat anti-mouse antibody for 1 h. The nucleus was stained by propidium iodide for 15 min. After a final wash, cell monolayers were mounted in Vectashield before examination on a Zeiss PASCAL confocal laser scanning system (Carl Zeiss Inc., Thornwood, NY). Control for nonspecific staining was the replacement of BXP-21 or BXP-34 with a nonspecific antibody from the same class, mouse IgG2a.

Transport Studies in Caco-2 Cells. Bidirectional transport studies were performed at 37°C in air. Prior to each experiment, the confluent cell monolayers on Transwell inserts were washed and equilibrated for 30 min with transport medium [Hanks’ balanced salt solution containing 10 mM N-2-hydroxyethyl-piperazine-N’-2-ethanesulfonic acid (HEPES) and 10 mM glucose, pH 7.4]. The experiment was initiated by adding a solution containing the test compound to either the apical (for A-to-B transport) or basolateral (for B-to-A transport) compartment. When applicable, inhibitors such as GF120918 (2 μM), prazocin (100 μM), MK571 (50 μM), Ko143 (1 μM), or LY335979 (5 μM) were present in the transport medium of the donor side from the preincubation period throughout the permeability study. At 15, 30, 45, and 60 min, the sample aliquots of receiving solutions were withdrawn from the basolateral side (for A-to-B transport) or the apical side (for B-to-A transport), and replaced immediately with an equal amount of fresh transport media except at the 60-min time point (the end of the incubation). The samples were mixed with 5 ml of scintillation cocktail and the radioactivity was determined in a liquid scintillation spectrophotometer (Beckman Coulter, Fullerton, CA).

Data Analysis. The cumulative amount of drug (Q) on the receiver side was plotted as a function of time. The steady-state flux J was then estimated from the slope (dQ/dt). The apparent permeability coefficient (P_app) of unidirectional flux for the test compound was estimated by normalizing the flux J (mols) against the nominal surface area A (0.33 cm²) and the initial drug concentration in the donor chamber C_D (mol/ml), or \( P_{app} = \frac{J}{A \cdot C_D} \).

Results

Expression and Localization of BCRP. BCRP protein expression was determined in Caco-2 cell lysate samples via Western blot analysis using BXP-21 as the primary antibody. As shown in Fig. 1A, both the half-transporter BCRP (~70 kDa) and its dimer (~140 kDa) were observed in lanes 1 and 4, in which the Caco-2 cell lysates were not treated with reducing agent (7–13% of dithiothreitol). Lanes 2, 3, 5, and 6 are the lysates treated with reducing agent (7–13% of dithiothreitol), which showed a strong band of BCRP monomer (~70 kDa). Lanes 6 depicts a trace amount of BCRP dimer when a high concentrated unheated sample was loaded.

When Caco-2 cells were cultured in cell culture T-flasks, the expression of BCRP monomer and dimer in Caco-2 cells was more than 3-fold higher in the late passage [passages 56 (p56) or 59 (p59)] than in the early passage [passages 33 (p33) or 36 (p36)] after normalization to the expression of β-actin (Fig. 1, B and C). However, the Caco-2 cells were grown on the Transwell plates, the BCRP monomer expression was about 3-fold higher and the dimer expression was about 10-fold higher in the early passage (p36) than in the late passage (p59). The expression level of BCRP, after normalization to the housekeeping protein β-actin, did not change on the 5th, 12th, and 19th days after the cell seeding on Transwell plates (Fig. 1B).

Caco-2 cells (p33) cultured on the Transwell plates showed positive staining of BCRP in the presence of BXP-21 (Fig. 2A, upper panel) or BXP-34 (Fig. 2A, lower panel) under confocal laser-scanning microscopy (Fig. 2A). Two days after cell seeding (prior to confluence), BCRP staining was observed not only on both apical and basolateral...
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membranes but also in the cytoplasm. On the 5-day-old culture, cells reached confluence and showed increased TEER values (indicative of the tight junction formation); BCRP staining was observed only on the apical membrane, and there was no detectable staining in the cytoplasm, indicating the localization of BCRP on the apical membrane. The same phenomena were also noted during the cell differentiation period (i.e., the 12th and 19th days after cell seeding). When the late passage cells (p59) were grown on the Transwell plates for 12 days, multiple layers were observed as demonstrated by the multiple nuclei staining on the x-z and y-z views (Fig. 2B). BCRP staining, however, was still preferentially shown on the apical side of cell membranes (Fig. 2B). The absence of any green staining in the cell monolayers with mouse IgG2a and FITC-conjugated second antibody served as a negative control.

Efflux Function of BCRP. The function of BCRP as an efflux transporter in Caco-2 cell monolayers was evaluated using a known substrate, E3S (Suzuki et al., 2002, 2003). As shown in Table 1, the \( P_{\text{app}} \) value of \(^{[3]H}\)E3S (0.02 \( \mu \)M) from B-to-A was 46.1 \( \times 10^{-6} \) cm/s and was 8-fold higher than the \( P_{\text{app}} \) value (5.7 \( \times 10^{-6} \) cm/s) for A-to-B in the 5-day culture (p33). In 12- and 19-day cultures (p33), the \( P_{\text{app}} \) values for A-to-B decreased to 4.1 and 4.5 \( \times 10^{-6} \) cm/s, and the \( P_{\text{app}} \) value for B-to-A increased to 63.0 and 58.0 \( \times 10^{-6} \) cm/s, respectively. In the presence of Ko143 (a BCRP inhibitor), the A-to-B and B-to-A transports of E3S were almost equal, with \( P_{\text{app}} \) values around 9 to 11 \( \times 10^{-6} \) cm/s. Figure 3 showed the comparison of BCRP-mediated \(^{[3]H}\)E3S (0.02 \( \mu \)M) efflux in both early passage (p36) and late passage (p59) Caco-2 cell monolayers. In p36, the \( P_{\text{app}} \) value for A-to-B transport of E3S was 2.3 \( \times 10^{-6} \) cm/s and was 32-fold less than the \( P_{\text{app}} \) value for B-to-A transport. The \( P_{\text{app}} \) value increased to 6.1 \( \times 10^{-6} \) cm/s in the presence of Ko143. However, \( P_{\text{app}} \) values of E3S for A-to-B and B-to-A transport in p59 were 16.2 and 37.1 \( \times 10^{-6} \) cm/s, respectively. The difference in the directional E3S transport was abolished by the BCRP inhibitor Ko143. The TEER value was 4-fold higher in p36 than in p59 in the present cell culture system (Fig. 3).

When \(^{[3]H}\)E3S (0.03 \( \mu \)M) was coincubated with Ko143 (1 \( \mu \)M), GF120918 (2 \( \mu \)M) (an inhibitor for BCRP and P-gp), or prazocin (100 \( \mu \)M) (an inhibitor for BCRP and P-gp) in Caco-2 cell monolayers, the \( P_{\text{app}} \) value of E3S for the A-to-B transport increased and the \( P_{\text{app}} \) value for B-to-A transport decreased, resulting in the B/A ratio dropping from 27 to almost unity (Fig. 4). In contrast, the P-gp-specific inhibitor LY335979 (5 \( \times 10^{-6} \)M) did not significantly change the \( P_{\text{app}} \) value for A-to-B nor the \( P_{\text{app}} \) value for B-to-A of MTX was significantly increased in the presence of MK571 (a MRP inhibitor; 50 \( \mu \)M), Ko143 (1 \( \mu \)M), and GF120918 (2 \( \mu \)M) \((p < 0.05)\). However, neither the \( P_{\text{app}} \) value for A-to-B nor the \( P_{\text{app}} \) value for B-to-A of MTX was significantly affected after coincubation with LY335979 (5 \( \mu \)M).
The B-to-A efflux of rhodamine 123, which is a substrate for P-gp and mutant BCRP, was significantly decreased in the presence of LY335979 and GF120918 ($p < 0.05$), but was not changed by coincubating with Ko143 (1 $\mu$M) (Fig. 7).

**Discussion**

Western blot analysis of Caco-2 cell lysates demonstrates the presence of both monomer ($\sim 70$ kDa) and homodimer ($\sim 140$ kDa) of BCRP in the Caco-2 cells (Fig. 1A). The 140-kDa BCRP complex dissociated to 70-kDa polypeptides in the presence of the reducing agent dithiothreitol, indicating that the BCRP dimer was linked by disulfide bonds. This finding is consistent with the recent discovery by Kage et al. (2002). The same group also demonstrates the necessity of homodimerization for the BCRP function when using a dominant-negative mutation of BCRP with a L554P alteration in the fifth transmembrane domain (Kage et al., 2002).

Immunohistochemical analysis of Caco-2 cells reveals the presence of BCRP on apical and basolateral plasma membranes as well as in the cytoplasm prior to cell confluence (Fig. 2). Upon cell confluence, BCRP was sorted to the “apical membrane” consistent with potential intracellular localization and redistribution of BCRP to the plasma membrane shown in BCRP-overexpressed MCF-7 AdVp3000 and S1-M1-80 drug-resistant cells (Litman et al., 2000). BCRP-transfected MDCK or LLC-PK epithelial cells also exhibited polarized apical localization of BCRP (Jonker et al., 2000; Maliepaard et al., 2001), implying that BCRP may contain a certain endogenous sorting signal, which was recognized by epithelial cells (Jonker et al., 2000; Imai et al., 2003). The polarized apical BCRP distribution in Caco-2 cells continues during the cell differentiation period, which is consistent with the proposed role for BCRP as a secretory detoxifying transporter, contributing to the gastrointestinal epithelial barrier. Results from transport function assays using E3S as the BCRP substrate agreed with immunohistochemical study results. The directional transport of E3S across Caco-2 monolayers was observed immediately after cells reached confluence (5-day culture) and continued throughout the differentiation phase (day 12–19 cell culture) (Table 1). In view of the fact that total BCRP protein expression had not changed over day 5, day 12, and day 19 (Fig. 1B), the higher A-to-B transport and lower B-to-A transport of E3S across Caco-2 cells on day 5, compared with those on day 12 and day 19, indicated that BCRP continuously redistributed to the apical membrane during the differentiation phase.

Similar to P-gp, MRP, and lung-resistant protein (Yu and Sinko, 1997), the expression of BCRP is passage-dependent. The expression
of BCRP increased approximately 3-fold from the early passage to the late passage when Caco-2 cells were cultured in T-flasks, but decreased about 3- to 10-fold for monomer and dimer, respectively, from the early passage to the late passage when Caco-2 cells were grown on Transwell plates (Fig. 1, B and C). Although our findings are novel, they are supported by the previous reports that culture conditions and cell-growing matrices may affect efflux pump protein expression (Yu and Sinko, 1997). The BCRP-mediated E3S efflux in Caco-2 cells, with the B/A ratio of 2 and 30, respectively (Fig. 3), are novel, they are supported by the previous reports that culture conditions and cell-growing matrices may affect efflux pump protein expression (Yu and Sinko, 1997). The BCRP-mediated E3S efflux in Caco-2 cells was 13.2 μM, which was close to the $K_v$ (16.6 μM) determined in BCRP vesicle studies. The $V_{max}$ was estimated as 10.8 pmol/s. (Data expressed as mean ± S.E.M., n = 3.)

The efflux activity of BCRP in Caco-2 cells was further investi-
implying that BCRP is present in Caco-2 cells. The decreased B-to-A transport of MTX in Caco-2 cells could be totally attributed to BCRP.

BCRP mutations have been found in some cancer cell lines (Honjo et al., 2001). The R482T or R482G mutation but not the wild-type expressed cell lines (Allen et al., 2002a). In Caco-2 cells, the efflux of rhodamine 123 could be totally abolished by P-gp inhibitors, LY335979 and GF120918, but not affected by the BCRP inhibitor, Ko143 (Fig. 7).

The conclusion is further demonstrated by the efflux of rhodamine 123, to which BCRP-confirmed resistance is observed in the R482T or R482G mutation but not the wild-type expressed cell lines (Allen et al., 2002a). In Caco-2 cells, the efflux of rhodamine 123 could be totally abolished by P-gp inhibitors, LY335979 and GF120918, but not affected by the BCRP inhibitor, Ko143 (Fig. 7).

In conclusion, both BCRP monomer and dimer are expressed in the Caco-2 cells. BCRP is polarized at the apical side of Caco-2 cells and can efficiently transport its substrate, such as E3S and MTX, out of cells. Therefore, besides BCRP-transfected cell lines, Caco-2 cells can also be used as an in vitro model to study the transport function of BCRP.

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**References**


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