Multidrug resistance-associated protein 4 is a determinant of arsenite resistance

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Abstract. Although arsenic trioxide (arsenite, As^III) has shown a remarkable efficacy in the treatment of acute promyelocytic leukemia patients, multidrug resistance is still a major concern for its clinical use. Multidrug resistance-associated protein 4 (MRP4), which belongs to the ATP-binding cassette (ABC) superfamily of transporters, is localized to the basolateral membrane of hepatocytes and the apical membrane of renal proximal tubule cells. Due to its characteristic localization, MRP4 is proposed as a candidate in the elimination of arsenic and may contribute to resistance to As^III. To test this hypothesis, stable HEK293 cells overexpressing MRP4 or MRP2 were used to establish the role of these two transporters in As^III resistance. The IC_{50} values of As^III in MRP4 cells were approximately 6-fold higher than those in MRP2 cells, supporting an important role for MRP4 in resistance to As^III. The capacity of MRP4 to confer resistance to As^III was further confirmed by a dramatic decrease in the IC_{50} values with the addition of MK571, an MRP4 inhibitor, and cyclosporine A, a well-known broad-spectrum inhibitor of ABC transporters. Surprisingly, the sensitivity of the MRP2 cells to As^III was similar to that of the parent cells, although insufficient formation of glutathione and/or Se conjugated arsenic compounds in the MRP2 cells might limit transport. Given that MRP4 is a major contributor to arsenic resistance in vitro, further investigation into the correlation between MRP4 expression and treatment outcome of leukemia patients treated with arsenic-based regimen is warranted.

Introduction

Arsenic and arsenic-containing compounds are widely distributed in the environment and exist in organic and inorganic forms. Although a well-known poison, arsenic has been used medicinally for over 2,000 years (1). In particular, administration of arsenic trioxide (arsenite, As^III), an arsenic derivative, has demonstrated a remarkable efficacy in the treatment of relapsed and refractory acute promyelocytic leukemia (APL) (1-4). Detailed pharmacokinetic studies of As^III in APL patients have been carried out to optimize treatment (3,5-8). Both inorganic arsenic and methylated arsenic metabolites accumulate in red blood cells during repeated administration of As^III to APL patients (7). Arsenic metabolites are also detected in cerebrospinal fluid (6) at concentrations of arsenic necessary for induction of differentiation (4,9). Recent data from our laboratory demonstrated that the profiles of arsenic species in peripheral blood (PB) plasma were very similar to those of bone marrow (BM) plasma, suggesting that the profiles of PB plasma could be predictive biomarkers for the treatment outcome of APL patients (8). These findings on the pharmacokinetics of As^III in APL patients provide new insight into clinical applications of As^III, and may contribute to designing better therapeutic protocols (1).

Multidrug resistance is a major concern for the clinical use of anticancer drugs. ATP-binding cassette (ABC) transporters contribute to drug resistance via ATP-dependent drug efflux (1,10). Multidrug resistance-associated proteins 1 and 2 (MRP1/2), and multidrug resistance protein 1 (MDR1; also known as P-glycoprotein, P-gp) have been implicated in the efflux of arsenic, and may contribute to resistance to arsenic therapy (1,11,12). Furthermore, we recently demonstrated the MRP2 and aquaporin-9 (AQP9), a member of the aquaporin superfamily, involvement in arsenic uptake (1,13-15), contributing to the differential sensitivity of primary human-derived normal cells to arsenite (14,16-18). Although these previous findings provide fundamental knowledge for understanding the cellular handling and elimination pathways of arsenic in...
cancer cells as well as normal cells, more studies are required to provide detailed information on the efficacy and safety of arsenic for clinical use.

MRP4 transports antiviral agents such as the nucleo-side/nucleotide analogs azidothymidine (AZT), adefovir [9-(2-phosphonylmethoxethyl) adenine or PMEA] and tenofovir (TFV) (19-21), and anticancer drugs such as campto-thecins and methotrexate (22-24). Of note, recent studies have demonstrated that hematopoietic progenitor cell differentiation affects expression and function of MRP4, and that MRP4 has a relevant role in tumor growth and apoptosis and in the eradication of leukemic stem cells, suggesting MRP4 as a new potential therapeutic target for acute myeloid leukemia (25,26).

It is noteworthy that MRP4 is localized to the basolateral membrane of hepatocytes and the apical membrane of renal proximal tubule cells, distinguishing itself from other members of the MRP family (21). Considering the localization of MRP4, biometylation of arsenic primarily in liver and elimination of arsenic by the kidney (1), MRP4 is an ideal candidate for the elimination of arsenic and may contribute to resistance to As\textsuperscript{III}. Only recently have studies been reported regarding the role of MRP4 in arsenic resistance (27,28). In the present study, stable human embryonic kidney epithelial (HEK)293 cells overexpressing MRP4 and MRP2 were created and used to investigate the cytotoxicity of As\textsuperscript{III} against both MRP overexpressing cells and reference cells transfected with an empty vector, in order to clarify whether MRP4 cells have the capacity to confer drug resistance to As\textsuperscript{III}.

Materials and methods

Reagents. Sodium arsenite (As\textsuperscript{III}) was purchased from Tri Chemical Laboratories (Yamanashi, Japan). Cyclosporin A (CsA), a broad-spectrum inhibitor of ABC transporters, was kindly provided by Professor Toshihiko Hirano, Department of Clinical Pharmacology, School of Pharmacy, Tokyo University of Pharmacy and Life Sciences. 3-[3-[2-(7-chloroquinolin-2-yl) vinyl]phenyl]-(2-dimethylcarbamoylethylsulfanyl)methyl-sulfanyl] propionic acid (MK571), an inhibitor of MRP4, was purchased from Merck (Darmstadt, Germany).

Construction of stable HEK293 cell lines expressing human MRP2 or MRP4 cDNA. Human ABCC2 or ABCC4 cDNA was subcloned into the pcDNA5/FRT vector (Invitrogen, Carlsbad, CA, USA) as described previously (19,20). Briefly, HEK293 Flp-In cells were seeded at 5x10\textsuperscript{4} cells/well in 6-well plates in Dulbecco's modified Eagle's medium (DMEM) with 10% fetal bovine serum (FBS) without antibiotics. After 24 h of cultivation, 0.4 µg MRP2 or MRP4 plasmid was transfected into HEK293 Flp-In cells using Lipofectamine 2000 (Invitrogen) according to the manufacturer's protocol. Empty pcDNA/FRT vector was transfected as a negative control (empty vector cells). DMEM fresh media were added at 6 h post-transfection. The selection of stable cell clones expressing MRP2 or MRP4 reference plasmid was started in 75 µg/ml hygromycin (Invitrogen) the following day. Media were changed every 2-3 days, and selection of stable transfecants generally took 10-14 days. HEK293 cells stably expressing MRP2 reference (MRP2 cells) or MRP4 reference (MRP4 cells) and empty vector cells were cultured in DMEM supplemented with 10% FBS, 100 U/ml of penicillin and 100 µg/ml of streptomycin at 37°C in a humidified atmosphere (5% CO\textsubscript{2} in air).

MRP2 protein isolation and western blot analysis. Proteins were isolated using PARIS™ kit (Ambion Life Technologies, Foster City, CA, USA) according to the manufacturer's protocol. Proteins (30 µg/lane) were separated on 4-15% Tris-HCI Criterion gels (Bio-Rad, Hercules, CA, USA) by SDS-PAGE at 80 V and then transferred onto nitrocellulose membranes at 250 mA for 2 h. Membranes were blocked in 2% skim milk for 1 h at room temperature and incubated overnight at 4°C with anti-MRP2 antibody (M2-III-6) at 1:250 dilution (Thermo Fisher Scientific, Rockford, IL, USA) or anti-GAPDH antibody (ab9483) at 1:100,000 dilution (Abcam, Cambridge, MA, USA). The following day, membranes were washed three times with phosphate-buffered saline (PBS) and then incubated with IRDye 800CW goat anti-mouse fluorescent secondary antibody (Li-COR, Lincoln, NE, USA) at 1:10,000 dilution. Membranes were washed with PBS and visualized on an Odyssey® Sa Infrared Imaging system (Li-COR).

Evaluation of MRP2 membrane expression by immunocytochemistry. Empty vector and MRP2 reference cells were seeded at 2.5x10\textsuperscript{4} cells/chamber on 4-chamber slides 24 h prior to staining. The following day cells were washed with cold PBS two times and permeabilized with cold acetone for 10 min on ice. An equal volume of 8% paraformaldehyde was added directly to the cells and incubated for 10 min at room temperature. Cells were washed with PBS two times and blocked with 2 mg/ml BSA solution for 30 min at room temperature. Primary anti-MRP2 (M2-III-6) antibody was added to BSA solution to the final dilution of 1:20. After three washes with PBS, the slides were incubated with a fluorescently conjugated secondary antibody (Alexa Fluor® 488; Thermo Fisher Scientific) for 1 h at room temperature protected from light. DAPI (Thermo Fisher Scientific) stain at 1:3,000 dilution was added 20 min before the end of incubation. After three washes with PBS, images were mounted and then captured using a Retiga CCD-cooled camera and associated QCapture Pro software (QImaging, Surrey, BC, Canada).

Functional assays of MRP2 cells. Evaluation of MRP2 transport activity was performed using a carboxyfluorescein (CF) fluorescent dye retention assay. CF was applied to cells in its di-acetate form (CFDA) to form (Sigma-Aldrich, St. Louis, MO, USA) which is non fluorescent and highly lipophilic; it freely enters the cells through passive diffusion. Once inside the cell the acetate groups of CFDA are cleaved by esterases yielding fluorescent CF, which has low permeability characteristics and is a substrate for ABC efflux transporters, including MRP2 (29). Briefly, empty vector and MRP2 overexpressing cells were trypsinized, washed with warm PBS and re-suspended at a cell density of 1x10\textsuperscript{5} cells/ml in fresh DMEM without FBS. Cells were then incubated for 30 min at 37°C with 10 µM of CFDA. Following accumulation, cells were pelleted, washed twice with warm PBS, re-suspended in DMEM media supplemented with 10% FBS and allowed to efflux for 30 min at 37°C. Following efflux, cells were pelleted, washed twice with ice-cold PBS and then re-suspended in ice-cold PBS supplemented with 10% FBS for analysis. Retention of CF was determined by
measuring fluorescence using flow cytometry performed on a BD FACSCalibur (BD Biosciences, Mississauga, ON, Canada). Briefly, 10,000 cell events were collected for each sample. Cells were co-stained with propidium iodide (PI; Thermo Fisher Scientific) to exclude non-viable cells from further analysis. CF fluorescence was measured in FL-1 channel (excitation wavelength 488 nm and emission wavelength 530 nm), and PI fluorescence was measured in FL-3 channel (excitation wavelength 488 nm and emission wavelength 600 nm). Each experiment was repeated four times. Data were normalized to empty vector transfected cells and a t-test was used on normalized data to test for differences in MRP2 overexpressing cells and reference cells.

**Functional assays of MRP4 cells.** Transport assays for MRP4 were carried out using two reported MRP4 substrates [adenine-8-3H-tenofovir disoproxil (TFV) (3.8 Ci/mmol, 98.1% purity); 3H-9-(2-phosphonylmethoxyethyl)-adenine (PMEA)] (Moravek Biochemicals, Brea, CA, USA) as described previously (19,20,30). Briefly, MRP4 and empty vector cells were seeded at 2.5x10^5 cells/well in triplicate in poly-D-lysine-coated 24-well plates (BD Biosciences, San Jose, CA, USA). After preincubation for 24 h, cells were incubated with 1 µM TFV or 100 nM PMEA in glucose-free DMEM supplemented with 10 µM NaN3 and 10 µM 2-deoxy-D-glucose, respectively, for 2 h at 37°C. After accumulation, cells were washed with ice-cold PBS and supplemented with complete DMEM. Supernatant fractions were collected at 0, 30 and 90 min, and cells were washed and lysed with 800 µl/well of an aqueous solution of 10% sodium dodecyl sulfate and 1 N NaOH. Supernatants were added to Ecolite scintillation fluid (MP Biomedicals, Santa Ana, CA, USA) and extracellular amounts of TFV and PMEA were analyzed by scintillation counting. The remaining cell lysate was used to determine the protein concentration with a BCA™ Protein Assay kit (Pierce Biotechnology, Rockford, IL, USA); TFV and PMEA levels were normalized to protein concentrations.

**Cell viability assay.** The cytotoxicity of AsIII to MRP4, MRP2 and empty vector cells was investigated by XTT dye-reduction assays according to the method previously described with slight modifications (14,31). Briefly, the cells were seeded in 96-well plates (Iwaki, Tokyo, Japan) at a density of 5x10^3 cells/well in 0.1 ml complete DMEM and cultivated for 24 h. Cultures in triplicate were treated with various concentrations of AsIII in the presence or absence of transporter inhibitors at the concentrations indicated. Cells were pre-incubated with transporter inhibitors at the indicated concentrations for 30 min. After treatment with AsIII for an additional 48 h, 2,3-bis(2-methoxy-4-nitro-5-sulfophenyl)-5-[(phenylaminocarbonyl]-2H-tetrazolium hydroxide (XTT; Sigma, MD, USA) and phenazine methosulfate (Wako Pure Chemical Industries, Osaka, Japan) were added into each well at final concentrations of 0.2 mg/ml and 1 mM, respectively.
After incubation at 37°C for 4 h, the plates were mixed, and the absorbance at 450 nm was measured with a microplate reader (Safire, Tecan, Switzerland). The relative cell viability was expressed as the ratio of the absorbance of each treatment group against those of the corresponding untreated control group. Data are shown as means ± SD from more than five independent experiments. The IC\textsubscript{50} values of As\textsuperscript{III} for all three cell types were calculated using GraphPad Prism® 5 software.

Statistical analysis. Data were analyzed using Student’s t-test and ANOVA with a Dunnett’s post test method. A p-value <0.05 was considered as statistically significant.

Results

Confirmation of MRP2 expression and transporter activity in the MRP2 stable cell line. Protein expression of MRP2 was confirmed using immunocytochemistry and western blotting. As shown in Fig. 1, MRP2 cells had significantly higher levels of membrane (Fig. 1A and B) and total (Fig. 1C) MRP2 expression compared to empty vector transfected cells. Functional activity of MRP2 was confirmed in MRP2 reference cells using a CF retention assay. As shown in Fig. 1D, retention of CF in MRP2 overexpressing cells was 4.5-fold lower than that in empty vector cells, indirectly confirming high efflux of fluorescent dye from MRP2 overexpressing cells and, therefore, providing strong evidence for MRP2 functional activity.

Confirmation of transporter activity in the MRP4 stable cell line. Accumulation of TFV or PMEA in the supernatant fractions of each cell type was assessed for confirmation of MRP4 function. As shown in Fig. 2, the accumulation of TFV or PMEA in the supernatant of MRP4 cells increased with time, and was much higher than that of empty vector cells. Compared to empty vector cells, an approximately 3-fold and 2.7-fold increase in the efflux of fluorescent dye from MRP2 overexpressing cells and, therefore, providing strong evidence for MRP2 functional activity.

As\textsuperscript{III}-induced cytotoxicity in MRP4 and MRP2 overexpressing cells. After treatment with 0-1,000 µM of As\textsuperscript{III} for 48 h, cell viability was investigated by XTT assay. A significant dose-dependent decrease in cell viability was observed in all three cell types (Fig. 3). As\textsuperscript{III} exhibited much lower cytotoxicity in MRP4 cells in comparison with MRP2 and empty vector cells. The IC\textsubscript{50} values of As\textsuperscript{III} in MRP4 cells was significantly greater than the corresponding value in empty vector cells (P<0.01).

Effect of MK571 on As\textsuperscript{III}-induced cytotoxicity in MRP4 and empty vector cells. Both MRP4 and empty vector cells were exposed to various concentrations of As\textsuperscript{III} (0-1,000 µM) in the presence or absence of 10 or 25 µM MK571 for 48 h, followed by the assessment of cell viability. A significant dose-dependent decrease in cell viability was observed in all three cell types (Fig. 3). As\textsuperscript{III} exhibited much lower cytotoxicity in MRP4 cells in comparison with MRP2 and empty vector cells. The IC\textsubscript{50} values of As\textsuperscript{III} were 36.6±8.1, 5.1±0.3 and 5.5±0.3 µM in MRP4, MRP2 and empty vector cells (MRP4 vs. empty vector, P<0.01; MRP2 vs. empty vector, P>0.05), respectively.

Effect of MK571 on As\textsuperscript{III}-induced cytotoxicity in MRP4 and empty vector cells. Both MRP4 and empty vector cells were exposed to various concentrations of As\textsuperscript{III} (0-1,000 µM) in the presence or absence of 10 or 25 µM MK571 for 48 h, followed by the assessment of cell viability. A significant dose-dependent decrease in cell viability was observed in all three cell types (Fig. 3). As\textsuperscript{III} exhibited much lower cytotoxicity in MRP4 cells in comparison with MRP2 and empty vector cells. The IC\textsubscript{50} values of As\textsuperscript{III} were 36.6±8.1, 5.1±0.3 and 5.5±0.3 µM in MRP4, MRP2 and empty vector cells (MRP4 vs. empty vector, P<0.01; MRP2 vs. empty vector, P>0.05), respectively.
Effect of CsA on As\textsuperscript{III}-induced cytotoxicity in MRP4 and empty vector cells. Both MRP4 and empty vector cells were exposed to various concentrations of As\textsuperscript{III} (0-1,000 µM) in the presence or absence of 1 or 3 µM CsA for 48 h, followed by the assessment of cell viability. In comparison to MK571, CsA, a broad-spectrum inhibitor of ABC transporters, more potently inhibited MRP4 and increased As\textsuperscript{III}-induced cytotoxicity (Fig. 5A). The IC\textsubscript{50} value of As\textsuperscript{III} in MRP4 cells decreased ≥90% after the addition of 1 and 3 µM CsA, respectively (Fig. 5A). There was some increase in As\textsuperscript{III} cytotoxicity as well in empty vector cells (5.3±0.2 vs 3.1±0.6 and 2.3±0.2 µM) (empty vector cells without CsA vs. empty vector cells with 1 or 3 µM CsA, P<0.05) (Fig. 5B).

Discussion

MRP2 and MRP4 overexpressing cells were used in the current study to evaluate the contribution of these ABC efflux transporters to As\textsuperscript{III} cytotoxicity. A 6-fold difference in IC\textsubscript{50} values of As\textsuperscript{III} between these cells and a significant reduction of the IC\textsubscript{50} value in MRP4 cells provide strong support that MRP4 is a major mediator of As\textsuperscript{III} efflux from cells and plays a major role in As\textsuperscript{III} resistance. MRP2 has been demonstrated to be involved in the efflux of arsenic, conferring resistance to As\textsuperscript{III} in other experimental systems (12,32,33). Wild-type and MRP2-deficient Wistar (TR\textsuperscript{−}) rats have been used to show that MRP2 is responsible for the biliary excretion of arsenic triglutathione [As(GS)\textsubscript{3}] and monomethyl arsenic diglutathione [CH\textsubscript{3}As(GS)\textsubscript{2}] (33). These in vivo findings are supported by cellular transport assays demonstrating that As(GS)\textsubscript{3} is also a substrate for human MRP2 (12). A recent in vitro study using MRP2-enriched membrane vesicles demonstrated that MRP2 transports seleno-bis(S-glutathionyl) arsinium ion [(GS)\textsubscript{2}AsSe\textsuperscript{+}] (32). These previous findings suggest that glutathione and/or the essential trace element selenium (Se) are required for the excretion and detoxification of arsenic. In contrast, using MRP2 overexpressing cells in the present study, there was no evidence that MRP2 plays a critical role in As\textsuperscript{III} transport and cytotoxicity. These conflicting results might reflect low levels of glutathione and/or Se conjugation in MRP2 overexpressing HEK293 cells, although more detailed analysis of the molecular events such as the amount of glutathione and its conjugated arsenic compounds is required to confirm this hypothesis.

Inhibition studies further support the role of MRP4 to confer resistance to As\textsuperscript{III}. MK571, an MRP4 inhibitor (21), was used to demonstrate that MRP4 confers resistance to a series of camptothecin analogs, including irinotecan and SN-38 (23,24). Consistent with these previous studies, the addition of MK571 significantly potentiated As\textsuperscript{III}-triggered cytotoxicity in the MRP4 cells in a dose-dependent manner. CsA, a well-known broad-spectrum inhibitor of ABC transporters, increased As\textsuperscript{III}-triggered cytotoxicity to a greater degree than MK571, supporting a role for multiple ABC transporters in As\textsuperscript{III} cytotoxicity. Collectively, our experimental results implicate MRP4 in resistance to As\textsuperscript{III}. Considering that multidrug resistance-reversing activity of CsA has been reported in phase II studies with myeloma and acute leukemia,
increased recognition of drug-drug interactions is necessary for optimal treatment of patients with arsenic-based regimens.

Our results are not completely in agreement with other studies of arsenic resistance. Increased resistance was not detected in MRP4-transfected NIH3T3 cells after exposure to sodium meta-arsenite for 72 h (28). Furthermore, a recent report demonstrated that after treatment with increasing concentrations of inorganic and methylated species of arsenic for 72 h, MRP4-transfected HEK293 cells conferred resist—ance to arsenate and methylated species of arsenic except for arsenite (27). Plausible explanations for the differences between our findings and previous reports are differences in cell lines, arsenite reagent, treatment durations and stable versus transient overexpression systems.

MRP4 has been proposed to contribute to arsenic elimination due to its characteristic localization in both the basolateral membrane of hepatocytes and the apical membrane of renal proximal tubule cells (21,27). Based on transport studies using MRP4-enriched membrane vesicles, the diglutathione conjugate of MMA_{III}, monomethylarsenic diglutathione [MMA(GS)_{2}], and DMA^+ are transported by MRP4 (27). It is noteworthy that MRP4 has been reported to regulate intracellular cyclic adenosine monophosphate (cAMP) levels, an endogenous substrate identified for MRP4 (21), in AML cell lines and contribute to cell proliferation and differentiation (34). Moreover, Copsel et al recently reported that MRP4 blockade strongly reduced tumor growth by inducing cell cycle arrest and apoptosis in U937 xenografted mice, and further demonstrated that increased cAMP levels and MRP4 inhibition resulted in leukemic stem cell differentiation (25). Of note, a rapid increase in intracellular cAMP has been linked to all-trans retinoic acid (ATRA)-induced differentiation in the human APL cell line NB4 and in fresh APL cells (35). More importantly, cAMP facilitates the degradation of As_{III}-mediated fusion protein promyelocytic leukemia (PML)-retinoic acid receptor α (RARα), a fusion gene generated by the t(15;17) translocation in APL and thought to play a central role in the initiation of leukemogenesis (1,4,36). These previous findings raised the possibility that MRP4 would be a novel promising target in APL therapy. The expression of MRP4 increased in CD34^+ cells differentiated toward megakaryocytes with thrombopoietin, and a similar increase was also observed in a megakaryoblastic cell line (M-07e) derived from a patient with megakaryoblastic leukemia, when differentiated toward megakaryocytes (26). However, the expression of MRP4 decreased in CD34^+ cells differentiated toward monocytes with G-CSF, suggesting a relevant role of MRP4 in hematopoietic progenitor cell differentiation (26). Therefore, considering the expression status of MRPs, including MRP4, is important for providing meaningful clinical benefits for patients with different types of hematological disorders.

It is noteworthy that the MRP4 gene is highly polymorphic, and numerous nonsynonymous single-nucleotide polymorphisms (SNPs) have been identified (21). Functional studies have shown that although there was no evidence for a complete loss of function allele, two variants (G187W and G487E) show...
a significantly reduced function compared to reference MRP4 as evidenced by higher intracellular accumulation of ATZ and PMEA, two antiviral substrates for MRP4 (19). Although no disease has so far been directly linked to altered MRP4 activity, evaluating the functional effects of high frequency variants on the disposition of arsenic has important implications for hematologic malignancy patients treated with arsenic-based regimens.

In conclusion, our results demonstrated the capacity of MRP4 to confer resistance to AsIII as evidenced by cell survival assays when treated with AsIII in the presence or absence of its two differential inhibitors. Given that MRP4 is widely distributed in the body, and plays a pivotal role in the drug concentrations achieved clinically, monitoring its expression levels may have important implications for predicting not only clinical efficacy but also side effects of arsenite and its metabolites. Obviously, clinical data are required to support the role of MRP4 in drug disposition and efficacy. Based on our findings and previous studies showing that MRP4 could be a major contributor to arsenic resistance, further investigation into the correlation between the expression level of MRP4 and treatment outcome of leukemia patients treated with arsenic-based regimens is warranted.

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