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To cite this version:
Stefan Harmsen, I. Meijerman, C. L. Febus, R. F. Maas-Bakker, J. H. Beijnen, et al.. PXR-mediated induction of P-glycoprotein by anticancer drugs in a human colon adenocarcinoma-derived cell line. Cancer Chemotherapy and Pharmacology, Springer Verlag, 2009, 66 (4), pp.765-771. <10.1007/s00280-009-1221-4>. <hal-00552487>
PXR-mediated induction of P-glycoprotein by anticancer drugs in a human colon adenocarcinoma-derived cell line

Stefan Harmsen · I. Meijerman · C. L. Febus · R. F. Maas-Bakker · J. H. Beijnen · J. H. M. Schellens

Abstract

**Purpose** The development of multidrug resistance (MDR) is one of the major limitations in the treatment of cancer. Induction of P-glycoprotein (Pgp) has been regarded as one of the main mechanisms underlying anticancer drug-induced MDR. Since the induction of Pgp is (in part) regulated by the pregnane X receptor (PXR), the ability of several widely used anticancer drugs to activate PXR-mediated Pgp induction was investigated.

**Methods** A Pgp-reporter gene assay was employed to determine the ability of a panel of widely used anticancer drugs to induce Pgp. To further assess whether PXR could be involved in the induction of Pgp by anticancer drugs, Pgp protein expression after treatment with the anticancer drugs was determined in both wild-type and PXR-knocked down LS180 cells. Furthermore, the effect of the anticancer drugs on the intracellular accumulation of the Pgp-probes rhodamine 123 and doxorubicin was determined.

**Results** Our study showed that vincristine, tamoxifen, vinblastine, docetaxel, cyclophosphamide, flutamide, ifosfamide and paclitaxel activate PXR-mediated Pgp induction, and were additionally shown to affect the intracellular accumulation of the Pgp probe rhodamine 123. Moreover, PXR activation was also shown to reduce the cytotoxic activity of the Pgp substrate doxorubicin in colon cancer cells.

**Conclusion** Our results indicate that several anticancer drugs can activate PXR-mediated induction of Pgp and affect the accumulation of Pgp substrates.

**Keywords** PXR · Induction · P-glycoprotein · Multidrug resistance · Anticancer drugs

**Abbreviations**

ABC ATP-binding cassette
ATP Adenosine triphosphate
DMSO Dimethylsulfoxide
HBSS Hank’s Balanced Salt Solution
HRP Horse radish peroxidase
MDR Multidrug resistance
NR Nuclear receptor
PBS phosphate buffer saline
Pgp P-glycoprotein
PXR Pregnan X receptor

Introduction

A major drawback in the successful treatment of advanced malignancies is the development of multidrug resistance (MDR). One of the main causes by which cancer cells acquire the multidrug resistant phenotype in response to chemotherapy is the induction of ATP-binding cassette (ABC) drug efflux transporter proteins. An important
mediator of MDR that has been associated with poor treatment response in various malignancies is the ABC-family member P-glycoprotein (Pgp; ABCB1; MDR1) [1–3]. Pgp is a transmembrane protein that protects the cell from xenobiotic stress by facilitating the extrusion of a broad spectrum of substrates, including anticancer drugs such as the anthracyclines, epipodophyllotoxins, Vinca alkaloids and taxanes. As a consequence, induction of Pgp affects the efficacy of these agents by decreasing their intracellular accumulation in cancer cells.

The pregnane X receptor (PXR; NR1I2) has been identified as a major regulator of Pgp induction [4], and, apart from expression in the liver and small intestines, has been shown to be expressed in several cancerous tissues such as breast, colon, bone, prostate and endometrial cancers [5–9]. PXR is a very promiscuous receptor that is activated by a wide variety of structurally unrelated ligands including rifampicin, hyperforin and the anticancer drug paclitaxel. Due to the promiscuity of PXR, possibly other widely used anticancer drugs could also activate PXR-mediated Pgp induction, and as a consequence induce MDR in cancer cells.

In the current study, a panel of widely used anticancer drugs was evaluated for their ability to activate PXR-mediated Pgp induction in the colon adenocarcinoma-derived cell line LS180. In addition, the effect of PXR activation on the intracellular accumulation of Pgp substrates was determined. Our results demonstrate that several widely used anticancer drugs can activate PXR-mediated induction of Pgp and as a consequence decrease the intracellular accumulation of Pgp substrates. In addition, evidence is presented that the cytotoxic activity of doxorubicin is reduced when cells are pretreated with the PXR activator rifampicin.

Materials and methods

Materials

All cell culture media and supplements were purchased from Invitrogen (Breda, The Netherlands). Carboplatin, ifosfamide, tamoxifen citrate and etoposide were obtained from Axxora (San Diego, CA, USA). Zosuquidar (LY335979) was obtained through Kanisa Pharmaceuticals Inc. (Irvine, CA, USA). All other chemicals were purchased from Sigma–Aldrich (Zwijndrecht, The Netherlands).

Plasmids

The pGL3-MDR1 (p-10224) luciferase reporter construct was generously provided by Dr. Oliver Burk (Institute of Clinical Pharmacology, Eberhard-Karls University, Tübingen, Germany). The pCDG-hPXR expression vector was generously provided by Dr. Ron Evans (Salk institute for biological studies, La Jolla, CA, USA). The pRL-TK control plasmid was obtained from Promega (Madison, WI, USA). Plasmids were checked by enzyme restriction and agarose gel electrophoresis and purified using Promega’s Pureyield Midi-prep (Madison, WI, USA) according to the instructions of the manufacturer.

Cell culture

The human colon adenocarcinoma-derived cell line LS180 was purchased from the ATCC (Manassas, VA, USA). The cell-line was maintained in Roswell Park Memorial Institute (RPMI) 1640 medium (with 25 mM HEPES and L-glutamine, supplemented with 10% (v/v) heat-inactivated fetal bovine serum, 100 U/ml penicillin and 100 µg/ml streptomycin), at 37°C under a humidified atmosphere of 5% CO2.

Cell viability

Cell viability was after 48 h, and the anticancer drug treatment was assessed with a 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide (MTT)-assay.

Pgp reporter gene assay

The Pgp reporter gene assay was performed in a similar manner as described previously [10]. In brief, LS180 cells were transfected with a pCDG-hPXR expression vector, a pGL3-MDR1 luciferase reporter construct, and a pRL-TK control plasmid for 24 h. After transfection, the transiently transfected LS180 cells were treated with the highest non-toxic concentration of each anticancer drug. After 48 h incubation, the reporter activity was determined.

Cell treatment

LS180 cells were plated at a density of 5 × 104 cells/well in 96 well plates (Pgp protein level determination) or 1 × 105 cells/well in 1 ml RPMI in 24 well plates (Pgp transport activity). After reaching 80–90% confluency, medium was replaced with medium containing the different anticancer drugs: carboplatin (10 µM), cyclophosphamide (100 µM), ifosfamide (100 µM), docetaxel (10 µM), paclitaxel (10 µM), flutamide (10 µM) and tamoxifen (10 µM). Rifampicin (10 µM) was used as a positive control and DMSO (0.1%) as a negative control. The cells were treated for 48 h with the drugs and the controls. At the end of each treatment, the cells were processed for immunoblotting or rhodamine 123-based Pgp activity assays.
RNA interference of PXR

The siRNA sequence targeting human PXR (sense: cguu-guguccccuucuggatt; antisense: cucaggaaggcaaaacagtg) and the negative control, which consisted of a non-complementary sequence, were purchased from Ambion (Austin, TX, USA). LS180 cells were reversely transfected 48 h prior to the Pgp induction experiment with 50 nM siRNA PXR or 50 nM negative control siRNA using LipoFectamine RNAiMax (Invitrogen, Breda, The Netherlands). Subsequently, PXR knockdown and control LS180 cells were treated with different anticancer drugs for 48 h.

Immunoblot analysis

After 48 h of culture, cells were washed with phosphate-buffered saline (PBS), and lysed in 1 ml/well MilliQ water that contained protease inhibitors (Roche, Basal, Switzerland). The lysate was centrifuged (5,000 rpm, 5 min, 4°C), and the remaining pellet was resuspended in 100 μL RIPA buffer. The lysate was centrifuged (10,000 rpm, 5 min, 4°C) and the supernatant was transferred to a new collection tube. Protein concentrations were determined by a Pierce BCA protein assay (Pierce, Rockford, IL, USA). Ten micrograms of total protein was separated by SDS–polyacrylamide gel electrophoresis using NuPage novex 4–12% bis–tris gradient gels (Invitrogen, Breda, The Netherlands). Proteins were electroblotted onto Immobilon P membranes (Millipore, Bedford, MA, USA). After blocking (1 h at RT) in 3% BSA in tris-buffered saline containing 0.1% Tween-20, the membranes were incubated overnight at 4°C with a murine monoclonal anti-Pgp primary antibody (C219, 1:500; Abcam, Cambridge, UK) followed by incubation with a goat anti-mouse IgG coupled to horseradish peroxidase (HRP) secondary antibody (1:1,000; Immunopure Peroxidase Conjugated Goat anti-Mouse IgG (H + L); Pierce, Rockford, IL, USA). β-actin (1:10,000; AC-15; Abcam, Cambridge, UK) was used as a loading control. The proteins were visualized by a chemiluminescence-based detection reagent (SuperSignal West Femto; Pierce, Rockford, IL, USA), and the intensities of the Pgp bands were determined on a Gel Doc XRS Imaging system with Quantity One analysis software (Bio-Rad, Hercules, CA, USA). The ratio of the Pgp signal and the β-actin signal were indicative for Pgp protein expression.

Rhodamine 123 accumulation assay

The efflux activity of Pgp was determined by measuring the accumulation of the fluorescent Pgp probe rhodamine 123 as described by Collett et al. [11]. In brief, LS180 cells were pretreated with the anticancer drugs as described earlier. After 48 h, the cells were washed with Hank’s Balanced Salt Solution (HBSS) and incubated at 37°C for 1 h with rhodamine 123 (10 μM) in the presence or absence of the Pgp-specific inhibitor zosuquidar (LY335979; 5 μM; [12]) in phenol red-free RPMI medium. After washing thrice with ice-cold HBSS, the cells were lysed with Triton X-100 (0.1%) in MilliQ water (Millipore, Bedford, MA, USA). The ratio of intracellular rhodamine 123 concentrations in the absence and presence of 5 μM zosuquidar is indicative for the efflux activity of Pgp.

Doxorubicin accumulation

LS180 cells (0.5 × 10^6 c/well) were plated in a 24-well plate. After a 48 h incubation with 0.1% DMSO or 10 μM rifampicin, cells were treated with 5 μM doxorubicin for 90 min. Cells were washed thrice with ice-cold PBS, lysed in 250 μl MilliQ water, and homogenized by ultrasonification Labsonic P (Braun; cycle 0.2 min en Amplitude 60%). Hundred microliters of the homogenate was processed for HPLC analysis. Daunorubicin was used as an internal standard. The samples and standard solutions were prepared as described by van Asperen et al. [13].

Doxorubicin HPLC analysis

Doxorubicin was quantified using a method as previously described by DeGregorio et al. [14]. In brief, samples were eluted isocratically with a mobile phase that consisted of 28% acetonitrile in 60 mM phosphoric acid. A Waters Symmetry C18 3.5 μM, 4.6 mm × 100 mm reverse-phase column was used. Fluorescence was detected using a JASCO FP-920 fluorometer (excitation: 455 nm; emission: 550 nm), Shimadzu LC10AT-VP pumps and a Shimadzu JASCO FP-920 spectrophotometer (excitation: 455 nm; emission: 550 nm). The flow rate was set at 1.0 ml/min. Doxorubicin and daunorubicin retention times were 2.38 and 5.01 min, respectively. A standard curve of doxorubicin (0.08, 0.4, 2, 10 and 50 μM) was prepared in milliQ water.

Doxorubicin cytotoxicity assay

LS180 cells were plated (2 × 10^4 cells/well; 96 well plates) and treated with 10 μM rifampicin or 0.1% DMSO. Following 72 h incubation, cells were exposed to doxorubicin (serially diluted 0–20 μM; dilution factor 2). Cell viability after 48 h doxorubicin exposure was assessed using a neutral red assay [15].

Statistical analysis

One-way ANOVA with Bonferroni post hoc testing was performed for statistical comparison of the obtained Pgp reporter gene and activity results and considered statistically
Results

Cell viability

Cell viability was assessed after 48 h treatment with different concentrations of the anticancer drugs. As shown in Fig. 1, the anticancer drugs did not significantly affect the LS180 cell viability at the concentrations used, while the doxorubicin (10 μM) did affect cell viability.

PXR-mediated Pgp reporter activation by anticancer drugs

The ability of widely used anticancer drugs to induce the expression of Pgp via activation of PXR was determined. LS180 cells were transfected with a Pgp reporter construct as well as a PXR expression plasmid. Treatment of the transiently transfected LS180 cells with cyclophosphamide, ifosfamide, docetaxel, paclitaxel, flutamide, and tamoxifen resulted in dose-dependent increases in Pgp reporter activity, while exposure to different concentrations of carboplatin and etoposide did not increase PXR-mediated Pgp reporter activity (Fig. 2). As a control, the same compounds were tested in LS180 cells that were only transfected with the Pgp reporter gene construct without co-transfection of a PXR expression plasmid. In these cells, none of the compounds was able to activate Pgp reporter activity (results not shown).

Induction of Pgp protein expression

To evaluate whether Pgp reporter activation is predictive for Pgp protein induction, the effect of the anticancer drugs on the protein expression levels of Pgp after 48 h treatment in LS180 cells was determined. As shown in Fig. 3, all compounds that were shown to activate PXR-mediated Pgp reporter activity (vincristine, tamoxifen, vinblastine, doce-
taxel, cyclophosphamide, flutamide, ifosfamide and paclitaxel) also induced Pgp protein levels in LS180 cells. In accordance with the Pgp-reporter gene assay, both etoposide and carboplatin did not induce Pgp protein expression. To further demonstrate that PXR plays an important role in the upregulation of Pgp, PXR was knocked down in the LS180 cells (knock down efficiency was 51% as was determined 48 h after siRNA transfection by Western blot (results not shown)). The cells were subsequently treated with the anticancer drugs. As shown in Fig. 3, Pgp protein induction was significantly lower in the PXR knockdown LS180 cells after treatment with the anticancer drugs compared to the control cells (LS180 cells transfected with non-targeting siRNA). However, Pgp induction was not completely abolished, most likely as a result of the knockdown efficiency of 51%.

Rhodamine 123 accumulation assay

To assess whether PXR-mediated Pgp induction affects the accumulation of Pgp substrates, the transport activity of Pgp was analyzed using a rhodamine 123 accumulation assay. Following 48 h pretreatment with the anticancer drugs, cells were exposed to rhodamine 123 in the absence or presence of the Pgp-specific inhibitor zosuquidar (LY335979; IC50 = 4nM) [16]. The ratio between the intracellular fluorescence of rhodamine 123 in the presence and absence of zosuquidar is indicative for the functionality of Pgp. The data are normalized to the DMSO controls and are presented as mean data (±SD) of three different experiments (*P < 0.05).
while etoposide and carboplatin did not affect rhodamine 123 accumulation. In contrast, the accumulation of rhodamine 123 in the cells that were treated with the Vinca alkaloids, cyclophosphamide, ifosfamide and tamoxifen did not correspond with Pgp protein expression.

Effect of PXR activation on doxorubicin cytotoxicity

To study whether activation of PXR affects the efficacy of a cytotoxic Pgp substrate, the effect of PXR activation on the cytotoxicity of doxorubicin in LS180 was assessed. Therefore, prior to doxorubicin exposure, LS180 cells were pretreated with the prototypical PXR agonist 10 μM rifampicin or 0.1% DMSO (solvent) for 48 h. As shown in Fig. 5a, rifampicin pretreatment affects the intracellular accumulation of doxorubicin. To determine whether reduced accumulation of doxorubicin also affects the efficacy of this agent, rifampicin- or solvent-pretreated cells were exposed to different doxorubicin concentrations (ranging from 0.3 to 20 μM; dilution factor 2). As shown in Fig. 5b, following doxorubicin exposure cell viability was significantly \( (P < 0.05) \) higher in the rifampicin-pretreated cells \( (IC_{50} = 2.5 \mu M) \) compared to the solvent (0.1% DMSO) pretreated cells \( (IC_{50} = 1.5 \mu M) \). This indicates that PXR activation indeed can reduce the activity of cytotoxic Pgp substrates as a result of decreased intracellular accumulation.

Discussion

The development of MDR in response to anticancer treatment is a major problem in the clinic. Since induction of Pgp is one of the main mechanisms underlying acquired MDR, we evaluated whether several widely used anticancer drugs can induce Pgp expression via activation of PXR, a nuclear receptor that is increasingly associated with acquired MDR [9, 17].

Indeed, several anticancer drugs (vincristine, vinblastine, cyclophosphamide, ifosfamide, docetaxel, flutamide and tamoxifen) were shown to increase Pgp reporter activity in cells co-transfected with a PXR expression plasmid. In accordance, the same anticancer drugs induced Pgp protein expression in LS180 cells. However, when PXR was knocked down in these cells with treatment with the indicated anticancer drugs did not result in Pgp protein induction. Together these results indicate that PXR plays a substantial role in anticancer drug-related Pgp induction.

Furthermore, since Pgp is a drug efflux transporter that is involved in the extrusion of a wide variety of substrates from the cell, the effect of Pgp protein induction following treatment with the indicated anticancer drugs on the intracellular accumulation of Pgp substrates was evaluated. Docetaxel, flutamide, paclitaxel and the positive control rifampicin that induced Pgp protein expression indeed affected the intracellular accumulation of the Pgp substrate rhodamine 123. In addition, carboplatin and etoposide, which had no effect on Pgp protein expression, did not affect rhodamine 123 accumulation. However, tamoxifen was shown to activate PXR-mediated Pgp protein induction, but did not affect the accumulation of rhodamine 123. This can be explained by the fact that tamoxifen is also a known inhibitor of Pgp and thus prevents the Pgp-mediated extrusion of rhodamine 123 [18]. The observed discrepancy between the Pgp protein expression and Pgp functionality

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after treatment with the Vinca alkaloids is difficult to explain and requires further investigation.

Since activation of PXR by anticancer drugs was shown to affect the accumulation of Pgp substrates, we further assessed whether PXR activation could also attenuate the cytotoxic effects of doxorubicin, a known cytotoxic Pgp substrate. However, to avoid potential pharmacodynamic interactions between the selected anticancer drugs and doxorubicin that could lead to increased cytotoxicity, we chose to pretreat the cells with the prototypical PXR agonist rifampicin. Indeed, pretreatment of LS180 cells with rifampicin decreased the intracellular accumulation of the cytotoxic Pgp substrate doxorubicin and additionally significantly reduced the cytotoxic efficacy of this agent. These results indicate that activation of PXR-mediated Pgp induction by anticancer drugs can underlie the development of acquired resistance. This is further supported by studies of Chen et al. [9] and Gupta et al. [17], who have shown that activation of PXR promotes the multidrug resistant phenotype of prostate and ovarian cancer cell lines, respectively.

Since Pgp is co-expressed with PXR in important barrier tissues, such as the intestines and the liver, activation of PXR-mediated Pgp induction could affect the pharmacokinetic profile of (oral) anticancer drugs. Indeed, drug-drug interactions as a result of PXR-mediated Pgp induction have been reported. For instance, intake of St. John’s wort, which contains the PXR agonist hyperforin, by healthy volunteers resulted in a 1.9-fold increase in the oral clearance of the Pgp substrate fexofenadine [19], while rifampicin pretreatment resulted in a 1.3- to 5.3-fold increase in the oral clearance of fexofenadine [20]. In the same manner, anticancer drugs that activate PXR might affect their own pharmacokinetics, but also that of other concomitantly administered (anticancer) drugs.

In conclusion, several widely used anticancer drugs were shown to activate PXR-mediated Pgp induction, and as a result affect the intracellular accumulation and cytotoxic activity of Pgp substrates.

Acknowledgments The authors would like to thank the Maurits and Anna de Kock foundation and the Nijbakker-Morra foundation for financially supporting this study.

Conflict of interest statement None.

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