



## OSI-930 analogues as novel reversal agents for ABCG2-mediated multidrug resistance

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### ABSTRACT

OSI-930, a dual c-Kit and KDR tyrosine kinase inhibitor, is reported to have undergone a Phase I dose escalation study in patients with advanced solid tumors. A series of fifteen pyridyl and phenyl analogues of OSI-930 were designed and synthesized. Extensive screening of these compounds led to the discovery that nitropyridyl and *ortho*-nitrophenyl analogues, VKJP1 and VKJP3, were effective in reversing ABC subfamily G member 2 (ABCG2) transporter-mediated multidrug resistance (MDR). VKJP1 and VKJP3 significantly sensitized ABCG2-expressing cells to established substrates of ABCG2 including mitoxantrone, SN-38, and doxorubicin in a concentration-dependent manner, but not to the non-ABCG2 substrate cisplatin. However, they were unable to reverse ABCB1- or ABCC1-mediated MDR indicating their selectivity for ABCG2. Western blotting analysis was performed to evaluate ABCG2 expression and it was found that neither VKJP1 nor VKJP3 significantly altered ABCG2 protein expression for up to 72 h. [<sup>3</sup>H]-mitoxantrone accumulation study demonstrated that VKJP1 and VKJP3 increased the intracellular accumulation of [<sup>3</sup>H]-mitoxantrone, a substrate of ABCG2. VKJP1 and VKJP3 also remarkably inhibited the transport of [<sup>3</sup>H]-methotrexate by ABCG2 membrane vesicles. Importantly, both VKJP1 and VKJP3 were efficacious in stimulating the activity of ATPase of ABCG2 and inhibited the photoaffinity labeling of this transporter by its substrate [<sup>125</sup>I]-iodoarylazidoprazosin. The results suggested that VKJP1 and VKJP3, specifically inhibit the function of ABCG2 through direct interaction with its substrate binding site(s). Thus VKJP1 and VKJP3 represent a new class of drugs for reducing MDR in ABCG2 over-expressing tumors.

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## 1. Introduction

Multidrug resistance (MDR) is a major obstacle to successful chemotherapy in cancer treatment. It is a phenomenon in which

cancer cells become resistant to a wide spectrum of anti cancer drugs that belong to different chemical or pharmacological classes and results in decreased cytotoxic activity of these anticancer drugs [1,2]. The mechanisms of MDR are multifactorial and include alteration in the permeability of lipid bilayer membrane, increased DNA repair in cancer cells, increased inactivation or detoxification of drugs, or inhibition of apoptosis [3,4]. Over-expression of multiple ATP binding cassette (ABC) transporters in cancer cells still remains the leading cause of MDR [5–7]. The term ABC transporter, is based on the fact that almost all members of the ABC superfamily, from yeast and bacteria to man, share a conserved consensus sequence of 90–110 amino acids, where ATP binds and is hydrolyzed by the ATPase enzyme. This consensus has Walker A and B motifs and another C region called ABC cassette. ABC transporters is the largest trans-membrane protein family encoded in the human genome, which are divided into 7 subfamilies ABC-A to G and subdivided into sub-subfamilies depending on their

**Abbreviations:** OSI-930, {N-(4-trifluoromethoxyphenyl)3-((quinolin-4-ylmethyl) amino) thiophene-2-carboxamide}; ABC, ATP binding cassette; ABCG2 (BCRP/MXR/ABCP), ABC subfamily G member 2; MDR, multidrug resistance; P-gp, P-glycoprotein; ABCCs (MRPs), ABC subfamily C member; RTK, receptor tyrosine kinases; KDR, kinase insert domain receptor; TKIs, tyrosine kinase inhibitors; IAAP, [<sup>125</sup>I]-iodoarylazidoprazosin; FTC, fumitremorgin C; MTT, 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide; DMSO, dimethylsulfoxide.

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structural similarity or difference in their trans-membrane domain. The prominent ABC transporters playing a role in MDR development are ABCB1, also called P-glycoprotein (P-gp) [8,9], multidrug resistance proteins (MRPs, ABCs) [10], and ABCG2 (BCRP/MXR/ABCP) [11,12]. This ABC cassette is involved in transportation of not only toxic metabolites or xenobiotics out of the cell, but also a multitude of anticancer drugs across cell membrane, thus decreasing their intracellular concentration and cytotoxicity. Therefore, their expression is believed to be the major mechanism by which MDR to anticancer drugs develops. Initially, verapamil was found by Tsuruo et al. to be the first modulator which could inhibit the function of ABCB1 and increase drug accumulation in the cancer cells [13]. Later, cyclosporine A (CsA), an immunosuppressant, was discovered to inhibit the function of ABCB1 and was used as another modulator of ABCB1 [14]. After that, many other compounds have been studied for their inhibitory effects to other members of ABC transporter family [15–18], raising hopes that ABC transporter mediated MDR could be reversed. At present, as the newly synthesized molecule-targeting agents such as tyrosine kinase inhibitors (TKIs) are being introduced into the clinic, research is being conducted to determine whether these novel compounds could represent a new class of ABC transporter inhibitors. TKIs such as imatinib, nilotinib, erlotinib, dasatinib and lapatinib have been extensively investigated as modulators of ABC transporters-mediated MDR, which most likely act as competitive inhibitors [19–23]. More recently, pre-clinical research and clinical trials have been conducted to study the efficacy of the combination of some TKIs with other anticancer drugs in improving the therapeutic outcome of cancer patients. Therefore, discovering novel inhibitors of ABC transporters to reverse the drug resistance can lead to improvement in the use of currently available anticancer drugs.

OSI-930 {N-(4-trifluoromethoxyphenyl)-3-((quinolin-4-ylmethyl) amino) thiophene-2-carboxamide} is a heterocyclic anthranilamide analogue synthesized by OSI Pharmaceuticals Inc. It is a potent and selective dual inhibitor of the closely related receptor tyrosine kinases (RTKs) Kit (c-Kit, a type III RTK over-expressed in seminoma, acute myeloid leukemia and in gastrointestinal stromal tumor) and kinase insert domain receptor (KDR, a type V RTK as a regulator of tumor angiogenesis) [24,25]. OSI-930 acts by these two distinct mechanisms in appropriate tumor types: i.e., direct effects on the tumor cell phenotype through inhibition of c-Kit and indirect effects via disruption of endothelial cell function by inhibition of KDR and is reported to be under phase I clinical trials. The quinoline domain of OSI-930 was modified with heteroatom substituted pyridyl and phenyl ring systems by our

group to obtain a series of fifteen compounds (Fig. 1) [26]. The effects of these changes on their activities and the binding site characteristics of c-Kit and KDR were evaluated using a kinase assay. In the present study, we report the screening of our newly designed and synthesized OSI 930 analogues to determine their ability to reverse ABC transporter-mediated MDR. The reversal mechanisms of the most efficacious analogues VKJP1 and VKJP3 were further evaluated by membrane vesicle transport assay, photolabeling affinity assay, intracellular drug accumulation assay and ATPase assay.

## 2. Materials and methods

### 2.1. Chemicals and reagents

[<sup>3</sup>H]-mitoxantrone (4 Ci/mmol) and [<sup>3</sup>H]-methotrexate (23 Ci/mmol) were purchased from Moravek Biochemicals, Inc. (Brea, CA). [<sup>125</sup>I]-iodoarylazidoprazosin (IAAP, 2200 Ci/mmol) were obtained from Perkin-Elmer Life Sciences. (Boston, MA). BXP-21 (against ABCG2) was acquired from Signet Laboratories, Inc. (Ded-Ham, MA). Anti-actin monoclonal antibody was obtained from Santa Cruz Biotechnology, Inc. (Santa Cruz, CA). OSI-930 was gift from Selleck Chemical (Houston, TX) OSI-930 analogues were successfully designed and synthesized by our group at St. John's University (New York, NY). PAK104P was a kind gift from Prof. Shin-Ichi Akiyama (Kagoshima University, Kagoshima, Japan) from Nissan Chemical Ind. Co. Ltd (Chiba, Japan). Fumitremorgin C (FTC) was synthesized by Thomas McCloud Developmental Therapeutics Program, Natural Products Extraction Laboratory, National Cancer Institute, NIH. (Bethesda, MD). Mitoxantrone, SN-38, doxorubicin, cisplatin, colchicine, vincristine, verapamil, penicillin/streptomycin, 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT), dimethylsulfoxide (DMSO), ATP, AMP and other chemicals were purchased from Sigma Chemical Co. (St. Louis, MO).

### 2.2. Cell lines and cell culture

HEK293/pcDNA3.1, ABCG2-482-R2, and ABCG2-482-T7 cells were established by selection with G418 after transfecting HEK293 with either the empty pcDNA3.1 vector or pcDNA3.1 vector containing the full length ABCG2 coding either arginine (R) or threonine (T) at amino acid 482, respectively [27]. They were cultured in medium with 2 mg/ml of G418 [27]. The parental human epidermoid carcinoma cell line KB-3-1 was selected in a stepwise manner using increasing concentration of colchicine to establish the ABCB1-overexpressing drug resistant cell line KB-C2

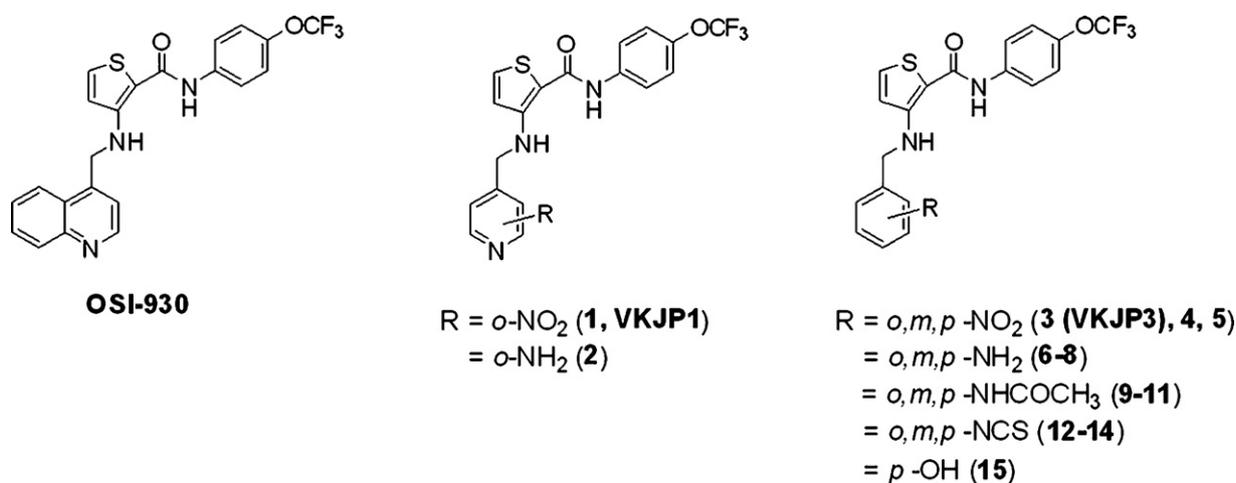


Fig. 1. Chemical structures of OSI-930 and VKJP compounds 1–15.

and was cultured in the medium with 2  $\mu\text{g/ml}$  of colchicine [28]. An ABCG2-overexpressing MDR cell line, KB-CV60, was also cloned from KB-3-1 cells and was maintained in medium with 1  $\mu\text{g/ml}$  cepharanthine and 60 ng/ml vincristine [29]. All the cell lines were grown in Dulbecco's modified Eagle's medium (DMEM) supplemented with 10% bovine serum, 100 units/ml penicillin, and 100 mg/ml streptomycin in a humidified incubator containing 5%  $\text{CO}_2$  at 37 °C.

### 2.3. Cell sensitivity to drugs by MTT assay

Cell sensitivity to drugs was analyzed using an MTT colorimetric assay [19]. Cells were harvested with trypsin and resuspended in a final concentration  $8 \times 10^3$  cells/ml for KB-C2 and  $5 \times 10^3$  for all the other cell lines. Cells were seeded evenly into (160  $\mu\text{l/well}$ ) 96-well multiplates. After 24 h of incubation, the test VKJP compound (20  $\mu\text{l/well}$ ) was added 1 h prior to different concentrations of chemotherapeutic drugs (20  $\mu\text{l/well}$ ) into designated wells. FTC, verapamil or PAK-104 (20  $\mu\text{l/well}$ ) was added as positive control inhibitors of ABCG2, ABCB1 or ABCG1, respectively. After 72 h of incubation, 20  $\mu\text{l}$  of MTT solution (4 mg/ml) was added to each well, further incubated for 4 h, the medium was discarded, and 100  $\mu\text{l}$  of DMSO was added into each well to dissolve the formazan crystals. The absorbance was determined at 570 nm by an OPSYS microplate Reader from DYNEX Technologies, Inc. (Chantilly, VA). The concentrations required to inhibit growth by 50% ( $\text{IC}_{50}$ ) were calculated from survival curves. The degree of resistance was calculated by dividing the  $\text{IC}_{50}$  of the MDR cells by that of the parental sensitive cells.

### 2.4. Preparation of cell lysates

Cells in T-25 flask treated with 10  $\mu\text{M}$  VKJP1 or VKJP3 for different time periods (0, 36, 72 h), then were harvested and rinsed twice with cold PBS. The cell extracts were prepared by incubating the cells with the radioimmunoprecipitation assay (RIPA) buffer [ $1 \times$  PBS, 1% Nonidet P-40, 0.5% sodium deoxycholate, 0.1% SDS, 100  $\mu\text{M}$  p-aminophenylmethylsulfonyl fluoride (p-APMSF), 10  $\mu\text{M}$  leupeptin, and 10  $\mu\text{M}$  aprotinin] for 30 min on ice with occasional rocking, followed by centrifugation at  $12,000 \times g$  at 4 °C for 15 min. The supernatant containing total cell lysates were collected and stored at  $-80$  °C until future experiments. The protein concentration was determined by bicinchoninic acid ( $\text{BCA}^{\text{TM}}$ )-based protein assay (Thermo Scientific, Rockford, IL).

### 2.5. Western blotting

Equal amounts of total cell lysates (40  $\mu\text{g}$ ) were resolved by 4–12% sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) and electrophoretically transferred onto PVDF membrane, then immersed in blocking solution (5% skim milk in Tris-buffered saline (TBST: 0.3% Tris, 0.8% NaCl, 0.02% KCl, 0.05% Tween 20) for 1 h at room temperature. The membrane was then immunoblotted overnight with monoclonal anti-ABCG2 antibody (BXP-21) at 1:500 dilution at 4 °C. The following day, the membrane was washed with TBST buffer three times and followed by 3 h incubation with horseradish peroxidase (HRP)-conjugated secondary anti-mouse IgG at 1:1000 for 3 h. Proteins were detected by enhanced chemiluminescence detection system (Amersham, NJ).  $\beta$ -Actin was used to confirm equal loading in each lane in the samples prepared from cell lysates.

### 2.6. [ $^3\text{H}$ ]-mitoxantrone accumulation

The effect of VKJP1 or VKJP3 on the intracellular accumulation of [ $^3\text{H}$ ]-mitoxantrone in HEK293/pcDNA3.1, ABCG2-482-R2, and ABCG2-482-T7 cells was determined as previously described [30]. Confluent cells in 24-well plates were preincubated with or without

VKJP1, VKJP3 (FTC as positive control) for 1 h at 37 °C. Intracellular drug accumulation was measured by incubating cells with 0.2  $\mu\text{M}$  [ $^3\text{H}$ ]-mitoxantrone for another 2 h. The cells were washed three times with ice-cold PBS, trypsinized and lysed in 10 mM lysis buffer (pH 7.4, containing 1% Triton X-100 and 0.2% SDS). Each sample was placed in scintillation fluid and radioactivity was measured in a Packard TRI-CARB<sup>®</sup> 1900CA liquid scintillation analyzer from Packard Instrument Company, Inc (Downers Grove, IL).

### 2.7. Preparation of membrane vesicles and in vitro transport assays

Membrane vesicles were prepared by using the nitrogen cavitation method [31]. Cells were rinsed twice with PBS and then scraped into PBS containing 1% aprotinin. Cells were then washed at 4 °C in PBS, collected by centrifugation ( $4000 \times g$  for 10 min), suspended in buffer A (10 mM Tris-HCl, pH 7.4, 0.25 M sucrose, 1 mM p-APMSF, and 0.2 mM  $\text{CaCl}_2$ ) and equilibrated at 4 °C for 15 min under a nitrogen pressure of 500 psi. EDTA was added to the suspension of lysed cells to make a final concentration of 1 mM, and the suspension was then diluted 1:4 with buffer B (10 mM Tris-HCl, pH 7.4, 0.25 M sucrose, and 1 mM p-APMSF) and centrifuged at  $4000 \times g$  for 10 min at 4 °C to remove nuclei and unlysed cells. The supernatant was layered onto a sucrose cushion (35% sucrose, 10 mM Tris-HCl, pH 7.4, and 1 mM EDTA) and centrifuged for 30 min at  $16,000 \times g$  at 4 °C. The interface was collected and centrifuged for 45 min at  $100,000 \times g$  at 4 °C. The pellet was resuspended in buffer B by repeated passage through a 25-gauge needle.

Transport assays were performed using the rapid filtration method as previously described [19]. Membrane vesicles were incubated with VKJP1 or VKJP3 (FTC as positive control) for 1 h on ice, and then transport experiments were carried out at 37 °C for 10 min in a total volume of 50  $\mu\text{l}$  medium (membrane vesicles 10  $\mu\text{g}$ , 0.25 M sucrose, 10 mM Tris-HCl, pH 7.4, 10 mM  $\text{MgCl}_2$ , 4 mM ATP or 4 mM AMP, 10 mM phosphocreatine, 100  $\mu\text{g/ml}$  creatine phosphokinase, and 0.5  $\mu\text{M}$  [ $^3\text{H}$ ]-methotrexate). Reactions were stopped by the addition of 3 ml of ice-cold stop solution (0.25 M sucrose, 100 mM NaCl, and 10 mM Tris-HCl, pH 7.4). During the rapid filtration step, samples were passed through 0.22  $\mu\text{m}$  GVWP filters (Millipore Corporation, Billerica, MA) that were presoaked in the stop solution. The filters were washed three times with 3 ml of ice-cold stop solution. Radioactivity was measured by the use of a liquid scintillation analyzer from Packard Instrument Company, Inc (Downers Grove, IL).

### 2.8. Effect of VKJP1 and VKJP3 on ATPase activity of ABCG2

ATPase activities of ABCG2 in High Five insect cell crude membranes were measured by endpoint inorganic phosphate  $\text{P}_i$  assay as described previously with minor modifications [32]. ABCG2-specific ATPase activity was recorded as BeFx-sensitive ATPase activity. Briefly, The membrane vesicles (100  $\mu\text{g}$  protein/ml) were incubated in ATPase assay buffer (50 mM MES-Tris, pH 6.8, 50 mM KCl, 5 mM  $\text{NaN}_3$ , 1 mM EGTA, 1 mM ouabain, 2 mM dithiothreitol and 10 mM  $\text{MgCl}_2$ ) for 20 min at 37 °C in the absence or presence of 0.2 mM beryllium sulfate and 2.5 mM sodium fluoride (BeFX), then incubated with different concentrations of VKJP1 or VKJP3 at 37 °C for 3 min. The reaction was initiated by the addition of 5 mM ATP and terminated by the addition of SDS (2.5% final concentration). The amount of  $\text{P}_i$  released was quantified using a colorimetric method.

### 2.9. Effect of VKJP1 and VKJP3 on photoaffinity labeling with [ $^{125}\text{I}$ ]-iodoarylazidoprazosin (IAAP) of ABCG2

Crude membranes of High Five insect cells (250  $\mu\text{g}$  protein/ml) were incubated with increasing concentrations of VKJP1 or VKJP3

at room temperature in 50 mM Tris–HCl, pH 7.5, for 3 min. [<sup>125</sup>I]-IAAP (3–6 nM) was added and further incubated for additional 5 min under subdued light. The samples were exposed to a UV lamp (365 nm) for 10 min at room temperature and processed as described previously [33]. The samples were separated on a 7% Tris–acetate gel at a constant voltage. Gel was dried and exposed to Bio-Max MR film (Eastman Kodak, Rochester, NY) for 9 h at –80 °C. The incorporation of [<sup>125</sup>I]-IAAP into ABCG2 band was quantified by estimating the radioactivity of this band using the STORM 860

phosphorimager (Molecular Dynamics, Sunnyvale, CA) and ImageQuaNT.

## 2.10. Statistical analysis

Unless otherwise indicated, all experiments were repeated at least three times and the differences were determined by the two-tailed Student's *t*-test. When statistical differences among more than 2 groups were analyzed, one-way ANOVA followed by Tukey's

**Table 1**  
Effects of the test compounds in reversing ABCG2-mediated resistance to mitoxantrone.

Treatment	HEK293/pcDNA3.1		ABCG2-482-R2	
	IC <sub>50</sub> (nM)	FR <sup>a</sup>	IC <sub>50</sub> (nM)	FR <sup>a</sup>
Mitoxantrone	17.77 ± 1.34	1.00	227.40 ± 14.95	12.80
.,OSI-930 10 μM	23.17 ± 3.37	1.30	42.24 ± 17.56	2.38
.,VKJP1 10 μM	11.28 ± 1.23	0.63	14.90 ± 8.70	0.84
.,VKJP2 10 μM	18.95 ± 7.45	1.07	61.81 ± 14.77	3.48
.,VKJP3 10 μM	10.68 ± 0.84	0.60	35.15 ± 11.96	1.98
.,VKJP4 10 μM	17.24 ± 2.96	0.97	58.95 ± 0.94	3.32
.,VKJP5 10 μM	14.93 ± 0.92	0.84	50.59 ± 5.57	2.85
.,VKJP6 10 μM	20.42 ± 0.69	1.15	61.88 ± 9.23	3.48
.,VKJP7 10 μM	15.64 ± 4.01	0.88	67.21 ± 11.78	3.78
.,VKJP8 10 μM	16.40 ± 3.35	0.92	79.48 ± 20.81	4.47
.,VKJP9 10 μM	18.32 ± 8.31	1.03	54.82 ± 24.55	3.08
.,VKJP10 10 μM	17.35 ± 8.45	0.98	53.93 ± 25.04	3.03
.,VKJP11 10 μM	15.72 ± 1.94	0.88	65.34 ± 25.35	3.68
.,VKJP12 10 μM	22.36 ± 6.90	1.26	54.66 ± 17.06	3.08
.,VKJP13 10 μM	17.96 ± 1.40	1.01	217.52 ± 18.65	12.20
.,VKJP14 10 μM	19.97 ± 5.74	1.12	257.63 ± 35.88	14.50
.,VKJP15 10 μM	14.51 ± 2.49	0.82	57.05 ± 26.69	3.21
.,FTC 2.5 μM	16.15 ± 1.19	0.91	20.17 ± 2.38	1.14

<sup>a</sup> Fold-resistance was determined by dividing the IC<sub>50</sub> value for mitoxantrone of ABCG2-482-R2 cells by IC<sub>50</sub> value of HEK293/pcDNA3.1 cells in the absence or presence of the test compounds or FTC.

**Table 2**  
Effect of VKJP1 and VKJP3 in reversing ABCG2-mediated MDR.

Treatment	HEK293/pcDNA3.1		ABCG2-482-R2		ABCG2-482-T7	
	IC <sub>50</sub> (nM)	FR <sup>a</sup>	IC <sub>50</sub> (nM)	FR <sup>a</sup>	IC <sub>50</sub> (nM)	FR <sup>a</sup>
Mitoxantrone	24.78 ± 0.68	1	277.91 ± 35.02	11.21	814.61 ± 3.05	32.87
.,VKJP1 1 μM	21.34 ± 4.65	0.86	109.68 ± 95.05	4.43	235.57 ± 97.66	9.51
.,VKJP1 3 μM	23.91 ± 0.93	0.96	28.52 ± 3.59	1.15	118.47 ± 67.94	4.78
.,VKJP1 10 μM	18.97 ± 5.41	0.77	16.75 ± 9.1	0.68	24.06 ± 0.49	0.97
.,VKJP3 1 μM	21.84 ± 7.17	0.88	108.58 ± 1.7	4.38	313.45 ± 194.86	12.65
.,VKJP3 3 μM	21.43 ± 7.75	0.86	46.94 ± 11.55	1.89	163.69 ± 37.21	6.6
.,VKJP3 10 μM	17.78 ± 6.88	0.72	53.19 ± 30.81	2.15	68.19 ± 0.08	2.75
.,FTC 2.5 μM	27.16 ± 3.43	1.1	9.03 ± 0.09	0.36	28.19 ± 1.97	1.14
SN-38	63.08 ± 0.94	1	2456.15 ± 376.77	38.94	2750.44 ± 180.13	43.6
.,VKJP1 1 μM	64.28 ± 7.45	1.02	641.69 ± 405.07	10.17	684.43 ± 40.57	10.85
.,VKJP1 3 μM	58.59 ± 21.93	0.93	310.36 ± 177.14	4.92	172.93 ± 16.78	2.74
.,VKJP1 10 μM	45.1 ± 18.13	0.72	213.17 ± 120.53	3.38	67.4 ± 5.01	1.07
.,VKJP3 1 μM	76.81 ± 17	1.22	931.17 ± 814.32	14.76	1164.95 ± 61.44	18.47
.,VKJP3 3 μM	73.12 ± 7.24	1.16	229.58 ± 23.21	3.64	403.42 ± 38.39	6.4
.,VKJP3 10 μM	68.49 ± 13.5	1.09	137.06 ± 39	2.17	191.8 ± 16.45	3.04
.,FTC 2.5 μM	61.88 ± 13.4	0.98	106.03 ± 63.17	1.68	61.33 ± 3.64	0.97
Doxorubicin	4.39 ± 0.21	1	15.02 ± 0.08	3.42	80.52 ± 5.96	18.33
.,VKJP1 1 μM	4.56 ± 0.69	1.04	10.5 ± 2.57	2.39	29.72 ± 3.35	6.77
.,VKJP1 3 μM	3.59 ± 0.12	0.82	4.78 ± 0.44	1.09	23.75 ± 7.78	5.41
.,VKJP1 10 μM	1.97 ± 0.89	0.45	4.9 ± 2.87	1.12	12.71 ± 6.55	2.89
.,VKJP3 1 μM	4.56 ± 0.2	1.04	13.85 ± 5.58	3.15	85.21 ± 34.22	19.4
.,VKJP3 3 μM	3.2 ± 0.46	0.73	9.35 ± 5.5	2.13	24.92 ± 10.13	5.67
.,VKJP3 10 μM	2.81 ± 0.2	0.64	6.51 ± 0.92	1.48	17.22 ± 8.58	3.92
.,FTC 2.5 μM	4.53 ± 0.57	1.03	5.05 ± 0.99	1.15	6.6 ± 3.92	1.5
Cisplatin	2380 ± 40	1	2410 ± 300	1.01	2460 ± 160	1.04
.,VKJP1 10 μM	2110 ± 240	0.89	2300 ± 530	0.97	2290 ± 170	0.96
.,VKJP3 10 μM	2330 ± 170	0.98	2610 ± 310	1.1	2570 ± 360	1.08
.,FTC 2.5 μM	2210 ± 50	0.93	2610 ± 410	1.1	2370 ± 260	1

<sup>a</sup> Fold-resistance was determined by dividing the IC<sub>50</sub> value for mitoxantrone, SN-38, doxorubicin, and cisplatin of ABCG2-482-R2 and ABCG2-482-T7 cells by IC<sub>50</sub> value of HEK293/pcDNA3.1 cells in the absence or presence of VKJP1, VKJP3 or FTC.

multiple comparison tests were performed. Results were presented as mean  $\pm$  standard deviations (SD). The statistical significance was determined to be  $P < 0.05$ .

### 3. Results

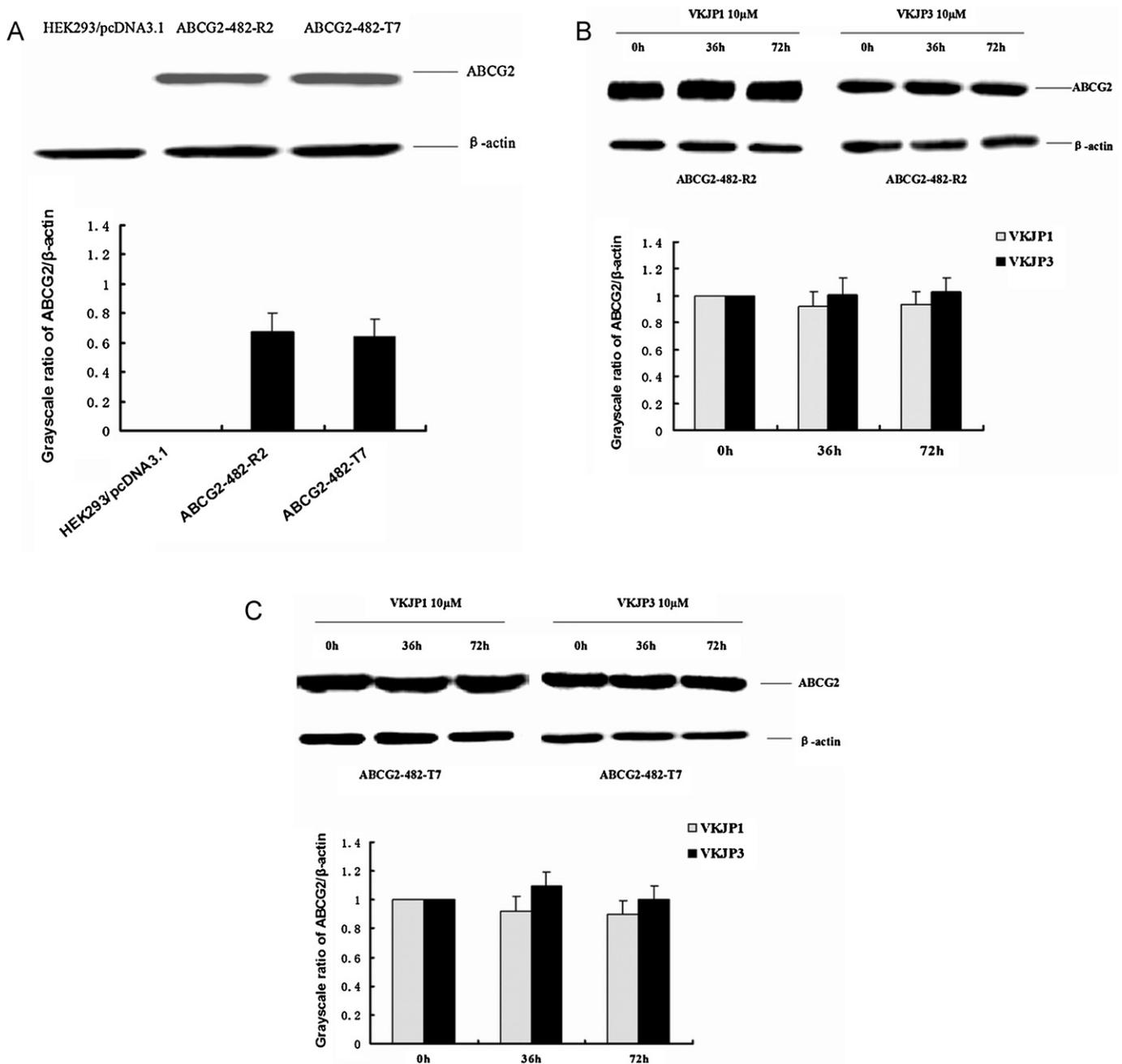
#### 3.1. Structure of VKJP1 and VKJP3

A series of OSI-930 analogues **1–15** (Fig. 1) were successfully designed, synthesized, and evaluated for their reversal effect of ABC transporter-mediated MDR.

#### 3.2. The effect of VKJP compounds on sensitivity of ABC-overexpressing cells to chemotherapeutic drugs

Prior to determining the effects of VKJP1 and VKJP3 in reversing the MDR, the effect of OSI-930 and its analogues on cell viability was examined. The  $IC_{50}$  values of these compounds in ABC-overexpressing cells were in the range of 100  $\mu$ M (data not shown). In order to avoid cellular toxicity, the highest non-toxic concentration of the test compounds used in subsequent reversal experiments was 10  $\mu$ M.

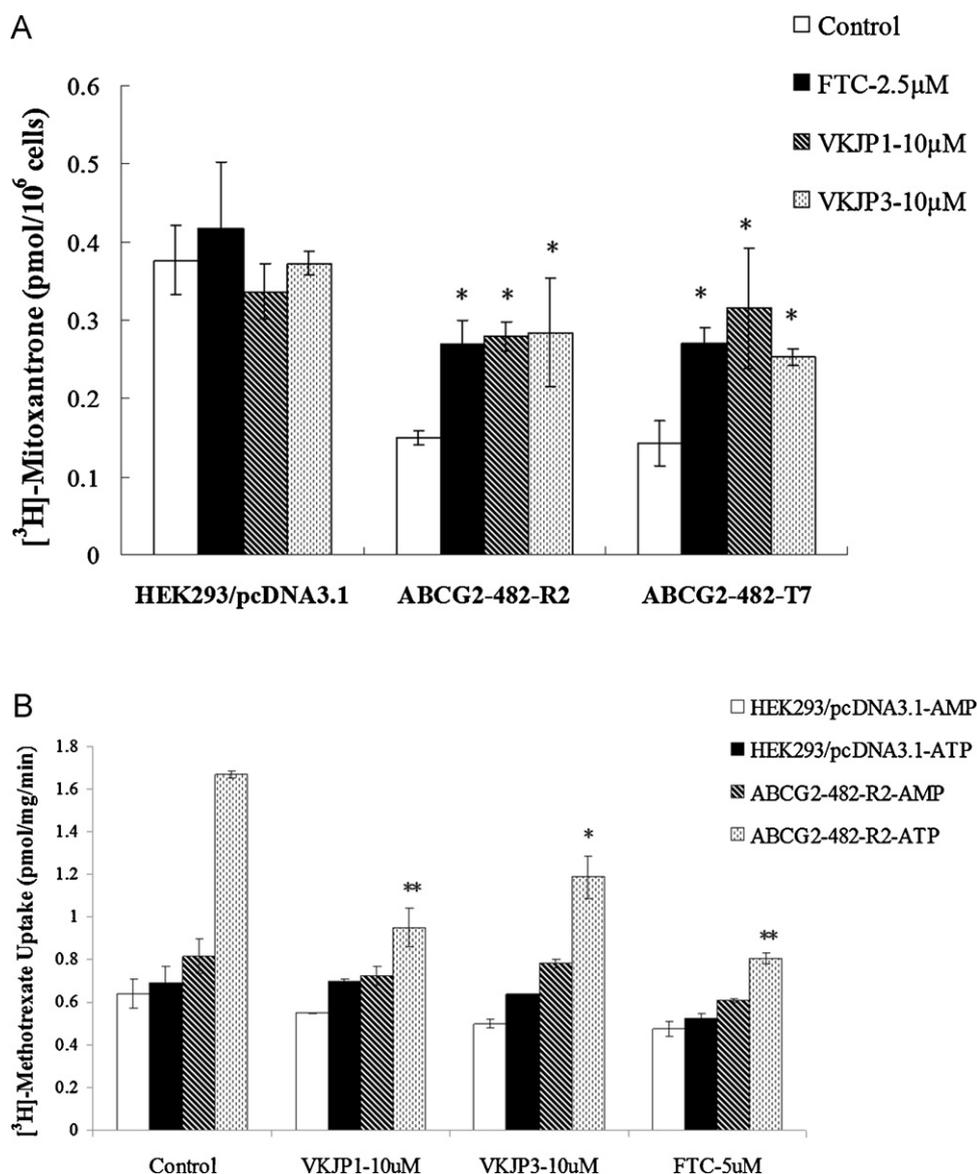
Subsequently, we screened the 15 compounds in the test series and determined whether they could sensitize the transfected



**Fig. 2.** Western blot detection of ABCG2 in cells. (A) The expression of ABCG2 in HEK293/pcDNA3.1, ABCG2-482-R2, ABCG2-482-T7 cells. Equal amounts (40  $\mu$ g protein) of total cell lysates were used for each sample. The membranes were immunoblotted with primary antibody against ABCG2 at 1:500 dilutions at 4 °C overnight, then incubated with HRP-conjugated secondary antibody at 1:1000 dilutions at room temperature for 3 h. Proteins were detected by enhanced chemiluminescence detection system (Amersham, NJ).  $\beta$ -Actin was used to confirm equal loading in each lane in the samples prepared from cell lysates. Representative result is shown out of three experiments. (B) Representative result showed the ABCG2 expression of ABCG2-482-R2 cells when treated with VKJP1 or VKJP3 (10  $\mu$ M) at 0, 36, and 72 h. C: Representative result showed the ABCG2 expression of ABCG2-482-T7 cells when treated with VKJP1 or VKJP3 (10  $\mu$ M) at 0, 36, and 72 h.

wild-type ABCG2-overexpressing MDR cells to certain chemotherapeutic drugs. VKJP1 and VKJP3 were the first two in the rank order of potencies for increasing the cytotoxicity of mitoxantrone among all of the members of the test series. For this reason, VKJP1 and VKJP3 were chosen for further study. Compound **13** had no significant reversing ability, while compound **14** was found to protect cells from cytotoxicity by mitoxantrone (Table 1). As shown in Table 2, VKJP1 or VKJP3 at 1, 3 and 10  $\mu\text{M}$ , concentration-dependently decreased the  $\text{IC}_{50}$  values of ABCG2 substrates including mitoxantrone, SN-38, and doxorubicin of the transfected wild-type (R2) or mutant (T7) ABCG2-overexpressing cells. VKJP1 was shown to be the most potent inhibitor of ABCG2 than VKJP3 at all concentrations. The magnitude of the sensitization produced by 10  $\mu\text{M}$  VKJP1 was similar to that induced by the known specific ABCG2 inhibitor FTC at 2.5  $\mu\text{M}$ . In addition, 10  $\mu\text{M}$

VKJP1 could significantly sensitize the parental HEK293-pcDNA3.1 cells to all the aforementioned chemotherapeutic drugs. However, this effect was significantly lower than measured in both ABCG2-482-R2 and ABCG2-482-T7 cells. In contrast, there was no significant effect in the  $\text{IC}_{50}$  values for all the cell lines when cisplatin (which is not a substrate of ABCG2) was incubated with VKJP1, VKJP3 or FTC (Table 2). Besides, VKJP1 or VKJP3 at 10  $\mu\text{M}$  did not significantly alter the cytotoxicity of colchicine (a substrate of ABCB1) on KB-C2 cells which is a drug selected cell line overexpressing ABCB1 (Supplemental Table 1) or vincristine (a substrate of ABCC1) on KB-CV60 cells which is a drug selected cell line overexpresses ABCC1 (Supplemental Table 2). These results suggest that VKJP1 and VKJP3 selectively sensitize chemotherapeutic drugs that are the substrates of ABCG2 but not ABCB1 or ABCC1.



**Fig. 3.** Effect of VKJP1 and VKJP3 on the intracellular accumulation of [<sup>3</sup>H]-mitoxantrone and ABCG2-mediated transport of [<sup>3</sup>H]-methotrexate. (A) The accumulation of [<sup>3</sup>H]-mitoxantrone was measured after cells were preincubated with or without VKJP1, VKJP3 (10  $\mu\text{M}$ ), or FTC (2.5  $\mu\text{M}$ ) for 1 h at 37 °C and then incubated with 0.2  $\mu\text{M}$  [<sup>3</sup>H]-mitoxantrone for another 2 h at 37 °C. Columns, mean of triplicate determinations; bars, SD; \*,  $P < 0.05$  versus the control group. Experiments were repeated at least three times and a representative experiment is shown. (B) Membrane vesicles (10  $\mu\text{g}$ ) were prepared from HEK293/pcDNA3.1 and ABCG2-482-R2 cells. The uptake rates of [<sup>3</sup>H]-methotrexate into membrane vesicles were measured for 10 min at 37 °C in uptake medium containing 4 mM of ATP or AMP. For inhibition experiments, membrane vesicles were incubated with VKJP1, VKJP3, or FTC for 1 h on ice, and then transport reactions were carried out for 10 min at 37 °C in uptake medium containing 4 mM ATP. Columns, mean of triplicate determinations; bars, SD; \*,  $P < 0.05$ ; \*\*,  $P < 0.01$ , versus the control group. Experiments were repeated at least three times and a representative experiment is shown.

### 3.3. Effects of VKJP1 and VKJP3 on the expression of ABCG2 in ABCG2-overexpressing MDR cell

The reversal of ABCG2-mediated MDR can be achieved either by decreasing ABCG2 expression or by inhibiting its efflux function. To study the effects of VKJP1 and VKJP3 on ABCG2 expression, ABCG2-482-R2 and ABCG2-482-T7 cells were treated with VKJP1 or VKJP3 at 10  $\mu\text{M}$  for different time periods (0, 36, and 72 h), and ABCG2 expression levels were examined by Western blotting analysis. As shown in Fig. 2B and C, the protein levels of ABCG2 in both cell lines were not significantly altered by VKJP1 or VKJP3 treatment, suggesting that the regulatory mechanisms of VKJP1 and VKJP3 are other than alteration of ABCG2 expression.

### 3.4. Effect of [ $^3\text{H}$ ]-mitoxantrone intracellular accumulation and [ $^3\text{H}$ ]-methotrexate transport by VKJP1 and VKJP3

In order to ascertain the mechanisms of VKJP1 and VKJP3 on the modulation of ABCG2 function, the intracellular levels of [ $^3\text{H}$ ]-mitoxantrone, a known substrate of ABCG2 was measured in the presence or absence of VKJP1 or VKJP3 in HEK293/pcDNA3.1, ABCG2-482-R2 and ABCG2-482-T7 cells (Fig. 3A). Both VKJP1 and VKJP3 at 10  $\mu\text{M}$ , were found to produce a significant increase in the intracellular levels of [ $^3\text{H}$ ]-mitoxantrone in both wild-type and mutant ABCG2 transfected cell lines. VKJP1 and VKJP3 at 10  $\mu\text{M}$  possess almost equal potency compared to the specific ABCG2 inhibitor FTC at 2.5  $\mu\text{M}$ . There was no significant change in the intracellular levels of [ $^3\text{H}$ ]-mitoxantrone in parental HEK293/pcDNA3.1 cells incubated either with VKJP1 or VKJP3. Due to the short time course, we did not see any difference between the potency of VKJP1 and VKJP3 on [ $^3\text{H}$ ]-mitoxantrone accumulation.

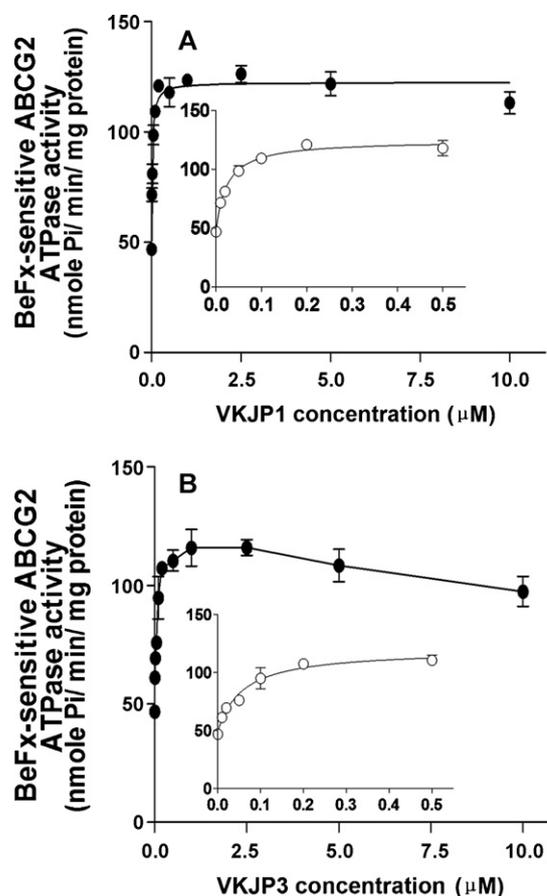
To further validate the ability of VKJP1 and VKJP3 to inhibit the transport activity of ABCG2, we performed transport experiments using membrane vesicles to determine the rate of ATP-dependent uptake of [ $^3\text{H}$ ]-methotrexate, another substrate of ABCG2. Previously, we have reported that only wild-type ABCG2 was able to robustly transport [ $^3\text{H}$ ]-methotrexate in the in vitro transport system [15,26]. Thus, in this study, we only used the membrane vesicles prepared from HEK293/pcDNA3.1 and ABCG2-482-R2 cell lines. Our data showed that VKJP1 and VKJP3 at 10  $\mu\text{M}$  produced a significant inhibition of the ATP-energized uptake rate of [ $^3\text{H}$ ]-methotrexate in ABCG2-482-R2 membrane vesicles (Fig. 3B), confirming that both VKJP1 and VKJP3 are able to directly inhibit the transport function of ABCG2-overexpressing membrane vesicles.

### 3.5. Effect of VKJP1 and VKJP3 on the ATPase activity of ABCG2

Since ABCG2 is an ATP-dependent membrane efflux pump which poses ATPase activity, we next measured the ABCG2-mediated ATP hydrolysis to investigate the effect of VKJP1 and VKJP3 on its ATPase activity. As shown in Fig. 4A and B, VKJP1 and VKJP3 enhanced the ATPase activity of ABCG2 in a concentration-dependent manner. The concentrations of VKJP1 or VKJP3 required for 50% stimulation of ATPase activity of ABCG2 were  $0.024 \pm 0.004$  and  $0.057 \pm 0.016$   $\mu\text{M}$ , respectively, indicating that VKJP1 and VKJP3 may be the substrates of ABCG2.

### 3.6. Effect of VKJP1 and VKJP3 on photoaffinity labeling of ABCG2 with [ $^{125}\text{I}$ ]-IAAP

ABCG2 is able to be photo-labeled by its substrate [ $^{125}\text{I}$ ]-IAAP, and its substrates as well as inhibitors can compete with [ $^{125}\text{I}$ ]-IAAP for binding to ABCG2. To understand whether there is physical interaction of VKJP1 or VKJP3 with the substrate interaction sites of ABCG2, we examined the photoaffinity labeling

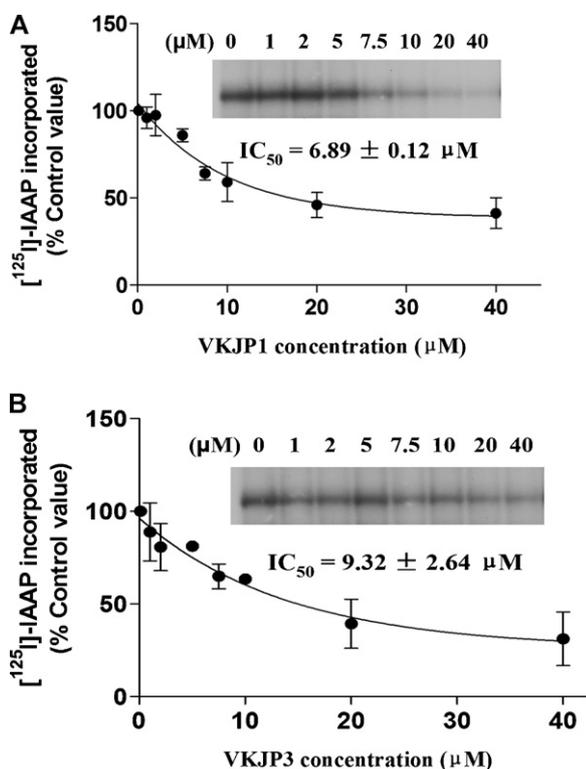


**Fig. 4.** Effect of VKJP1 and VKJP3 on the ATPase activity of ABCG2. ATPase activity of ABCG2 in membrane vesicles was measured with different concentrations of VKJP1 (panel A) and VKJP3 (panel B) as described in "Experimental Procedures". The lines represent the best fit for the data either by linear or non-linear least squares regression analysis using GraphPad Prism version 2.0. Spot, mean of triplicate determinations; bars, SD. In both panels A and B, inset shows ATP hydrolysis in the presence of 0–0.5  $\mu\text{M}$  VKJP1 and VKJP3. Experiments were repeated at least three times and a representative experiment is shown.

of ABCG2 with [ $^{125}\text{I}$ ]-IAAP by using the membranes in the presence of various concentrations of VKJP1 and VKJP3. As indicated in Fig. 5A and B, VKJP1 and VKJP3 significantly inhibited the photoaffinity labeling of ABCG2 with [ $^{125}\text{I}$ ]-IAAP in a concentration-dependent manner. The concentrations of VKJP1 and VKJP3 required for 50% inhibition of photoaffinity labeling were  $6.89 \pm 0.12$   $\mu\text{M}$  and  $9.32 \pm 2.64$   $\mu\text{M}$ , respectively. These results suggest that VKJP1 and VKJP3 could bind to ABCG2 substrate-binding sites with different affinity.

## 4. Discussion

ABCG2 is a 655-amino acid membrane protein, located on 4q22 chromosome, with a molecular weight of 72 kDa. It is a half transporter in since it has one single N-terminal nucleotide binding domain and one trans-membrane domain. Its unique structure indicates that it might have to dimerize with itself (homodimerize) or other members of ABC-G subfamily (heterodimerize) to function as an efflux transporter [34,35]. ABCG2 is expressed endogenously in many tissues, including placental syncytiotrophoblast cells, epithelial cells of gastrointestinal tract and liver, and endothelial cells of blood brain barrier [36]. It is also over-expressed in many kinds of MDR cancer tissues or established MDR cell lines. In recent years, ABCG2 has been shown to confer resistance to a diverse group of chemotherapeutic drugs, including nucleoside analogues,



**Fig. 5.** Effect of VKJP1 and VKJP3 on the photoaffinity labeling of ABCG2 by [<sup>125</sup>I]-IAAP. The photoaffinity labeling of ABCG2 with [<sup>125</sup>I]-IAAP was performed with different concentration of VKJP1 or VKJP3. The radioactivity incorporated into ABCG2 was determined by exposing the gel to an X-ray film at  $-70^{\circ}C$ . An autoradiogram and quantification of incorporation of IAAP into the ABCG2 band from at least three independent experiments.

anthracyclines, camptothecin-derived indolocarbazole topoisomerase I inhibitors, methotrexate, and flavopiridols [35]. Several compounds have been identified that serve as ABCG2 inhibitors and have been shown to overcome ABCG2-mediated MDR [37]. Additionally, many newly designed compounds are under investigation for their inhibitory effects in *in vitro* experiments. Several tyrosine kinase inhibitors (TKIs), including imatinib (Gleevec, STI571), AG1478, erlotinib (Tarceva, OSI-774), and lapatinib (Tykerb, GW572016) [19–23] have been identified as novel ABCG2 reversal agents by our group and others. These TKIs interact with ABCG2 and block its efflux function; increasing the intracellular accumulation of anticancer drugs and reverse ABCG2-mediated MDR.

In the current study, we found that nearly all the OSI-930 and its pyridyl and phenyl analogues were shown to reverse drug resistance in wild-type ABCG2 transfected MDR ABCG2-482-R2 cells. Among which, VKJP1 and VKJP3 had the most potency for increasing the cytotoxicity of ABCG2 substrate mitoxantrone. For this reason, we chose to conduct further mechanistic studies on these analogues. We found that meta-isothiocyanate derivative **13** had no significant reversing ability, while the para isothiocyanate derivative **14** interestingly had an opposite effect of protecting the cells from cytotoxicity by mitoxantrone (Table 1). We assume that these 15 OSI-930 analogues resulted in variant reversal or protective effects due to their structural differences. The isothiocyanate analogues in phenyl series were less potent in reversing the MDR. Based on the present results, OSI-930 analogues such as VKJP1, VKJP3 also showed the enhancement of cytotoxicity of mitoxantrone in the parental HEK293/pCDNA3.1 cells. One explanation could be that the parental cells might also have other endogenous drug transporters that we could not detect, or some

other mechanisms, which were interfered by the compounds, we tested.

Our previous data showed that amino acid residues at position 482 of ABCG2 is essential for substrate recognition [30]. In order to further study whether the reversal abilities of VKJP1 and VKJP3 is related to the amino acid residue, we used wild-type and mutant transfectants, ABCG2-482-R2 and ABCG2-482-T7 cells, in our experiments. Our results showed that VKJP1 and VKJP3 markedly increased the cytotoxicity of several ABCG2 anticancer substrates including mitoxantrone, SN-38, and doxorubicin in a concentration-dependent manner in both cell lines implying that the amino acid residues at ABCG2 482 position did not play an important role in the reversal abilities of VKJP1 and VKJP3. The reversal effect of VKJP1 on ABCG2-mediated MDR is more potent than that of VKJP3. The sensitization produced by 10 μM VKJP1 was similar to that induced by the known specific ABCG2 inhibitor FTC at 2.5 μM, showing that the use of VKJP1 as an ABCG2 modulator is very promising. Besides, both VKJP1 and VKJP3 were found to be ABCG2 selective inhibitors as demonstrated by their inability to significantly increase the sensitivity of all cell lines to a non ABCG2 substrate, cisplatin (Table 2) and inability to reverse ABCB1 or ABCC1 mediated drug resistance (Supplemental Tables 1 and 2).

Neither VKJP1 nor VKJP3 at 10 μM altered the expression levels of ABCG2 in the ABCG2-482-R2 and ABCG2-482-T7 cells suggesting that VKJP1 and VKJP3 exerted their inhibitory abilities by directly interacting with ABCG2 transporter. Consistent with the above hypothesis, we found that when ABCG2-482-R2 and ABCG2-482-T7 MDR cells were incubated concomitantly with [<sup>3</sup>H]-mitoxantrone and VKJP1 or VKJP3 at 10 μM, a higher intracellular drug accumulation was seen than when the cells were incubated with [<sup>3</sup>H]-mitoxantrone alone (Fig. 3A). A similar result was obtained when we examined the transport of [<sup>3</sup>H]-methotrexate using wild-type ABCG2 overexpressing membrane vesicles, the rates of [<sup>3</sup>H]-methotrexate uptake were significantly inhibited by VKJP1 or VKJP3 (Fig. 3B). These results were similar to some of the ABCG2 inhibitors that were previously reported by our research group. For example, we previously discovered that several TKIs such as AG1478, erlotinib, lapatinib could selectively modulate MDR protein-ATPase activity, by competitively inhibiting the efflux function of ABCG2 and ABCB1 and thus increase the intracellular accumulation of certain substrates into the MDR cells [19,22,23]. We then examined whether VKJP1 and VKJP3 have the ability to interact with the ABCG2 transporter substrate binding sites and stimulate ATP hydrolysis. Indeed this was confirmed by the finding that VKJP1 and VKJP3 significantly stimulated the ATP hydrolysis and inhibited the binding of [<sup>125</sup>I]-IAAP to the substrate binding sites of ABCG2 (Figs. 4 and 5).

In conclusion, our study shows that VKJP1 and VKJP3 have specific and the most potential reversal activity on ABCG2-mediated MDR by directly inhibiting the drug efflux function, resulting in an increase in the intracellular drug accumulation. The mechanistic study found that VKJP1 and VKJP3 stimulated the activity of ATPase and inhibited the photoaffinity labeling of ABCG2 with [<sup>125</sup>I]-IAAP. The pyridyl and phenyl analogues of OSI-930 are novel reversal agents of ABCG2 transporter with a potential to increase clinical response when combined with conventional chemotherapeutic agents.

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## Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at <http://dx.doi.org/10.1016/j.bcp.2012.06.019>.

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