

Martina Ceckova

List of Publications by Year in descending order

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52
papers

2,216
citations

218677

26
h-index

223800

46
g-index

52
all docs

52
docs citations

52
times ranked

2533
citing authors

#	ARTICLE	IF	CITATIONS
1	Dabrafenib inhibits ABCG2 and cytochrome P450 isoenzymes; potential implications for combination anticancer therapy. <i>Toxicology and Applied Pharmacology</i> , 2022, 434, 115797.	2.8	4
2	ABCB1 as a potential beneficial target of midostaurin in acute myeloid leukemia. <i>Biomedicine and Pharmacotherapy</i> , 2022, 150, 112962.	5.6	4
3	Interplay of drug transporters P-glycoprotein (MDR1), MRP1, OATP1A2 and OATP1B3 in passage of maraviroc across human placenta. <i>Biomedicine and Pharmacotherapy</i> , 2020, 129, 110506.	5.6	6
4	In vitro function and in situ localization of Multidrug Resistance-associated Protein (MRP)1 (ABCC1) suggest a protective role against methyl mercury-induced oxidative stress in the human placenta. <i>Archives of Toxicology</i> , 2020, 94, 3799-3817.	4.2	14
5	Targeting Pharmacokinetic Drug Resistance in Acute Myeloid Leukemia Cells with CDK4/6 Inhibitors. <i>Cancers</i> , 2020, 12, 1596.	3.7	13
6	Interactions between Maraviroc and the ABCB1, ABCG2, and ABCC2 Transporters: An Important Role in Transplacental Pharmacokinetics. <i>Drug Metabolism and Disposition</i> , 2019, 47, 954-960.	3.3	13
7	Brivanib Exhibits Potential for Pharmacokinetic Drug-Drug Interactions and the Modulation of Multidrug Resistance through the Inhibition of Human ABCG2 Drug Efflux Transporter and CYP450 Biotransformation Enzymes. <i>Molecular Pharmaceutics</i> , 2019, 16, 4436-4450.	4.6	22
8	Transport of ribavirin across the rat and human placental barrier: Roles of nucleoside and ATP-binding cassette drug efflux transporters. <i>Biochemical Pharmacology</i> , 2019, 163, 60-70.	4.4	11
9	Interactions of Alectinib with Human ATP-Binding Cassette Drug Efflux Transporters and Cytochrome P450 Biotransformation Enzymes: Effect on Pharmacokinetic Multidrug Resistance. <i>Drug Metabolism and Disposition</i> , 2019, 47, 699-709.	3.3	15
10	Cyclin-dependent kinase inhibitors AZD5438 and R547 show potential for enhancing efficacy of daunorubicin-based anticancer therapy: Interaction with carbonyl-reducing enzymes and ABC transporters. <i>Biochemical Pharmacology</i> , 2019, 163, 290-298.	4.4	9
11	The inhibitory effect of antiretroviral drugs on the L-carnitine uptake in human placenta. <i>Toxicology and Applied Pharmacology</i> , 2019, 368, 18-25.	2.8	10
12	Ribociclib shows potential for pharmacokinetic drug-drug interactions being a substrate of ABCB1 and potent inhibitor of ABCB1, ABCG2 and CYP450 isoforms in vitro. <i>Biochemical Pharmacology</i> , 2018, 154, 10-17.	4.4	41
13	LC-MS/MS method for determination of cyclin-dependent kinase inhibitors, BP-14 and BP-20, and its application in pharmacokinetic study in rat. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2018, 1089, 24-32.	2.3	0
14	Equilibrative Nucleoside Transporter 1 (ENT1, <i>SLC29A1</i>) Facilitates Transfer of the Antiretroviral Drug Abacavir across the Placenta. <i>Drug Metabolism and Disposition</i> , 2018, 46, 1817-1826.	3.3	25
15	Efavirenz reduces renal excretion of lamivudine in rats by inhibiting organic cation transporters (OCT, Oct) and multidrug and toxin extrusion proteins (MATE, Mate). <i>PLoS ONE</i> , 2018, 13, e0202706.	2.5	11
16	In vitro and in silico Evaluation of Non-Quaternary Reactivators of AChE as Antidotes of Organophosphorus Poisoning - a New Hope or a Blind Alley?. <i>Medicinal Chemistry</i> , 2018, 14, 281-292.	1.5	19
17	Emtricitabine is a substrate of MATE1 but not of OCT1, OCT2, P-gp, BCRP or MRP2 transporters. <i>Xenobiotica</i> , 2017, 47, 77-85.	1.1	27
18	Universal efavirenz determination in transport study, rat placenta perfusion and placenta lysate by HPLC-UV. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2017, 137, 70-77.	2.8	1

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19	Role of nucleoside transporters in transplacental pharmacokinetics of nucleoside reverse transcriptase inhibitors zidovudine and emtricitabine. <i>Placenta</i> , 2017, 60, 86-92.	1.5	12
20	MDR1 and BCRP Transporter-Mediated Drug-Drug Interaction between Rilpivirine and Abacavir and Effect on Intestinal Absorption. <i>Antimicrobial Agents and Chemotherapy</i> , 2017, 61, .	3.2	23
21	Placental passage of olomoucine II, but not purvalanol A, is affected by p-glycoprotein (ABCB1), breast cancer resistance protein (ABCG2) and multidrug resistance-associated proteins (ABCCs). <i>Xenobiotica</i> , 2016, 46, 416-423.	1.1	1
22	Etravirine inhibits ABCG2 drug transporter and affects transplacental passage of tenofovir disoproxil fumarate. <i>Placenta</i> , 2016, 47, 124-129.	1.5	13
23	Role of ABC and Solute Carrier Transporters in the Placental Transport of Lamivudine. <i>Antimicrobial Agents and Chemotherapy</i> , 2016, 60, 5563-5572.	3.2	19
24	Role of ABCB1, ABCG2, ABCC2 and ABCC5 transporters in placental passage of zidovudine. <i>Biopharmaceutics and Drug Disposition</i> , 2016, 37, 28-38.	1.9	24
25	Regulation of drug transporter expression and function in the placenta. <i>Expert Opinion on Drug Metabolism and Toxicology</i> , 2015, 11, 533-555.	3.3	40
26	Effect of drug efflux transporters on placental transport of antiretroviral agent abacavir. <i>Reproductive Toxicology</i> , 2015, 57, 176-182.	2.9	29
27	Boldine enhances bile production in rats via osmotic and Farnesoid X receptor dependent mechanisms. <i>Toxicology and Applied Pharmacology</i> , 2015, 285, 12-22.	2.8	19
28	Interactions of cyclin-dependent kinase inhibitors AT-7519, flavopiridol and SNS-032 with ABCB1, ABCG2 and ABCC1 transporters and their potential to overcome multidrug resistance in vitro. <i>Cancer Chemotherapy and Pharmacology</i> , 2015, 76, 105-116.	2.3	28
29	Dinaciclib, a cyclin-dependent kinase inhibitor, is a substrate of human ABCB1 and ABCG2 and an inhibitor of human ABCC1 in vitro. <i>Biochemical Pharmacology</i> , 2015, 98, 465-472.	4.4	27
30	Interactions of tenofovir and tenofovir disoproxil fumarate with drug efflux transporters ABCB1, ABCG2, and ABCC2; role in transport across the placenta. <i>Aids</i> , 2014, 28, 9-17.	2.2	68
31	Multidrug and toxin extrusion proteins (MATE/SLC47); role in pharmacokinetics. <i>International Journal of Biochemistry and Cell Biology</i> , 2013, 45, 2007-2011.	2.8	61
32	Tetratricopeptide Repeat Motifs in the World of Bacterial Pathogens: Role in Virulence Mechanisms. <i>Infection and Immunity</i> , 2013, 81, 629-635.	2.2	156
33	Organic Cation Transporter 3 (OCT3/SLC22A3) and Multidrug and Toxin Extrusion 1 (MATE1/SLC47A1) Transporter in the Placenta and Fetal Tissues: Expression Profile and Fetus Protective Role at Different Stages of Gestation1. <i>Biology of Reproduction</i> , 2013, 88, 55.	2.7	58
34	Olomoucine II, but Not Purvalanol A, Is Transported by Breast Cancer Resistance Protein (ABCG2) and P-Glycoprotein (ABCB1). <i>PLoS ONE</i> , 2013, 8, e75520.	2.5	6
35	Purvalanol A, Olomoucine II and Roscovitine Inhibit ABCB1 Transporter and Synergistically Potentiate Cytotoxic Effects of Daunorubicin In Vitro. <i>PLoS ONE</i> , 2013, 8, e83467.	2.5	27
36	Synchronized Activity of Organic Cation Transporter 3 (OCT3/SLC22A3) and Multidrug and Toxin Extrusion 1 (MATE1/SLC47A1) Transporter in Transplacental Passage of MPP+ in Rat. <i>Toxicological Sciences</i> , 2012, 128, 471-481.	3.1	38

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37	Pharmacotherapy in pregnancy; effect of ABC and SLC transporters on drug transport across the placenta and fetal drug exposure. <i>Journal of Drug Targeting</i> , 2012, 20, 736-763.	4.4	99
38	Olomoucine II and purvalanol A inhibit ABCG2 transporter in vitro and in situ and synergistically potentiate cytostatic effect of mitoxantrone. <i>Pharmacological Research</i> , 2012, 65, 312-319.	7.1	23
39	Fetoprotective activity of breast cancer resistance protein (BCRP, ABCG2): expression and function throughout pregnancy. <i>Drug Metabolism Reviews</i> , 2011, 43, 53-68.	3.6	42
40	Expression and Function of P-Glycoprotein in Normal Tissues: Effect on Pharmacokinetics. <i>Methods in Molecular Biology</i> , 2010, 596, 199-222.	0.9	74
41	Transplacental Pharmacokinetics of Glyburide, Rhodamine 123, and BODIPY FL Prazosin: Effect of Drug Efflux Transporters and Lipid Solubility. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2009, 331, 1118-1125.	2.5	64
42	Variation of Drug Kinetics in Pregnancy. <i>Current Drug Metabolism</i> , 2009, 10, 520-529.	1.2	93
43	Role of breast cancer resistance protein (Bcrp/Abcg2) in fetal protection during gestation in rat. <i>Toxicology Letters</i> , 2008, 178, 176-180.	0.8	44
44	Effect of ABCG2 on cytotoxicity of platinum drugs: Interference of EGFP. <i>Toxicology in Vitro</i> , 2008, 22, 1846-1852.	2.4	28
45	Salicylanilide Acetates: Synthesis and Antibacterial Evaluation. <i>Molecules</i> , 2007, 12, 1-12.	3.8	40
46	EXPRESSION AND FUNCTIONAL ACTIVITY OF BREAST CANCER RESISTANCE PROTEIN (BCRP, ABCG2) TRANSPORTER IN THE HUMAN CHORIOCARCINOMA CELL LINE BEWO. <i>Clinical and Experimental Pharmacology and Physiology</i> , 2006, 33, 58-65.	1.9	74
47	Synthesis and antimicrobial evaluation of new 2-substituted 5,7-di-tert-butylbenzoxazoles. <i>Bioorganic and Medicinal Chemistry</i> , 2006, 14, 5850-5865.	3.0	100
48	P-glycoprotein in the placenta: Expression, localization, regulation and function. <i>Reproductive Toxicology</i> , 2006, 22, 400-410.	2.9	187
49	Expression and Transport Activity of Breast Cancer Resistance Protein (Bcrp/Abcg2) in Dually Perfused Rat Placenta and HRP-1 Cell Line. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2006, 319, 53-62.	2.5	79
50	Human Breast Cancer Resistance Protein: Interactions with Steroid Drugs, Hormones, the Dietary Carcinogen 2-Amino-1-methyl-6-phenylimidazo(4,5- <i>b</i>)pyridine, and Transport of Cimetidine. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2005, 312, 144-152.	2.5	258
51	P-glycoprotein expression and distribution in the rat placenta during pregnancy. <i>Reproductive Toxicology</i> , 2004, 18, 785-792.	2.9	63
52	Examination of the Functional Activity of P-glycoprotein in the Rat Placental Barrier Using Rhodamine 123. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2003, 305, 1239-1250.	2.5	54