List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Conjugate export pumps of the multidrug resistance protein (MRP) family: localization, substrate specificity, and MRP2-mediated drug resistance. Biochimica Et Biophysica Acta - Biomembranes, 1999, 1461, 377-394.	2.6	681
2	cDNA Cloning of the Hepatocyte Canalicular Isoform of the Multidrug Resistance Protein, cMrp, Reveals a Novel Conjugate Export Pump Deficient in Hyperbilirubinemic Mutant Rats. Journal of Biological Chemistry, 1996, 271, 15091-15098.	3.4	580
3	A novel human organic anion transporting polypeptide localized to the basolateral hepatocyte membrane. American Journal of Physiology - Renal Physiology, 2000, 278, G156-G164.	3.4	479
4	Transporters and Drug-Drug Interactions: Important Determinants of Drug Disposition and Effects. Pharmacological Reviews, 2013, 65, 944-966.	16.0	475
5	Localization and Genomic Organization of a New Hepatocellular Organic Anion Transporting Polypeptide. Journal of Biological Chemistry, 2000, 275, 23161-23168.	3.4	462
6	Hepatic Uptake of Bilirubin and Its Conjugates by the Human Organic Anion Transporter SLC21A6. Journal of Biological Chemistry, 2001, 276, 9626-9630.	3.4	458
7	Characterization of the human multidrug resistance protein isoform MRP3 localized to the basolateral hepatocyte membrane. Hepatology, 1999, 29, 1156-1163.	7.3	430
8	Pharmacogenomics of human OATP transporters. Naunyn-Schmiedeberg's Archives of Pharmacology, 2006, 372, 432-443.	3.0	308
9	ABCC Drug Efflux Pumps and Organic Anion Uptake Transporters in Human Gliomas and the Blood-Tumor Barrier. Cancer Research, 2005, 65, 11419-11428.	0.9	266
10	Expression and localization of the conjugate export pump encoded by the <i>MRP2 (cMRP/cMOAJ)</i> gene in liver. FASEB Journal, 1997, 11, 509-515.	0.5	265
11	Hepatic Secretion of Conjugated Drugs and Endogenous Substances. Seminars in Liver Disease, 2000, Volume 20, 265-272.	3.6	224
12	Expression of the MRP2 Gene-Encoded Conjugate Export Pump in Human Kidney Proximal Tubules and in Renal Cell Carcinoma. Journal of the American Society of Nephrology: JASN, 1999, 10, 1159-1169.	6.1	224
13	Vectorial Transport by Double-Transfected Cells Expressing the Human Uptake Transporter SLC21A8 and the Apical Export Pump ABCC2. Molecular Pharmacology, 2001, 60, 934-943.	2.3	209
14	Human Hepatobiliary Transport of Organic Anions Analyzed by Quadruple-Transfected Cells. Molecular Pharmacology, 2005, 68, 1031-1038.	2.3	193
15	Deletion of the Mammalian INDY Homolog Mimics Aspects of Dietary Restriction and Protects against Adiposity and Insulin Resistance in Mice. Cell Metabolism, 2011, 14, 184-195.	16.2	193
16	The Influence of Macrolide Antibiotics on the Uptake of Organic Anions and Drugs Mediated by OATP1B1 and OATP1B3. Drug Metabolism and Disposition, 2007, 35, 779-786.	3.3	175
17	Mutations in the SLCO1B3 gene affecting the substrate specificity of the hepatocellular uptake transporter OATP1B3 (OATP8). Pharmacogenetics and Genomics, 2004, 14, 441-452.	5.7	170
18	Expression and localization of human multidrug resistance protein (ABCC) family members in pancreatic carcinoma, International Journal of Cancer, 2005. 115. 359-367.	5.1	165

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19	Expression of the multidrug resistance proteins MRP2 and MRP3 in human hepatocellular carcinoma. International Journal of Cancer, 2001, 94, 492-499.	5.1	163
20	The multidrug resistance protein MRP1 mediates the release of glutathione disulfide from rat astrocytes during oxidative stress. Journal of Neurochemistry, 2001, 76, 627-636.	3.9	153
21	Transport of monoglucuronosyl and bisglucuronosyl bilirubin by recombinant human and rat multidrug resistance protein 2. Hepatology, 1999, 30, 485-490.	7.3	151
22	Changes in the expression and localization of hepatocellular transporters and radixin in primary biliary cirrhosis. Journal of Hepatology, 2003, 39, 693-702.	3.7	149
23	Exon-intron organization of the human multidrug-resistance protein 2 (MRP2) gene mutated in Dubin–Johnson syndrome. Gastroenterology, 1999, 117, 653-660.	1.3	148
24	A Naturally Occurring Mutation in the SLC21A6Gene Causing Impaired Membrane Localization of the Hepatocyte Uptake Transporter. Journal of Biological Chemistry, 2002, 277, 43058-43063.	3.4	127
25	Influence of the flavonoids apigenin, kaempferol, and quercetin on the function of organic anion transporting polypeptides 1A2 and 2B1. Biochemical Pharmacology, 2010, 80, 1746-1753.	4.4	121
26	Expression of mRNAs of multidrug resistance proteins (Mrps) in cultured rat astrocytes, oligodendrocytes, microglial cells and neurones. Journal of Neurochemistry, 2002, 82, 716-719.	3.9	120
27	Interaction of Oral Antidiabetic Drugs With Hepatic Uptake Transporters. Diabetes, 2008, 57, 1463-1469.	0.6	111
28	Uptake Transporters of the Human OATP Family. Handbook of Experimental Pharmacology, 2011, , 1-28.	1.8	107
29	Detection of the Human Organic Anion Transporters SLC21A6 (OATP2) and SLC21A8 (OATP8) in Liver and Hepatocellular Carcinoma. Laboratory Investigation, 2003, 83, 527-538.	3.7	105
30	Role of P-Glycoprotein Inhibition forÂDrug Interactions. Clinical Pharmacokinetics, 2007, 46, 1039-1049.	3.5	101
31	Structural determinants of inhibitor interaction with the human organic cation transporter OCT2 (SLC22A2). Naunyn-Schmiedeberg's Archives of Pharmacology, 2009, 379, 337-348.	3.0	101
32	Expression and localization of the multidrug resistance proteins MRP2 and MRP3 in human gallbladder epithelia. Gastroenterology, 2001, 121, 1203-1208.	1.3	99
33	Influence of Non-Steroidal Anti-Inflammatory Drugs on Organic Anion Transporting Polypeptide (OATP) 1B1- and OATP1B3-Mediated Drug Transport. Drug Metabolism and Disposition, 2011, 39, 1047-1053.	3.3	94
34	Hepatic OATP and OCT uptake transporters: their role for drug-drug interactions and pharmacogenetic aspects. Drug Metabolism Reviews, 2010, 42, 380-401.	3.6	93
35	Clinical Aspects of Transporterâ€Mediated Drug–Drug Interactions. Clinical Pharmacology and Therapeutics, 2019, 105, 1386-1394.	4.7	88
36	Characterization of the 5′-flanking region of the human multidrug resistance protein 2 (MRP2) gene and its regulation in comparison withthe multidrug resistance protein 3 (MRP3) gene. FEBS Journal, 2000, 267, 1347-1358.	0.2	87

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37	Disposition of ezetimibe is influenced by polymorphisms of the hepatic uptake carrier OATP1B1. Pharmacogenetics and Genomics, 2008, 18, 559-568.	1.5	87
38	Export pumps for anionic conjugates encoded by MRP genes. Advances in Enzyme Regulation, 1999, 39, 237-246.	2.6	86
39	Role of organic cation transporter OCT2 and multidrug and toxin extrusion proteins MATE1 and MATE2-K for transport and drug interactions of the antiviral lamivudine. Biochemical Pharmacology, 2013, 86, 808-815.	4.4	85
40	Identification and functional characterization of the natural variant MRP3-Arg1297His of human multidrug resistance protein 3 (MRP3/ABCC3). Pharmacogenetics and Genomics, 2004, 14, 213-223.	5.7	84
41	The canalicular multidrug resistance protein, cMRP/MRP2, a novel conjugate export pump expressed in the apical membrane of hepatocytes. Advances in Enzyme Regulation, 1997, 37, 321-333.	2.6	82
42	Functional Characterization of the Human Organic Cation Transporter 2 Variant p.270Ala>Ser. Drug Metabolism and Disposition, 2009, 37, 1312-1318.	3.3	80
43	Localization, substrate specificity, and drug resistance conferred by conjugate export pumps of the MRP family. Advances in Enzyme Regulation, 2000, 40, 339-349.	2.6	71
44	<i>In vitro</i> evidence for the role of OATP and OCT uptake transporters in drug–drug interactions. Expert Opinion on Drug Metabolism and Toxicology, 2009, 5, 489-500.	3.3	71
45	Structural requirements for the apical sorting of human multidrug resistance protein 2 (ABCC2). FEBS Journal, 2002, 269, 1866-1876.	0.2	64
46	Molecular Mechanism of Renal Tubular Secretion of the Antimalarial Drug Chloroquine. Antimicrobial Agents and Chemotherapy, 2011, 55, 3091-3098.	3.2	64
47	Inhibitory Effects of Green Tea and (–)-Epigallocatechin Gallate on Transport by OATP1B1, OATP1B3, OCT1, OCT2, MATE1, MATE2-K and P-Glycoprotein. PLoS ONE, 2015, 10, e0139370.	2.5	64
48	Biomarkers for In Vivo Assessment of Transporter Function. Pharmacological Reviews, 2018, 70, 246-277.	16.0	59
49	The human longevity gene homolog INDY and interleukinâ€6 interact in hepatic lipid metabolism. Hepatology, 2017, 66, 616-630.	7.3	55
50	Expression and localization of the uptake transporters OATP2B1, OATP3A1 and OATP5A1 in non-malignant and malignant breast tissue. Cancer Biology and Therapy, 2011, 11, 584-591.	3.4	54
51	Genome-Wide Association Study of <scp>l</scp> -Arginine and Dimethylarginines Reveals Novel Metabolic Pathway for Symmetric Dimethylarginine. Circulation: Cardiovascular Genetics, 2014, 7, 864-872.	5.1	53
52	Inhibition of Mouse Erythroid Band 3-Mediated Chloride Transport by Site-Directed Mutagenesis of Histidine Residues and Its Reversal by Second Site Mutation of Lys 558, the Locus of Covalent H2DIDS Binding. Biochemistry, 1995, 34, 9315-9324.	2.5	52
53	Interaction of the cardiovascular risk marker asymmetric dimethylarginine (ADMA) with the human cationic amino acid transporter 1 (CAT1). Journal of Molecular and Cellular Cardiology, 2012, 53, 392-400.	1.9	52
54	Gender Is an Important Determinant of the Disposition of the Loop Diuretic Torasemide. Journal of Clinical Pharmacology, 2010, 50, 160-168.	2.0	49

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55	The Prostaglandin Transporter OATP2A1 Is Expressed in Human Ocular Tissues and Transports the Antiglaucoma Prostanoid Latanoprost. , 2010, 51, 2504.		48
56	<i>SLCO1B1</i> genetic polymorphism influences mycophenolic acid tolerance in renal transplant recipients. Pharmacogenomics, 2010, 11, 1703-1713.	1.3	48
57	ATP-Binding Cassette Transporters in the Heart. Trends in Cardiovascular Medicine, 2006, 16, 7-15.	4.9	44
58	The functional consequences of genetic variations in transporter genes encoding human organic anion-transporting polypeptide family members. Expert Opinion on Drug Metabolism and Toxicology, 2008, 4, 51-64.	3.3	44
59	Transport of asymmetric dimethylarginine (ADMA) by cationic amino acid transporter 2 (CAT2), organic cation transporter 2 (OCT2) and multidrug and toxin extrusion protein 1 (MATE1). Amino Acids, 2013, 45, 989-1002.	2.7	41
60	Role of Organic Anion-Transporting Polypeptides for Cellular Mesalazine (5-Aminosalicylic Acid) Uptake. Drug Metabolism and Disposition, 2011, 39, 1097-1102.	3.3	40
61	Purification of the human apical conjugate export pump MRP2. Reconstitution and functional characterization as substrate-stimulated ATPase. FEBS Journal, 1999, 265, 281-289.	0.2	39
62	In vivo evidence that Agxt2 can regulate plasma levels of dimethylarginines in mice. Biochemical and Biophysical Research Communications, 2013, 430, 84-89.	2.1	36
63	Alanine-glyoxylate aminotransferase 2 (AGXT2) Polymorphisms Have Considerable Impact on Methylarginine and β-aminoisobutyrate Metabolism in Healthy Volunteers. PLoS ONE, 2014, 9, e88544.	2.5	33
64	The Nonmetabolized β-Blocker Nadolol Is a Substrate of OCT1, OCT2, MATE1, MATE2-K, and P-Glycoprotein, but Not of OATP1B1 and OATP1B3. Molecular Pharmaceutics, 2016, 13, 512-519.	4.6	33
65	MRP2, THE APICAL EXPORT PUMP FOR ANIONIC CONJUGATES. , 2003, , 423-443.		29
66	Transporter-Mediated Drug–Drug Interactions with Oral Antidiabetic Drugs. Pharmaceutics, 2011, 3, 680-705.	4.5	29
67	Tropane alkaloids as substrates and inhibitors of human organic cation transporters of the SLC22 (OCT) and the SLC47 (MATE) families. Biological Chemistry, 2017, 398, 237-249.	2.5	28
68	Functional analysis of the polymorphism â^211C>T in the regulatory region of the human ABCC3 gene. Life Sciences, 2007, 80, 1490-1494.	4.3	27
69	The prognostic biomarker L-homoarginine is a substrate of the cationic amino acid transporters CAT1, CAT2A and CAT2B. Scientific Reports, 2017, 7, 4767.	3.3	27
70	Human Mast Cells Secreting Leukotriene C4 Express the MRP1 Gene-Encoded Conjugate Export Pump. Biological Chemistry, 1998, 379, 1121-6.	2.5	25
71	Characterization of Ursodeoxycholic and Norursodeoxycholic Acid as Substrates of the Hepatic Uptake Transporters <scp>OATP</scp> 1B1, <scp>OATP</scp> 1B3, <scp>OATP</scp> 2B1 and <scp>NTCP</scp> . Basic and Clinical Pharmacology and Toxicology, 2012, 111, 81-86.	2.5	24
72	Analysis of naturally occurring mutations in the human uptake transporter NaCT important for bone and brain development and energy metabolism. Scientific Reports, 2018, 8, 11330.	3.3	24

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73	The Influence of Oral Antidiabetic Drugs on Cellular Drug Uptake Mediated by Hepatic <scp>OATP</scp> Family Members. Basic and Clinical Pharmacology and Toxicology, 2013, 112, 244-250.	2.5	23
74	Importance of OCT2 and MATE1 for the Cimetidine–Metformin Interaction: Insights from Investigations of Polarized Transport in Single- And Double-Transfected MDCK Cells with a Focus on Perpetrator Disposition. Molecular Pharmaceutics, 2018, 15, 3425-3433.	4.6	23
75	Contribution of MATE1 to Renal Secretion of the NMDA Receptor Antagonist Memantine. Molecular Pharmaceutics, 2017, 14, 2991-2998.	4.6	22
76	Transport of L-Arginine Related Cardiovascular Risk Markers. Journal of Clinical Medicine, 2020, 9, 3975.	2.4	21
77	The renal transport protein OATP4C1 mediates uptake of the uremic toxin asymmetric dimethylarginine (ADMA) and efflux of cardioprotective L-homoarginine. PLoS ONE, 2019, 14, e0213747.	2.5	17
78	Interplay of the Organic Cation Transporters OCT1 and OCT2 with the Apically Localized Export Protein MATE1 for the Polarized Transport of Trospium. Molecular Pharmaceutics, 2019, 16, 510-517.	4.6	14
79	Interaction of Remdesivir with Clinically Relevant Hepatic Drug Uptake Transporters. Pharmaceutics, 2021, 13, 369.	4.5	14
80	Reconstitution of Transport-Active Multidrug Resistance Protein 2 (MRP2; ABCC2) in Proteoliposomes. Biological Chemistry, 2002, 383, 1001-9.	2.5	13
81	Impact of the CYP3A5 genotype on midazolam pharmacokinetics and pharmacodynamics during intensive care sedation. European Journal of Clinical Pharmacology, 2007, 63, 1129-1133.	1.9	13
82	Esters of Bendamustine Are by Far More Potent Cytotoxic Agents than the Parent Compound against Human Sarcoma and Carcinoma Cells. PLoS ONE, 2015, 10, e0133743.	2.5	12
83	Transport of Drugs and Endogenous Compounds Mediated by Human OCT1: Studies in Single- and Double-Transfected Cell Models. Frontiers in Pharmacology, 2021, 12, 662535.	3.5	11
84	Non-synonymous polymorphisms in the human SLCO1B1 gene: an in vitro analysis of SNPÂc.1929A>C. Molecular Genetics and Genomics, 2008, 279, 149-157.	2.1	9
85	Deletion of the diabetes candidate gene Slc16a13 in mice attenuates diet-induced ectopic lipid accumulation and insulin resistance. Communications Biology, 2021, 4, 826.	4.4	6
86	Vectorial transport of the arginine derivatives asymmetric dimethylarginine (ADMA) and l-homoarginine by OATP4C1 and P-glycoprotein studied in double-transfected MDCK cells. Amino Acids, 2020, 52, 975-985.	2.7	5
87	Assays for Analyzing the Role of Transport Proteins in the Uptake and the Vectorial Transport of Substances Affecting Cell Viability. Methods in Molecular Biology, 2017, 1601, 123-135.	0.9	5
88	Molecular analysis of regulation of gene expression of the human erythroid anion exchanger (AE) 1. FEBS Letters, 1998, 438, 315-320.	2.8	4
89	Deletion of the Mammalian INDY Homolog Mimics Aspects of Dietary Restriction and Protects against Adiposity and Insulin Resistance in Mice. Cell Metabolism, 2011, 14, 567.	16.2	4
90	Analysis of amino acid residues in the predicted transmembrane pore influencing transport kinetics of the hepatic drug transporter organic anion transporting polypeptide 1B1 (OATP1B1). Biochimica Et Biophysica Acta - Biomembranes, 2016, 1858, 2894-2902.	2.6	4

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91	Exposure of Fexofenadine, but Not Pseudoephedrine, Is Markedly Decreased by Green Tea Extract in Healthy Volunteers. Clinical Pharmacology and Therapeutics, 2022, 112, 627-634.	4.7	4
92	Role of Uptake Transporters OAT4, OATP2A1, and OATP1A2 in Human Placental Bio-disposition of Pravastatin. Journal of Pharmaceutical Sciences, 2022, 111, 505-516.	3.3	3
93	Carbamazepine efficacy in a severe electroâ€clinical presentation of <scp><i>SLC13A5</i></scp> â€epilepsy. Annals of Clinical and Translational Neurology, 2022, 9, 1095-1099.	3.7	3
94	The human hepatocyte-specific organic anion transporter encoded by the SLC21A8 gene. Gastroenterology, 2002, 122, 1545-1546.	1.3	2
95	Screening of commonly prescribed drugs for effects on the CAT1-mediated transport of l-arginine and arginine derivatives. Amino Acids, 2022, 54, 1101-1108.	2.7	2
96	L-Arginine and Cardioactive Arginine Derivatives as Substrates and Inhibitors of Human and Mouse NaCT/Nact. Metabolites, 2022, 12, 273.	2.9	2
97	The Role of ABCC Family Members in the Disposition of Endogenous Compounds and Drugs. , 2011, , 209-245.		0
98	Transport of Bilirubin Conjugates across Hepatocellular Membrane Domains and the Conjugated Hyperbilirubinemia of Dubin-Johnson Syndrome. , 2004, , 195-210.		0
99	Zebrafish Oatp1d1 Acts as a Cellular Efflux Transporter of the Anionic Herbicide Bromoxynil. Chemical Research in Toxicology, 2022, , .	3.3	0