Kathleen M Giacomini

List of Publications by Year in Descending Order

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ext. papers

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

 142
 10,303
 46
 100

 papers
 citations
 h-index
 g-index

 151
 11,800
 7.6
 5.91

ext. citations

avg, IF

L-index

#	Paper	IF	Citations
142	High Throughput Screening of a Prescription Drug Library for Inhibitors of Organic Cation Transporter 3, OCT3 <i>Pharmaceutical Research</i> , 2022 , 1	4.5	2
141	The Clinical Pharmacogenetics Implementation Consortium (CPIC) guideline for SLCO1B1, ABCG2, and CYP2C9 and statin-associated musculoskeletal symptoms <i>Clinical Pharmacology and Therapeutics</i> , 2022 ,	6.1	9
140	Mechanisms and genetics of drug transport 2022 , 213-239		O
139	Pharmacogenomic mechanisms of drug toxicity 2022 , 303-322		
138	Response to Comment on Dawed et al. Genome-Wide Meta-analysis Identifies Genetic Variants Associated With Glycemic Response to Sulfonylureas. Diabetes Care 2021;44:2673-2682 <i>Diabetes Care</i> , 2022 , 45, e82-e83	14.6	
137	ORGANIC CATION AND ZWITTERION TRANSPORTERS 2022 , 9-32		
136	A Critical Overview of the Biological Effects of Excipients (Part I): Impact on Gastrointestinal Absorption <i>AAPS Journal</i> , 2022 , 24, 60	3.7	O
135	Drug Metabolism and Disposition, 2021 ,	4	4
134	Genome-Wide Meta-analysis Identifies Genetic Variants Associated With Glycemic Response to Sulfonylureas. <i>Diabetes Care</i> , 2021 , 44, 2673-2682	14.6	5
133	The Effects of Genetic Mutations and Drugs on the Activity of the Thiamine Transporter, SLC19A2. <i>AAPS Journal</i> , 2021 , 23, 35	3.7	1
132	Opportunities and challenges for the computational interpretation of rare variation in clinically important genes. <i>American Journal of Human Genetics</i> , 2021 , 108, 535-548	11	10
131	Oxypurinol pharmacokinetics and pharmacodynamics in healthy volunteers: Influence of BCRP Q141K polymorphism and patient characteristics. <i>Clinical and Translational Science</i> , 2021 , 14, 1431-1443	4.9	2
130	Drugs in COVID-19 Clinical Trials: Predicting Transporter-Mediated Drug-Drug Interactions Using In Vitro Assays and Real-World Data. <i>Clinical Pharmacology and Therapeutics</i> , 2021 , 110, 108-122	6.1	3
129	Drug Metabolites Potently Inhibit Renal Organic Anion Transporters, OAT1 and OAT3. <i>Journal of Pharmaceutical Sciences</i> , 2021 , 110, 347-353	3.9	4
128	Characterization of cytochrome P450 (CYP) 2D6 drugs as substrates of human organic cation transporters and multidrug and toxin extrusion proteins. <i>British Journal of Pharmacology</i> , 2021 , 178, 1459-1474	8.6	3
127	A New Era in Pharmacovigilance: Toward Real-World Data and Digital Monitoring. <i>Clinical Pharmacology and Therapeutics</i> , 2021 , 109, 1197-1202	6.1	8
126	Advancing Precision Medicine Through the New Pharmacogenomics Global Research Network. <i>Clinical Pharmacology and Therapeutics</i> , 2021 , 110, 559-562	6.1	4

125	Interaction of Commonly Used Oral Molecular Excipients with P-glycoprotein. <i>AAPS Journal</i> , 2021 , 23, 106	3.7	1
124	Bacterial metabolism rescues the inhibition of intestinal drug absorption by food and drug additives. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020 , 117, 16009-16018	11.5	15
123	GenEpi: gene-based epistasis discovery using machine learning. <i>BMC Bioinformatics</i> , 2020 , 21, 68	3.6	10
122	Interactions of Oral Molecular Excipients with Breast Cancer Resistance Protein, BCRP. <i>Molecular Pharmaceutics</i> , 2020 , 17, 748-756	5.6	12
121	Global Pharmacogenomics Within Precision Medicine: Challenges and Opportunities. <i>Clinical Pharmacology and Therapeutics</i> , 2020 , 107, 57-61	6.1	17
120	Drug-nutrient interactions: discovering prescription drug inhibitors of the thiamine transporter ThTR-2 (SLC19A3). <i>American Journal of Clinical Nutrition</i> , 2020 , 111, 110-121	7	15
119	Expanding Precompetitive Multisector Collaborations to Advance Drug Development and Pharmacogenomics. <i>Clinical Pharmacology and Therapeutics</i> , 2020 , 107, 96-101	6.1	3
118	The activities of drug inactive ingredients on biological targets. <i>Science</i> , 2020 , 369, 403-413	33.3	34
117	Scientific considerations for global drug development. Science Translational Medicine, 2020, 12,	17.5	6
116	Neural production of kynurenic acid in requires the AAT-1 transporter. <i>Genes and Development</i> , 2020 , 34, 1033-1038	12.6	1
115	Deorphaning a solute carrier 22 family member, SLC22A15, through functional genomic studies. <i>FASEB Journal</i> , 2020 , 34, 15734-15752	0.9	13
114	Unraveling the functional role of the orphan solute carrier, SLC22A24 in the transport of steroid conjugates through metabolomic and genome-wide association studies. <i>PLoS Genetics</i> , 2019 , 15, e1008	208	14
113	Impact of Pharmaceutical Excipients on Oral Drug Absorption: A Focus on Intestinal Drug Transporters. <i>Clinical Pharmacology and Therapeutics</i> , 2019 , 105, 323-325	6.1	7
112	l-Type amino acid transporter 1 activity of 1,2,3-triazolyl analogs of l-histidine and l-tryptophan. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2019 , 29, 2254-2258	2.9	10
111	A Comprehensive Analysis of Ontogeny of Renal Drug Transporters: mRNA Analyses, Quantitative Proteomics, and Localization. <i>Clinical Pharmacology and Therapeutics</i> , 2019 , 106, 1083-1092	6.1	45
110	Organic Anion Transporter Polypeptide 1B1 Polymorphism Modulates the Extent of Drug-Drug Interaction and Associated Biomarker Levels in Healthy Volunteers. <i>Clinical and Translational Science</i> , 2019 , 12, 388-399	4.9	26
109	Genome-Wide Association and Functional Studies Reveal Novel Pharmacological Mechanisms for Allopurinol. <i>Clinical Pharmacology and Therapeutics</i> , 2019 , 106, 623-631	6.1	15
108	Functional and structural analysis of rare SLC2A2 variants associated with Fanconi-Bickel syndrome and metabolic traits. <i>Human Mutation</i> , 2019 , 40, 983-995	4.7	6

107	Research Projects Supported by the University of California, San Francisco-Stanford Center of Excellence in Regulatory Science and Innovation. <i>Clinical Pharmacology and Therapeutics</i> , 2019 , 105, 81.	5-818	3
106	A conserved role of the insulin-like signaling pathway in diet-dependent uric acid pathologies in Drosophila melanogaster. <i>PLoS Genetics</i> , 2019 , 15, e1008318	6	20
105	In Vitro Evaluation of Excipients as Inhibitors of Human Intestinal P-glycoprotein. <i>FASEB Journal</i> , 2019 , 33, 814.3	0.9	1
104	Organic cation transporter 3 (Oct3) is a distinct catecholamines clearance route in adipocytes mediating the beiging of white adipose tissue. <i>PLoS Biology</i> , 2019 , 17, e2006571	9.7	21
103	The Role of Transporters in Drug Accumulation and Mitochondrial Toxicity 2018, 15-24		
102	Influence of Transporter Polymorphisms on Drug Disposition and Response: A Perspective From the International Transporter Consortium. <i>Clinical Pharmacology and Therapeutics</i> , 2018 , 104, 803-817	6.1	60
101	Genetic Variants in and Are Associated With Variation in Response to Metformin in Individuals With Type 2 Diabetes. <i>Diabetes</i> , 2018 , 67, 1428-1440	0.9	18
100	Molecular Mechanisms for Species Differences in Organic Anion Transporter 1, OAT1: Implications for Renal Drug Toxicity. <i>Molecular Pharmacology</i> , 2018 , 94, 689-699	4.3	29
99	Reverse Translational Research of ABCG2 (BCRP) in Human Disease and Drug Response. <i>Clinical Pharmacology and Therapeutics</i> , 2018 , 103, 233-242	6.1	17
98	Reevaluating the Substrate Specificity of the L-Type Amino Acid Transporter (LAT1). <i>Journal of Medicinal Chemistry</i> , 2018 , 61, 7358-7373	8.3	38
97	Organic cation transporter 1 (OCT1) modulates multiple cardiometabolic traits through effects on hepatic thiamine content. <i>PLoS Biology</i> , 2018 , 16, e2002907	9.7	29
96	Transporters in Drug Development: 2018 ITC Recommendations for Transporters of Emerging Clinical Importance. <i>Clinical Pharmacology and Therapeutics</i> , 2018 , 104, 890-899	6.1	113
95	Emerging Clinical Importance of Hepatic Organic Cation Transporter 1 (OCT1) in Drug Pharmacokinetics, Dynamics, Pharmacogenetic Variability, and Drug Interactions. <i>Clinical Pharmacology and Therapeutics</i> , 2018 , 103, 758-760	6.1	31
94	Clinical Probes and Endogenous Biomarkers as Substrates for Transporter Drug-Drug Interaction Evaluation: Perspectives From the International Transporter Consortium. <i>Clinical Pharmacology and Therapeutics</i> , 2018 , 104, 836-864	6.1	77
93	Pharmacogenetics of Antidiabetic Drugs. <i>Advances in Pharmacology</i> , 2018 , 83, 361-389	5.7	9
92	ITC Commentary on Metformin Clinical Drug-Drug Interaction Study Design That Enables an Efficacy- and Safety-Based Dose Adjustment Decision. <i>Clinical Pharmacology and Therapeutics</i> , 2018 , 104, 781-784	6.1	14
91	Discovery of Competitive and Noncompetitive Ligands of the Organic Cation Transporter 1 (OCT1; SLC22A1). <i>Journal of Medicinal Chemistry</i> , 2017 , 60, 2685-2696	8.3	39
90	The Effect of Uremic Solutes on the Organic Cation Transporter 2. <i>Journal of Pharmaceutical Sciences</i> , 2017 , 106, 2551-2557	3.9	17

(2015-2017)

89	Transporters Involved in Metformin Pharmacokinetics and Treatment Response. <i>Journal of Pharmaceutical Sciences</i> , 2017 , 106, 2245-2250	3.9	72
88	Computational Discovery and Experimental Validation of Inhibitors of the Human Intestinal Transporter OATP2B1. <i>Journal of Chemical Information and Modeling</i> , 2017 , 57, 1402-1413	6.1	13
87	PharmGKB summary: very important pharmacogene information for ABCG2. <i>Pharmacogenetics and Genomics</i> , 2017 , 27, 420-427	1.9	14
86	Human Concentrative Nucleoside Transporter 3 (hCNT3, SLC28A3) Forms a Cyclic Homotrimer. <i>Biochemistry</i> , 2017 , 56, 3475-3483	3.2	9
85	Genome-wide association studies of drug response and toxicity: an opportunity for genome medicine. <i>Nature Reviews Drug Discovery</i> , 2017 , 16, 1	64.1	59
84	LAT1 activity of carboxylic acid bioisosteres: Evaluation of hydroxamic acids as substrates. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016 , 26, 5000-5006	2.9	29
83	A research roadmap for next-generation sequencing informatics. <i>Science Translational Medicine</i> , 2016 , 8, 335ps10	17.5	29
82	The Effect of Nizatidine, a MATE2K Selective Inhibitor, on the Pharmacokinetics and Pharmacodynamics of Metformin in Healthy Volunteers. <i>Clinical Pharmacokinetics</i> , 2016 , 55, 495-506	6.2	22
81	Rapid Method To Determine Intracellular Drug Concentrations in Cellular Uptake Assays: Application to Metformin in Organic Cation Transporter 1-Transfected Human Embryonic Kidney 293 Cells. <i>Drug Metabolism and Disposition</i> , 2016 , 44, 356-64	4	46
80	The Effect of Famotidine, a MATE1-Selective Inhibitor, on the Pharmacokinetics and Pharmacodynamics of Metformin. <i>Clinical Pharmacokinetics</i> , 2016 , 55, 711-21	6.2	37
79	Genomic Characterization of Metformin Hepatic Response. <i>PLoS Genetics</i> , 2016 , 12, e1006449	6	30
78	Pharmacometabolomic Assessment of Metformin in Non-diabetic, African Americans. <i>Frontiers in Pharmacology</i> , 2016 , 7, 135	5.6	20
77	LAT-1 activity of meta-substituted phenylalanine and tyrosine analogs. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016 , 26, 2616-2621	2.9	30
76	Identification and Quantitative Assessment of Uremic Solutes as Inhibitors of Renal Organic Anion Transporters, OAT1 and OAT3. <i>Molecular Pharmaceutics</i> , 2016 , 13, 3130-40	5.6	61
75	Variation in the glucose transporter gene SLC2A2 is associated with glycemic response to metformin. <i>Nature Genetics</i> , 2016 , 48, 1055-1059	36.3	108
74	Targeted disruption of organic cation transporter 3 attenuates the pharmacologic response to metformin. <i>Molecular Pharmacology</i> , 2015 , 88, 75-83	4.3	62
73	SLC transporters as therapeutic targets: emerging opportunities. <i>Nature Reviews Drug Discovery</i> , 2015 , 14, 543-60	64.1	363
72	A pharmacogenetic candidate gene study of tenofovir-associated Fanconi syndrome. <i>Pharmacogenetics and Genomics</i> , 2015 , 25, 82-92	1.9	23

71	Prediction and validation of enzyme and transporter off-targets for metformin. <i>Journal of Pharmacokinetics and Pharmacodynamics</i> , 2015 , 42, 463-75	2.7	32
70	Metformin Is a Substrate and Inhibitor of the Human Thiamine Transporter, THTR-2 (SLC19A3). <i>Molecular Pharmaceutics</i> , 2015 , 12, 4301-10	5.6	61
69	Unmet needs: Research helps regulators do their jobs. <i>Science Translational Medicine</i> , 2015 , 7, 315ps22	17.5	11
68	Towards quantitation of the effects of renal impairment and probenecid inhibition on kidney uptake and efflux transporters, using physiologically based pharmacokinetic modelling and simulations. <i>Clinical Pharmacokinetics</i> , 2014 , 53, 283-293	6.2	67
67	OCT1 is a high-capacity thiamine transporter that regulates hepatic steatosis and is a target of metformin. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014 , 111, 9983-8	11.5	155
66	Metformin pharmacogenomics: current status and future directions. <i>Diabetes</i> , 2014 , 63, 2590-9	0.9	90
65	Genome-wide discovery of drug-dependent human liver regulatory elements. <i>PLoS Genetics</i> , 2014 , 10, e1004648	6	30
64	A genome-wide association study of bronchodilator response in Latinos implicates rare variants. Journal of Allergy and Clinical Immunology, 2014 , 133, 370-8	11.5	84
63	Gene expression profiling of transporters in the solute carrier and ATP-binding cassette superfamilies in human eye substructures. <i>Molecular Pharmaceutics</i> , 2013 , 10, 650-63	5.6	41
62	Discovery of potent, selective multidrug and toxin extrusion transporter 1 (MATE1, SLC47A1) inhibitors through prescription drug profiling and computational modeling. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 781-795	8.3	100
61	Renal transporters in drug development. Annual Review of Pharmacology and Toxicology, 2013, 53, 503-	29 7.9	226
60	The Pharmacogenomics of Membrane Transporters Project 2013 , 73-108		
59	OCT (SLC22A) and OCTN Family 2013 , 171-208		5
58	Reduced renal clearance of cefotaxime in asians with a low-frequency polymorphism of OAT3 (SLC22A8). <i>Journal of Pharmaceutical Sciences</i> , 2013 , 102, 3451-7	3.9	41
57	Structure-based ligand discovery for the Large-neutral Amino Acid Transporter 1, LAT-1. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013 , 110, 5480-5	11.5	129
56	Molecular modeling and ligand docking for solute carrier (SLC) transporters. <i>Current Topics in Medicinal Chemistry</i> , 2013 , 13, 843-56	3	65
55	High selectivity of the Eminobutyric acid transporter 2 (GAT-2, SLC6A13) revealed by structure-based approach. <i>Journal of Biological Chemistry</i> , 2012 , 287, 37745-56	5.4	41
54	Pharmacogenomics and patient care: one size does not fit all. <i>Science Translational Medicine</i> , 2012 , 4, 153ps18	17.5	39

53	The role of ATM in response to metformin treatment and activation of AMPK. <i>Nature Genetics</i> , 2012 , 44, 359-60	36.3	44
52	Metformin pathways: pharmacokinetics and pharmacodynamics. <i>Pharmacogenetics and Genomics</i> , 2012 , 22, 820-7	1.9	256
51	Germline Genetic Polymorphisms Are Associated with Disease-Free Survival in Adults with Acute Myeloid Leukemia (AML): A Genomewide Association Study From the Pgrn-Riken Global Alliance <i>Blood</i> , 2012 , 120, 2548-2548	2.2	
50	Profiling of a prescription drug library for potential renal drug-drug interactions mediated by the organic cation transporter 2. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 4548-58	8.3	117
49	Role of organic cation transporter 1, OCT1 in the pharmacokinetics and toxicity of cis-diammine(pyridine)chloroplatinum(II) and oxaliplatin in mice. <i>Pharmaceutical Research</i> , 2011 , 28, 610) -2 5	36
48	Structure-based discovery of prescription drugs that interact with the norepinephrine transporter, NET. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011 , 108, 15810-5	5 ^{11.5}	101
47	Interactions of tyrosine kinase inhibitors with organic cation transporters and multidrug and toxic compound extrusion proteins. <i>Molecular Cancer Therapeutics</i> , 2011 , 10, 531-9	6.1	145
46	SLCO1B1 Variation and Methotrexate Disposition in Children with Acute Lymphoblastic Leukemia: The Importance of Rare Variants in Pharmacogenetics. <i>Blood</i> , 2011 , 118, 571-571	2.2	
45	Membrane transporters in drug development. <i>Nature Reviews Drug Discovery</i> , 2010 , 9, 215-36	64.1	2464
44	Organic cation transporters modulate the uptake and cytotoxicity of picoplatin, a third-generation platinum analogue. <i>Molecular Cancer Therapeutics</i> , 2010 , 9, 1058-69	6.1	66
43	Genetic variants of human organic anion transporter 4 demonstrate altered transport of endogenous substrates. <i>American Journal of Physiology - Renal Physiology</i> , 2010 , 299, F767-75	4.3	18
42	Role of organic cation transporter 3 (SLC22A3) and its missense variants in the pharmacologic action of metformin. <i>Pharmacogenetics and Genomics</i> , 2010 , 20, 687-99	1.9	145
41	Effect of genetic variation in the organic cation transporter 2 on the renal elimination of metformin. <i>Pharmacogenetics and Genomics</i> , 2009 , 19, 497-504	1.9	184
40	Genetic variants in multidrug and toxic compound extrusion-1, hMATE1, alter transport function. <i>Pharmacogenomics Journal</i> , 2009 , 9, 127-36	3.5	87
39	Identification and characterization of novel polymorphisms in the basal promoter of the human transporter, MATE1. <i>Pharmacogenetics and Genomics</i> , 2009 , 19, 770-80	1.9	52
38	Genetic variation in the proximal promoter of ABC and SLC superfamilies: liver and kidney specific expression and promoter activity predict variation. <i>PLoS ONE</i> , 2009 , 4, e6942	3.7	33
37	cis-Diammine(pyridine)chloroplatinum(II), a monofunctional platinum(II) antitumor agent: Uptake, structure, function, and prospects. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2008 , 105, 8902-7	11.5	198
36	Organic anion transporter 2 (SLC22A7) is a facilitative transporter of cGMP. <i>Molecular Pharmacology</i> , 2008 , 73, 1151-8	4.3	91

35	Genetic variation in human aquaporins and effects on phenotypes of water homeostasis. <i>Human Mutation</i> , 2008 , 29, 1108-17	4.7	18
34	Transport of paraquat by human organic cation transporters and multidrug and toxic compound extrusion family. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2007 , 322, 695-700	4.7	88
33	Functional effects of protein sequence polymorphisms in the organic cation/ergothioneine transporter OCTN1 (SLC22A4). <i>Pharmacogenetics and Genomics</i> , 2007 , 17, 773-82	1.9	39
32	Effect of genetic variation in the organic cation transporter 1 (OCT1) on metformin action. <i>Journal of Clinical Investigation</i> , 2007 , 117, 1422-31	15.9	673
31	The human organic anion transporter 3 (OAT3; SLC22A8): genetic variation and functional genomics. <i>American Journal of Physiology - Renal Physiology</i> , 2006 , 290, F905-12	4.3	78
30	Organic cation transporters are determinants of oxaliplatin cytotoxicity. Cancer Research, 2006, 66, 884	17 <u>+</u> 57 <u>1</u>	339
29	Functional genetic diversity in the high-affinity carnitine transporter OCTN2 (SLC22A5). <i>Molecular Pharmacology</i> , 2006 , 70, 1602-11	4.3	46
28	Functional analysis of polymorphisms in the organic anion transporter, SLC22A6 (OAT1). <i>Pharmacogenetics and Genomics</i> , 2005 , 15, 201-9	1.9	81
27	The concentrative nucleoside transporter family, SLC28. <i>Pflugers Archiv European Journal of Physiology</i> , 2004 , 447, 728-34	4.6	304
26	Natural variation in human membrane transporter genes reveals evolutionary and functional constraints. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2003 , 100, 5896-901	11.5	208
25	Sorting of rat SPNT in renal epithelium is independent of N-glycosylation. <i>Pharmaceutical Research</i> , 2003 , 20, 319-23	4.5	14
24	Evolutionary conservation predicts function of variants of the human organic cation transporter, OCT1. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2003 , 100, 5902-	7 ^{11.5}	241
23	Polymorphisms in a human kidney xenobiotic transporter, OCT2, exhibit altered function. <i>Pharmacogenetics and Genomics</i> , 2002 , 12, 395-405		169
22	Arginine 454 and lysine 370 are essential for the anion specificity of the organic anion transporter, rOAT3. <i>Biochemistry</i> , 2001 , 40, 5511-20	3.2	67
21	Molecular determinants of substrate selectivity in Na+-dependent nucleoside transporters. <i>Journal of Biological Chemistry</i> , 1997 , 272, 28845-8	5.4	46
20	Cloning and functional expression of a human liver organic cation transporter. <i>Molecular Pharmacology</i> , 1997 , 51, 913-21	4.3	343
19	Taurine transport in cultured choroid plexus. <i>Pharmaceutical Research</i> , 1997 , 14, 406-9	4.5	8
18	Mechanisms of 5-fluorouracil (5-FU) transport in isolated rabbit choroid plexus tissue slices. <i>Pharmaceutical Research</i> , 1996 , 13, 1276-8	4.5	5

LIST OF PUBLICATIONS

17	Expression of a renal Na(+)-nucleoside cotransport system (N2) in Xenopus laevis oocytes. <i>Pflugers Archiv European Journal of Physiology</i> , 1994 , 427, 381-3	4.6	12
16	Interaction of nucleoside analogues with the sodium-nucleoside transport system in brush border membrane vesicles from human kidney. <i>Pharmaceutical Research</i> , 1993 , 10, 423-6	4.5	55
15	Stereoselective interactions of organic cations with the organic cation transporter in OK cells. <i>Pharmaceutical Research</i> , 1993 , 10, 1169-73	4.5	16
14	Formycin B elimination from the cerebrospinal fluid of the rat. <i>Pharmaceutical Research</i> , 1993 , 10, 611-5	4.5	10
13	Effect of probenecid on the pharmacokinetics and pharmacodynamics of procainamide. <i>Journal of Clinical Pharmacology</i> , 1991 , 31, 429-32	2.9	4
12	The pharmacokinetics and pharmacodynamics of diltiazem and its metabolites in healthy adults after a single oral dose. <i>Clinical Pharmacology and Therapeutics</i> , 1989 , 46, 408-19	6.1	41
11	The pharmacokinetics of the enantiomers of atenolol. <i>Clinical Pharmacology and Therapeutics</i> , 1989 , 45, 403-10	6.1	38
10	The effect of probenecid on the renal elimination of cimetidine. <i>Clinical Pharmacology and Therapeutics</i> , 1989 , 45, 444-52	6.1	44
9	Renal transport of drugs: an overview of methodology with application to cimetidine. <i>Pharmaceutical Research</i> , 1988 , 5, 465-71	4.5	7
8	Cimetidine elimination from the cerebrospinal fluid of the rat. <i>Pharmaceutical Research</i> , 1988 , 5, 628-33	4.5	3
7	Stereoselective binding of disopyramide to plasma proteins. <i>Pharmaceutical Research</i> , 1988 , 5, 316-8	4.5	7
6	Verapamil interacts stereoselectively with the muscarinic receptor. <i>Pharmaceutical Research</i> , 1985 , 2, 94-5	4.5	
5	Correction for Volume Shift during Equilibrium Dialysis by Measurement of Protein Concentration. <i>Pharmaceutical Research</i> , 1984 , 1, 179-81	4.5	14
4	Effect of concentration-dependent binding to plasma proteins on the pharmacokinetics and pharmacodynamics of disopyramide. <i>Clinical Pharmacokinetics</i> , 1984 , 9 Suppl 1, 42-8	6.2	14
3	Propoxyphene and norpropoxyphene plasma concentrations after oral propoxyphene in cirrhotic patients with and without surgically constructed portacaval shunt. <i>Clinical Pharmacology and Therapeutics</i> , 1980 , 28, 417-24	6.1	24
2	Effect of hemodialysis on propoxyphene and norpropoxyphene concentrations in blood of anephric patients. <i>Clinical Pharmacology and Therapeutics</i> , 1980 , 27, 508-14	6.1	27
1	Propoxyphene and norpropoxyphene plasma concentrations in the anephric patient. <i>Clinical Pharmacology and Therapeutics</i> , 1980 , 27, 665-70	6.1	54