Ingolf Cascorbi

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#	Paper	IF	Citations
228	Functional polymorphisms of the human multidrug-resistance gene: multiple sequence variations and correlation of one allele with P-glycoprotein expression and activity in vivo. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2000 , 97, 3473-8	11.5	1017
227	Frequency of single nucleotide polymorphisms in the P-glycoprotein drug transporter MDR1 gene in white subjects. <i>Clinical Pharmacology and Therapeutics</i> , 2001 , 69, 169-74	6.1	541
226	Role of pharmacogenetics of ATP-binding cassette transporters in the pharmacokinetics of drugs 2006 , 112, 457-73		274
225	Deposition of Alzheimer® beta-amyloid is inversely correlated with P-glycoprotein expression in the brains of elderly non-demented humans. <i>Pharmacogenetics and Genomics</i> , 2002 , 12, 535-41		270
224	Association between the C3435T MDR1 gene polymorphism and susceptibility for ulcerative colitis. <i>Gastroenterology</i> , 2003 , 124, 26-33	13.3	266
223	Modulation of steady-state kinetics of digoxin by haplotypes of the P-glycoprotein MDR1 gene. <i>Clinical Pharmacology and Therapeutics</i> , 2002 , 72, 584-94	6.1	243
222	Mental and physical distress is modulated by a polymorphism in the 5-HT transporter gene interacting with social stressors and chronic disease burden. <i>Molecular Psychiatry</i> , 2005 , 10, 220-4	15.1	232
221	MDR1-P-Glycoprotein (ABCB1) Mediates Transport of Alzheimer@amyloid-beta peptidesimplications for the mechanisms of Abeta clearance at the blood-brain barrier. <i>Brain Pathology</i> , 2007 , 17, 347-53	6	189
220	Polymorphisms of drug-metabolizing enzymes CYP2C9, CYP2C19, CYP2D6, CYP1A1, NAT2 and of P-glycoprotein in a Russian population. <i>European Journal of Clinical Pharmacology</i> , 2003 , 59, 303-12	2.8	186
219	The effects of the human MDR1 genotype on the expression of duodenal P-glycoprotein and disposition of the probe drug talinolol. <i>Clinical Pharmacology and Therapeutics</i> , 2002 , 72, 572-83	6.1	161
218	Should we use N-acetyltransferase type 2 genotyping to personalize isoniazid doses?. <i>Antimicrobial Agents and Chemotherapy</i> , 2005 , 49, 1733-8	5.9	157
217	Glyburide and glimepiride pharmacokinetics in subjects with different CYP2C9 genotypes. <i>Clinical Pharmacology and Therapeutics</i> , 2002 , 72, 326-32	6.1	156
216	CYP1A1 and GSTM1 genetic polymorphisms and lung cancer risk in Caucasian non-smokers: a pooled analysis. <i>Carcinogenesis</i> , 2003 , 24, 875-82	4.6	152
215	Functional gene variants of CYP3A4. Clinical Pharmacology and Therapeutics, 2014, 96, 340-8	6.1	146
214	Association of Liver Injury From Specific Drugs, or Groups of Drugs, With Polymorphisms in HLA and Other Genes in a Genome-Wide Association Study. <i>Gastroenterology</i> , 2017 , 152, 1078-1089	13.3	137
213	Influence of polymorphisms of ABCB1 and ABCC2 on mRNA and protein expression in normal and cancerous kidney cortex. <i>Pharmacogenomics Journal</i> , 2007 , 7, 56-65	3.5	136
212	P-glycoprotein: tissue distribution, substrates, and functional consequences of genetic variations. <i>Handbook of Experimental Pharmacology</i> , 2011 , 261-83	3.2	135

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211	Carbamazepine regulates intestinal P-glycoprotein and multidrug resistance protein MRP2 and influences disposition of talinolol in humans. <i>Clinical Pharmacology and Therapeutics</i> , 2004 , 76, 192-200	6.1	134
210	CYP1A1 and GSTM1 genotypes affect benzo[a]pyrene DNA adducts in smokersQung: comparison with aromatic/hydrophobic adduct formation. <i>Carcinogenesis</i> , 2002 , 23, 1969-77	4.6	133
209	The role of P-glycoprotein in cerebral amyloid angiopathy; implications for the early pathogenesis of Alzheimer@ disease. <i>Current Alzheimer Research</i> , 2004 , 1, 121-5	3	131
208	Variable expression of MRP2 (ABCC2) in human placenta: influence of gestational age and cellular differentiation. <i>Drug Metabolism and Disposition</i> , 2005 , 33, 896-904	4	129
207	CYP1A1 T3801 C polymorphism and lung cancer: a pooled analysis of 2451 cases and 3358 controls. <i>International Journal of Cancer</i> , 2003 , 104, 650-7	7.5	121
206	Clozapine-induced agranulocytosis is associated with rare HLA-DQB1 and HLA-B alleles. <i>Nature Communications</i> , 2014 , 5, 4757	17.4	118
205	MDR1 genotypes do not influence the absorption of a single oral dose of 1 mg digoxin in healthy white males. <i>British Journal of Clinical Pharmacology</i> , 2002 , 54, 610-6	3.8	114
204	Modulation of benzo[a]pyrene diolepoxide-DNA adduct levels in human white blood cells by CYP1A1, GSTM1 and GSTT1 polymorphism. <i>Carcinogenesis</i> , 2000 , 21, 35-41	4.6	113
203	CYP3A5 genotype markedly influences the pharmacokinetics of tacrolimus and sirolimus in kidney transplant recipients. <i>Clinical Pharmacology and Therapeutics</i> , 2007 , 81, 228-34	6.1	105
202	CYP2D6 genotype and induction of intestinal drug transporters by rifampin predict presystemic clearance of carvedilol in healthy subjects. <i>Clinical Pharmacology and Therapeutics</i> , 2004 , 75, 213-22	6.1	102
201	Transient receptor potential channel polymorphisms are associated with the somatosensory function in neuropathic pain patients. <i>PLoS ONE</i> , 2011 , 6, e17387	3.7	102
200	Polymorphisms in CYP1A1, GSTM1, GSTT1 and lung cancer below the age of 45 years. <i>International Journal of Epidemiology</i> , 2003 , 32, 60-3	7.8	98
199	Candidate gene analysis identifies a polymorphism in HLA-DQB1 associated with clozapine-induced agranulocytosis. <i>Journal of Clinical Psychiatry</i> , 2011 , 72, 458-63	4.6	98
198	Differential metabolism of benzo[a]pyrene and benzo[a]pyrene-7,8-dihydrodiol by human CYP1A1 variants. <i>Carcinogenesis</i> , 2001 , 22, 453-9	4.6	93
197	Polymorphisms of the drug transporters ABCB1, ABCG2, ABCC2 and ABCC3 and their impact on drug bioavailability and clinical relevance. <i>Expert Opinion on Drug Metabolism and Toxicology</i> , 2014 , 10, 1337-54	5.5	89
196	Correlation between genotype and phenotype of the human arylamine N-acetyltransferase type 1 (NAT1). <i>Biochemical Pharmacology</i> , 1999 , 58, 1759-64	6	89
195	Expression and localization of P-glycoprotein in human heart: effects of cardiomyopathy. <i>Journal of Histochemistry and Cytochemistry</i> , 2002 , 50, 1351-6	3.4	87
194	Pooled analysis of the CYP1A1 exon 7 polymorphism and lung cancer (United States). <i>Cancer Causes and Control</i> , 2003 , 14, 339-46	2.8	83

193	Association of cyclophosphamide pharmacokinetics to polymorphic cytochrome P450 2C19. <i>Pharmacogenomics Journal</i> , 2005 , 5, 365-73	3.5	82
192	Drug interactionsprinciples, examples and clinical consequences. <i>Deutsches A&#x0308;rzteblatt International</i> , 2012 , 109, 546-55; quiz 556	2.5	81
191	Intestinal expression of P-glycoprotein (ABCB1), multidrug resistance associated protein 2 (ABCC2), and uridine diphosphate-glucuronosyltransferase 1A1 predicts the disposition and modulates the effects of the cholesterol absorption inhibitor ezetimibe in humans. <i>Clinical Pharmacology and</i>	6.1	81
190	Therapeutics, 200 6, 79, 206-17 Expression and localization of the multidrug resistance protein 5 (MRP5/ABCC5), a cellular export pump for cyclic nucleotides, in human heart. <i>American Journal of Pathology</i> , 2003 , 163, 1567-77	5.8	81
189	Non-response to antiepileptic pharmacotherapy is associated with the ABCC2 -24C>T polymorphism in young and adult patients with epilepsy. <i>Pharmacogenetics and Genomics</i> , 2009 , 19, 353	3- 62	79
188	The ATP-binding cassette transporter ABCG2 (BCRP), a marker for side population stem cells, is expressed in human heart. <i>Journal of Histochemistry and Cytochemistry</i> , 2006 , 54, 215-21	3.4	77
187	A1/A2 polymorphism of glycoprotein IIIa and association with excess procedural risk for coronary catheter interventions: a case-controlled study. <i>Lancet, The,</i> 1999 , 353, 708-12	40	77
186	Arylamine N-acetyltransferase activity in man. <i>Drug Metabolism Reviews</i> , 1999 , 31, 489-502	7	75
185	Polymorphisms in the human CYP1A1 gene as susceptibility factors for lung cancer: exon-7 mutation (4889 A to G), and a T to C mutation in the 3G lanking region. <i>The Clinical Investigator</i> , 1994 , 72, 240-8		74
184	Influence of genetic polymorphisms on intestinal expression and rifampicin-type induction of ABCC2 and on bioavailability of talinolol. <i>Pharmacogenetics and Genomics</i> , 2008 , 18, 357-65	1.9	73
183	Modulation of multidrug resistance P-glycoprotein 1 (ABCB1) expression in human heart by hereditary polymorphisms. <i>Pharmacogenetics and Genomics</i> , 2004 , 14, 381-5		71
182	MicroRNAs and their relevance to ABC transporters. <i>British Journal of Clinical Pharmacology</i> , 2014 , 77, 587-96	3.8	70
181	Down-regulation of ATP-binding cassette C2 protein expression in HepG2 cells after rifampicin treatment is mediated by microRNA-379. <i>Molecular Pharmacology</i> , 2011 , 80, 314-20	4.3	66
180	Genetic basis of toxic reactions to drugs and chemicals. <i>Toxicology Letters</i> , 2006 , 162, 16-28	4.4	66
179	A European Spectrum of Pharmacogenomic Biomarkers: Implications for Clinical Pharmacogenomics. <i>PLoS ONE</i> , 2016 , 11, e0162866	3.7	66
178	Impact of ABCC2 haplotypes on transcriptional and posttranscriptional gene regulation and function. <i>Pharmacogenomics Journal</i> , 2011 , 11, 25-34	3.5	62
177	MicroRNA profiling in K-562 cells under imatinib treatment: influence of miR-212 and miR-328 on ABCG2 expression. <i>Pharmacogenetics and Genomics</i> , 2012 , 22, 198-205	1.9	62
176	A LC-MS/MS method to quantify the novel cholesterol lowering drug ezetimibe in human serum, urine and feces in healthy subjects genotyped for SLCO1B1. <i>Journal of Chromatography B:</i> Analytical Technologies in the Biomedical and Life Sciences, 2006 , 830, 143-50	3.2	60

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175	A Missense Variant in PTPN22 is a Risk Factor for Drug-induced Liver Injury. <i>Gastroenterology</i> , 2019 , 156, 1707-1716.e2	13.3	59	
174	Molecular genetics of cancer susceptibility. <i>Pharmacology</i> , 2000 , 61, 212-27	2.3	57	
173	Pitfalls in N-acetyltransferase 2 genotyping. <i>Pharmacogenetics and Genomics</i> , 1999 , 9, 123-7		57	
172	miRNAs as mediators of drug resistance. <i>Epigenomics</i> , 2012 , 4, 369-81	4.4	56	
171	High DNA damage by benzo[a]pyrene 7,8-diol-9,10-epoxide in bronchial epithelial cells from patients with lung cancer: comparison with lung parenchyma. <i>Cancer Letters</i> , 2004 , 207, 157-63	9.9	56	
170	Genetic determinants of clozapine-induced agranulocytosis: recent results of HLA subtyping in a non-jewish caucasian sample. <i>Archives of General Psychiatry</i> , 2001 , 58, 93-4		55	
169	Decreased levels of dopamine D3 receptor mRNA in schizophrenic and bipolar patients. <i>Neuropsychobiology</i> , 2004 , 50, 305-10	4	54	
168	High CA repeat numbers in intron 13 of the endothelial nitric oxide synthase gene and increased risk of coronary artery disease. <i>Pharmacogenetics and Genomics</i> , 2000 , 10, 133-40		53	
167	Hematopoietic stem cell involvement in positive ALL as a potential mechanism of resistance to blinatumomab therapy. <i>Blood</i> , 2017 , 130, 2027-2031	2.2	52	
166	Pharmacogenetics of ATP-binding cassette transporters and clinical implications. <i>Methods in Molecular Biology</i> , 2010 , 596, 95-121	1.4	52	
165	Clozapine-induced agranulocytosis in schizophrenic Caucasians: confirming clues for associations with human leukocyte class I and II antigens. <i>Pharmacogenomics Journal</i> , 2007 , 7, 325-32	3.5	52	
164	HLA-DRB1*16: 01-DQB1*05: 02 is a novel genetic risk factor for flupirtine-induced liver injury. <i>Pharmacogenetics and Genomics</i> , 2016 , 26, 218-24	1.9	50	
163	Allelic variants of human cytochrome P450 1A1 (CYP1A1): effect of T461N and I462V substitutions on steroid hydroxylase specificity. <i>Pharmacogenetics and Genomics</i> , 2000 , 10, 519-30		49	
162	Further evidence of human leukocyte antigen-encoded susceptibility to clozapine-induced agranulocytosis independent of ancestry. <i>Pharmacogenetics and Genomics</i> , 2001 , 11, 135-41		46	
161	Cellular uptake of imatinib into leukemic cells is independent of human organic cation transporter 1 (OCT1). <i>Clinical Cancer Research</i> , 2014 , 20, 985-94	12.9	45	
160	Impact of myeloperoxidase and NADPH-oxidase polymorphisms in drug-induced agranulocytosis. Journal of Clinical Psychopharmacology, 2004 , 24, 613-7	1.7	45	
159	Antimicrobial peptides and proteins of the horseinsights into a well-armed organism. <i>Veterinary Research</i> , 2011 , 42, 98	3.8	44	
158	Metabolic gene polymorphisms and lung cancer risk in non-smokers. An update of the GSEC study. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 2005 , 592, 45-57	3.3	43	

157	Polymorphisms in xenobiotic conjugation and disease predisposition. <i>Toxicology Letters</i> , 1998 , 102-103, 173-83	4.4	43
156	Mutations in the human paraoxonase 1 gene: frequencies, allelic linkages, and association with coronary artery disease. <i>Pharmacogenetics and Genomics</i> , 1999 , 9, 755-61		43
155	Myeloperoxidase G-463A polymorphism and lung cancer: a HuGE genetic susceptibility to environmental carcinogens pooled analysis. <i>Genetics in Medicine</i> , 2007 , 9, 67-73	8.1	42
154	Effect of levothyroxine administration on intestinal P-glycoprotein expression: consequences for drug disposition. <i>Clinical Pharmacology and Therapeutics</i> , 2002 , 72, 256-64	6.1	42
153	Identification of six methylenetetrahydrofolate reductase (MTHFR) genotypes resulting from common polymorphisms: impact on plasma homocysteine levels and development of coronary artery disease. <i>Atherosclerosis</i> , 2001 , 154, 651-8	3.1	41
152	Impact of ABCC2 genotype on antiepileptic drug response in Caucasian patients with childhood epilepsy. <i>Pharmacogenetics and Genomics</i> , 2011 , 21, 624-30	1.9	40
151	Clinical trial: a novel high-dose 1 g mesalamine suppository (Salofalk) once daily is as efficacious as a 500-mg suppository thrice daily in active ulcerative proctitis. <i>Inflammatory Bowel Diseases</i> , 2010 , 16, 1947-56	4.5	40
150	Myeloperoxidase463A variant reduces benzo[a]pyrene diol epoxide DNA adducts in skin of coal tar treated patients. <i>Carcinogenesis</i> , 2001 , 22, 1015-8	4.6	40
149	Association of arylamine N-acetyltransferases NAT1 and NAT2 genotypes to laryngeal cancer risk. <i>Pharmacogenetics and Genomics</i> , 1999 , 9, 103???112		40
148	Decreased sigmoidal ABCB1 (P-glycoprotein) expression in ulcerative colitis is associated with disease activity. <i>Pharmacogenomics</i> , 2009 , 10, 1941-53	2.6	39
147	Elimination half-life of anti-Mllerian hormone. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2012 , 97, 2160-3	5.6	37
146	Population frequency, mutation linkage and analytical methodology for the Arg16Gly, Gln27Glu and Thr164Ile polymorphisms in the beta2-adrenergic receptor among Turks. <i>British Journal of Clinical Pharmacology</i> , 1999 , 48, 761-4	3.8	37
145	Occupational history and genetic N-acetyltransferase polymorphism in urothelial cancer patients of Leverkusen, Germany. <i>Scandinavian Journal of Work, Environment and Health</i> , 1996 , 22, 332-8	4.3	37
144	Which Genetic Determinants Should be Considered for Tacrolimus Dose Optimization in Kidney Transplantation? A Combined Analysis of Genes Affecting the CYP3A Locus. <i>Therapeutic Drug Monitoring</i> , 2015 , 37, 288-95	3.2	36
143	Epigenetic modulation of the drug resistance genes MGMT, ABCB1 and ABCG2 in glioblastoma multiforme. <i>BMC Cancer</i> , 2013 , 13, 617	4.8	35
142	Gender and smoking-related risk reduction of periodontal disease with variant myeloperoxidase alleles. <i>Genes and Immunity</i> , 2002 , 3, 102-6	4.4	35
141	Determination and allelic allocation of seven nucleotide transitions within the arylamine N-acetyltransferase gene in the Polish population. <i>Clinical Pharmacology and Therapeutics</i> , 1996 , 59, 37	6-8 1	35
140	The Nomenclature, Definition and Distinction of Types of Shock. <i>Deutsches A&#x0308;rzteblatt International</i> , 2018 , 115, 757-768	2.5	35

139	Association of ATP-binding cassette transporter variants with the risk of Alzheimer@disease. <i>Pharmacogenomics</i> , 2013 , 14, 485-94	2.6	34
138	High frequency of CYP1A1 mutations in a Turkish population. <i>Archives of Toxicology</i> , 1998 , 72, 215-8	5.8	34
137	Simvastatin does not influence the intestinal P-glycoprotein and MPR2, and the disposition of talinolol after chronic medication in healthy subjects genotyped for the ABCB1, ABCC2 and SLCO1B1 polymorphisms. <i>British Journal of Clinical Pharmacology</i> , 2006 , 61, 440-50	3.8	34
136	Functional significance of a hereditary adenine insertion variant in the 5GUTR of the endothelin-1 gene. <i>Pharmacogenetics and Genomics</i> , 2003 , 13, 445-51		34
135	SOX11 identified by target gene evaluation of miRNAs differentially expressed in focal and non-focal brain tissue of therapy-resistant epilepsy patients. <i>Neurobiology of Disease</i> , 2015 , 77, 127-40	7.5	33
134	Association of ABCB1 genetic variants 3435C>T and 2677G>T to ABCB1 mRNA and protein expression in brain tissue from refractory epilepsy patients. <i>Epilepsia</i> , 2008 , 49, 1555-61	6.4	33
133	Clozapine-induced agranulocytosis and hereditary polymorphisms of clozapine metabolizing enzymes: no association with myeloperoxidase and cytochrome P4502D6. <i>Pharmacopsychiatry</i> , 2000 , 33, 218-20	2	33
132	Pharmacogenomics education in medical and pharmacy schools: conclusions of a global survey. <i>Pharmacogenomics</i> , 2019 , 20, 643-657	2.6	31
131	Identification and characterization of a defective CYP3A4 genotype in a kidney transplant patient with severely diminished tacrolimus clearance. <i>Clinical Pharmacology and Therapeutics</i> , 2014 , 95, 416-22	6.1	30
130	Endothelial nitric oxide synthase Glu298Asp gene polymorphism, blood pressure and hypertension in a general population sample. <i>Journal of Hypertension</i> , 2005 , 23, 1361-6	1.9	30
129	Endothelial NO synthase polymorphisms and postural tachycardia syndrome. <i>Hypertension</i> , 2005 , 46, 1103-10	8.5	30
128	Key Learning Outcomes for Clinical Pharmacology and Therapeutics Education in Europe: A Modified Delphi Study. <i>Clinical Pharmacology and Therapeutics</i> , 2018 , 104, 317-325	6.1	30
127	Paraoxonase 1 mutations in a Turkish population. <i>Toxicology and Applied Pharmacology</i> , 1999 , 157, 174-	· 7 4.6	29
126	ABC transporters in drug-refractory epilepsy: limited clinical significance of pharmacogenetics?. <i>Clinical Pharmacology and Therapeutics</i> , 2010 , 87, 15-8	6.1	28
125	Influence of CYP3A4, CYP3A5, and ABCB1 genotype and expression on budesonide pharmacokinetics: a possible role of intestinal CYP3A4 expression. <i>Clinical Pharmacology and Therapeutics</i> , 2008 , 84, 43-6	6.1	28
124	Genetic determinants of drug-induced agranulocytosis: potential risk of olanzapine?. <i>Pharmacopsychiatry</i> , 1999 , 32, 110-2	2	28
123	NAT2*12A (803A>G) codes for rapid arylamine n-acetylation in humans. <i>Pharmacogenetics and Genomics</i> , 1996 , 6, 257-9		28
122	Genetic variants may play an important role in mRNA-miRNA interaction: evidence for haplotype-dependent downregulation of ABCC2 (MRP2) by miRNA-379. <i>Pharmacogenetics and Genomics</i> 2014 24 283-91	1.9	27

121	Arylamine N-acetyltransferase (NAT2) genotypes in a Turkish population. <i>Pharmacogenetics and Genomics</i> , 1997 , 7, 327-31		27
120	Pharmacogenetics-based new therapeutic concepts. <i>Drug Metabolism Reviews</i> , 2004 , 36, 617-38	7	27
119	Co-expression of human cytochrome P4501A1 (CYP1A1) variants and human NADPH-cytochrome P450 reductase in the baculovirus/insect cell system. <i>Xenobiotica</i> , 2001 , 31, 345-56	2	27
118	How to manage individualized drug therapy: application of pharmacogenetic knowledge of drug metabolism and transport. <i>Clinical Chemistry and Laboratory Medicine</i> , 2000 , 38, 869-76	5.9	27
117	Role of Kozak sequence polymorphism of platelet glycoprotein Ibalpha as a risk factor for coronary artery disease and catheter interventions. <i>Journal of the American College of Cardiology</i> , 2001 , 38, 1023-	.7 5.1	27
116	Polymorphic enzymes of xenobiotic metabolism as modulators of acquired P53 mutations in bladder cancer. <i>Pharmacogenetics and Genomics</i> , 1996 , 6, 535-45		27
115	Differential expression and functionality of TRPA1 protein genetic variants in conditions of thermal stimulation. <i>Journal of Biological Chemistry</i> , 2012 , 287, 27087-94	5.4	26
114	Lack of association between arylamine N-acetyltransferase 2 (NAT2) polymorphism and systemic lupus erythematosus. <i>Pharmacogenetics and Genomics</i> , 2002 , 12, 559-63		26
113	The Neurophysiology and Treatment of Motion Sickness. <i>Deutsches A&#x0308;rzteblatt International</i> , 2018 , 115, 687-696	2.5	26
112	Interaction of herbal products with prescribed medications: A systematic review and meta-analysis. <i>Pharmacological Research</i> , 2019 , 141, 397-408	10.2	25
111	OpenVigilfree eyeballs on AERS pharmacovigilance data. <i>Nature Biotechnology</i> , 2012 , 30, 137-8	44.5	25
110	CYP1A1 mutations 4887A, 4889G, 5639C and 6235C in the Polish population and their allelic linkage, determined by peptide nucleic acid-mediated PCR clamping. <i>Pharmacogenetics and Genomics</i> , 1997 , 7, 303-7		25
109	Clinically Relevant Multidrug Transporters Are Regulated by microRNAs along the Human Intestine. <i>Molecular Pharmaceutics</i> , 2017 , 14, 2245-2253	5.6	24
108	Relationship of drug metabolizing enzyme genotype to plasma levels as well as myelotoxicity of cyclophosphamide in breast cancer patients. <i>European Journal of Clinical Pharmacology</i> , 2012 , 68, 389-9	5 ^{2.8}	23
107	N-Acetyltransferase 2 (NAT2) and Glutathione S-Transferase $\bar{\mu}$ (GSTM1) in Bladder-cancer Patients in a Highly Industrialized Area. <i>International Journal of Occupational and Environmental Health</i> , 1997 , 3, 105-110		23
106	Dysregulation of Mucosal Membrane Transporters and Drug-Metabolizing Enzymes in Ulcerative Colitis. <i>Journal of Pharmaceutical Sciences</i> , 2019 , 108, 1035-1046	3.9	23
105	Dihydropyrimidine Dehydrogenase Testing prior to Treatment with 5-Fluorouracil, Capecitabine, and Tegafur: A Consensus Paper. <i>Oncology Research and Treatment</i> , 2020 , 43, 628-636	2.8	22
104	Endothelial nitric oxide synthase Glu(298)>Asp polymorphism, carotid atherosclerosis and intima-media thickness in a general population sample. <i>Clinical Science</i> , 2005 , 109, 475-81	6.5	21

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103	Meta-analysis on outcome-worsening comorbidities of COVID-19 and related potential drug-drug interactions. <i>Pharmacological Research</i> , 2020 , 161, 105250	10.2	21
102	Pharmacogenetics and Predictive Testing of Drug Hypersensitivity Reactions. <i>Frontiers in Pharmacology</i> , 2016 , 7, 396	5.6	21
101	Association of metabolic gene polymorphisms with tobacco consumption in healthy controls. <i>International Journal of Cancer</i> , 2004 , 110, 266-70	7.5	20
100	Interaction of xenobiotics on the glucose-transport system and the Na+/K(+)-ATPase of human skin fibroblasts. <i>Ecotoxicology and Environmental Safety</i> , 1991 , 21, 38-46	7	20
99	©omparison of extremes approach provides evidence against the modifying role of NAT2 polymorphism in lung cancer susceptibility. <i>Cancer Letters</i> , 2005 , 221, 177-83	9.9	19
98	Elevated serum leptin in patients with coronary artery disease: no association with the Trp64Arg polymorphism of the beta3-adrenergic receptor. <i>International Journal of Obesity</i> , 2000 , 24, 369-75	5.5	19
97	Correlation between the lipophilicity of substituted phenols and their inhibition of the Na+/K+-ATPase of Chinese hamster ovary cells. <i>Toxicology</i> , 1989 , 58, 197-210	4.4	18
96	Drug Hypersensitivity. <i>Deutsches A&#x0308;rzteblatt International</i> , 2018 , 115, 501-512	2.5	18
95	Effects of CYP2B6 genetic polymorphisms in patients receiving cyclophosphamide combination chemotherapy for breast cancer. <i>Cancer Chemotherapy and Pharmacology</i> , 2015 , 75, 207-14	3.5	17
94	Length variants of the ABCB1 3GUTR and loss of miRNA binding sites: possible consequences in regulation and pharmacotherapy resistance. <i>Pharmacogenomics</i> , 2016 , 17, 327-40	2.6	17
93	Pharmacogenomics of heart failure focus on drug disposition and action. <i>Cardiovascular Research</i> , 2004 , 64, 32-9	9.9	17
92	Reduced procedural risk for coronary catheter interventions in carriers of the coagulation factor VII-Gln353 gene. <i>Journal of the American College of Cardiology</i> , 2000 , 36, 1520-5	15.1	17
91	Cyclophosphamide treatment-induced leukopenia rates in ANCA-associated vasculitis are influenced by variant CYP450 2C9 genotypes. <i>Pharmacogenomics</i> , 2016 , 17, 367-74	2.6	16
90	Genotype frequencies of selected drug metabolizing enzymes and ABC drug transporters among breast cancer patients on FAC chemotherapy. <i>Basic and Clinical Pharmacology and Toxicology</i> , 2010 , 107, 570-6	3.1	16
89	Phenotyping of N-acetyltransferase type 2 by caffeine from uncontrolled dietary exposure. <i>European Journal of Clinical Pharmacology</i> , 2004 , 60, 17-21	2.8	16
88	Association between the N-acetylation genetic polymorphism and bronchial asthma. <i>British Journal of Clinical Pharmacology</i> , 2002 , 54, 671-4	3.8	16
87	Effects of a heterogeneous set of xenobiotics on growth and plasma membranes of mammalian and fungal cell cultures. <i>Ecotoxicology and Environmental Safety</i> , 1993 , 26, 113-26	7	16
86	Pharmacogenomic or -epigenomic biomarkers in drug treatment: Two sides of the same medal?. <i>Clinical Pharmacology and Therapeutics</i> , 2016 , 99, 478-80	6.1	16

85	Implementation and obstacles of pharmacogenetics in clinical practice: An international survey. <i>British Journal of Clinical Pharmacology</i> , 2019 , 85, 2076-2088	3.8	14
84	Transcriptional and Post-Transcriptional Regulation of Duodenal P-Glycoprotein and MRP2 in Healthy Human Subjects after Chronic Treatment with Rifampin and Carbamazepine. <i>Molecular Pharmaceutics</i> , 2019 , 16, 3823-3830	5.6	14
83	Expression of N-acetyltransferases in periodontal granulation tissue. <i>Journal of Dental Research</i> , 2002 , 81, 349-53	8.1	14
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