Ann K Daly

List of Publications by Year in descending order

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		4370	5806
280	28,911	86	161
papers	citations	h-index	g-index
324	324	324	20108
all docs	docs citations	times ranked	citing authors

ANN K DALV

#	Article	IF	CITATIONS
1	A genetic risk score and diabetes predict development of alcohol-related cirrhosis in drinkers. Journal of Hepatology, 2022, 76, 275-282.	1.8	33
2	Increased serum miR-193a-5p during non-alcoholic fatty liver disease progression: Diagnostic and mechanistic relevance. JHEP Reports, 2022, 4, 100409.	2.6	20
3	Macrophage scavenger receptor 1 mediates lipid-induced inflammation in non-alcoholic fatty liver disease. Journal of Hepatology, 2022, 76, 1001-1012.	1.8	54
4	Metabolic signatures across the full spectrum of non-alcoholic fatty liver disease. JHEP Reports, 2022, 4, 100477.	2.6	31
5	A multiancestry genome-wide association study of unexplained chronic ALT elevation as a proxy for nonalcoholic fatty liver disease with histological and radiological validation. Nature Genetics, 2022, 54, 761-771.	9.4	68
6	Human Leukocyte Antigen B*14:01 and B*35:01 Are Associated With Trimethoprim‣ulfamethoxazole Induced Liver Injury. Hepatology, 2021, 73, 268-281.	3.6	43
7	PharmVar GeneFocus: <i>CYP2C19</i> . Clinical Pharmacology and Therapeutics, 2021, 109, 352-366.	2.3	72
8	Genetic Risk Factors in Drugâ€Induced Liver Injury Due to Isoniazidâ€Containing Antituberculosis Drug Regimens. Clinical Pharmacology and Therapeutics, 2021, 109, 1125-1135.	2.3	31
9	rs641738C>T near MBOAT7 is associated with liver fat, ALT and fibrosis in NAFLD: A meta-analysis. Journal of Hepatology, 2021, 74, 20-30.	1.8	77
10	Genomeâ€wide Association Study and Metaâ€analysis on Alcoholâ€Associated Liver Cirrhosis Identifies Genetic Risk Factors. Hepatology, 2021, 73, 1920-1931.	3.6	54
11	NASH limits anti-tumour surveillance in immunotherapy-treated HCC. Nature, 2021, 592, 450-456.	13.7	649
12	A PDCD1 Role in the Genetic Predisposition to NAFLD-HCC?. Cancers, 2021, 13, 1412.	1.7	26
13	Pharmacogenomics spotlight commentary: From the United Kingdom to global populations. British Journal of Clinical Pharmacology, 2021, 87, 4546-4548.	1.1	4
14	PharmVar GeneFocus: <i>CYP2C9</i> . Clinical Pharmacology and Therapeutics, 2021, 110, 662-676.	2.3	34
15	Transcriptomics Identify Thrombospondinâ€2 as a Biomarker for NASH and Advanced Liver Fibrosis. Hepatology, 2021, 74, 2452-2466.	3.6	71
16	Diagnostic accuracy of elastography and magnetic resonance imaging in patients with NAFLD: A systematic review and meta-analysis. Journal of Hepatology, 2021, 75, 770-785.	1.8	149
17	Obesity, Diabetes, Coffee, Tea, and Cannabis Use Alter Risk for Alcohol-Related Cirrhosis in 2 Large Cohorts of High-Risk Drinkers. American Journal of Gastroenterology, 2021, 116, 106-115.	0.2	25
18	Peptide-based urinary monitoring of fibrotic nonalcoholic steatohepatitis by mass-barcoded activity-based sensors. Science Translational Medicine, 2021, 13, eabe8939.	5.8	17

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19	Genome-wide association study identifies susceptibility loci for acute myeloid leukemia. Nature Communications, 2021, 12, 6233.	5.8	17
20	Investigation of Oxidative Stress-Related Candidate Genes as Risk Factors for Drug-Induced Liver Injury due to Co-Amoxiclav. DNA and Cell Biology, 2020, 39, 349-354.	0.9	10
21	Pharmacogenomics of Drug-Induced Liver Injury. Advances in Molecular Pathology, 2020, 3, 107-115.	0.2	3
22	Transcriptomic profiling across the nonalcoholic fatty liver disease spectrum reveals gene signatures for steatohepatitis and fibrosis. Science Translational Medicine, 2020, 12, .	5.8	205
23	HLA DRB1*15:01-DQB1*06:02-Restricted Human CD4+ T Cells Are Selectively Activated With Amoxicillin-Peptide Adducts. Toxicological Sciences, 2020, 178, 115-126.	1.4	14
24	Polygenic architecture informs potential vulnerability to drug-induced liver injury. Nature Medicine, 2020, 26, 1541-1548.	15.2	55
25	HLA associations with infliximab-induced liver injury. Pharmacogenomics Journal, 2020, 20, 681-686.	0.9	17
26	Enhanced liver fibrosis test for the non-invasive diagnosis of fibrosis in patients with NAFLD: A systematic review and meta-analysis. Journal of Hepatology, 2020, 73, 252-262.	1.8	170
27	Genome-wide association study of non-alcoholic fatty liver and steatohepatitis in a histologically characterised cohortâ^†. Journal of Hepatology, 2020, 73, 505-515.	1.8	279
28	Genetic Polymorphisms Implicated in Nonalcoholic Liver Disease or Selected Other Disorders Have No Influence on Drugâ€Induced Liver Injury. Hepatology Communications, 2019, 3, 1032-1035.	2.0	7
29	Drugâ€Induced Liver Injury due to Flucloxacillin: Relevance of Multiple Human Leukocyte Antigen Alleles. Clinical Pharmacology and Therapeutics, 2019, 106, 245-253.	2.3	58
30	Endocytosis and Lack of Cytotoxicity of Alkyl-Capped Silicon Quantum Dots Prepared from Porous Silicon. Materials, 2019, 12, 1702.	1.3	7
31	Shared Genetic Risk Factors Across Carbamazepineâ€Induced Hypersensitivity Reactions. Clinical Pharmacology and Therapeutics, 2019, 106, 1028-1036.	2.3	52
32	Next-Generation Sequencing of PTGS Genes Reveals an Increased Frequency of Non-synonymous Variants Among Patients With NSAID-Induced Liver Injury. Frontiers in Genetics, 2019, 10, 134.	1.1	10
33	A Missense Variant in PTPN22 is a Risk Factor for Drug-induced Liver Injury. Gastroenterology, 2019, 156, 1707-1716.e2.	0.6	97
34	Paracetamol metabolism, hepatotoxicity, biomarkers and therapeutic interventions: a perspective. Toxicology Research, 2018, 7, 347-357.	0.9	70
35	Human Leukocyte Antigen (HLA) and Other Genetic Risk Factors in Drug-Induced Liver Injury (DILI). Methods in Pharmacology and Toxicology, 2018, , 497-509.	0.1	1
36	Evaluation of laboratory tests for cirrhosis and for alcohol use, in the context of alcoholic cirrhosis. Alcohol, 2018, 66, 1-7.	0.8	13

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37	Genetic and Clinical Factors Are Associated With Statinâ€Related Myotoxicity of Moderate Severity: A Case–Control Study. Clinical Pharmacology and Therapeutics, 2018, 104, 178-187.	2.3	14
38	HLA-A*33:03-Restricted Activation of Ticlopidine-Specific T-Cells from Human Donors. Chemical Research in Toxicology, 2018, 31, 1022-1024.	1.7	9
39	Pharmacogenomics of CYP2C9: Functional and Clinical Considerations. Journal of Personalized Medicine, 2018, 8, 1.	1.1	136
40	Pharmacogenetics of Adverse Drug Reactions. Advances in Pharmacology, 2018, 83, 155-190.	1.2	32
41	Genetic risk factors for DILI-recent findings from large international networks. Proceedings for Annual Meeting of the Japanese Pharmacological Society, 2018, WCP2018, SY76-2.	0.0	0
42	Dosing algorithms for vitamin K antagonists across VKORC1 and CYP2C9 genotypes. Journal of Thrombosis and Haemostasis, 2017, 15, 465-472.	1.9	8
43	Drug-Induced Cholestasis: Mechanisms and Importance. , 2017, , 117-128.		0
44	Minocycline hepatotoxicity: Clinical characterization and identification of HLA-Bâ^—35:02 as a risk factor. Journal of Hepatology, 2017, 67, 137-144.	1.8	100
45	Are Polymorphisms in Genes Relevant to Drug Disposition Predictors of Susceptibility to Drug-Induced Liver Injury?. Pharmaceutical Research, 2017, 34, 1564-1569.	1.7	33
46	Association of Liver Injury From Specific Drugs, or Groups ofÂDrugs, With Polymorphisms in HLA and Other Genes in aÂGenome-Wide Association Study. Gastroenterology, 2017, 152, 1078-1089.	0.6	174
47	Ageâ€stratified outcome of a genotypeâ€guided dosing algorithm for acenocoumarol and phenprocoumon. Journal of Thrombosis and Haemostasis, 2017, 15, 454-464.	1.9	4
48	Pharmacogenetics: a general review on progress to date. British Medical Bulletin, 2017, 124, 1-15.	2.7	48
49	Telomerase reverse transcriptase germline mutations and hepatocellular carcinoma in patients with nonalcoholic fatty liver disease. Cancer Medicine, 2017, 6, 1930-1940.	1.3	43
50	Defining drug response for stratified medicine. Drug Discovery Today, 2017, 22, 173-179.	3.2	24
51	HLA-DRB1*16. Pharmacogenetics and Genomics, 2016, 26, 218-224.	0.7	63
52	Pharmacogenetic allele nomenclature: International workgroup recommendations for test result reporting. Clinical Pharmacology and Therapeutics, 2016, 99, 172-185.	2.3	146
53	The Effect of the <i>CYP1A1*2A</i> Allele on Colorectal Cancer Susceptibility in a British Population. Genetic Testing and Molecular Biomarkers, 2016, 20, 475-477.	0.3	3
54	Variants in the LGALS9 Gene Are Associated With Development of Liver Disease in Heavy Consumers of Alcohol. Clinical Gastroenterology and Hepatology, 2016, 14, 762-768.e1.	2.4	9

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55	Weight Loss Decreases Excess Pancreatic Triacylglycerol Specifically in Type 2 Diabetes. Diabetes Care, 2016, 39, 158-165.	4.3	135
56	A multi-factorial analysis of response to warfarin in a UK prospective cohort. Genome Medicine, 2016, 8, 2.	3.6	41
57	Transcriptional regulation of PNPLA3 and its impact on susceptibility to nonalcoholic fatty liver Disease (NAFLD) in humans. Aging, 2016, 9, 26-40.	1.4	11
58	Genetics of Alcoholic Liver Disease. Seminars in Liver Disease, 2015, 35, 361-374.	1.8	39
59	Brief Report: Genetics of Alcoholic Cirrhosis— <scp>G</scp> enom <scp>ALC</scp> Multinational Study. Alcoholism: Clinical and Experimental Research, 2015, 39, 836-842.	1.4	29
60	Characterization of amoxicillin―and clavulanic acidâ€specific T cells in patients with amoxicillinâ€clavulanate–induced liver injury. Hepatology, 2015, 62, 887-899.	3.6	83
61	Quality of life in patients with venous thromboembolism and atrial fibrillation treated with coumarin anticoagulants. Thrombosis Research, 2015, 136, 69-75.	0.8	28
62	PNPLA3 Gene Polymorphism Is Associated With Predisposition to and Severity of Alcoholic Liver Disease. American Journal of Gastroenterology, 2015, 110, 846-856.	0.2	120
63	TM6SF2 as a genetic risk factor for fibrosis. Hepatology, 2015, 62, 1321-1321.	3.6	3
64	Oral anticoagulation: a critique of recent advances and controversies. Trends in Pharmacological Sciences, 2015, 36, 153-163.	4.0	70
65	Pharmacogenetics of drug metabolizing enzymes in the United Kingdom population: review of current knowledge and comparison with selected European populations. Drug Metabolism and Personalized Therapy, 2015, 30, 165-174.	0.3	18
66	Promiscuous T-cell responses to drugs and drug-haptens. Journal of Allergy and Clinical Immunology, 2015, 136, 474-476.e8.	1.5	41
67	Comparison of dosing algorithms for acenocoumarol and phenprocoumon using clinical factors with the standard care in the Netherlands. Thrombosis Research, 2015, 136, 94-100.	0.8	2
68	TM6SF2: Catch-22 in the Fight Against Nonalcoholic Fatty Liver Disease and Cardiovascular Disease?. Gastroenterology, 2015, 148, 679-684.	0.6	75
69	Polymorphic Variants of Cytochrome P450. Advances in Pharmacology, 2015, 74, 85-111.	1.2	34
70	Pharmacogenomics of Warfarin. , 2014, , 497-507.		0
71	Direct-to-consumer pharmacogenomic testing assessed in a US-based study. Journal of the Royal College of Physicians of Edinburgh, The, 2014, 44, 212-213.	0.2	0
72	Gene polymorphisms of cellular senescence marker p21 and disease progression in non-alcohol-related fatty liver disease. Cell Cycle, 2014, 13, 1489-1494.	1.3	54

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73	Genetic Basis of Drug-Induced Liver Injury: Present and Future. Seminars in Liver Disease, 2014, 34, 123-133.	1.8	101
74	Pharmacogeneticâ€guided dosing of coumarin anticoagulants: algorithms for warfarin, acenocoumarol and phenprocoumon. British Journal of Clinical Pharmacology, 2014, 77, 626-641.	1.1	113
75	Patients Benefit From Genetics-Guided Coumarin Anticoagulant Therapy. Clinical Pharmacology and Therapeutics, 2014, 96, 15-17.	2.3	15
76	Is There a Need to Teach Pharmacogenetics?. Clinical Pharmacology and Therapeutics, 2014, 95, 245-247.	2.3	17
77	Identification of susceptible HLA class II coâ€amoxiclav genotypes based on the analysis of drugâ€specific Tâ€cells from patients with liver injury. Clinical and Translational Allergy, 2014, 4, O3.	1.4	2
78	Characterization of amoxicillin and clavulanicâ€acidâ€responsive CD4+ And CD8+ Tâ€cells in patients with coâ€amoxiclavâ€induced liver injury. Clinical and Translational Allergy, 2014, 4, P42.	1.4	0
79	N-acetyltransferase 2 (NAT2) genotype as a risk factor for development of drug-induced liver injury relating to antituberculosis drug treatment in a mixed-ethnicity patient group. European Journal of Clinical Pharmacology, 2014, 70, 1079-1086.	0.8	56
80	Opportunities and limitations: the value of pharmacogenetics in clinical practice. British Journal of Clinical Pharmacology, 2014, 77, 583-586.	1.1	7
81	TM6SF2 rs58542926 influences hepatic fibrosis progression in patients with non-alcoholic fatty liver disease. Nature Communications, 2014, 5, 4309.	5.8	478
82	Carriage of the PNPLA3 rs738409 C >G polymorphism confers an increased risk of non-alcoholic fatty liver disease associated hepatocellular carcinoma. Journal of Hepatology, 2014, 61, 75-81.	1.8	431
83	Point of care testing for improving risk- benefit ratio of aspirin and warfarin. Molecular Cytogenetics, 2014, 7, 154.	0.4	0
84	Human Leukocyte Antigen (HLA) Pharmacogenomic Tests: Potential and Pitfalls. Current Drug Metabolism, 2014, 15, 196-201.	0.7	20
85	Optimal dosing of warfarin and other coumarin anticoagulants: the role of genetic polymorphisms. Archives of Toxicology, 2013, 87, 407-420.	1.9	50
86	Relevance of CYP2E1 to Non-alcoholic Fatty Liver Disease. Sub-Cellular Biochemistry, 2013, 67, 165-175.	1.0	17
87	A Randomized Trial of Genotype-Guided Dosing of Warfarin. New England Journal of Medicine, 2013, 369, 2294-2303.	13.9	735
88	Pharmacogenomics of adverse drug reactions. Genome Medicine, 2013, 5, 5.	3.6	87
89	Efavirenz induced acute liver failure requiring liver transplantation in a slow drug metaboliser. Journal of Clinical Virology, 2013, 58, 331-333.	1.6	17

90 Genetic Factors in the Pathogenesis of Drug-Induced Liver Injury. , 2013, , 215-225.

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91	Lung cancer risk in relation to nicotinic acetylcholine receptor, CYP2A6 and CYP1A1 genotypes in the Bangladeshi population. Clinica Chimica Acta, 2013, 416, 11-19.	0.5	58
92	Adaptive Dosing Approaches to the Individualization of 13- <i>Cis</i> -Retinoic Acid (Isotretinoin) Treatment for Children with High-Risk Neuroblastoma. Clinical Cancer Research, 2013, 19, 469-479.	3.2	45
93	Human leukocyte antigen (HLA)-B*57:01-restricted activation of drug-specific T cells provides the immunological basis for flucloxacillin-induced liver injury. Hepatology, 2013, 57, 727-739.	3.6	212
94	Institutional Profile: Pharmacogenomics research at Newcastle University. Pharmacogenomics, 2012, 13, 1333-1338.	0.6	0
95	CYP2D6 update. Pharmacogenetics and Genomics, 2012, 22, 692-694.	0.7	19
96	Limited contribution of common genetic variants to risk for liver injury due to a variety of drugs. Pharmacogenetics and Genomics, 2012, 22, 784-795.	0.7	108
97	Genetic Polymorphisms Affecting Drug Metabolism. Advances in Pharmacology, 2012, 63, 137-167.	1.2	35
98	Cost–effectiveness of pharmacogenetics in anticoagulation: international differences in healthcare systems and costs. Pharmacogenomics, 2012, 13, 1405-1417.	0.6	17
99	The SOD2 C47T polymorphism influences NAFLD fibrosis severity: Evidence from case-control and intra-familial allele association studies. Journal of Hepatology, 2012, 56, 448-454.	1.8	156
100	Genetic association studies in drug-induced liver injury. Drug Metabolism Reviews, 2012, 44, 116-126.	1.5	100
101	VKORC1 and CYP2C9 genotype and patient characteristics explain a large proportion of the variability in warfarin dose requirement among children. Blood, 2012, 119, 868-873.	0.6	99
102	Using Genome-Wide Association Studies to Identify Genes Important in Serious Adverse Drug Reactions. Annual Review of Pharmacology and Toxicology, 2012, 52, 21-35.	4.2	100
103	The population pharmacokinetics of <i>R</i> ―and <i>S</i> â€warfarin: effect of genetic and clinical factors. British Journal of Clinical Pharmacology, 2012, 73, 66-76.	1.1	70
104	Longâ€ŧerm anticoagulant effects of the CYP2C9 and VKORC1 genotypes in acenocoumarol users. Journal of Thrombosis and Haemostasis, 2012, 10, 606-614.	1.9	22
105	An evaluation of gene–gene interaction between the CYP2C9 and VKORC1 genotypes affecting the anticoagulant effect of phenprocoumon and acenocoumarol. Journal of Thrombosis and Haemostasis, 2012, 10, 767-772.	1.9	15
106	Polymorphism in the Farnesyl Diphosphate Farnesyl Transferase 1 Gene and Nonalcoholic Fatty Liver Disease Severity. Gastroenterology, 2011, 140, 1694-1695.	0.6	14
107	Susceptibility to Amoxicillin-Clavulanate-Induced Liver Injury Is Influenced by Multiple HLA Class I and II Alleles. Gastroenterology, 2011, 141, 338-347.	0.6	412
108	A Genome-Wide Association Study Identifies Potential Susceptibility Loci for Hepatotoxicity Due to Various Drugs. Gastroenterology, 2011, 140, S-886.	0.6	3

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109	Genotypes and phenotypes of CYP3A in Bangladeshi population. Clinica Chimica Acta, 2011, 412, 531-536.	0.5	9
110	Genotyping for CYP2C9 and VKORC1 alleles by a novel point of care assay with HyBeacon® probes. Clinica Chimica Acta, 2011, 412, 2063-2069.	0.5	33
111	Genetic modifiers of non-alcoholic fatty liver disease progression. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2011, 1812, 1557-1566.	1.8	59
112	The APOC3 T-455C and C-482T promoter region polymorphisms are not associated with the severity of liver damage independently of PNPLA3 I148M genotype in patients with nonalcoholic fatty liver. Journal of Hepatology, 2011, 55, 1409-1414.	1.8	74
113	Genetic determinants of susceptibility and severity in nonalcoholic fatty liver disease. Expert Review of Gastroenterology and Hepatology, 2011, 5, 253-263.	1.4	53
114	Family History of Cancer and Tobacco Exposure in Index Cases of Pancreatic Ductal Adenocarcinoma. Journal of Oncology, 2011, 2011, 1-7.	0.6	11
115	Cyclooxygenase-2 Polymorphisms and Pancreatic Cancer Susceptibility. Pancreas, 2011, 40, 1289-1294.	0.5	11
116	Characterization of the metabolism of fenretinide by human liver microsomes, cytochrome P450 enzymes and UDPâ€glucuronosyltransferases. British Journal of Pharmacology, 2011, 162, 989-999.	2.7	14
117	The Phenotype Standardization Project: Improving Pharmacogenetic Studies of Serious Adverse Drug Reactions. Clinical Pharmacology and Therapeutics, 2011, 89, 784-785.	2.3	61
118	Case Definition and Phenotype Standardization in Drug-Induced Liver Injury. Clinical Pharmacology and Therapeutics, 2011, 89, 806-815.	2.3	773
119	Loading and maintenance dose algorithms for phenprocoumon and acenocoumarol using patient characteristics and pharmacogenetic data. European Heart Journal, 2011, 32, 1909-1917.	1.0	86
120	Genetics of Alcoholic and Nonalcoholic Fatty Liver Disease. Seminars in Liver Disease, 2011, 31, 128-146.	1.8	101
121	A common polymorphism in the <i>ABCB11</i> gene is associated with advanced fibrosis in hepatitis C but not in non-alcoholic fatty liver disease. Clinical Science, 2011, 120, 287-296.	1.8	44
122	Clinical and Pharmacogenetic Influences on Response to Hydroxychloroquine in Discoid Lupus Erythematosus: A Retrospective Cohort Study. Journal of Investigative Dermatology, 2011, 131, 1981-1986.	0.3	84
123	Inter-individual variation in DNA damage and base excision repair in young, healthy non-smokers: effects of dietary supplementation and genotype. British Journal of Nutrition, 2010, 103, 1585-1593.	1.2	40
124	A systematic review of cost–effectiveness analyses of pharmacogenetic-guided dosing in treatment with coumarin derivatives. Pharmacogenomics, 2010, 11, 989-1002.	0.6	26
125	A role for the pregnane X receptor in flucloxacillin-induced liver injury. Hepatology, 2010, 51, 1656-1664.	3.6	55
126	Patatin-like phospholipase domain containing 3: A case in point linking genetic susceptibility for alcoholic and nonalcoholic liver disease. Hepatology, 2010, 51, 1463-1465.	3.6	26

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127	Homozygosity for the patatin-like phospholipase-3/adiponutrin I148M polymorphism influences liver fibrosis in patients with nonalcoholic fatty liver disease. Hepatology, 2010, 51, 1209-1217.	3.6	563
128	Reversal of warfarin-induced over-anticoagulation with individualised dosing of oral vitamin k: a pilot study. Journal of Thrombosis and Haemostasis, 2010, 8, 1123-5.	1.9	6
129	Preempting and preventing drug-induced liver injury. Nature Genetics, 2010, 42, 650-651.	9.4	25
130	Genome-wide association studies in pharmacogenomics. Nature Reviews Genetics, 2010, 11, 241-246.	7.7	238
131	Role of UDP-Glucuronosyltransferase Isoforms in 13-cis Retinoic Acid Metabolism in Humans. Drug Metabolism and Disposition, 2010, 38, 1211-1217.	1.7	15
132	Relevance of Nonsynonymous CYP2C8 Polymorphisms to 13-cis Retinoic Acid and Paclitaxel Hydroxylation. Drug Metabolism and Disposition, 2010, 38, 1261-1266.	1.7	21
133	Polymorphisms in the Tropomyosin TPM1 Short Isoform Promoter Alter Gene Expression and Are Associated With Increased Risk of Metabolic Syndrome. American Journal of Hypertension, 2010, 23, 399-404.	1.0	5
134	Genetic variants regulating insulin receptor signalling are associated with the severity of liver damage in patients with non-alcoholic fatty liver disease. Gut, 2010, 59, 267-273.	6.1	148
135	Association between anti-tumour necrosis factor treatment response and genetic variants within the TLR and NFÂB signalling pathways. Annals of the Rheumatic Diseases, 2010, 69, 1315-1320.	0.5	74
136	Human leucocyte antigen class II genotype in susceptibility and resistance to co-amoxiclav-induced liver injury. Journal of Hepatology, 2010, 53, 1049-1053.	1.8	137
137	Pharmacogenetics and human genetic polymorphisms. Biochemical Journal, 2010, 429, 435-449.	1.7	91
138	Drug-induced liver injury: past, present and future. Pharmacogenomics, 2010, 11, 607-611.	0.6	69
139	Genotype-guided dosing of coumarin derivatives: the European pharmacogenetics of anticoagulant therapy (EU-PACT) trial design. Pharmacogenomics, 2009, 10, 1687-1695.	0.6	131
140	Genetic Association Studies in Drug-Induced Liver Injury. Seminars in Liver Disease, 2009, 29, 400-411.	1.8	98
141	Inter-individual variation in nucleotide excision repair in young adults: effects of age, adiposity, micronutrient supplementation and genotype. British Journal of Nutrition, 2009, 101, 1316.	1.2	40
142	HLA-B*5701 genotype is a major determinant of drug-induced liver injury due to flucloxacillin. Nature Genetics, 2009, 41, 816-819.	9.4	950
143	Pharmacogenomics of anticoagulants: steps toward personal dosage. Genome Medicine, 2009, 1, 10.	3.6	33
144	Estimation of the Warfarin Dose with Clinical and Pharmacogenetic Data. New England Journal of Medicine, 2009, 360, 753-764.	13.9	1,375

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145	Nomenclature for alleles of the cytochrome P450 oxidoreductase gene. Pharmacogenetics and Genomics, 2009, 19, 565-566.	0.7	30
146	Genetic and environmental factors determining clinical outcomes and cost of warfarin therapy: a prospective study. Pharmacogenetics and Genomics, 2009, 19, 800-812.	0.7	57
147	Genetic variants of hepatic transporters and susceptibility to drug induced liver injury. Toxicology, 2008, 253, 10.	2.0	8
148	Relevance of PXR to Flucloxacillin-induced Liver Injury. Toxicology, 2008, 253, 2.	2.0	0
149	Flucloxacillin-induced liver injury. Toxicology, 2008, 254, 158-163.	2.0	29
150	Pharmacogenomics Applications in Drug Metabolism. Methods in Pharmacology and Toxicology, 2008, , 109-120.	0.1	0
151	The Relationship between PON1 Phenotype and PON1-192 Genotype in Detoxification of Three Oxons by Human Liver. Drug Metabolism and Disposition, 2007, 35, 315-320.	1.7	20
152	Genetic Susceptibility to Diclofenac-Induced Hepatotoxicity: Contribution of UGT2B7, CYP2C8, and ABCC2 Genotypes. Gastroenterology, 2007, 132, 272-281.	0.6	318
153	Genetic susceptibility in pancreatic ductal adenocarcinoma. British Journal of Surgery, 2007, 95, 22-32.	0.1	19
154	Factors Affecting Drug Concentrations and QT Interval During Thioridazine Therapy. Clinical Pharmacology and Therapeutics, 2007, 82, 555-565.	2.3	32
155	Evidence that a polymorphism within the 3′UTR of glutathione peroxidase 4 is functional and is associated with susceptibility to colorectal cancer. Genes and Nutrition, 2007, 2, 225-232.	1.2	83
156	Individualized drug therapy. Current Opinion in Drug Discovery & Development, 2007, 10, 29-36.	1.9	9
157	Genotyping for Cytochrome P450 Polymorphisms. , 2006, 320, 193-208.		23
158	Contribution of CYP2C9 to variability in vitamin K antagonist metabolism. Expert Opinion on Drug Metabolism and Toxicology, 2006, 2, 3-15.	1.5	13
159	Significance of the Minor Cytochrome P450 3A Isoforms. Clinical Pharmacokinetics, 2006, 45, 13-31.	1.6	205
160	APOE genotype makes a small contribution to warfarin dose requirements. Pharmacogenetics and Genomics, 2006, 16, 609-611.	0.7	46
161	A rapid genotyping method for the vitamin K epoxide reductase complex subunit 1 (VKORC1) gene. Journal of Thrombosis and Haemostasis, 2006, 4, 1158-1159.	1.9	9
162	The impact of simvastatin on warfarin disposition and dose requirements. Journal of Thrombosis and Haemostasis, 2006, 4, 1422-1424.	1.9	21

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163	Polymorphic organic anion transporting polypeptide 1B1 is a major determinant of repaglinide pharmacokinetics. Clinical Pharmacology and Therapeutics, 2005, 77, 468-478.	2.3	320
164	Genetic polymorphism of manganese superoxide dismutase (MnSOD) and breast cancer susceptibility. Cell Biochemistry and Function, 2005, 23, 73-76.	1.4	35
165	Influence of IL-6, COL1A1, and VDR gene polymorphisms on bone mineral density in Crohn's disease. Gut, 2005, 54, 1579-1584.	6.1	35
166	The impact of CYP2C9 and VKORC1 genetic polymorphism and patient characteristics upon warfarin dose requirements: proposal for a new dosing regimen. Blood, 2005, 106, 2329-2333.	0.6	894
167	Pharmacogenetics of oral anticoagulants. Personalized Medicine, 2005, 2, 23-27.	0.8	3
168	Warfarin and celecoxib interaction in the setting of cytochrome P450 (CYP2C9) polymorphism with bleeding complication. Postgraduate Medical Journal, 2004, 80, 107-109.	0.9	33
169	N-Acetyltransferase (NAT2) Polymorphism and Breast Cancer Susceptibility: A Lack of Association in a Case-Control Study of Turkish Population. International Journal of Toxicology, 2004, 23, 25-31.	0.6	18
170	CYP3A5 phenotype-genotype correlations in a British population. British Journal of Clinical Pharmacology, 2004, 57, 664-664.	1.1	0
171	Development of analytical technology in pharmacogenetic research. Naunyn-Schmiedeberg's Archives of Pharmacology, 2004, 369, 133-140.	1.4	17
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