

Mark McCormack

List of Publications by Year in descending order

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Version: 2024-02-01

29
papers

2,881
citations

516215

16
h-index

476904

29
g-index

30
all docs

30
docs citations

30
times ranked

5831
citing authors

#	ARTICLE	IF	CITATIONS
1	A pharmacogenomic assessment of psychiatric adverse drug reactions to levetiracetam. <i>Epilepsia</i> , 2022, 63, 1563-1570.	2.6	11
2	Symptomatology of carbamazepine and oxcarbazepine induced hyponatremia in people with epilepsy. <i>Epilepsia</i> , 2021, 62, 778-784.	2.6	9
3	Assessing the role of rare genetic variants in drug-resistant, non-lesional focal epilepsy. <i>Annals of Clinical and Translational Neurology</i> , 2021, 8, 1376-1387.	1.7	16
4	Role of Common Genetic Variants for Drug-Resistance to Specific Anti-Seizure Medications. <i>Frontiers in Pharmacology</i> , 2021, 12, 688386.	1.6	6
5	Pharmacoresponse in genetic generalized epilepsy: a genome-wide association study. <i>Pharmacogenomics</i> , 2020, 21, 325-335.	0.6	21
6	Analysis of shared common genetic risk between amyotrophic lateral sclerosis and epilepsy. <i>Neurobiology of Aging</i> , 2020, 92, 153.e1-153.e5.	1.5	4
7	Testing association of rare genetic variants with resistance to three common antiseizure medications. <i>Epilepsia</i> , 2020, 61, 657-666.	2.6	22
8	A comparison of genomic diagnostics in adults and children with epilepsy and comorbid intellectual disability. <i>European Journal of Human Genetics</i> , 2020, 28, 1066-1077.	1.4	30
9	De-novo mutations in patients with chronic ultra-refractory epilepsy with onset after age five years. <i>European Journal of Medical Genetics</i> , 2020, 63, 103625.	0.7	9
10	Ultra-Rare Genetic Variation in the Epilepsies: A Whole-Exome Sequencing Study of 17,606 Individuals. <i>American Journal of Human Genetics</i> , 2019, 105, 267-282.	2.6	237
11	Development of a genomics module within an epilepsy-specific electronic health record: Toward genomic medicine in epilepsy care. <i>Epilepsia</i> , 2019, 60, 1670-1677.	2.6	7
12	Comparative effectiveness of antiepileptic drugs in juvenile myoclonic epilepsy. <i>Epilepsia Open</i> , 2019, 4, 420-430.	1.3	34
13	Genomic and clinical predictors of lacosamide response in refractory epilepsies. <i>Epilepsia Open</i> , 2019, 4, 563-571.	1.3	12
14	Influence of common SCN1A promoter variants on the severity of SCN1A related phenotypes. <i>Molecular Genetics & Genomic Medicine</i> , 2019, 7, e00727.	0.6	13
15	A genome-wide association study of sodium levels and drug metabolism in an epilepsy cohort treated with carbamazepine and oxcarbazepine. <i>Epilepsia Open</i> , 2019, 4, 102-109.	1.3	9
16	Genetic variation in <i>CFH</i> predicts phenytoin-induced maculopapular exanthema in European-descent patients. <i>Neurology</i> , 2018, 90, e332-e341.	1.5	43
17	Distinct HLA associations of LGI1 and CASPR2-antibody diseases. <i>Brain</i> , 2018, 141, 2263-2271.	3.7	100
18	Rare coding variants in genes encoding GABAA receptors in genetic generalised epilepsies: an exome-based case-control study. <i>Lancet Neurology</i> , The, 2018, 17, 699-708.	4.9	67

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19	The impact of ERBB-family germline single nucleotide polymorphisms on survival response to adjuvant trastuzumab treatment in HER2-positive breast cancer. <i>Oncotarget</i> , 2016, 7, 75518-75525.	0.8	12
20	Association of CYP3A variants with kidney transplant outcomes. <i>Renal Failure</i> , 2015, 37, 562-566.	0.8	9
21	Tibetans living at sea level have a hyporesponsive hypoxia-inducible factor system and blunted physiological responses to hypoxia. <i>Journal of Applied Physiology</i> , 2014, 116, 893-904.	1.2	97
22	TDP2 protects transcription from abortive topoisomerase activity and is required for normal neural function. <i>Nature Genetics</i> , 2014, 46, 516-521.	9.4	122
23	Genetic determinants of common epilepsies: a meta-analysis of genome-wide association studies. <i>Lancet Neurology</i> , The, 2014, 13, 893-903.	4.9	264
24	A genome-wide association study of recipient genotype and medium-term kidney allograft function. <i>Clinical Transplantation</i> , 2013, 27, 379-387.	0.8	39
25	Genome-wide mapping for clinically relevant predictors of lamotrigine- and phenytoin-induced hypersensitivity reactions. <i>Pharmacogenomics</i> , 2012, 13, 399-405.	0.6	38
26	Exome Sequencing Followed by Large-Scale Genotyping Fails to Identify Single Rare Variants of Large Effect in Idiopathic Generalized Epilepsy. <i>American Journal of Human Genetics</i> , 2012, 91, 293-302.	2.6	95
27	HLA-A*3101 and Carbamazepine-Induced Hypersensitivity Reactions in Europeans. <i>New England Journal of Medicine</i> , 2011, 364, 1134-1143.	13.9	815
28	Pharmacogenomics and epilepsy: the road ahead. <i>Pharmacogenomics</i> , 2011, 12, 1429-1447.	0.6	31
29	Natural selection on <i>EPAS1</i> (<i>HIF2α</i>) associated with low hemoglobin concentration in Tibetan highlanders. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010, 107, 11459-11464.	3.3	708