Stefan Wolking

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/3094677/publications.pdf

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24	954	686830 13	525886 27
papers	citations	h-index	g-index

29	29	29	2344
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Ultra-Rare Genetic Variation in the Epilepsies: A Whole-Exome Sequencing Study of 17,606 Individuals. American Journal of Human Genetics, 2019, 105, 267-282.	2.6	237
2	Impact of Genetic Polymorphisms of ABCB1 (MDR1, P-Glycoprotein) on Drug Disposition and Potential Clinical Implications: Update of the Literature. Clinical Pharmacokinetics, 2015, 54, 709-735.	1.6	207
3	Frequent genes in rare diseases: panelâ€based next generation sequencing to disclose causal mutations in hereditary neuropathies. Journal of Neurochemistry, 2017, 143, 507-522.	2.1	68
4	Rare coding variants in genes encoding GABAA receptors in genetic generalised epilepsies: an exome-based case-control study. Lancet Neurology, The, 2018, 17, 699-708.	4.9	67
5	Genetic variation in CFH predicts phenytoin-induced maculopapular exanthema in European-descent patients. Neurology, 2018, 90, e332-e341.	1.5	43
6	Clinical spectrum of STX1B -related epileptic disorders. Neurology, 2019, 92, e1238-e1249.	1.5	43
7	Early-onset familial hemiplegic migraine due to a novel SCN1A mutation. Cephalalgia, 2016, 36, 1238-1247.	1.8	36
8	Comparative effectiveness of antiepileptic drugs in juvenile myoclonic epilepsy. Epilepsia Open, 2019, 4, 420-430.	1.3	34
9	Comparative effectiveness of antiepileptic drugs in patients with mesial temporal lobe epilepsy with hippocampal sclerosis. Epilepsia, 2017, 58, 1734-1741.	2.6	26
10	Focal epilepsy in Glucose transporter type 1 (Glut1) defects: case reports and a review of literature. Journal of Neurology, 2014, 261, 1881-1886.	1.8	24
11	Testing association of rare genetic variants with resistance to three common antiseizure medications. Epilepsia, 2020, 61, 657-666.	2.6	22
12	Pharmacoresponse in genetic generalized epilepsy: a genome-wide association study. Pharmacogenomics, 2020, 21, 325-335.	0.6	21
13	Polygenic risk scores of several subtypes of epilepsies in a founder population. Neurology: Genetics, 2020, 6, e416.	0.9	17
14	Assessing the role of rare genetic variants in drugâ€resistant, nonâ€lesional focal epilepsy. Annals of Clinical and Translational Neurology, 2021, 8, 1376-1387.	1.7	16
15	Episodic itch in a case of spinal glioma. BMC Neurology, 2013, 13, 124.	0.8	15
16	Genomic and clinical predictors of lacosamide response in refractory epilepsies. Epilepsia Open, 2019, 4, 563-571.	1.3	12
17	The role of common genetic variation in presumed monogenic epilepsies. EBioMedicine, 2022, 81, 104098.	2.7	12
18	Genetic testing before epilepsy surgery â€“ An exploratory survey and case collection from German epilepsy centers. Seizure: the Journal of the British Epilepsy Association, 2022, 95, 4-10.	0.9	11

#	Article	IF	Citations
19	Reduced REM sleep: a potential biomarker for epilepsy â€“ a retrospective case-control study. Seizure: the Journal of the British Epilepsy Association, 2022, 98, 27-33.	0.9	11
20	Rare gene deletions in genetic generalized and Rolandic epilepsies. PLoS ONE, 2018, 13, e0202022.	1.1	6
21	Role of Common Genetic Variants for Drug-Resistance to Specific Anti-Seizure Medications. Frontiers in Pharmacology, 2021, 12, 688386.	1.6	6
22	A case of DRESS (drug reaction with eosinophilia and systemic symptoms) under treatment with eslicarbazepine. Seizure: the Journal of the British Epilepsy Association, 2019, 72, 11-12.	0.9	3
23	Macrodactylia lipomatosa with fibrolipomatous hamartomas: Macroscopic and ultrasound clues. Clinical Neurophysiology, 2017, 128, 1315-1316.	0.7	2
24	Assessment of burden and segregation profiles of <scp>CNVs</scp> in patients with epilepsy. Annals of Clinical and Translational Neurology, 2022, 9, 1050-1058.	1.7	2