

# Stefan Wolking

## List of Publications by Year in descending order

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Version: 2024-02-01

24  
papers

954  
citations

686830

13  
h-index

525886

27  
g-index

29  
all docs

29  
docs citations

29  
times ranked

2344  
citing authors

#	ARTICLE	IF	CITATIONS
1	Genetic testing before epilepsy surgery – An exploratory survey and case collection from German epilepsy centers. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2022, 95, 4-10.	0.9	11
2	Reduced REM sleep: a potential biomarker for epilepsy – a retrospective case-control study. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2022, 98, 27-33.	0.9	11
3	The role of common genetic variation in presumed monogenic epilepsies. <i>EBioMedicine</i> , 2022, 81, 104098.	2.7	12
4	Assessment of burden and segregation profiles of CNVs in patients with epilepsy. <i>Annals of Clinical and Translational Neurology</i> , 2022, 9, 1050-1058.	1.7	2
5	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
6	Role of Common Genetic Variants for Drug-Resistance to Specific Anti-Seizure Medications. <i>Frontiers in Pharmacology</i> , 2021, 12, 688386.	1.6	6
7	Pharmacoresponse in genetic generalized epilepsy: a genome-wide association study. <i>Pharmacogenomics</i> , 2020, 21, 325-335.	0.6	21
8	Testing association of rare genetic variants with resistance to three common antiseizure medications. <i>Epilepsia</i> , 2020, 61, 657-666.	2.6	22
9	Polygenic risk scores of several subtypes of epilepsies in a founder population. <i>Neurology: Genetics</i> , 2020, 6, e416.	0.9	17
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	Comparative effectiveness of antiepileptic drugs in juvenile myoclonic epilepsy. <i>Epilepsia Open</i> , 2019, 4, 420-430.	1.3	34
12	Genomic and clinical predictors of lacosamide response in refractory epilepsies. <i>Epilepsia Open</i> , 2019, 4, 563-571.	1.3	12
13	A case of DRESS (drug reaction with eosinophilia and systemic symptoms) under treatment with eslicarbazepine. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2019, 72, 11-12.	0.9	3
14	Clinical spectrum of STX1B-related epileptic disorders. <i>Neurology</i> , 2019, 92, e1238-e1249.	1.5	43
15	Genetic variation in CFH predicts phenytoin-induced maculopapular exanthema in European-descent patients. <i>Neurology</i> , 2018, 90, e332-e341.	1.5	43
16	Rare gene deletions in genetic generalized and Rolandic epilepsies. <i>PLoS ONE</i> , 2018, 13, e0202022.	1.1	6
17	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
18	Comparative effectiveness of antiepileptic drugs in patients with mesial temporal lobe epilepsy with hippocampal sclerosis. <i>Epilepsia</i> , 2017, 58, 1734-1741.	2.6	26

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19	Frequent genes in rare diseases: panel-based next generation sequencing to disclose causal mutations in hereditary neuropathies. <i>Journal of Neurochemistry</i> , 2017, 143, 507-522.	2.1	68
20	Macrodactylia lipomatosa with fibrolipomatous hamartomas: Macroscopic and ultrasound clues. <i>Clinical Neurophysiology</i> , 2017, 128, 1315-1316.	0.7	2
21	Early-onset familial hemiplegic migraine due to a novel <i>SCN1A</i> mutation. <i>Cephalalgia</i> , 2016, 36, 1238-1247.	1.8	36
22	Impact of Genetic Polymorphisms of ABCB1 (MDR1, P-Glycoprotein) on Drug Disposition and Potential Clinical Implications: Update of the Literature. <i>Clinical Pharmacokinetics</i> , 2015, 54, 709-735.	1.6	207
23	Focal epilepsy in Glucose transporter type 1 (Glut1) defects: case reports and a review of literature. <i>Journal of Neurology</i> , 2014, 261, 1881-1886.	1.8	24
24	Episodic itch in a case of spinal glioma. <i>BMC Neurology</i> , 2013, 13, 124.	0.8	15