## Simona Balestrini

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/305762/publications.pdf

Version: 2024-02-01

69 papers 2,390 citations

236925 25 h-index 233421 45 g-index

76 all docs

76 docs citations

76 times ranked 3990 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.	6.2	237
2	The landscape of epilepsy-related GATOR1 variants. Genetics in Medicine, 2019, 21, 398-408.	2.4	137
3	Clinical spectrum and genotype–phenotype associations of KCNA2-related encephalopathies. Brain, 2017, 140, 2337-2354.	7.6	117
4	Pharmacogenomics in epilepsy. Neuroscience Letters, 2018, 667, 27-39.	2.1	109
5	Severe carotid stenosis and impaired cerebral hemodynamics can influence cognitive deterioration. Neurology, 2013, 80, 2145-2150.	1.1	103
6	Intronic ATTTC repeat expansions in STARD7 in familial adult myoclonic epilepsy linked to chromosome 2. Nature Communications, 2019, 10, 4920.	12.8	99
7	<i>TBC1D24</i> genotype–phenotype correlation. Neurology, 2016, 87, 77-85.	1.1	97
8	Neurologic phenotypes associated with <i>COL4A1</i> /i>/ <i>2</i> /i> mutations. Neurology, 2018, 91, e2078-e2088.	1.1	97
9	Structural imaging biomarkers of sudden unexpected death in epilepsy. Brain, 2015, 138, 2907-2919.	7.6	95
10	Methodology for classification and definition of epilepsy syndromes with list of syndromes: Report of the ILAE Task Force on Nosology and Definitions. Epilepsia, 2022, 63, 1333-1348.	5.1	84
11	Genome-wide Polygenic Burden of Rare Deleterious Variants in Sudden Unexpected Death in Epilepsy. EBioMedicine, 2015, 2, 1063-1070.	6.1	74
12	The challenges of treating epilepsy with 25 antiepileptic drugs. Pharmacological Research, 2016, 107, 211-219.	7.1	72
13	Multimodal responses induced by cortical stimulation of the parietal lobe: a stereo-electroencephalography study. Brain, 2015, 138, 2596-2607.	7.6	64
14	Cerebellar, limbic, and midbrain volume alterations in sudden unexpected death in epilepsy. Epilepsia, 2019, 60, 718-729.	5.1	54
15	De novo mutations of i>KIAA2022 / i> in females cause intellectual disability and intractable epilepsy. Journal of Medical Genetics, 2016, 53, 850-858.	3.2	47
16	The <scp>ENIGMAâ€Epilepsy</scp> working group: Mapping disease from large data sets. Human Brain Mapping, 2022, 43, 113-128.	3.6	47
17	Epilepsy subtype-specific copy number burden observed in a genome-wide study of 17 458 subjects. Brain, 2020, 143, 2106-2118.	7.6	47
18	Markers for the Risk of Progression from Mild Cognitive Impairment to Alzheimer's Disease. Journal of Alzheimer's Disease, 2015, 45, 883-890.	2.6	44

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19	Clinical spectrum of <i>STX1B</i> -related epileptic disorders. Neurology, 2019, 92, e1238-e1249.	1.1	43
20	Applying a perceptions and practicalities approach to understanding nonadherence to antiepileptic drugs. Epilepsia, 2015, 56, 1398-1407.	5.1	38
21	Monogenic Epilepsies. Neurology, 2021, 97, 817-831.	1.1	38
22	One-Year Progression of Moderate Asymptomatic Carotid Stenosis Predicts the Risk of Vascular Events. Stroke, 2013, 44, 792-794.	2.0	36
23	Audit of use of stiripentol in adults with Dravet syndrome. Acta Neurologica Scandinavica, 2017, 135, 73-79.	2.1	36
24	The aetiologies of epilepsy. Epileptic Disorders, 2021, 23, 1-16.	1.3	35
25	Sub-genic intolerance, ClinVar, and the epilepsies: A whole-exome sequencing study of 29,165 individuals. American Journal of Human Genetics, 2021, 108, 965-982.	6.2	35
26	Comparative effectiveness of antiepileptic drugs in juvenile myoclonic epilepsy. Epilepsia Open, 2019, 4, 420-430.	2.4	34
27	Perampanel Confirms to Be Effective and Well-Tolerated as an Add-On Treatment in Patients With Brain Tumor-Related Epilepsy (PERADET Study). Frontiers in Neurology, 2020, 11, 592.	2.4	32
28	Climate change and epilepsy: Insights from clinical and basic science studies. Epilepsy and Behavior, 2021, 116, 107791.	1.7	30
29	Real-life survey of pitfalls and successes of precision medicine in genetic epilepsies. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 1044-1052.	1.9	30
30	Genomeâ€wide association study: Exploring the genetic basis for responsiveness to ketogenic dietary therapies for drugâ€resistant epilepsy. Epilepsia, 2018, 59, 1557-1566.	5.1	23
31	Psychiatric comorbidities in patients from seven families with autosomal dominant cortical tremor, myoclonus, and epilepsy. Epilepsy and Behavior, 2016, 56, 38-43.	1.7	21
32	Diagnostic delay and prognosis in primary central nervous system lymphoma compared with glioblastoma multiforme. Neurological Sciences, 2016, 37, 23-29.	1.9	20
33	Diagnostic Biomarkers of Epilepsy. Current Pharmaceutical Biotechnology, 2018, 19, 440-450.	1.6	20
34	Cardiac phenotype in <i>ATP1A3</i> -related syndromes. Neurology, 2020, 95, e2866-e2879.	1.1	19
35	Retinal nerve fibre layer thinning is associated with drug resistance in epilepsy. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, 396-401.	1.9	18
36	K.Vita: a feasibility study of a blend of medium chain triglycerides to manage drug-resistant epilepsy. Brain Communications, 2021, 3, fcab160.	3.3	17

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37	The impact of SARS-CoV-2 vaccination in Dravet syndrome: A UK survey. Epilepsy and Behavior, 2021, 124, 108258.	1.7	15
38	Electroclinical Features and Long-term Seizure Outcome in Patients With Eyelid Myoclonia With Absences. Neurology, 2022, 98, .	1.1	15
39	Sex-Associated Differences in the Modulation of Vascular Risk in Patients with Asymptomatic Carotid Stenosis. Journal of Cerebral Blood Flow and Metabolism, 2015, 35, 684-688.	4.3	13
40	Muscle and brain sodium channelopathies: genetic causes, clinical phenotypes, and management approaches. The Lancet Child and Adolescent Health, 2020, 4, 536-547.	5.6	13
41	Expanding the genetic and phenotypic spectrum of <scp><i>CHD2</i></scp> â€related disease: From early neurodevelopmental disorders to adultâ€onset epilepsy. American Journal of Medical Genetics, Part A, 2022, 188, 522-533.	1.2	13
42	Increased Common Carotid Artery Wall Thickness Is Associated with Rapid Progression of Asymptomatic Carotid Stenosis. Journal of Neuroimaging, 2014, 24, 473-478.	2.0	11
43	Clinical outcomes of COVID-19 in long-term care facilities for people with epilepsy. Epilepsy and Behavior, 2021, 115, 107602.	1.7	11
44	Postictal Psychosis in Epilepsy: A Clinicogenetic Study. Annals of Neurology, 2021, 90, 464-476.	5.3	11
45	Two-center experience of cannabidiol use in adults with Dravet syndrome. Seizure: the Journal of the British Epilepsy Association, 2021, 91, 5-8.	2.0	11
46	Intracerebral electrical stimulations of the temporal lobe: A stereoelectroencephalography study. European Journal of Neuroscience, 2021, 54, 5368-5383.	2.6	10
47	Emergency room access for recurring seizures: when and why. European Journal of Neurology, 2013, 20, 1411-1416.	3.3	9
48	Transcranial magnetic stimulation as a tool to understand genetic conditions associated with epilepsy. Epilepsia, 2020, 61, 1818-1839.	5.1	9
49	Late diagnoses of Dravet syndrome: How many individuals are we missing?. Epilepsia Open, 2021, 6, 770-776.	2.4	9
50	10.2174/1381612823666170809115827. Current Pharmaceutical Design, 2018, 23, 5667-5690.	1.9	9
51	Percutaneous transluminal angioplasty for chronic cerebrospinal venous insufficiency in multiple sclerosis: dichotomy between subjective and objective outcome scores. Neurological Sciences, 2013, 34, 2205-2210.	1.9	8
52	From Cannabis to Cannabidiol to Treat Epilepsy, Where Are We?. Current Pharmaceutical Design, 2017, 22, 6426-6433.	1.9	8
53	Coasting, embryo development and outcomes of blastocyst transfer: a case–control study. Reproductive BioMedicine Online, 2014, 29, 231-238.	2.4	7
54	Ring Chromosome 17 Not Involving the Miller-Dieker Region: A Case with Drug-Resistant Epilepsy. Molecular Syndromology, 2018, 9, 38-44.	0.8	7

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55	The impact of COVIDâ€19 in Dravet syndrome: A UK survey. Acta Neurologica Scandinavica, 2021, 143, 389-395.	2.1	7
56	Efficacy and Safety of Long-Term Treatment with Stiripentol in Children and Adults with Drug-Resistant Epilepsies: A Retrospective Cohort Study of 196 Patients. Drugs - Real World Outcomes, 2022, 9, 451-461.	1.6	6
57	Cortical myoclonus and epilepsy in a family with a new SLC20A2 mutation. Journal of Neurology, 2020, 267, 2221-2227.	3.6	5
58	Increased facial asymmetry in focal epilepsies associated with unilateral lesions. Brain Communications, 2021, 3, fcab068.	3.3	5
59	Rare and Complex Epilepsies from Childhood to Adulthood: Requirements for Separate Management or Scope for a Lifespan Holistic Approach?. Current Neurology and Neuroscience Reports, 2021, 21, 65.	4.2	4
60	Generalized epilepsy in a patient with myotonic dystrophy type 2. Neurological Sciences, 2014, 35, 489-490.	1.9	3
61	Safe use of perampanel in a carrier of variegate porphyria. Practical Neurology, 2016, 16, 217-219.	1.1	3
62	Personalized treatment in the epilepsies: challenges and opportunities. Expert Review of Precision Medicine and Drug Development, 2018, 3, 237-247.	0.7	3
63	Complex epilepsy: it's all in the history. Practical Neurology, 2021, 21, 153-156.	1.1	3
64	Reply: The dorsal cingulate cortex as a critical gateway in the network supporting conscious awareness. Brain, 2016, 139, e24-e24.	7.6	2
65	Ammonia: what adult neurologists need to know. Practical Neurology, 2020, , practneurol-2020-002654.	1.1	2
66	Transcranial magnetic stimulation as a biomarker of treatment response in children with epilepsy. Developmental Medicine and Child Neurology, 2020, 62, 770-770.	2.1	1
67	Nonâ€Stationary Outcome of Alternating Hemiplegia of Childhood into Adulthood. Movement Disorders Clinical Practice, 2022, 9, 206-211.	1.5	1
68	Burden of uncontrolled epilepsy in patients requiring an emergency room visit or hospitalization. Neurology, 2013, 80, 2170-2170.	1.1	0
69	Drug-resistant epilepsy, early-onset hypertension and white matter lesions: a hidden paraganglioma. BMJ Case Reports, 2019, 12, e228348.	0.5	O