Miriam Kessi

List of Publications by Year
in descending order

21	251	1163117 8	1058476 14
papers	citations	h-index	g-index

24	24	24	312
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Genetic etiologies of the electrical status epilepticus during slow wave sleep: systematic review. BMC Genetics, 2018, 19, 40.	2.7	39
2	Calcium channelopathies and intellectual disability: a systematic review. Orphanet Journal of Rare Diseases, 2021, 16, 219.	2.7	33
3	Myelin Oligodendrocyte Glycoprotein (MOG) Antibody Diseases in Children in Central South China: Clinical Features, Treatments, Influencing Factors, and Outcomes. Frontiers in Neurology, 2019, 10, 868.	2.4	29
4	Intellectual Disability and Potassium Channelopathies: A Systematic Review. Frontiers in Genetics, 2020, 11, 614.	2.3	28
5	Rare Copy Number Variations and Predictors in Children With Intellectual Disability and Epilepsy. Frontiers in Neurology, 2018, 9, 947.	2.4	23
6	The Contribution of HCN Channelopathies in Different Epileptic Syndromes, Mechanisms, Modulators, and Potential Treatment Targets: A Systematic Review. Frontiers in Molecular Neuroscience, 2022, 15, .	2.9	15
7	Efficacy of different treatment modalities for acute and chronic phases of the febrile infection-related epilepsy syndrome: A systematic review. Seizure: the Journal of the British Epilepsy Association, 2020, 79, 61-68.	2.0	13
8	The Study of Genetic Susceptibility and Mitochondrial Dysfunction in Mesial Temporal Lobe Epilepsy. Molecular Neurobiology, 2020, 57, 3920-3930.	4.0	11
9	Familial SYN1 variants related neurodevelopmental disorders in Asian pediatric patients. BMC Medical Genomics, 2021, 14, 182.	1.5	9
10	Treatment for the Benign Childhood Epilepsy With Centrotemporal Spikes: A Monocentric Study. Frontiers in Neurology, 2021, 12, 670958.	2.4	8
11	De Novo KCNQ2 Mutation in One Case of Epilepsy of Infancy With Migrating Focal Seizures That Evolved to Infantile Spasms. Child Neurology Open, 2018, 5, 2329048X1876773.	1.1	7
12	Immunotherapies for Anti-N-M-methyl-D-aspartate Receptor Encephalitis: Multicenter Retrospective Pediatric Cohort Study in China. Frontiers in Pediatrics, 2021, 9, 691599.	1.9	7
13	Etiologic Classification of 541 Infantile Spasms Cases: A Cohort Study. Frontiers in Pediatrics, 2022, 10, 774828.	1.9	7
14	A Case With 4 de Novo Copy Number Variations With Clinical Features That Overlap 1q43q44 Microdeletion and 3q29 Microduplication Syndromes. Child Neurology Open, 2018, 5, 2329048X1879820.	1.1	4
15	Correlation Analyses of Clinical Manifestations and Variant Effects in KCNB1-Related Neurodevelopmental Disorder. Frontiers in Pediatrics, 2021, 9, 755344.	1.9	4
16	Urine Organic Acids as Metabolic Indicators for Global Developmental Delay/Intellectual Disability in Chinese Children. Frontiers in Molecular Biosciences, 2021, 8, 792319.	3.5	4
17	The Recommendations for the Management of Chinese Children With Epilepsy During the COVID-19 Outbreak. Frontiers in Pediatrics, 2020, 8, 495.	1.9	3
18	Observed and self-reported hand hygiene compliance and associated factors among healthcare workers at a county referral hospital in Kenya. Scientific African, 2021, 14, e00984.	1.5	2

#	Article	IF	Citations
19	Restoration of Sarco/Endoplasmic Reticulum Ca2+-ATPase Activity Functions as a Pivotal Therapeutic Target of Anti-Glutamate-Induced Excitotoxicity to Attenuate Endoplasmic Reticulum Ca2+ Depletion. Frontiers in Pharmacology, 2022, 13, 877175.	3.5	2
20	The patterns of response of 11 regimens for infantile spasms. Scientific Reports, 2020, 10, 11509.	3.3	1
21	A survey on pediatric anti-N-methyl-D-aspartate-receptor encephalitis treatment strategies in China. Chinese Medical Journal, 2021, 134, 1498-1499.	2.3	1