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	Etiologic Classification of 541 Infantile Spasms Cases: A Cohort Study. Frontiers in Pediatrics, 2022, 10, 774828.	1.9	7
2	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
3	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
4	Treatment for the Benign Childhood Epilepsy With Centrotemporal Spikes: A Monocentric Study. Frontiers in Neurology, 2021, 12, 670958.	2.4	8
5	Calcium channelopathies and intellectual disability: a systematic review. Orphanet Journal of Rare Diseases, 2021, 16, 219.	2.7	33
6	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
7	Familial SYN1 variants related neurodevelopmental disorders in Asian pediatric patients. BMC Medical Genomics, 2021, 14, 182.	1.5	9
8	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
9	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
10	Correlation Analyses of Clinical Manifestations and Variant Effects in KCNB1-Related Neurodevelopmental Disorder. Frontiers in Pediatrics, 2021, 9, 755344.	1.9	4
11	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
12	The Study of Genetic Susceptibility and Mitochondrial Dysfunction in Mesial Temporal Lobe Epilepsy. Molecular Neurobiology, 2020, 57, 3920-3930.	4.0	11
13	The patterns of response of 11 regimens for infantile spasms. Scientific Reports, 2020, 10, 11509.	3.3	1
14	The Recommendations for the Management of Chinese Children With Epilepsy During the COVID-19 Outbreak. Frontiers in Pediatrics, 2020, 8, 495.	1.9	3
15	Intellectual Disability and Potassium Channelopathies: A Systematic Review. Frontiers in Genetics, 2020, 11, 614.	2.3	28
16	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
17	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
18	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

#	Article	IF	Citations
19	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
20	Rare Copy Number Variations and Predictors in Children With Intellectual Disability and Epilepsy. Frontiers in Neurology, 2018, 9, 947.	2.4	23
21	Genetic etiologies of the electrical status epilepticus during slow wave sleep: systematic review. BMC Genetics, 2018, 19, 40.	2.7	39