

Anita N Datta

List of Publications by Year in descending order

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Version: 2024-02-01

21
papers

334
citations

1307594

7
h-index

839539

18
g-index

21
all docs

21
docs citations

21
times ranked

822
citing authors

#	ARTICLE	IF	CITATIONS
1	Loss-of-Function and Gain-of-Function Mutations in KCNQ5 Cause Intellectual Disability or Epileptic Encephalopathy. <i>American Journal of Human Genetics</i> , 2017, 101, 65-74.	6.2	99
2	Diagnostic Yield and Treatment Impact of Targeted Exome Sequencing in Early-Onset Epilepsy. <i>Frontiers in Neurology</i> , 2019, 10, 434.	2.4	70
3	Lack of response to quinidine in <i>KCNT1</i> -related neonatal epilepsy. <i>Epilepsia</i> , 2018, 59, 1889-1898.	5.1	53
4	mTOR Inhibitors as a New Therapeutic Strategy in Treatment Resistant Epilepsy in Hemimegalencephaly: A Case Report. <i>Journal of Child Neurology</i> , 2019, 34, 132-138.	1.4	24
5	An Infant With Epilepsy and Recurrent Hemiplegia due to Compound Heterozygous Variants in ATP1A2. <i>Pediatric Neurology</i> , 2017, 75, 87-90.	2.1	21
6	Clinical Experience With Perampanel for Refractory Pediatric Epilepsy in One Canadian Center. <i>Journal of Child Neurology</i> , 2017, 32, 834-839.	1.4	15
7	Manifestations of Tuberous Sclerosis Complex: The Experience of a Provincial Clinic. <i>Canadian Journal of Neurological Sciences</i> , 2017, 44, 35-43.	0.5	15
8	Two Patients With KCNT1-Related Epilepsy Responding to Phenobarbital and Potassium Bromide. <i>Journal of Child Neurology</i> , 2019, 34, 728-734.	1.4	7
9	Co-existence of Rolandic and 3 Hz Spike-Wave Discharges on EEG in Children with Epilepsy. <i>Canadian Journal of Neurological Sciences</i> , 2019, 46, 64-70.	0.5	6
10	Case Report. <i>Journal of Child Neurology</i> , 2017, 32, 403-407.	1.4	5
11	Childhood Small Vessel Primary Angiitis of the Central Nervous System: A Treatable Cause of Super-refractory Status Epilepticus. <i>Journal of Child Neurology</i> , 2020, 35, 31-36.	1.4	4
12	Resilience of adolescents and teenagers with self-limited and genetic-generalized epilepsy during the COVID-19 pandemic. <i>Epilepsy and Behavior Reports</i> , 2022, 17, 100520.	1.0	4
13	Pediatric Occipital Spikes at a Single Center Over 26 Years and the Significance of Tangential Dipole. <i>Journal of Child Neurology</i> , 2021, 36, 530-536.	1.4	3
14	Infantile Spasms and Trisomy 21: Unfavorable Outcomes with First-line Vigabatrin Therapy. <i>Canadian Journal of Neurological Sciences</i> , 2021, , 1-6.	0.5	2
15	Hypnagogic Frontal EEG Bursts in Children. <i>Journal of Clinical Neurophysiology</i> , 2020, Publish Ahead of Print, 536-541.	1.7	2
16	Predictive Value of Midline Spikes on Pediatric EEG for Seizure and Developmental Outcome. <i>Journal of Clinical Neurophysiology</i> , 2018, 35, 490-495.	1.7	1
17	Clinical Significance of Incidental Rolandic Spikes in Children With Absence Epilepsy. <i>Journal of Child Neurology</i> , 2019, 34, 631-638.	1.4	1
18	At What Point Should We Discontinue Treatment in Super-Refractory Status Epilepticus?. <i>Journal of Pediatric Epilepsy</i> , 2020, 9, 100-101.	0.2	1

#	ARTICLE	IF	CITATIONS
19	School performance in children at the time of new-onset seizures and at long-term follow-up: A retrospective cohort study. <i>Journal of International Medical Research</i> , 2022, 50, 030006052210810.	1.0	1
20	Children With Trisomy 21 and Lennox-Gastaut Syndrome With Predominant Myoclonic Seizures. <i>Journal of Child Neurology</i> , 2021, 36, 1027-1033.	1.4	0
21	Effect of Training on Visual Identification of High Frequency Oscillationsâ€™A Delphi-Style Intervention. <i>Frontiers in Neurology</i> , 2022, 13, 794668.	2.4	0