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List of Publications by Year
in descending order

39 papers	815 citations	687363 13 h-index	526287 27 g-index
40 all docs	40 docs citations	40 times ranked	1695 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	PRRT2 Mutations are the major cause of benign familial infantile seizures. Human Mutation, 2012, 33, 1439-1443.	2.5	93
3	A prevalence study of restless legs syndrome in Turkish children and adolescents. Sleep Medicine, 2011, 12, 315-321.	1.6	83
4	Epilepsy subtype-specific copy number burden observed in a genome-wide study of 17â€‰458 subjects. Brain, 2020, 143, 2106-2118.	7.6	47
5	Characteristic and Overlapping Features of Migraine and Tension-Type Headache. Headache, 2006, 46, 461-468.	3.9	41
6	Lipid Peroxidation and Antioxidative Enzyme Activities in Childhood Epilepsy. Journal of Child Neurology, 2002, 17, 673-676.	1.4	38
7	Subdural EEG Patterns in Children With Taylor-Type Cortical Dysplasia: Comparison With Nondysplastic Lesions. Journal of Clinical Neurophysiology, 2005, 22, 37-42.	1.7	38
8	Genomeâ€wide linkage metaâ€analysis identifies susceptibility loci at 2q34 and 13q31.3 for genetic generalized epilepsies. Epilepsia, 2012, 53, 308-318.	5.1	32
9	Auditory neuropathy in hyperbilirubinemia: is there a correlation between serum bilirubin, neuron-specific enolase levels and auditory neuropathy?. International Journal of Audiology, 2004, 43, 516-522.	1.7	23
10	Visual and auditory event related potentials in epileptic children: a comparison with normal and abnormal MRI findings. Brain and Development, 2003, 25, 396-400.	1.1	17
11	Ketogenic diet as a successful early treatment modality for SCN2A mutation. Brain and Development, 2019, 41, 389-391.	1.1	17
12	Deep brain stimulation as treatment for dystonic storm in pantothenate kinase-associated neurodegeneration syndrome: case report of a patient with homozygous C.628 2ÂTâ€‰>â€‰G mutation of the PANK2 gene. Acta Neurochirurgica, 2015, 157, 1513-1517.	1.7	16
13	Involvement of sympathetic reflex activity in patients with acute and chronic stroke: a comparison with functional motor capacity. Archives of Physical Medicine and Rehabilitation, 2004, 85, 470-473.	0.9	14
14	The frequency of late-onset Pompe disease in pediatric patients with limb-girdle muscle weakness and nonspecific hyperCKemia: A multicenter study. Neuromuscular Disorders, 2016, 26, 796-800.	0.6	14
15	SCN1A gene sequencing in 46 Turkish epilepsy patients disclosed 12 novel mutations. Seizure: the Journal of the British Epilepsy Association, 2016, 39, 34-43.	2.0	13
16	Familial early infantile epileptic encephalopathy and cardiac conduction disorder: A rare cause of SUDEP in infancy. Seizure: the Journal of the British Epilepsy Association, 2017, 50, 171-172.	2.0	13
17	Glial fibrillary acidic protein (GFAP)-antibody in children with focal seizures of undetermined cause. Acta Neurologica Belgica, 2021, 121, 1275-1280.	1.1	9
18	The Effects of Neurodevelopmental Therapy on Feeding and Swallowing Activities in Children with Cerebral Palsy. Dysphagia, 2022, 37, 800-811.	1.8	9

#	ARTICLE	IF	CITATIONS
19	When Is EEG Indicated in Attention-Deficit/Hyperactivity Disorder?. Journal of Child Neurology, 2015, 30, 1785-1793.	1.4	8
20	An association analysis at 2q36 reveals a new candidate susceptibility gene for juvenile absence epilepsy and/or absence seizures associated with generalized tonic-clonic seizures. Epilepsia, 2011, 52, 975-983.	5.1	6
21	Anti-N-Methyl-<scp>d</scp>-Aspartate (Anti-NMDA) Receptor Encephalitis. Journal of Child Neurology, 2014, 29, 684-687.	1.4	6
22	The Burden of Primary Caregivers of Spinal Muscular Atrophy Patients and Their Needs. , 2021, 56, 366-373.		6
23	Possible role of SCN4A skeletal muscle mutation in apnea during seizure. Epilepsia Open, 2019, 4, 498-503.	2.4	5
24	Evaluation of cases with cerebral thrombosis in children. Turk Pediatri Arsivi, 2016, 51, 87-93.	0.9	5
25	Transient Splenial Lesion of the Corpus Callosum Related to Migraine with Aura in a Pediatric Patient. Acta Medica (Hradec Kralove), 2016, 59, 64-66.	0.5	5
26	De novo FZR1 loss-of-function variants cause developmental and epileptic encephalopathies. Brain, 2022, 145, 1684-1697.	7.6	5
27	A novel truncating mutation of DOCK7 gene with an early-onset non-encephalopathic epilepsy. Seizure: the Journal of the British Epilepsy Association, 2019, 66, 12-14.	2.0	4
28	Awareness of sudden unexpected death in epilepsy among parents of children with epilepsy in a tertiary center. Epilepsy and Behavior, 2020, 111, 107125.	1.7	4
29	Demographic and Clinical Findings of Cerebral Palsy Patients in Istanbul: A Multicenter Study. FTR - Turkiye Fiziksel Tip Ve Rehabilitasyon Dergisi, 2014, 60, 134-138.	0.1	3
30	The effect of neck-trunk stabilization exercises in cerebral palsy: randomized controlled trial. Minerva Pediatrics, 2021, , .	0.4	2
31	Neurodevelopmental Findings and Epilepsy in Malformations of Cortical Development. , 2021, 56, 356-356.		1
32	Overlapping features of restless legs syndrome and growing pains in Turkish children and adolescents. Brain and Development, 2022, 44, 372-379.	1.1	1
33	Continuous muscle fiber activity syndrome. Pediatrics International, 1995, 37, 94-96.	0.5	0
34	Global aphasia without hemiparesis may be caused by blunt head trauma: An adolescent boy with transient aphasia. Journal of Clinical Neuroscience, 2017, 39, 84-86.	1.5	0
35	Relationships Among 3 Movement Analysis Tests in Preterm Infants. Pediatric Physical Therapy, 2019, 31, 251-256.	0.6	0
36	CLB add-on treatment in patients with epileptic encephalopathy: a single center experience with long-term follow-up. Acta Neurologica Belgica, 2022, 122, 51-57.	1.1	0

#	ARTICLE	IF	CITATIONS
37	Chromosomal microarray and exome sequencing in unexplained early infantile epileptic encephalopathies in a highly consanguineous population. International Journal of Neuroscience, 2023, 133, 683-700.	1.6	0
38	How do presentation age and CSF opening pressure level affect long-term prognosis of pseudotumor cerebri syndrome in children? Experience of a single tertiary clinic. Child's Nervous System, 2021, , 1.	1.1	0
39	Effect of Nusinersen treatment on motor functions in children and adolescents with spinal muscular atrophy who gave a break to physiotherapy during COVID-19 pandemic. Turkish Journal of Physical Medicine and Rehabilitation, 2022, 68, 157-158.	0.9	0