Morteza Heidari

List of Publications by Year in descending order

35 papers	276 citations	1170033 9 h-index	1113639 15 g-index
41 all docs	41 docs citations	41 times ranked	391 citing authors

#	Article	IF	Citations
1	A novel homozygous missense variant in the NAXE gene in an Iranian family with progressive encephalopathy with brain edema and leukoencephalopathy. Acta Neurologica Belgica, 2022, 122, 1201-1210.	0.5	10
2	Primary and Secondary Microcephaly, Global Developmental Delay, and Seizure in Two Siblings Caused by a Novel Missense Variant in the ZNF335 Gene. Journal of Molecular Neuroscience, 2022, 72, 719-729.	1.1	8
3	Genetic Analysis of Forty MLPA-Negative Duchenne Muscular Dystrophy Patients by Whole-Exome Sequencing. Journal of Molecular Neuroscience, 2022, , 1.	1.1	1
4	The safety and efficacy of umbilical cord blood mononuclear cells in individuals with spastic cerebral palsy: a randomized double-blind sham-controlled clinical trial. BMC Neurology, 2022, 22, 123.	0.8	7
5	Epilepsia Partialis Continua a Clinical Feature of a Missense Variant in the ADCK3 Gene and Poor Response to Therapy. Journal of Molecular Neuroscience, 2022, 72, 1125-1132.	1.1	6
6	Followâ€up of 25 patients with treatable ataxia: AÂcomprehensive case series study. Clinical Case Reports (discontinued), 2022, 10, e05777.	0.2	1
7	Characteristics of disease progression and genetic correlation in ambulatory Iranian boys with Duchenne muscular dystrophy. BMC Neurology, 2022, 22, 162.	0.8	4
8	A novel pathogenic variant of BRAT1 gene causes rigidity and multifocal seizure syndrome, lethal neonatal. International Journal of Neuroscience, 2021, 131, 875-878.	0.8	8
9	Identification of a novel missense c.386Gâ€‰>â€‰A variant in a boy with the POMGNT1-related muscular dystrophy-dystroglycanopathy. Acta Neurologica Belgica, 2021, 121, 143-151.	0.5	11
10	Managing Status Epilepticus in a Child with Dravet Syndrome: How Difficult It Could Be?. Journal of Pediatric Epilepsy, 2021, 10, 128-134.	0.1	1
11	The Association between Behavioral Problems with SelfEsteem and Self-Concept in Pediatric Patients with Thalassemia. Iranian Journal of Psychiatry, 2021, 16, 36-42.	0.4	1
12	Clinical and Paraclinical Screening for Celiac Disease in Children with Intractable Epilepsy. Neurology Research International, 2021, 2021, 1-4.	0.5	0
13	Mitochondrial <scp>D</scp><scp>NA</scp> Analysis from Exome Sequencing Data Improves Diagnostic Yield in Neurological Diseases. Annals of Neurology, 2021, 89, 1240-1247.	2.8	12
14	ACER3-related leukoencephalopathy: expanding the clinical and imaging findings spectrum due to novel variants. Human Genomics, 2021, 15, 45.	1.4	1
15	Clinical and imaging outcomes after intrathecal injection of umbilical cord tissue mesenchymal stem cells in cerebral palsy: a randomized double-blind sham-controlled clinical trial. Stem Cell Research and Therapy, 2021, 12, 439.	2.4	18
16	Defective complex III mitochondrial respiratory chain due to a novel variant in CYC1 gene masquerades acute demyelinating syndrome or Leber hereditary optic neuropathy. Mitochondrion, 2021, 60, 12-20.	1.6	4
17	Cinnarizine and sodium valproate as the preventive agents of pediatric migraine: A randomized double-blind placebo-controlled trial. Cephalalgia, 2020, 40, 665-674.	1.8	7
18	An update on clinical, pathological, diagnostic, and therapeutic perspectives of childhood leukodystrophies. Expert Review of Neurotherapeutics, 2020, 20, 65-84.	1.4	47

#	Article	IF	Citations
19	Letter to the Editor on â€œPerspective: Therapeutic Potential of Flavonoids as Alternative Medicines in Epilepsyâ€• Advances in Nutrition, 2020, 11, 741.	2.9	0
20	The First Comprehensive Cohort of the Duchenne Muscular Dystrophy in Iranian Population: Mutation Spectrum of 314 Patients and Identifying Two Novel Nonsense Mutations. Journal of Molecular Neuroscience, 2020, 70, 1565-1573.	1.1	7
21	Clinical Characteristics and Electrodiagnostic Features of Guillain-BarrÃ© Syndrome Among the Pediatric Population. Journal of Child Neurology, 2020, 35, 448-455.	0.7	4
22	Children with Epilepsy in the Era of COVID-19 Pandemic. Journal of Pediatric Epilepsy, 2020, 09, 053-054.	0.1	0
23	Coronavirus, Its Neurologic Manifestations, and Complications. Iranian Journal of Pediatrics, 2020, 30, .	0.1	10
24	Levetiracetam for prophylactic treatment of pediatric migraine: A randomized double-blind placebo-controlled trial. Cephalalgia, 2019, 39, 1509-1517.	1.8	11
25	The First Report of Relative Incidence of Inherited White Matter Disorders in an Asian Country Based on an Iranian Bioregistry System. Journal of Child Neurology, 2018, 33, 255-259.	0.7	13
26	The Efficacy of Cerebrolysin in Improvement of Spasticity in Children with Cerebral Palsy: A Clinical Trial. Iranian Journal of Pediatrics, 2018, 28, .	0.1	1
27	The efficacy of the ketogenic diet in infants and young children with refractory epilepsies using a formula-based powder. Acta Neurologica Belgica, 2017, 117, 175-182.	0.5	12
28	The quality of life in boys with Duchenne muscular dystrophy. Neuromuscular Disorders, 2016, 26, 423-427.	0.3	19
29	Efficacy of Potassium Bromide (As an Out-of-Date Drug) in Epilepsia Partialis Continua: A Brief Report. Iranian Journal of Pediatrics, 2016, In Press, .	0.1	0
30	Association of IL6 single nucleotide polymorphisms with febrile seizures. Journal of the Neurological Sciences, 2014, 342, 25-28.	0.3	20
31	Efficacy and Safety of Cinnarizine in the Prophylaxis of Migraine in Children: A Double-Blind Placebo-Controlled Randomized Trial. Pediatric Neurology, 2014, 51, 503-508.	1.0	28
32	Diagnosis of an abnormality in brain MRI. Iranian Journal of Pediatrics, 2014, 24, 779-80.	0.1	0
33	Impact of N-myc Amplification on Median Survival in Children With Neuroblastoma. Journal of Comprehensive Pediatrics, 2012, 3, 29-33.	0.1	3
34	We need well-designed multicenter studies to investigate neurologic manifestations of coronavirus disease 2019. Current Journal of Neurology, 0, , .	0.0	0
35	Clinical and Molecular Findings of Autosomal Recessive Spastic Ataxia of Charlevoix Saguenay: an Iranian Case Series Expanding the Genetic and Neuroimaging Spectra. Cerebellum, 0, , .	1.4	1