Sareh Hosseinpour

List of Publications by Year
in descending order

9 papers	52 citations	1937685 4 h-index	1720034 7 g-index
12 all docs	12 docs citations	12 times ranked	143 citing authors

#	Article	IF	Citations
1	Primary creatine deficiency syndrome as a potential missed diagnosis in children with psychomotor delay and seizure: case presentation with two novel variants and literature review. Acta Neurologica Belgica, 2020, 120, 511-516.	1.1	12
2	Homozygous inâ€frame variant of SCL6A3 causes dopamine transporter deficiency syndrome in a consanguineous family. Annals of Human Genetics, 2020, 84, 315-323.	0.8	12
3	Hypomyelinating Leukodystrophy with Spinal Cord Involvement Caused by a Novel Variant in RARS: Report of Two Unrelated Patients. Neuropediatrics, 2019, 50, 130-134.	0.6	8
4	Deep brain stimulation in status dystonicus caused by anti-NMDA receptor encephalitis. Parkinsonism and Related Disorders, 2019, 66, 255-257.	2.2	6
5	Leukoencephalopathy in RIN2 syndrome: Novel mutation and expansion of clinical spectrum. European Journal of Medical Genetics, 2020, 63, 103629.	1.3	5
6	Persistent dystonia and basal ganglia involvement following metronidazole induced encephalopathy. Neurological Sciences, 2020, 41, 957-959.	1.9	4
7	Characteristics of disease progression and genetic correlation in ambulatory Iranian boys with Duchenne muscular dystrophy. BMC Neurology, 2022, 22, 162.	1.8	4
8	Genetic Analysis of Forty MLPA-Negative Duchenne Muscular Dystrophy Patients by Whole-Exome Sequencing. Journal of Molecular Neuroscience, 2022, , 1.	2.3	1
9	An Asymptomatic Case of Megalencephalic Leukoencephalopathy with Subcortical Cysts. Iranian Journal of Pediatrics, 2019, 29, .	0.3	0