Matthias Eckenweiler

List of Publications by Year
in descending order

14 papers	271 citations	1163117 8 h-index	1125743 13 g-index
14 all docs	14 docs citations	14 times ranked	558 citing authors

#	Article	IF	Citations
1	A Novel Variant of <scp>ATP5MC3</scp> Associated with Both Dystonia and Spastic Paraplegia. Movement Disorders, 2022, 37, 375-383.	3.9	10
2	Variants in Mitochondrial <scp>ATP</scp> Synthase Cause Variable Neurologic Phenotypes. Annals of Neurology, 2022, 91, 225-237.	5.3	12
3	Quality of Life After Deep Brain Stimulation of Pediatric Patients with Dyskinetic Cerebral Palsy: A Prospective, Singleâ€Arm, Multicenter Study with a Subsequent Randomized Doubleâ€Blind Crossover (<scp>STIMâ€CP</scp>). Movement Disorders, 2022, 37, 799-811.	3.9	10
4	Postâ€dural puncture headacheâ€”a singleâ€centre analysis in paediatric patients with and without SMA. Acta Paediatrica, International Journal of Paediatrics, 2021, 110, 1895-1901.	1.5	0
5	The spectrum of peripheral neuropathy in disorders of the mitochondrial trifunctional protein. Journal of Inherited Metabolic Disease, 2021, 44, 893-902.	3.6	12
6	Agammaglobulinemia with normal B-cell numbers in a patient lacking Bob1. Journal of Allergy and Clinical Immunology, 2021, 147, 1977-1980.	2.9	12
7	Scoring Algorithmâ€Based Genomic Testing in Dystonia: A Prospective Validation Study. Movement Disorders, 2021, 36, 1959-1964.	3.9	7
8	Facilitation of drug-resistant epilepsy and catastrophic status epilepticus in children with combined pituitary hormone deficiency. European Journal of Paediatric Neurology, 2021, 33, 99-105.	1.6	1
9	Thiamine Treatment and Favorable Outcome in an Infant with Biallelic TPK1 Variants. Neuropediatrics, 2021, 52, 123-125.	0.6	4
10	Severe Locked-In-Like Guillainâ€“BarrÃ©'s Syndrome: Dilemmas in Diagnosis and Treatment. Neuropediatrics, 2021, 52, 019-026.	0.6	5
11	Monogenic variants in dystonia: an exome-wide sequencing study. Lancet Neurology, The, 2020, 19, 908-918.	10.2	139
12	Cerebrospinal fluid findings in patients with myelin oligodendrocyte glycoprotein (MOG) antibodies. Part 2: Results from 108 lumbar punctures in 80 pediatric patients. Journal of Neuroinflammation, 2020, 17, 262.	7.2	44
13	De novo variant in SCN4A causes neonatal sodium channel myotonia with general muscle stiffness and respiratory failure. Neuromuscular Disorders, 2019, 29, 907-909.	0.6	5
14	Novel mutation in two brothers with Hermansky Pudlak syndrome type 3. Blood Cells, Molecules, and Diseases, 2017, 67, 75-80.	1.4	10