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	Monogenic variants in dystonia: an exome-wide sequencing study. Lancet Neurology, The, 2020, 19, 908-918.	10.2	139
2	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
3	The spectrum of peripheral neuropathy in disorders of the mitochondrial trifunctional protein. Journal of Inherited Metabolic Disease, 2021, 44, 893-902.	3.6	12
4	Agammaglobulinemia with normal B-cell numbers in a patient lacking Bob1. Journal of Allergy and Clinical Immunology, 2021, 147, 1977-1980.	2.9	12
5	Variants in Mitochondrial <scp>ATP</scp> Synthase Cause Variable Neurologic Phenotypes. Annals of Neurology, 2022, 91, 225-237.	5.3	12
6	Novel mutation in two brothers with Hermansky Pudlak syndrome type 3. Blood Cells, Molecules, and Diseases, 2017, 67, 75-80.	1.4	10
7	A Novel Variant of <scp>ATP5MC3</scp> Associated with Both Dystonia and Spastic Paraplegia. Movement Disorders, 2022, 37, 375-383.	3.9	10
8	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
9	Scoring Algorithmâ€Based Genomic Testing in Dystonia: A Prospective Validation Study. Movement Disorders, 2021, 36, 1959-1964.	3.9	7
10	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
11	Severe Locked-In-Like Guillainâ€“BarrÃ©'s Syndrome: Dilemmas in Diagnosis and Treatment. Neuropediatrics, 2021, 52, 019-026.	0.6	5
12	Thiamine Treatment and Favorable Outcome in an Infant with Biallelic TPK1 Variants. Neuropediatrics, 2021, 52, 123-125.	0.6	4
13	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
14	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