Astrid Blaschek

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/8578311/publications.pdf

Version: 2024-02-01

32	1,559	430874	454955
papers	citations	18 h-index	30 g-index

33	33	33	1908
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Prognostic relevance of MOG antibodies in children with an acquired demyelinating syndrome. Neurology, 2017, 89, 900-908.	1.1	278
2	Antiâ€“Myelin Oligodendrocyte Glycoprotein Antibodies in Pediatric Patients With Optic Neuritis. Archives of Neurology, 2012, 69, 752-6.	4.5	181
3	Autoantibodies to MOG in a distinct subgroup of adult multiple sclerosis. Neurology: Neuroimmunology and NeuroInflammation, 2016, 3, e257.	6.0	178
4	One Year of Newborn Screening for SMA â€“ Results of a German Pilot Project. Journal of Neuromuscular Diseases, 2019, 6, 503-515.	2.6	105
5	Antibodies to MOG and AQP4 in children with neuromyelitis optica and limited forms of the disease. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, 897-905.	1.9	98
6	Newborn screening for spinal muscular atrophy in Germany: clinical results after 2 years. Orphanet Journal of Rare Diseases, 2021, 16, 153.	2.7	81
7	MRI of the first event in pediatric acquired demyelinating syndromes with antibodies to myelin oligodendrocyte glycoprotein. Journal of Neurology, 2018, 265, 845-855.	3.6	68
8	Oligoclonal bands predict multiple sclerosis in children with optic neuritis. Annals of Neurology, 2015, 77, 1076-1082.	5.3	61
9	Optical coherence tomography in myelin-oligodendrocyte-glycoprotein antibody-seropositive patients: a longitudinal study. Journal of Neuroinflammation, 2019, 16, 154.	7.2	61
10	Gene replacement therapy with onasemnogene abeparvovec in children with spinal muscular atrophy aged 24 months or younger and bodyweight up to 15 kg: an observational cohort study. The Lancet Child and Adolescent Health, 2022, 6, 17-27.	5.6	57
11	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
12	Self-reported neck pain is associated with migraine but not with tension-type headache in adolescents. Cephalalgia, 2014, 34, 895-903.	3.9	39
13	Infants Diagnosed with Spinal Muscular Atrophy and 4 SMN2 Copies through Newborn Screening â€“ Opportunity or Burden?1. Journal of Neuromuscular Diseases, 2020, 7, 109-117.	2.6	39
14	Patientâ€specific determinants of responsiveness to robotâ€enhanced treadmill therapy in children and adolescents with cerebral palsy. Developmental Medicine and Child Neurology, 2014, 56, 1172-1179.	2.1	38
15	Childhood multiple sclerosis is associated with reduced brain volumes at first clinical presentation and brain growth failure. Multiple Sclerosis Journal, 2019, 25, 927-936.	3.0	32
16	Fatigue and depression predict health-related quality of life in patients with pediatric-onset multiple sclerosis. Multiple Sclerosis and Related Disorders, 2019, 36, 101368.	2.0	31
17	SACS variants are a relevant cause of autosomal recessive hereditary motor and sensory neuropathy. Human Genetics, 2018, 137, 911-919.	3.8	29
18	Endocrine and Growth Abnormalities in 4H Leukodystrophy Caused by Variants in POLR3A, POLR3B, and POLR1C. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e660-e674.	3.6	26

#	Article	IF	Citations
19	Six-minute walk test versus two-minute walk test in children with Duchenne muscular dystrophy: Is more time more information?. European Journal of Paediatric Neurology, 2015, 19, 640-646.	1.6	18
20	Neuropsychological Aspects of Childhood Multiple Sclerosis: An Overview. Neuropediatrics, 2012, 43, 176-183.	0.6	16
21	Delayed-Release Dimethyl Fumarate Safety and Efficacy in Pediatric Patients With Relapsing-Remitting Multiple Sclerosis. Frontiers in Neurology, 2020, 11, 606418.	2.4	16
22	Newborn Screening for SMA â€“ Can a Wait-and-See Strategy be Responsibly Justified in Patients With Four SMN2 Copies?. Journal of Neuromuscular Diseases, 2022, 9, 597-605.	2.6	16
23	A homozygous splice variant in AP4S1 mimicking neurodegeneration with brain iron accumulation. Movement Disorders, 2017, 32, 797-799.	3.9	14
24	De novo stop-loss variants in CLDN11 cause hypomyelinating leukodystrophy. Brain, 2021, 144, 411-419.	7.6	12
25	Spinal Muscular Atrophy â€“ Is Newborn Screening Too Late for Children with Two SMN2 Copies?. Journal of Neuromuscular Diseases, 2022, 9, 389-396.	2.6	10
26	Jumping Mechanography as a Complementary Testing Tool for Motor Function in Children with Hereditary Motor and Sensory Neuropathy. Neuropediatrics, 2017, 48, 420-425.	0.6	4
27	Quantitative Motion Measurements Based on Markerless 3D Full-Body Tracking in Children with SMA Highly Correlate with Standardized Motor Assessments. Journal of Neuromuscular Diseases, 2022, 9, 121-128.	2.6	3
28	Clinical and magnetic resonance imaging features of children, adolescents, and adults with a clinically isolated syndrome. European Journal of Paediatric Neurology, 2018, 22, 1087-1094.	1.6	2
29	Is Exercise-Induced Fatigue a Problem in Children with Duchenne Muscular Dystrophy?. Neuropediatrics, 2020, 51, 342-348.	0.6	1
30	Jumping Mechanography is a Suitable Complementary Method to Assess Motor Function in Ambulatory Boys with Duchenne Muscular Dystrophy. Neuropediatrics, 2021, 52, 455-461.	0.6	1
31	Intelligence Quotient and Cognitive Fatigue are Independent Predictors of Cognitive Deficit in Pediatric MS Patients. , 2019, 50, .		0
32	Failure of Expected Brain Growth in Children with ADEM. , 2019, 50, .		0