Annette Hackenberg

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/9924862/publications.pdf

Version: 2024-02-01

40 papers	1,229 citations	567281 15 h-index	395702 33 g-index
41 all docs	41 docs citations	41 times ranked	2250 citing authors

#	Article	IF	Citations
1	Blood DNA methylation provides an accurate biomarker of KMT2B-related dystonia and predicts onset. Brain, 2022, 145, 644-654.	7.6	18
2	Long-Term Efficacy of T3 Analogue Triac in Children and Adults With MCT8 Deficiency: A Real-Life Retrospective Cohort Study. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e1136-e1147.	3.6	15
3	Homozygosity for a Novel DOCK7 Variant Due to Segmental Uniparental Isodisomy of Chromosome 1 Associated with Early Infantile Epileptic Encephalopathy (EIEE) and Cortical Visual Impairment. International Journal of Molecular Sciences, 2022, 23, 7382.	4.1	2
4	Preoperative neurodevelopment of children with moyamoya angiopathy. Developmental Medicine and Child Neurology, 2021, 63, 218-225.	2.1	4
5	<scp>MED27</scp> Variants Cause Developmental Delay, Dystonia, and Cerebellar Hypoplasia. Annals of Neurology, 2021, 89, 828-833.	5.3	14
6	Multiple Sklerose bei Kindern und Jugendlichen â€“ Neue Therapien und wichtige Differentialdiagnosen. Paediatrica, 2021, 32, .	0.1	0
7	SclÃ©rose en plaques de lâ€™enfant et de lâ€™adolescent â€“ nouvelles thÃ©rapies et diagnostics diffÃ©rentiels importants. Paediatrica, 2021, 32, .	0.0	0
8	Management of Acute Demyelinating Attacks in the Pediatric Population: A Swiss Consensus Statement. Clinical and Translational Neuroscience, 2021, 5, 17.	0.9	1
9	Preoperative clinical symptomatology and stroke burden in pediatric moyamoya angiopathy: Defining associated risk variables. European Journal of Paediatric Neurology, 2021, 35, 130-136.	1.6	8
10	Management of Acute Demyelinating Attacks in the Pediatric Population: A Swiss Consensus Statement. Neuropediatrics, 2021, 52, .	0.6	0
11	Impact of stroke volume on motor outcome in neonatal arterial ischemic stroke. European Journal of Paediatric Neurology, 2020, 25, 97-105.	1.6	21
12	Monogenic variants in dystonia: an exome-wide sequencing study. Lancet Neurology, The, 2020, 19, 908-918.	10.2	139
13	Disease characteristics of MCT8 deficiency: an international, retrospective, multicentre cohort study. Lancet Diabetes and Endocrinology,the, 2020, 8, 594-605.	11.4	50
14	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
15	Risk Factors for Postprocedural Arterial Ischemic Stroke in Children With Cardiac Disease. Stroke, 2020, 51, e242-e245.	2.0	7
16	High association of MOG-IgG antibodies in children with bilateral optic neuritis. European Journal of Paediatric Neurology, 2020, 27, 86-93.	1.6	22
17	Health related quality of life and manual ability 5 years after neonatal ischemic stroke. European Journal of Paediatric Neurology, 2019, 23, 716-722.	1.6	15
18	Further corroboration of distinct functional features in SCN2A variants causing intellectual disability or epileptic phenotypes. Molecular Medicine, 2019, 25, 6.	4.4	42

#	Article	IF	Citations
19	The role of recessive inheritance in early-onset epileptic encephalopathies: a combined whole-exome sequencing and copy number study. European Journal of Human Genetics, 2019, 27, 408-421.	2.8	52
20	Feasibility, safety, and outcome of recanalization treatment in childhood stroke. Annals of Neurology, 2018, 83, 1125-1132.	5.3	73
21	Cutis laxa, exocrine pancreatic insufficiency and altered cellular metabolomics as additional symptoms in a new patient with ATP6AP1-CDG. Molecular Genetics and Metabolism, 2018, 123, 364-374.	1.1	23
22	Case report. Medicine (United States), 2018, 97, e11490.	1.0	1
23	Infantile Basal Ganglia Stroke after Mild Head Trauma Associated with Mineralizing Angiopathy of Lenticulostriate Arteries: An Under Recognized Entity. Neuropediatrics, 2018, 49, 262-268.	0.6	8
24	Clinical and functional characterization of two novel ZBTB20 mutations causing Primrose syndrome. Human Mutation, 2018, 39, 959-964.	2.5	11
25	Torcular Pseudomass. Neuropediatrics, 2018, 49, 225-226.	0.6	2
26	Low Voice, Spasmodic Dysphonia, and Hand Dystonia as Clinical Clues for KMT2B-Associated Early-Onset Dystonia. Neuropediatrics, 2018, 49, 356-356.	0.6	6
27	Horizontal Gaze Palsy in Two Brothers with Compound Heterozygous ROBO3 Gene Mutations. Neuropediatrics, 2017, 48, 057-058.	0.6	4
28	Ommaya reservoir â€œoffâ€dutyâ€•causing major lateâ€onset complications in a child with medulloblastoma. Pediatric Blood and Cancer, 2017, 64, e26384.	1.5	1
29	Tubulin-related cerebellar dysplasia: definition of a distinct pattern of cerebellar malformation. European Radiology, 2017, 27, 5080-5092.	4.5	36
30	Inflammatory markers in pediatric stroke: An attempt to better understanding the pathophysiology. European Journal of Paediatric Neurology, 2016, 20, 252-260.	1.6	23
31	Intrathecal Anti-GalC Antibodies in Bickerstaff Brain Stem Encephalitis. Neuropediatrics, 2015, 46, e1-e1.	0.6	0
32	Severe childhood Guillainâ€BarrÃ© syndrome associated with Mycoplasma pneumoniae infection: a case series. Journal of the Peripheral Nervous System, 2015, 20, 72-78.	3.1	17
33	Intrathecal Anti-GalC Antibodies in Bickerstaff Brain Stem Encephalitis. Neuropediatrics, 2015, 46, 428-430.	0.6	6
34	Long-term outcome after arterial ischemic stroke in children and young adults. Neurology, 2015, 84, 1941-1947.	1.1	117
35	Factors affecting cognitive outcome in early pediatric stroke. Neurology, 2014, 82, 784-792.	1.1	99
36	Infantile Epileptic Encephalopathy, Transient Choreoathetotic Movements, and Hypersomnia due to a De Novo Missense Mutation in the SCN2A Gene. Neuropediatrics, 2014, 45, 261-264.	0.6	30

#	Article	IF	Citations
37	Mycoplasma pneumoniae Intrathecal Antibody Responses in Bickerstaff Brain Stem Encephalitis. Neuropediatrics, 2014, 45, 061-063.	0.6	14
38	Fatal Outcome of Rhino-orbital-cerebral Mucormycosis Due to Bilateral Internal Carotid Occlusion in a Child After Hematopoietic Stem Cell Transplantation. Pediatric Infectious Disease Journal, 2013, 32, 1149-1150.	2.0	10
39	Infection-Triggered Familial or Recurrent Cases of Acute Necrotizing Encephalopathy Caused by Mutations in a Component of the Nuclear Pore, RANBP2. American Journal of Human Genetics, 2009, 84, 44-51.	6.2	291
40	Neuroimaging in cerebellar ataxia in childhood: A review. Journal of Neuroimaging, 0, , .	2.0	0