

Annette Hackenberg

List of Publications by Year in descending order

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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	Infection-Triggered Familial or Recurrent Cases of Acute Necrotizing Encephalopathy Caused by Mutations in a Component of the Nuclear Pore, RANBP2. <i>American Journal of Human Genetics</i> , 2009, 84, 44-51.	6.2	291
2	Monogenic variants in dystonia: an exome-wide sequencing study. <i>Lancet Neurology</i> , The, 2020, 19, 908-918.	10.2	139
3	Long-term outcome after arterial ischemic stroke in children and young adults. <i>Neurology</i> , 2015, 84, 1941-1947.	1.1	117
4	Factors affecting cognitive outcome in early pediatric stroke. <i>Neurology</i> , 2014, 82, 784-792.	1.1	99
5	Feasibility, safety, and outcome of recanalization treatment in childhood stroke. <i>Annals of Neurology</i> , 2018, 83, 1125-1132.	5.3	73
6	The role of recessive inheritance in early-onset epileptic encephalopathies: a combined whole-exome sequencing and copy number study. <i>European Journal of Human Genetics</i> , 2019, 27, 408-421.	2.8	52
7	Disease characteristics of MCT8 deficiency: an international, retrospective, multicentre cohort study. <i>Lancet Diabetes and Endocrinology</i> , the, 2020, 8, 594-605.	11.4	50
8	Cerebrospinal fluid findings in patients with myelin oligodendrocyte glycoprotein (MOG) antibodies. Part 2: Results from 108 lumbar punctures in 80 pediatric patients. <i>Journal of Neuroinflammation</i> , 2020, 17, 262.	7.2	44
9	Further corroboration of distinct functional features in SCN2A variants causing intellectual disability or epileptic phenotypes. <i>Molecular Medicine</i> , 2019, 25, 6.	4.4	42
10	Tubulin-related cerebellar dysplasia: definition of a distinct pattern of cerebellar malformation. <i>European Radiology</i> , 2017, 27, 5080-5092.	4.5	36
11	Infantile Epileptic Encephalopathy, Transient Choreoathetotic Movements, and Hypersomnia due to a De Novo Missense Mutation in the SCN2A Gene. <i>Neuropediatrics</i> , 2014, 45, 261-264.	0.6	30
12	Inflammatory markers in pediatric stroke: An attempt to better understanding the pathophysiology. <i>European Journal of Paediatric Neurology</i> , 2016, 20, 252-260.	1.6	23
13	Cutis laxa, exocrine pancreatic insufficiency and altered cellular metabolomics as additional symptoms in a new patient with ATP6AP1-CDG. <i>Molecular Genetics and Metabolism</i> , 2018, 123, 364-374.	1.1	23
14	High association of MOG-IgG antibodies in children with bilateral optic neuritis. <i>European Journal of Paediatric Neurology</i> , 2020, 27, 86-93.	1.6	22
15	Impact of stroke volume on motor outcome in neonatal arterial ischemic stroke. <i>European Journal of Paediatric Neurology</i> , 2020, 25, 97-105.	1.6	21
16	Blood DNA methylation provides an accurate biomarker of <i>KMT2B</i> -related dystonia and predicts onset. <i>Brain</i> , 2022, 145, 644-654.	7.6	18
17	Severe childhood Guillain-Barré syndrome associated with <i>Mycoplasma pneumoniae</i> infection: a case series. <i>Journal of the Peripheral Nervous System</i> , 2015, 20, 72-78.	3.1	17
18	Health related quality of life and manual ability 5 years after neonatal ischemic stroke. <i>European Journal of Paediatric Neurology</i> , 2019, 23, 716-722.	1.6	15

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19	Long-Term Efficacy of T3 Analogue Triac in Children and Adults With MCT8 Deficiency: A Real-Life Retrospective Cohort Study. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, e1136-e1147.	3.6	15
20	<i>Mycoplasma pneumoniae</i> Intrathecal Antibody Responses in Bickerstaff Brain Stem Encephalitis. <i>Neuropediatrics</i> , 2014, 45, 061-063.	0.6	14
21	<sc><i>MED27</i></sc> Variants Cause Developmental Delay, Dystonia, and Cerebellar Hypoplasia. <i>Annals of Neurology</i> , 2021, 89, 828-833.	5.3	14
22	Clinical and functional characterization of two novel <i>ZBTB20</i> mutations causing Primrose syndrome. <i>Human Mutation</i> , 2018, 39, 959-964.	2.5	11
23	Fatal Outcome of Rhino-orbital-cerebral Mucormycosis Due to Bilateral Internal Carotid Occlusion in a Child After Hematopoietic Stem Cell Transplantation. <i>Pediatric Infectious Disease Journal</i> , 2013, 32, 1149-1150.	2.0	10
24	Infantile Basal Ganglia Stroke after Mild Head Trauma Associated with Mineralizing Angiopathy of Lenticulostriate Arteries: An Under Recognized Entity. <i>Neuropediatrics</i> , 2018, 49, 262-268.	0.6	8
25	Preoperative clinical symptomatology and stroke burden in pediatric moyamoya angiopathy: Defining associated risk variables. <i>European Journal of Paediatric Neurology</i> , 2021, 35, 130-136.	1.6	8
26	Risk Factors for Postprocedural Arterial Ischemic Stroke in Children With Cardiac Disease. <i>Stroke</i> , 2020, 51, e242-e245.	2.0	7
27	Intrathecal Anti-GalC Antibodies in Bickerstaff Brain Stem Encephalitis. <i>Neuropediatrics</i> , 2015, 46, 428-430.	0.6	6
28	Low Voice, Spasmodic Dysphonia, and Hand Dystonia as Clinical Clues for KMT2B-Associated Early-Onset Dystonia. <i>Neuropediatrics</i> , 2018, 49, 356-356.	0.6	6
29	Horizontal Gaze Palsy in Two Brothers with Compound Heterozygous ROBO3 Gene Mutations. <i>Neuropediatrics</i> , 2017, 48, 057-058.	0.6	4
30	Preoperative neurodevelopment of children with moyamoya angiopathy. <i>Developmental Medicine and Child Neurology</i> , 2021, 63, 218-225.	2.1	4
31	Torcular Pseudomass. <i>Neuropediatrics</i> , 2018, 49, 225-226.	0.6	2
32	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. <i>International Journal of Molecular Sciences</i> , 2022, 23, 7382.	4.1	2
33	Ommaya reservoir â€œoffâ€dutyâ€ causing major lateâ€onset complications in a child with medulloblastoma. <i>Pediatric Blood and Cancer</i> , 2017, 64, e26384.	1.5	1
34	Case report. <i>Medicine (United States)</i> , 2018, 97, e11490.	1.0	1
35	Management of Acute Demyelinating Attacks in the Pediatric Population: A Swiss Consensus Statement. <i>Clinical and Translational Neuroscience</i> , 2021, 5, 17.	0.9	1
36	Intrathecal Anti-GalC Antibodies in Bickerstaff Brain Stem Encephalitis. <i>Neuropediatrics</i> , 2015, 46, e1-e1.	0.6	0

#	ARTICLE	IF	CITATIONS
37	Multiple Sklerose bei Kindern und Jugendlichen â€“ Neue Therapien und wichtige Differentialdiagnosen. Paediatrica, 2021, 32, .	0.1	0
38	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
39	Management of Acute Demyelinating Attacks in the Pediatric Population: A Swiss Consensus Statement. Neuropediatrics, 2021, 52, .	0.6	0
40	Neuroimaging in cerebellar ataxia in childhood: A review. Journal of Neuroimaging, 0, , .	2.0	0