Christian M Korff

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/3989492/publications.pdf

Version: 2024-02-01

91	2,607	230014	242451
papers	citations	27	47
		h-index	g-index

92	92	92	4313
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Childhood-Onset Movement Disorders Can Mask a Primary Immunodeficiency: 6 Cases of Classical Ataxia-Telangiectasia and Variant Forms. Frontiers in Immunology, 2022, 13, 791522.	2.2	4
2	Pyridoxine or pyridoxalâ€5â€phosphate treatment for seizures in glycosylphosphatidylinositol deficiency: A cohort study. Developmental Medicine and Child Neurology, 2022, 64, 789-798.	1.1	6
3	A Review of Targeted Therapies for Monogenic Epilepsy Syndromes. Frontiers in Neurology, 2022, 13, 829116.	1.1	15
4	Selected Abstracts of the 48th Annual Meeting of the SENP (SociÃ©tÃ© EuropÃ©enne de Neurologie) Tj ETQq0 0 0 rgBT /Overlock 10	0.3	0
5	Natural History Study of STXBP1-Developmental and Epileptic Encephalopathy Into Adulthood. Neurology, 2022, 99, .	1.5	23
6	NEXMIF encephalopathy: an X-linked disorder with male and female phenotypic patterns. Genetics in Medicine, 2021, 23, 363-373.	1.1	28
7	Ketogenic diet treatment in diffuse intrinsic pontine glioma in children: Retrospective analysis of feasibility, safety, and survival data. Cancer Reports, 2021, 4, e1383.	0.6	10
8	<scp>ZMYND11</scp> variants are a novel cause of centrotemporal and generalised epilepsies with neurodevelopmental disorder. Clinical Genetics, 2021, 100, 412-429.	1.0	5
9	Structural brain abnormalities in epilepsy with myoclonic atonic seizures. Epilepsy Research, 2021, 177, 106771.	0.8	2
10	The phenotypic spectrum of Xâ€linked, infantile onset ALG13â€related developmental and epileptic encephalopathy. Epilepsia, 2021, 62, 325-334.	2.6	10
11	Localizing non-epileptiform abnormal brain function in children using high density EEG: Electric Source Imaging of focal slowing. Epilepsy Research, 2020, 159, 106245.	0.8	12
12	Status Epilepticus in Children. Journal of Clinical Neurophysiology, 2020, 37, 429-433.	0.9	10
13	Lessons learned from 40 novel PIGA patients and a review of the literature. Epilepsia, 2020, 61, 1142-1155.	2.6	32
14	Acute monophasic erythromelalgia pain in five children diagnosed as small-fiber neuropathy. European Journal of Paediatric Neurology, 2020, 28, 198-204.	0.7	11
15	SCN8A heterozygous variants are associated with anoxicâ€epileptic seizures. American Journal of Medical Genetics, Part A, 2020, 182, 1209-1216.	0.7	7
16	DIPG-25. KETOGENIC DIET IN DIFFUSE INTRINSIC PONTINE GLIOMA IN CHILDREN: A RETROSPECTIVE STUDY INVESTIGATING THE FEASIBILITY. Neuro-Oncology, 2020, 22, iii291-iii292.	0.6	0
17	Infantile-Onset Paroxysmal Movement Disorder and Episodic Ataxia Associated with a TBC1D24 Mutation. Neuropediatrics, 2019, 50, 308-312.	0.3	12
18	Neurodevelopmental problems of unaccompanied refugee and migrant children: a new challenge for pediatric neurologists. Developmental Medicine and Child Neurology, 2019, 61, 1348-1348.	1.1	2

#	Article	IF	Citations
19	The spectrum of intermediate <scp>SCN</scp>8Aâ€related epilepsy. Epilepsia, 2019, 60, 830-844.	2.6	70
20	Comment on: Ketogenic diet treatment in recurrent diffuse intrinsic pontine glioma in children: A safety and feasibility study. Pediatric Blood and Cancer, 2019, 66, e27664.	0.8	2
21	Circulating neural antibodies in unselected children with new-onset seizures. European Journal of Paediatric Neurology, 2018, 22, 396-403.	0.7	6
22	Focal cortical malformations in children with early infantile epilepsy and <scp>PCDH</scp>19 mutations: case report. Developmental Medicine and Child Neurology, 2018, 60, 100-105.	1.1	56
23	A triad of infantile spasms, nystagmus and a focal tonic seizure. Epileptic Disorders, 2018, 20, 295-300.	0.7	6
24	Why the TimeToStop trial failed to recruit: a survey on antiepileptic drug withdrawal after paediatric epilepsy surgery. Epileptic Disorders, 2018, 20, 374-385.	0.7	12
25	NBEA: Developmental disease gene with early generalized epilepsy phenotypes. Annals of Neurology, 2018, 84, 788-795.	2.8	44
26	Desaturation During Cry in the Neonatal Period. Global Pediatric Health, 2018, 5, 2333794X1876451.	0.3	1
27	Characterization of glycosylphosphatidylinositol biosynthesis defects by clinical features, flow cytometry, and automated image analysis. Genome Medicine, 2018, 10, 3.	3.6	67
28	GLUT1-DS in a Girl with Transitory Abnormal Eye Movements and Seizures Responding to Carbamazepine. Neuropediatrics, 2018, 49, S1-S12.	0.3	0
29	An epidemic of dystonic reactions in central Africa. The Lancet Global Health, 2017, 5, e137-e138.	2.9	20
30	GRIN2Bencephalopathy: novel findings on phenotype, variant clustering, functional consequences and treatment aspects. Journal of Medical Genetics, 2017, 54, 460-470.	1.5	190
31	Genetic and phenotypic dissection of 1q43q44 microdeletion syndrome and neurodevelopmental phenotypes associated with mutations in ZBTB18 and HNRNPU. Human Genetics, 2017, 136, 463-479.	1.8	66
32	Drug-Level Monitoring on Admission for Presurgical Epilepsy Evaluation. European Neurology, 2017, 78, 105-110.	0.6	1
33	SCN1A Does Not Determine Seizure Duration in Children Unaffected by Dravet's Syndrome. Journal of Pediatric Epilepsy, 2017, 06, 111-114.	0.1	0
34	The Immune System in Pediatric Seizures and Epilepsies. Pediatrics, 2017, 140, e20163534.	1.0	26
35	Clinical spectrum and genotypeâ€“phenotype associations of KCNA2-related encephalopathies. Brain, 2017, 140, 2337-2354.	3.7	117
36	SERPINI1 pathogenic variants: An emerging cause of childhoodâ€onset progressive myoclonic epilepsy. American Journal of Medical Genetics, Part A, 2017, 173, 2456-2460.	0.7	13

#	Article	IF	Citations
37	Epileptic networks are strongly connected with and without the effects of interictal discharges. Epilepsia, 2016, 57, 1086-1096.	2.6	36
38	Genetic and neurodevelopmental spectrum ofSYNGAP1-associated intellectual disability and epilepsy. Journal of Medical Genetics, 2016, 53, 511-522.	1.5	135
39	Epileptic spasms in epilepsy with myoclonic-atonic seizures (Doose syndrome). Epileptic Disorders, 2016, 18, 289-296.	0.7	3
40	Effectiveness of antiepileptic therapy in patients with PCDH19 mutations. Seizure: the Journal of the British Epilepsy Association, 2016, 35, 106-110.	0.9	61
41	Pediatric epilepsy surgery: could age be a predictor of outcomes?. Journal of Neurosurgery: Pediatrics, 2016, 18, 235-241.	0.8	24
42	Seizures and Epilepsies due to Channelopathies and Neurotransmitter Receptor Dysfunction: A Parallel between Genetic and Immune Aspects. Molecular Syndromology, 2016, 7, 197-209.	0.3	17
43	Diabetic Striatopathy in Childhood: A Case Report. Pediatrics, 2016, 137, .	1.0	18
44	Combined VIth and VIIth nerve palsy: Consider idiopathic intracranial hypertension!. European Journal of Paediatric Neurology, 2016, 20, 336-338.	0.7	7
45	Riboflavin in cyclic vomiting syndrome: efficacy in three children. European Journal of Pediatrics, 2016, 175, 131-135.	1.3	12
46	Yield of MRI, high-density electric source imaging (HD-ESI), SPECT and PET in epilepsy surgery candidates. Clinical Neurophysiology, 2016, 127, 150-155.	0.7	97
47	Editorial commentary on â€œRefractory absence seizures: An Italian multicenter retrospective studyâ€•by E. Franzoni etÂal.. European Journal of Paediatric Neurology, 2015, 19, 617-618.	0.7	0
48	Epileptic activity is a surrogate for an underlying etiology and stopping the activity has a limited impact on developmental outcome. Epilepsia, 2015, 56, 1477-1481.	2.6	31
49	A child with ictal vocalizations and generalized epilepsy. Epileptic Disorders, 2015, 17, 67-71.	0.7	5
50	Severe phenotypic spectrum of biallelic mutations inPRRT2gene. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 782-785.	0.9	72
51	Widespread intracranial calcifications in the follow-up of a patient with cartilage-hair hypoplasia â€“ Anauxetic dysplasia spectrum disorder: A coincidental finding?. European Journal of Paediatric Neurology, 2015, 19, 367-371.	0.7	0
52	When should clinicians search for GLUT1 deficiency syndrome in childhood generalized epilepsies?. European Journal of Paediatric Neurology, 2015, 19, 170-175.	0.7	9
53	A practical, simple, and useful method of categorizing interictal EEG features in children. Neurology, 2015, 85, 471-478.	1.5	11
54	De novo loss- or gain-of-function mutations in KCNA2 cause epileptic encephalopathy. Nature Genetics, 2015, 47, 393-399.	9.4	224

#	Article	IF	Citations
55	Clinical Variability of GLUT1DS. Pediatric Neurology Briefs, 2015, 29, 14.	0.2	1
56	Vaccinations and Dravet Syndrome. Pediatric Neurology Briefs, 2015, 29, 85.	0.2	1
57	Autoimmunity and inflammation in status epilepticus: from concepts to therapies. Expert Review of Neurotherapeutics, 2014, 14, 1181-1202.	1.4	27
58	Positive Outcome following Early Diagnosis and Treatment of Pyridoxal-5â€²-Phosphate Oxidase Deficiency: A Case Report. Neuropediatrics, 2014, 45, 064-068.	0.3	27
59	When is a child with status epilepticus likely to have Dravet syndrome?. Epilepsy Research, 2014, 108, 740-747.	0.8	16
60	Towards the identification of a genetic basis for <scp>L</scp>andauâ€<scp>K</scp>leffner <scp>s</scp>yndrome. Epilepsia, 2014, 55, 858-865.	2.6	44
61	Epilepsy: Old Syndromes, New Genes. Current Neurology and Neuroscience Reports, 2014, 14, 447.	2.0	17
62	Localization of the epileptogenic tuber with electric source imaging in patients with tuberous sclerosis. Epilepsy Research, 2014, 108, 267-279.	0.8	30
63	Tracking the source of cerebellar epilepsy: Hemifacial seizures associated with cerebellar cortical dysplasia. Epilepsy Research, 2013, 105, 245-249.	0.8	19
64	Hashimoto's encephalopathy: Identification and long-term outcome in children. European Journal of Paediatric Neurology, 2013, 17, 280-287.	0.7	40
65	Epilepsy classification. Current Opinion in Neurology, 2013, 26, 163-167.	1.8	24
66	Reduction of seizure frequency after epilepsy surgery in a patient with <scp>STXBP1</scp> encephalopathy and clinical description of six novel mutation carriers. Epilepsia, 2013, 54, e74-80.	2.6	59
67	Head stereotypies in <scp>STXBP</scp>1 encephalopathy. Developmental Medicine and Child Neurology, 2013, 55, 769-772.	1.1	17
68	Postoperative EEG in Hemimegalencephaly. European Neurology, 2012, 68, 358-360.	0.6	4
69	Anti-NMDA receptor encephalitis: The importance of early diagnosis and aggressive immunotherapy in tumor negative pediatric patients. European Journal of Paediatric Neurology, 2012, 16, 764-765.	0.7	9
70	Ohtahara syndrome or early-onset West syndrome? A case with overlapping features and favorable response to vigabatrin. European Journal of Paediatric Neurology, 2012, 16, 753-757.	0.7	8
71	Partial Rhombencephalosynapsis and Chiari Type II Malformation in a Child: a True Association Supported by DTI Tractography. Cerebellum, 2012, 11, 227-232.	1.4	5
72	Benzodiazepines in the acute management of seizures with autonomic manifestations: Anticipate complications!. Epilepsia, 2011, 52, e156-e159.	2.6	10

#	Article	IF	Citations
73	Encephalitis Associated With Glutamic Acid Decarboxylase Autoantibodies in a Child. Archives of Neurology, 2011, 68, 1065.	4.9	42
74	ABCB1 polymorphisms and neuropsychiatric adverse events in oseltamivir-treated children during influenza H1N1/09 pandemia. Pharmacogenomics, 2011, 12, 1493-1501.	0.6	13
75	Severe childhood encephalopathy with dyskinesia and prolonged cognitive disturbances: evidence for antiâ€Nâ€methylâ€<scp>d</scp>â€aspartate receptor encephalitis. Developmental Medicine and Child Neurology, 2010, 52, e78-82.	1.1	48
76	A case of SUDEP in a patient with Dravet syndrome with SCN1A mutation. Epilepsia, 2010, 51, 1915-1918.	2.6	57
77	Status epilepticus in fragile X syndrome. Epilepsia, 2010, 51, 2470-2473.	2.6	15
78	West Syndrome. , 2010, , 885-889.		0
79	Legends forÂvideo sequences. Epileptic Disorders, 2009, 11, 132-135.	0.7	8
80	Alexander disease: Early presence of cerebral MRI criteria. European Journal of Paediatric Neurology, 2009, 13, 556-558.	0.7	9
81	FOOD POISONING AS A CAUSE OF ACUTE LIVER FAILURE. Pediatric Infectious Disease Journal, 2008, 27, 846-847.	1.1	72
82	Diagnosis and management of nonconvulsive status epilepticus in children. Nature Clinical Practice Neurology, 2007, 3, 505-516.	2.7	30
83	Dravet Syndrome (Severe Myoclonic Epilepsy in Infancy): A Retrospective Study of 16 Patients. Journal of Child Neurology, 2007, 22, 185-194.	0.7	76
84	Cryptogenic Late-Onset Epileptic Spasms or Late Infantile Epileptogenic Encephalopathy?. Epilepsia, 2007, 48, 206-8.	2.6	18
85	Epilepsy Syndromes in Infancy. Pediatric Neurology, 2006, 34, 253-263.	1.0	50
86	Epilepsy syndromes undetermined whether focal or generalized in infants. Epilepsy Research, 2006, 70, 105-109.	0.8	6
87	The clinical-electrographic expression of infantile seizures. Epilepsy Research, 2006, 70, 116-131.	0.8	18
88	Notched Delta, Phenotype, and Angelman Syndrome. Journal of Clinical Neurophysiology, 2005, 22, 238-243.	0.9	37
89	Paroxysmal Events in Infants: Persistent Eye Closure Makes Seizures Unlikely. Pediatrics, 2005, 116, e485-e486.	1.0	10
90	Do generalized tonic-clonic seizures in infancy exist?. Neurology, 2005, 65, 1750-1753.	1.5	67

#	Article	IF	Citations
91	Isolated Absence of Optic Chiasm Revealed by Congenital Nystagmus, MRI and VEPs. Neuropediatrics, 2003, 34, 219-223.	0.3	25