Elaine C Wirrell

List of Publications by Year
in descending order

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Version: 2024-02-01

229 papers	13,287 citations	22099 59 h-index	30010 103 g-index
237 all docs	237 docs citations	237 times ranked	10239 citing authors

#	Article	IF	Citations
1	Dravet Syndrome as an Example of Precision Medicine in Epilepsy. Epilepsy Currents, 2023, 23, 4-7.	0.4	5
2	New and emerging pharmacologic treatments for developmental and epileptic encephalopathies. Current Opinion in Neurology, 2022, 35, 145-154.	1.8	8
3	A tale of two cohorts: Differing outcomes in infantileâ€onset focal epilepsy. Epilepsia, 2022, 63, 950-960.	2.6	6
4	Evaluation of First Seizure and Newly Diagnosed Epilepsy. CONTINUUM Lifelong Learning in Neurology, 2022, 28, 230-260.	0.4	6
5	COVIDâ€19 vaccine in patients with Dravet syndrome: Observations and realâ€world experiences. Epilepsia, 2022, 63, 1778-1786.	2.6	13
6	Safety and efficacy of responsive neurostimulation in the pediatric population: Evidence from institutional review and patient-level meta-analysis. Epilepsy and Behavior, 2022, 129, 108646.	0.9	17
7	Inequities in Therapy for Infantile Spasms: A Call to Action. Annals of Neurology, 2022, 92, 32-44.	2.8	7
8	Safety and efficacy of ganaxolone in patients with CDKL5 deficiency disorder: results from the double-blind phase of a randomised, placebo-controlled, phase 3 trial. Lancet Neurology, The, 2022, 21, 417-427.	4.9	63
9	International League Against Epilepsy classification and definition of epilepsy syndromes with onset in childhood: Position paper by the ILAE Task Force on Nosology and Definitions. Epilepsia, 2022, 63, 1398-1442.	2.6	263
10	Introduction to the epilepsy syndrome papers. Epilepsia, 2022, 63, 1330-1332.	2.6	23
11	Methodology for classification and definition of epilepsy syndromes with list of syndromes: Report of the ILAE Task Force on Nosology and Definitions. Epilepsia, 2022, 63, 1333-1348.	2.6	84
12	International League Against Epilepsy classification and definition of epilepsy syndromes with onset at a variable age: position statement by the ILAE Task Force on Nosology and Definitions. Epilepsia, 2022, 63, 1443-1474.	2.6	81
13	Value of genetic testing for pediatric epilepsy: Driving earlier diagnosis of ceroid lipofuscinosis type 2 Batten disease. Epilepsia, 2022, 63, .	2.6	6
14	ILAE classification and definition of epilepsy syndromes with onset in neonates and infants: Position statement by the ILAE Task Force on Nosology and Definitions. Epilepsia, 2022, 63, 1349-1397.	2.6	237
15	ILAE definition of the Idiopathic Generalized Epilepsy Syndromes: Position statement by the ILAE Task Force on Nosology and Definitions. Epilepsia, 2022, 63, 1475-1499.	2.6	148
16	International consensus on diagnosis and management of Dravet syndrome. Epilepsia, 2022, 63, 1761-1777.	2.6	62
17	Impact of Antiseizure Medications on Appetite and Weight in Children. Paediatric Drugs, 2022, 24, 335-363.	1.3	10
18	The gain of function SCN1A disorder spectrum: novel epilepsy phenotypes and therapeutic implications. Brain, 2022, 145, 3816-3831.	3.7	43

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19	Considerations for determining the efficacy of new antiseizure medications in children age 1Âmonth to younger than 2â€‰years. Epilepsia, 2022, 63, 2664-2670.	2.6	4
20	Proposal to optimize evaluation and treatment of Febrile infectionâ€related epilepsy syndrome (FIRES): A Report from FIRES workshop. Epilepsia Open, 2021, 6, 62-72.	1.3	35
21	The role of new medical treatments for the management of developmental and epileptic encephalopathies: Novel concepts and results. Epilepsia, 2021, 62, 857-873.	2.6	26
22	Results of an international Delphi consensus in epilepsy with myoclonic atonic seizures/ Doose syndrome. Seizure: the Journal of the British Epilepsy Association, 2021, 85, 12-18.	0.9	9
23	Add-on Cannabidiol Treatment for Drug-Resistant Seizures in Tuberous Sclerosis Complex. JAMA Neurology, 2021, 78, 285.	4.5	139
24	Time to onset of cannabidiol (CBD) treatment effect in Lennoxâ€“Gastaut syndrome: Analysis from two randomized controlled trials. Epilepsia, 2021, 62, 1130-1140.	2.6	20
25	Management of CLN1 Disease: International Clinical Consensus. Pediatric Neurology, 2021, 120, 38-51.	1.0	10
26	Comparative Effectiveness of Initial Treatment for Infantile Spasms in a Contemporary US Cohort. Neurology, 2021, 97, .	1.5	19
27	Cognitive outcome in children with infantile spasms using a standardized treatment protocol. A five-year longitudinal study. Seizure: the Journal of the British Epilepsy Association, 2021, 89, 73-80.	0.9	6
28	Discontinuing Antiseizure Medication in Neonates With Acute Symptomatic Seizuresâ€”Primum non nocere. JAMA Neurology, 2021, 78, 797.	4.5	0
29	Improved everyday executive functioning following profound reduction in seizure frequency with fenfluramine: Analysis from a phase 3 long-term extension study in children/young adults with Dravet syndrome. Epilepsy and Behavior, 2021, 121, 108024.	0.9	31
30	Dravet syndrome: A quick transition guide for the adult neurologist. Epilepsy Research, 2021, 177, 106743.	0.8	11
31	Monogenic Epilepsies. Neurology, 2021, 97, 817-831.	1.5	38
32	Treatment with fenfluramine in patients with Dravet syndrome has no long-term effects on weight and growth. Epilepsy and Behavior, 2021, 122, 108212.	0.9	9
33	DRAVET ENGAGE. Parent caregivers of children with Dravet syndrome: Perspectives, needs, and opportunities for clinical research. Epilepsy and Behavior, 2021, 122, 108198.	0.9	17
34	What is the optimal duration for vigabatrin monotherapy in patients with infantile spasms: 6 months or longer?. Seizure: the Journal of the British Epilepsy Association, 2021, 91, 503-506.	0.9	0
35	Centromedian thalamic nucleus with or without anterior thalamic nucleus deep brain stimulation for epilepsy in children and adults: A retrospective case series. Seizure: the Journal of the British Epilepsy Association, 2021, 84, 101-107.	0.9	39
36	Impact of fenfluramine on the expected SUDEP mortality rates in patients with Dravet syndrome. Seizure: the Journal of the British Epilepsy Association, 2021, 93, 154-159.	0.9	41

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37	PURA- Related Developmental and Epileptic Encephalopathy. Neurology: Genetics, 2021, 7, e613.	0.9	15
38	Rational Approach to Children with Drug-Resistant Epilepsy. Journal of International Child Neurology Association, 2021, 1, .	0.0	1
39	Fenfluramine for Treatment-Resistant Seizures in Patients With Dravet Syndrome Receiving Stiripentol-Inclusive Regimens. JAMA Neurology, 2020, 77, 300.	4.5	152
40	Care Delivery for Children With Epilepsy During the COVID-19 Pandemic: An International Survey of Clinicians. Journal of Child Neurology, 2020, 35, 924-933.	0.7	48
41	Selfâ€injurious and suicidal behavior in young adults, teens, and children with epilepsy: A populationâ€based study. Epilepsia, 2020, 61, 1919-1930.	2.6	13
42	An accelerated shift in the use of remote systems in epilepsy due to the COVID-19 pandemic. Epilepsy and Behavior, 2020, 112, 107376.	0.9	29
43	Anakinra usage in febrile infection related epilepsy syndrome: an international cohort. Annals of Clinical and Translational Neurology, 2020, 7, 2467-2474.	1.7	80
44	Seizures in Children. Pediatrics in Review, 2020, 41, 321-347.	0.2	57
45	Extinguishing Febrile Infection-Related Epilepsy Syndrome: Pipe Dream or Reality?. Seminars in Neurology, 2020, 40, 263-272.	0.5	10
46	Post-placement Lead Deformation Secondary to Cerebrospinal Fluid Loss in Transventricular Trajectory During Responsive Neurostimulation Surgery. Cureus, 2020, 12, e6823.	0.2	1
47	The co-occurrence of epilepsy and autism: A systematic review. Epilepsy and Behavior, 2019, 98, 238-248.	0.9	120
48	Classifications of Seizures and Epilepsies. , 2019, , 58-76.		2
49	The Genetic Landscape of Epilepsy of Infancy with Migrating Focal Seizures. Annals of Neurology, 2019, 86, 821-831.	2.8	96
50	Recent Advances in the Drug Treatment of Dravet Syndrome. CNS Drugs, 2019, 33, 867-881.	2.7	64
51	Correspondence Reply to Madaan etÂal.. Pediatric Neurology, 2019, 95, 92-93.	1.0	0
52	Seizure Rescue Medication Use among US Pediatric Epilepsy Providers: A Survey of the Pediatric Epilepsy Research Consortium. Journal of Pediatrics, 2019, 212, 111-116.	0.9	11
53	Immediate outcomes in early life epilepsy: A contemporary account. Epilepsy and Behavior, 2019, 97, 44-50.	0.9	27
54	The Ketogenic and Modified Atkins Diet Therapy for Children With Refractory Epilepsy of Genetic Etiology. Pediatric Neurology, 2019, 94, 32-37.	1.0	24

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55	How can transition to adult care be best orchestrated for adolescents with epilepsy?. Epilepsy and Behavior, 2019, 93, 138-147.	0.9	39
56	Impact of Prior Authorization of Antiepileptic Drugs in Children With Epilepsy. Pediatric Neurology, 2018, 83, 38-41.	1.0	14
57	Ketogenic Diet Therapy in Infants: Efficacy and Tolerability. Pediatric Neurology, 2018, 82, 13-18.	1.0	29
58	Comparative Effectiveness of Levetiracetam vs Phenobarbital for Infantile Epilepsy. JAMA Pediatrics, 2018, 172, 352.	3.3	30
59	How do we diagnose and treat epilepsy with myoclonic-atonic seizures (Doose syndrome)? Results of the Pediatric Epilepsy Research Consortium survey. Epilepsy Research, 2018, 144, 14-19.	0.8	14
60	Treatment Strategies for Dravet Syndrome. CNS Drugs, 2018, 32, 335-350.	2.7	43
61	Systematic review of the screening, diagnosis, and management of <scp>ADHD</scp> in children with epilepsy. Consensus paper of the Task Force on Comorbidities of the <scp>ILAE</scp> Pediatric Commission. Epilepsia, 2018, 59, 1867-1880.	2.6	68
62	Challenges in managing epilepsy associated with focal cortical dysplasia in children. Epilepsy Research, 2018, 145, 1-17.	0.8	25
63	Optimal clinical management of children receiving dietary therapies for epilepsy: Updated recommendations of the International Ketogenic Diet Study Group. Epilepsia Open, 2018, 3, 175-192.	1.3	412
64	Effect of Cannabidiol on Drop Seizures in the Lennoxâ€“Gastaut Syndrome. New England Journal of Medicine, 2018, 378, 1888-1897.	13.9	682
65	A Novel Phenotype in a Previously Described Epilepsyâ€”Aphasia Disorder. Seminars in Pediatric Neurology, 2018, 26, 21-24.	1.0	1
66	LGI1 and CASPR2 neurological autoimmunity in children. Annals of Neurology, 2018, 84, 473-480.	2.8	53
67	Jeavons Syndrome: Clinical Features and Response to Treatment. Pediatric Neurology, 2018, 86, 46-51.	1.0	40
68	Neuroimaging of Early Life Epilepsy. Pediatrics, 2018, 142, .	1.0	23
69	De Novo DNM1L Variant in a Teenager With Progressive Paroxysmal Dystonia and Lethal Super-refractory Myoclonic Status Epilepticus. Journal of Child Neurology, 2018, 33, 651-658.	0.7	25
70	Why West? Comparisons of clinical, genetic and molecular features of infants with and without spasms. PLoS ONE, 2018, 13, e0193599.	1.1	28
71	Understanding Death in Children With Epilepsy. Pediatric Neurology, 2017, 70, 7-15.	1.0	29
72	Optimizing the Diagnosis and Management of Dravet Syndrome: Recommendations From a North American Consensus Panel. Pediatric Neurology, 2017, 68, 18-34.e3.	1.0	207

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73	Epidemiology and Outcomes of Arterial Ischemic Stroke in Children: The Canadian Pediatric Ischemic Stroke Registry. Pediatric Neurology, 2017, 69, 58-70.	1.0	213
74	Stiripentol in the Management of Epilepsy. CNS Drugs, 2017, 31, 405-416.	2.7	37
75	Predictive models in the diagnosis and treatment of autoimmune epilepsy. Epilepsia, 2017, 58, 1181-1189.	2.6	120
76	Cognitive and Social Outcomes of Epileptic Encephalopathies. Seminars in Pediatric Neurology, 2017, 24, 264-275.	1.0	17
77	Early-Life Epilepsies and the Emerging Role of Genetic Testing. JAMA Pediatrics, 2017, 171, 863.	3.3	125
78	Not all SCN1A epileptic encephalopathies are Dravet syndrome. Neurology, 2017, 89, 1035-1042.	1.5	97
79	The impact of hypsarrhythmia on infantile spasms treatment response: Observational cohort study from the National Infantile Spasms Consortium. Epilepsia, 2017, 58, 2098-2103.	2.6	55
80	Initial Treatment for Nonsyndromic Early-Life Epilepsy: An Unexpected Consensus. Pediatric Neurology, 2017, 75, 73-79.	1.0	18
81	Prevalence and Risk Factors of Peri-ictal Autonomic Changes in Children With Temporal Lobe Seizures. Pediatric Neurology, 2017, 67, 36-39.	1.0	7
82	Response to treatment in a prospective national infantile spasms cohort. Annals of Neurology, 2016, 79, 475-484.	2.8	182
83	A microRNAâ€328 binding site inPAX6is associated with centrotemporal spikes of rolandic epilepsy. Annals of Clinical and Translational Neurology, 2016, 3, 512-522.	1.7	27
84	False ictal-appearing EEG from a frontal sinus pseudomeningocele. Neurology, 2016, 87, 2600-2601.	1.5	2
85	Focal photoparoxysmal response in the Heidenhain variant of CJD. Neurology, 2016, 86, 1647-1648.	1.5	3
86	Patient and caregiver view on seizure detection devices: A survey study. Seizure: the Journal of the British Epilepsy Association, 2016, 41, 179-181.	0.9	40
87	Usefulness of repeat review of head magnetic resonance images during presurgical epilepsy conferences. Epilepsy Research, 2016, 126, 106-108.	0.8	6
88	Response to second treatment after initial failed treatment in a multicenter prospective infantile spasms cohort. Epilepsia, 2016, 57, 1834-1842.	2.6	58
89	Cognitive and neurodevelopmental comorbidities in paediatric epilepsy. Nature Reviews Neurology, 2016, 12, 465-476.	4.9	152
90	Febrile infectionâ€related epilepsy syndrome treated with anakinra. Annals of Neurology, 2016, 80, 939-945.	2.8	208

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91	Mortality in Dravet syndrome. Epilepsy Research, 2016, 128, 43-47.	0.8	218
92	Treatment of Dravet Syndrome. Canadian Journal of Neurological Sciences, 2016, 43, S13-S18.	0.3	84
93	Do Patients Require Inpatient Admission to Receive Adrenocorticotropic Hormone (ACTH)? A Survey of US-Based Prescribers. Journal of Child Neurology, 2016, 31, 164-169.	0.7	4
94	Routine vs extended outpatient EEG for the detection of interictal epileptiform discharges. Neurology, 2016, 86, 1524-1530.	1.5	48
95	Pharmacotherapy for Dravet Syndrome. Paediatric Drugs, 2016, 18, 197-208.	1.3	64
96	Infantile, Childhood, and Adolescent Epilepsies. CONTINUUM Lifelong Learning in Neurology, 2016, 22, 60-93.	0.4	9
97	Does treatment have an impact on incidence and risk factors for autism spectrum disorders in children with infantile spasms?. Epilepsia, 2015, 56, 856-863.	2.6	23
98	Acetazolamide for electrical status epilepticus in slowâ€wave sleep. Epilepsia, 2015, 56, e134-8.	2.6	23
99	How should children with West syndrome be efficiently and accurately investigated? Results from the National Infantile Spasms Consortium. Epilepsia, 2015, 56, 617-625.	2.6	130
100	Seizure Freedom in Children With Pathology-Confirmed Focal Cortical Dysplasia. Pediatric Neurology, 2015, 53, 513-518.	1.0	16
101	Predictors of Length of Stay in Children Admitted forÂPresurgicalÂEvaluation for Epilepsy Surgery. Pediatric Neurology, 2015, 53, 207-210.	1.0	8
102	Repeat resective surgery in complex pediatric refractory epilepsy: lessons learned. Journal of Neurosurgery: Pediatrics, 2015, 16, 94-100.	0.8	21
103	Yield and Predictors of Epilepsy Surgery Candidacy in Children Admitted for Surgical Evaluation. Pediatric Neurology, 2015, 53, 58-64.	1.0	5
104	CNKSR2 deletions: A novel cause of Xâ€linked intellectual disability and seizures. American Journal of Medical Genetics, Part A, 2015, 167, 1668-1670.	0.7	23
105	Patient considerations in the management of focal seizures in children and adolescents. Adolescent Health, Medicine and Therapeutics, 2014, 5, 49.	0.7	3
106	Transition issues for benign epilepsy with centrotemporal spikes, nonlesional focal epilepsy in otherwise normal children, childhood absence epilepsy, and juvenile myoclonic epilepsy. Epilepsia, 2014, 55, 16-20.	2.6	22
107	Factors contributing to the yield of asymmetric bilateral implantation of intracranial electrodes. Epilepsia, 2014, 55, 1620-1625.	2.6	0
108	Sleep Abnormalities in Children With Dravet Syndrome. Pediatric Neurology, 2014, 50, 474-478.	1.0	27

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109	Early-Onset Focal Seizures and Spasmsâ€"Is it Surgical?. Seminars in Pediatric Neurology, 2014, 21, 101-103.	1.0	2
110	Predictors of Recording an Event During Prolonged Inpatient Video Electroencephalogram Monitoring in Children. Pediatric Neurology, 2014, 50, 458-463.	1.0	7
111	Intractable Epilepsy, Hemispheric Malformation, and Generalized Electroencephalography Abnormalities. Seminars in Pediatric Neurology, 2014, 21, 73-75.	1.0	3
112	Chronic subthreshold subdural cortical stimulation for the treatment of focal epilepsy originating from eloquent cortex. Epilepsia, 2014, 55, e18-21.	2.6	47
113	DPPX potassium channel antibody. Neurology, 2014, 83, 1797-1803.	1.5	255
114	Editorial Comment: The Janus of Rasmussen Encephalitis: Never a Friendly Face. Seminars in Pediatric Neurology, 2014, 21, 137-138.	1.0	0
115	Bizarre Semiology and Medically Intractable Seizures. Seminars in Pediatric Neurology, 2014, 21, 154-159.	1.0	0
116	Early Efficacy of the Ketogenic Diet Is Not Affected by Initial Body Mass Index Percentile. Pediatric Neurology, 2014, 50, 469-473.	1.0	3
117	Outcome of intracranial electroencephalography monitoring and surgery in magnetic resonance imaging-negative temporal lobe epilepsy. Epilepsy Research, 2014, 108, 937-944.	0.8	17
118	Seizure outcome after AED failure in pediatric focal epilepsy: Impact of underlying etiology. Epilepsy and Behavior, 2014, 34, 20-24.	0.9	15
119	Long-term Outcomes After Nonlesional Extratemporal Lobe Epilepsy Surgery. JAMA Neurology, 2013, 70, 1003.	4.5	145
120	CDKL5 and ARX Mutations in Males With Early-Onset Epilepsy. Pediatric Neurology, 2013, 48, 367-377.	1.0	53
121	Autonomic epileptic seizures, autonomic effects of seizures, and SUDEP. Epilepsy and Behavior, 2013, 26, 375-385.	0.9	78
122	Role of the sodium channel <scp>SCN</scp>9A in genetic epilepsy with febrile seizures plus and Dravet syndrome. Epilepsia, 2013, 54, e122-6.	2.6	62
123	Predicting pharmacoresistance in pediatric epilepsy. Epilepsia, 2013, 54, 19-22.	2.6	116
124	Stiripentol in <scp>D</scp>ravet syndrome: Results of a retrospective <scp>U</scp>.<scp>S</scp>. study. Epilepsia, 2013, 54, 1595-1604.	2.6	84
125	Mortality Risks in New-Onset Childhood Epilepsy. Pediatrics, 2013, 132, 124-131.	1.0	141
126	Novel de novo SCN2A Mutation in a Child With Migrating Focal Seizures of Infancy. Pediatric Neurology, 2013, 49, 486-488.	1.0	58

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127	Early onset epilepsy is associated with increased mortality: A population-based study. Epilepsy Research, 2013, 105, 410-414.	0.8	30
128	Characteristics of postictal generalized EEG suppression in children. Epilepsy Research, 2013, 106, 123-127.	0.8	68
129	What predicts enduring intractability in children who appear medically intractable in the first 2Âyears after diagnosis?. Epilepsia, 2013, 54, 1056-1064.	2.6	49
130	Ketogenic Diet. Canadian Journal of Neurological Sciences, 2013, 40, 158-167.	0.3	61
131	Seizure Outcomes After Corpus Callosotomy for Drop Attacks. Neurosurgery, 2013, 73, 993-1000.	0.6	53
132	DIETARY TREATMENT OF EPILEPSY. PRACTICAL IMPLEMENTATION OF KETOGENIC THERAPY. 2012. Edited by Elizabeth Neal. Published by Wiley-Blackwell. 241 pages. C$80 approx.. Canadian Journal of Neurological Sciences, 2013, 40, 440-440.	0.3	1
133	Autoimmune Epilepsy. Archives of Neurology, 2012, 69, 582.	4.9	324
134	The Cessation of Continuous Spike Wave in Slow-Wave Sleep Following a Temporal Lobectomy. Journal of Child Neurology, 2012, 27, 113-116.	0.7	8
135	Epilepsy in Childrenâ€"When Should We Think Neurometabolic Disease?. Journal of Child Neurology, 2012, 27, 663-671.	0.7	13
136	Surgical Outcomes for Intractable Epilepsy in Children With Epileptic Spasms. Journal of Child Neurology, 2012, 27, 713-720.	0.7	20
137	Temporal Lobe Epilepsy in Children. Epilepsy Research & Treatment, 2012, 2012, 1-16.	1.4	21
138	Autoimmune Encephalopathies and Epilepsies in Children and Teenagers. Canadian Journal of Neurological Sciences, 2012, 39, 134-144.	0.3	20
139	What's New in the Medical Management of Pediatric Epilepsy?. Canadian Journal of Neurological Sciences, 2012, 39, S30-S34.	0.3	0
140	Predictors and course of medically intractable epilepsy in young children presenting before 36â€ƒmonths of age: A retrospective, populationâ€based study. Epilepsia, 2012, 53, 1563-1569.	2.6	97
141	A randomized controlled trial of flunarizine as addâ€on therapy and effect on cognitive outcome in children with infantile spasms. Epilepsia, 2012, 53, 1570-1576.	2.6	17
142	Historic, Clinical, and Prognostic Features of Epileptic Encephalopathies Caused by CDKL5 Mutations. Pediatric Neurology, 2012, 46, 101-105.	1.0	55
143	Epilepsyâ€related mortality is low in children: A 30â€year populationâ€based study in Olmsted County, MN. Epilepsia, 2012, 53, 2164-2171.	2.6	81
144	Specific safety and tolerability considerations in the use of anticonvulsant medications in children. Drug, Healthcare and Patient Safety, 2012, 4, 39.	1.0	11

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145	Social skills in children with epilepsy: How do they compare to healthy and chronic disease controls?. Epilepsy and Behavior, 2011, 21, 238-241.	0.9	24
146	Neuronal Voltage-Gated Potassium Channel Complex Autoimmunity in Children. Pediatric Neurology, 2011, 44, 275-281.	1.0	56
147	Microdeletion of Chromosome 15q26.1 in a Child With Intractable Generalized Epilepsy. Pediatric Neurology, 2011, 45, 60-62.	1.0	16
148	SISCOM in Children with Tuberous Sclerosis Complex-Related Epilepsy. Pediatric Neurology, 2011, 45, 83-88.	1.0	39
149	Outcome following decompressive craniectomy for malignant middle cerebral artery infarction in children. Developmental Medicine and Child Neurology, 2011, 53, 29-33.	1.1	59
150	A populationâ€based study of longâ€term outcomes of cryptogenic focal epilepsy in childhood: Cryptogenic epilepsy is probably not symptomatic epilepsy. Epilepsia, 2011, 52, 738-745.	2.6	38
151	A populationâ€based study of longâ€term outcome of epilepsy in childhood with a focal or hemispheric lesion on neuroimaging. Epilepsia, 2011, 52, 1522-1526.	2.6	32
152	Incidence and classification of new-onset epilepsy and epilepsy syndromes in children in Olmsted County, Minnesota from 1980 to 2004: A population-based study. Epilepsy Research, 2011, 95, 110-118.	0.8	160
153	Electrocardiographic and oximetric changes during partial complex and generalized seizures. Epilepsy Research, 2011, 95, 237-245.	0.8	76
154	Valproate-Induced Worsening of Seizures. Journal of Child Neurology, 2011, 26, 1319-1321.	0.7	14
155	Generalized Periodic Epileptiform Discharges in a Child With Dravet Syndrome. Journal of Child Neurology, 2011, 26, 907-910.	0.7	7
156	Prognostic Significance of Interictal Epileptiform Discharges in Newly Diagnosed Seizure Disorders. Journal of Clinical Neurophysiology, 2010, 27, 239-248.	0.9	72
157	PEDIATRIC EPILEPSY SYNDROMES. CONTINUUM Lifelong Learning in Neurology, 2010, 16, 57-85.	0.4	2
158	How common is ictal hypoxemia and bradycardia in children with partial complex and generalized convulsive seizures?. Epilepsia, 2010, 51, 1219-1224.	2.6	82
159	The incidence of injuries in persons with and without epilepsyâ€"A populationâ€based study. Epilepsia, 2010, 51, 2247-2253.	2.6	46
160	Epilepsy in Children With Attention-Deficit/Hyperactivity Disorder. Pediatric Neurology, 2010, 42, 325-330.	1.0	71
161	Vitamin D and Bone Health in Children With Epilepsy: Fad or Fact?. Pediatric Neurology, 2010, 42, 394-395.	1.0	10
162	Bilateral Striopallidodentate Calcinosis With Paroxysmal Kinesigenic Dyskinesia. Pediatric Neurology, 2010, 43, 46-48.	1.0	18

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163	EEG and Neuroimaging Studies in Young Children Having Epilepsy Surgery. Pediatric Neurology, 2010, 43, 335-340.	1.0	9
164	Monographs in Human Genetics Neurofibromatoses. Volume 16. 2008. Edited by Dieter Kaufmann. Published by Karger. 192 pages. Price C$190.. Canadian Journal of Neurological Sciences, 2009, 36, 130-130.	0.3	0
165	Controversies in Neonatal Seizure Management. Journal of Child Neurology, 2009, 24, 591-599.	0.7	58
166	Centrotemporal sharp wave EEG trait in rolandic epilepsy maps to Elongator Protein Complex 4 (ELP4). European Journal of Human Genetics, 2009, 17, 1171-1181.	1.4	176
167	Optimal clinical management of children receiving the ketogenic diet: Recommendations of the International Ketogenic Diet Study Group. Epilepsia, 2009, 50, 304-317.	2.6	505
168	Genome wide high density SNP-based linkage analysis of childhood absence epilepsy identifies a susceptibility locus on chromosome 3p23-p14. Epilepsy Research, 2009, 87, 247-255.	0.8	29
169	Are children with epilepsy at greater risk for bullying than their peers?. Epilepsy and Behavior, 2009, 15, 500-505.	0.9	50
170	Mesial Temporal Sclerosis After Posterior Reversible Encephalopathy Syndrome. Pediatric Neurology, 2009, 41, 226-228.	1.0	10
171	Ketogenic ratio, calories, and fluids: Do they matter?. Epilepsia, 2008, 49, 17-19.	2.6	24
172	Electrical Status Epilepticus in Sleep. Seminars in Pediatric Neurology, 2008, 15, 50-60.	1.0	163
173	Self-Concept in Adolescents with Epilepsy: Biological and Social Correlates. Pediatric Neurology, 2008, 38, 335-339.	1.0	33
174	Peri-Ictal Headache in Children: Prevalence and Character. Pediatric Neurology, 2008, 39, 91-96.	1.0	40
175	Anxiety and Depressive Symptoms in Children Presenting With a First Seizure. Pediatric Neurology, 2008, 39, 236-240.	1.0	27
176	Maternal Depression: The Cost of Caring for a Child With Intractable Epilepsy. Pediatric Neurology, 2008, 39, 418-422.	1.0	45
177	Family function in cognitively normal children with epilepsy: Impact on competence and problem behaviors. Epilepsy and Behavior, 2008, 12, 90-95.	0.9	27
178	Health behavior in teens with epilepsy: How do they compare with controls?. Epilepsy and Behavior, 2008, 13, 90-95.	0.9	20
179	Parenting stress in mothers of children with intractable epilepsy. Epilepsy and Behavior, 2008, 13, 169-173.	0.9	58
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