

Alexis Arzimanoglou

List of Publications by Year in descending order

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Version: 2024-02-01

143
papers

15,416
citations

53939

47
h-index

20625

120
g-index

153
all docs

153
docs citations

153
times ranked

14710
citing authors

#	ARTICLE	IF	CITATIONS
1	CNTNAP1-encephalopathy: Six novel patients surviving the neonatal period. <i>European Journal of Paediatric Neurology</i> , 2022, 37, 98-104.	0.7	4
2	The p.Glu787Lys variant in the GRIA3 gene causes developmental and epileptic encephalopathy mimicking structural epilepsy in a female patient. <i>European Journal of Medical Genetics</i> , 2022, 65, 104442.	0.7	4
3	Sleep disorders and ADHD symptoms in children and adolescents with typical absence seizures: An observational study. <i>Epilepsy and Behavior</i> , 2022, 128, 108513.	0.9	2
4	Electroclinical Features and Long-term Seizure Outcome in Patients With Eyelid Myoclonia With Absences. <i>Neurology</i> , 2022, 98, .	1.5	15
5	Gain-of-function and loss-of-function GABRB3 variants lead to distinct clinical phenotypes in patients with developmental and epileptic encephalopathies. <i>Nature Communications</i> , 2022, 13, 1822.	5.8	32
6	The gain of function <i>SCN1A</i> disorder spectrum: novel epilepsy phenotypes and therapeutic implications. <i>Brain</i> , 2022, 145, 3816-3831.	3.7	43
7	Epileptic spasms are associated with increased stereo-electroencephalography derived functional connectivity in tuberous sclerosis complex. <i>Epilepsia</i> , 2022, 63, 2359-2370.	2.6	4
8	Basal Ganglia Dismorphism in Patients With Aicardi Syndrome. <i>Neurology</i> , 2021, 96, e1319-e1333.	1.5	6
9	A survey of the European Reference Network EpiCARE on clinical practice for selected rare epilepsies. <i>Epilepsia Open</i> , 2021, 6, 160-170.	1.3	3
10	The aetiologies of epilepsy. <i>Epileptic Disorders</i> , 2021, 23, 1-16.	0.7	35
11	Clinical spectrum of MTOR-related hypomelanosis of Ito with neurodevelopmental abnormalities. <i>Genetics in Medicine</i> , 2021, 23, 1484-1491.	1.1	14
12	Revision of the diagnostic criteria of alternating hemiplegia of childhood. <i>European Journal of Paediatric Neurology</i> , 2021, 32, A4-A5.	0.7	16
13	Alternating hemiplegia of childhood: evolution over time and mouse model corroboration. <i>Brain Communications</i> , 2021, 3, fcab128.	1.5	8
14	Safety and efficacy of rufinamide in children and adults with Lennox-Gastaut syndrome: A post hoc analysis from Study 022. <i>Epilepsy and Behavior</i> , 2021, 124, 108275.	0.9	2
15	Hypothalamic Hamartomas. <i>Neurology</i> , 2021, 97, 864-873.	1.5	12
16	Trends in pediatric epilepsy surgery in Europe between 2008 and 2015: Country-, center-, and age-specific variation. <i>Epilepsia</i> , 2020, 61, 216-227.	2.6	44
17	Epileptogenicity in tuberous sclerosis complex: A stereoelectroencephalographic study. <i>Epilepsia</i> , 2020, 61, 81-95.	2.6	31
18	Neural correlates of verbal working memory in children with epilepsy with centro-temporal spikes. <i>NeuroImage: Clinical</i> , 2020, 28, 102392.	1.4	4

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19	Corticosteroids versus clobazam in epileptic encephalopathy with ESES: a European multicentre randomised controlled clinical trial (RESCUE ESES*). <i>Trials</i> , 2020, 21, 957.	0.7	8
20	Establishing criteria for pediatric epilepsy surgery center levels of care: Report from the ILAE Pediatric Epilepsy Surgery Task Force. <i>Epilepsia</i> , 2020, 61, 2629-2642.	2.6	19
21	An accelerated shift in the use of remote systems in epilepsy due to the COVID-19 pandemic. <i>Epilepsy and Behavior</i> , 2020, 112, 107376.	0.9	29
22	Interrater agreement of classification of photoparoxysmal electroencephalographic response. <i>Epilepsia</i> , 2020, 61, e124-e128.	2.6	6
23	Did the COVID-19 pandemic silence the needs of people with epilepsy?. <i>Epileptic Disorders</i> , 2020, 22, 439-442.	0.7	46
24	Cardiac phenotype in <i>ATP1A3</i> -related syndromes. <i>Neurology</i> , 2020, 95, e2866-e2879.	1.5	19
25	Seizure outcome and use of antiepileptic drugs after epilepsy surgery according to histopathological diagnosis: a retrospective multicentre cohort study. <i>Lancet Neurology</i> , The, 2020, 19, 748-757.	4.9	177
26	Exome sequencing in 57 patients with self-limited focal epilepsies of childhood with typical or atypical presentations suggests novel candidate genes. <i>European Journal of Paediatric Neurology</i> , 2020, 27, 104-110.	0.7	17
27	Movement disorders in patients with alternating hemiplegia. <i>Neurology</i> , 2020, 94, e1378-e1385.	1.5	14
28	Brain volumetrics in alternating hemiplegia of childhood. <i>European Journal of Paediatric Neurology</i> , 2020, 26, 1.	0.7	1
29	Epilepsy in <i>LAMA2</i> -related muscular dystrophy: An electro-clinico-radiological characterization. <i>Epilepsia</i> , 2020, 61, 971-983.	2.6	12
30	Meta-analysis of drug efficacy in adult vs pediatric trials of patients with PGTC seizures. <i>Neurology</i> , 2020, 94, e1845-e1852.	1.5	2
31	Early-onset epileptic encephalopathy with migrating focal seizures associated with a <i>FARS2</i> homozygous nonsense variant. <i>Epileptic Disorders</i> , 2020, 22, 327-335.	0.7	6
32	The COVID-19 outbreak and approaches to performing EEG in Europe. <i>Epileptic Disorders</i> , 2020, 22, 548-554.	0.7	12
33	BLAST paradigm: A new test to assess brief attentional fluctuations in children with epilepsy, ADHD, and normally developing children. <i>Epilepsy and Behavior</i> , 2019, 99, 106470.	0.9	3
34	Novel study design to assess the efficacy and tolerability of antiseizure medications for focal-onset seizures in infants and young children: A consensus document from the regulatory task force and the pediatric commission of the International League against Epilepsy (ILAE), in collaboration with the Pediatric Epilepsy Research Consortium (PERC). <i>Epilepsia Open</i> , 2019, 4, 537-543.	1.3	20
35	The evaluation and costs of transition programs for youth with epilepsy. <i>Epilepsy and Behavior</i> , 2019, 93, 133-137.	0.9	12
36	Resective surgery in tuberous Sclerosis complex, from Penfield to 2018: A critical review. <i>Revue Neurologique</i> , 2019, 175, 163-182.	0.6	19

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37	Role of observational studies in supporting extrapolation of efficacy data from adults to children with epilepsy – A systematic review of the literature using lacosamide as an example. <i>European Journal of Paediatric Neurology</i> , 2019, 23, 589-603.	0.7	5
38	Disappearance of symptomatic generalized 3-Hz discharges after focal surgery in a patient with tuberous sclerosis. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2019, 67, 71-72.	0.9	0
39	<i>GRIN2A</i> -related disorders: genotype and functional consequence predict phenotype. <i>Brain</i> , 2019, 142, 80-92.	3.7	143
40	Evaluation of long-term safety, tolerability, and behavioral outcomes with adjunctive rufinamide in pediatric patients (1 to 4 years old) with Lennox-Gastaut syndrome: Final results from randomized study 303. <i>European Journal of Paediatric Neurology</i> , 2019, 23, 126-135.	0.7	18
41	Classification of paroxysmal events and the four-dimensional epilepsy classification system. <i>Epileptic Disorders</i> , 2019, 21, 1-29.	0.7	20
42	Roadmap for a competency-based educational curriculum in epileptology: report of the Epilepsy Education Task Force of the International League Against Epilepsy. <i>Epileptic Disorders</i> , 2019, 21, 129-140.	0.7	50
43	Predictive factors and prognostic value for status epilepticus in newborns. <i>European Journal of Paediatric Neurology</i> , 2019, 23, 270-279.	0.7	2
44	Epilepsy and EEG Patterns in Children Diagnosed with Hyperinsulinism. Report of Two Cases Presented as Atypical Generalized Epilepsy and Review of EEGs of 15 Supplementary Cases. , 2019, 50, .		0
45	Cognitive impairment and behavioral disorders in Encephalopathy related to Status Epilepticus during slow Sleep: diagnostic assessment and outcome. <i>Epileptic Disorders</i> , 2019, 21, 71-75.	0.7	9
46	A Review of the New Antiepileptic Drugs for Focal-Onset Seizures in Pediatrics: Role of Extrapolation. <i>Paediatric Drugs</i> , 2018, 20, 249-264.	1.3	35
47	Movement disorders in children: The need to observe, describe in detail and integrate your findings to the global clinical picture. <i>European Journal of Paediatric Neurology</i> , 2018, 22, 217-218.	0.7	1
48	Individualized prediction of seizure relapse and outcomes following antiepileptic drug withdrawal after pediatric epilepsy surgery. <i>Epilepsia</i> , 2018, 59, e28-e33.	2.6	23
49	Neonatal tremor episodes and hyperekplexia-like presentation at onset in a child with <i>SCN8A</i> developmental and epileptic encephalopathy. <i>Epileptic Disorders</i> , 2018, 20, 289-294.	0.7	13
50	Identifying the educational needs of physicians in pediatric epilepsy in order to improve care: results from a needs assessment in Germany, Spain, and the United States. <i>Epileptic Disorders</i> , 2018, 20, 239-256.	0.7	7
51	Why the TimeToStop trial failed to recruit: a survey on antiepileptic drug withdrawal after paediatric epilepsy surgery. <i>Epileptic Disorders</i> , 2018, 20, 374-385.	0.7	12
52	Frameless robot-assisted stereoelectroencephalography for refractory epilepsy in pediatric patients: accuracy, usefulness, and technical issues. <i>Acta Neurochirurgica</i> , 2018, 160, 2489-2500.	0.9	20
53	Epilepsy surgery near or in eloquent cortex in children – Practice patterns and recommendations for minimizing and reporting deficits. <i>Epilepsia</i> , 2018, 59, 1484-1491.	2.6	18
54	Electrical status epilepticus in sleep, a constitutive feature of Christianson syndrome?. <i>European Journal of Paediatric Neurology</i> , 2018, 22, 1124-1132.	0.7	11

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55	European trends in epilepsy surgery. <i>Neurology</i> , 2018, 91, e96-e106.	1.5	108
56	SEEG in ... Family. <i>Neuropediatrics</i> , 2018, 49, S1-S12.	0.3	1
57	Treatment issues for children with epilepsy transitioning to adult care. <i>Epilepsy and Behavior</i> , 2017, 69, 153-160.	0.9	33
58	Histopathological Findings in Brain Tissue Obtained during Epilepsy Surgery. <i>New England Journal of Medicine</i> , 2017, 377, 1648-1656.	13.9	621
59	Extrapolating evidence of antiepileptic drug efficacy in adults to children 2 years of age with focal seizures: The case for disease similarity. <i>Epilepsia</i> , 2017, 58, 1686-1696.	2.6	49
60	Behavioral and fMRI responses to fearful faces are altered in benign childhood epilepsy with centrotemporal spikes (BCECTS). <i>Epilepsia</i> , 2017, 58, 1716-1727.	2.6	19
61	How long for epilepsy remission in the ILAE definition?. <i>Epilepsia</i> , 2017, 58, 1486-1487.	2.6	4
62	The role of EEG in the diagnosis and classification of the epilepsy syndromes: a tool for clinical practice by the ILAE Neurophysiology Task Force (Part 1). <i>Epileptic Disorders</i> , 2017, 19, 233-298.	0.7	79
63	The role of EEG in the diagnosis and classification of the epilepsy syndromes: a tool for clinical practice by the ILAE Neurophysiology Task Force (Part 2). <i>Epileptic Disorders</i> , 2017, 19, 385-437.	0.7	48
64	Expert Opinion on the Management of Lennox-Gastaut Syndrome: Treatment Algorithms and Practical Considerations. <i>Frontiers in Neurology</i> , 2017, 8, 505.	1.1	129
65	ADHD in childhood epilepsy: Clinical determinants of severity and of the response to methylphenidate. <i>Epilepsia</i> , 2016, 57, 1069-1077.	2.6	31
66	Safety and pharmacokinetic profile of rufinamide in pediatric patients aged less than 4 years with Lennox-Gastaut syndrome: An interim analysis from a multicenter, randomized, active-controlled, open-label study. <i>European Journal of Paediatric Neurology</i> , 2016, 20, 393-402.	0.7	29
67	Cost-effectiveness analysis of epilepsy surgery in a controlled cohort of adult patients with intractable partial epilepsy: A 5-year follow-up study. <i>Epilepsia</i> , 2016, 57, 1669-1679.	2.6	90
68	Partial validation of a French version of the ADHD-rating scale IV on a French population of children with ADHD and epilepsy. Factorial structure, reliability, and responsiveness. <i>Epilepsy and Behavior</i> , 2016, 58, 1-6.	0.9	18
69	Safety of levetiracetam among infants younger than 12 months Results from a European multicenter observational study. <i>European Journal of Paediatric Neurology</i> , 2016, 20, 368-375.	0.7	11
70	Clinical profile of patients with ATP1A3 mutations in Alternating Hemiplegia of Childhood—a study of 155 patients. <i>Orphanet Journal of Rare Diseases</i> , 2015, 10, 123.	1.2	117
71	Cognitive consequences of early versus late antiepileptic drug withdrawal after pediatric epilepsy surgery, the TimeToStop (TTS) trial: study protocol for a randomized controlled trial. <i>Trials</i> , 2015, 16, 482.	0.7	18
72	Faulty cardiac repolarization reserve in alternating hemiplegia of childhood broadens the phenotype. <i>Brain</i> , 2015, 138, 2859-2874.	3.7	30

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73	Intelligence quotient improves after antiepileptic drug withdrawal following pediatric epilepsy surgery. <i>Annals of Neurology</i> , 2015, 78, 104-114.	2.8	97
74	Vagus nerve stimulation for drug-resistant epilepsy: A European long-term study up to 24 months in 347 children. <i>Epilepsia</i> , 2014, 55, 1576-1584.	2.6	185
75	Time to relapse after epilepsy surgery in children: AED withdrawal policies are a contributing factor. <i>Epileptic Disorders</i> , 2014, 16, 305-311.	0.7	5
76	A subset of genomic alterations detected in rolandic epilepsies contains candidate or known epilepsy genes including <i>GRIN2A</i> and <i>PRRT2</i> . <i>Epilepsia</i> , 2014, 55, 370-378.	2.6	69
77	Intrinsular functional connectivity in human. <i>Human Brain Mapping</i> , 2014, 35, 2779-2788.	1.9	46
78	The administration of rescue medication to children with prolonged acute convulsive seizures in a non-hospital setting: an exploratory survey of healthcare professionals' perspectives. <i>European Journal of Pediatrics</i> , 2014, 173, 773-779.	1.3	8
79	ILAE Official Report: A practical clinical definition of epilepsy. <i>Epilepsia</i> , 2014, 55, 475-482.	2.6	3,770
80	Functional connectivity of insular efferences. <i>Human Brain Mapping</i> , 2014, 35, 5279-5294.	1.9	66
81	Distinct neurological disorders with ATP1A3 mutations. <i>Lancet Neurology</i> , The, 2014, 13, 503-514.	4.9	206
82	Visual and auditory socio-cognitive perception in unilateral temporal lobe epilepsy in children and adolescents: a prospective controlled study. <i>Epileptic Disorders</i> , 2014, 16, 456-470.	0.7	19
83	The outcome of childhood epilepsy: what improvements are needed?. <i>Epileptic Disorders</i> , 2013, 15, 101-104.	0.7	9
84	GRIN2A mutations in acquired epileptic aphasia and related childhood focal epilepsies and encephalopathies with speech and language dysfunction. <i>Nature Genetics</i> , 2013, 45, 1061-1066.	9.4	380
85	The administration of rescue medication to children with prolonged acute convulsive seizures in the community: What happens in practice?. <i>European Journal of Paediatric Neurology</i> , 2013, 17, 14-23.	0.7	39
86	Caveats and pitfalls of "enduring value" publications: Is only the first author responsible?. <i>Epilepsy and Behavior</i> , 2013, 28, 533-534.	0.9	0
87	Therapeutic approach to epileptic encephalopathies. <i>Epilepsia</i> , 2013, 54, 45-50.	2.6	74
88	Are we failing to provide adequate rescue medication to children at risk of prolonged convulsive seizures in schools?. <i>Archives of Disease in Childhood</i> , 2013, 98, 777-780.	1.0	21
89	Epileptic encephalopathies of the Landau-Kleffner and continuous spike and waves during slow-wave sleep types: Genomic dissection makes the link with autism. <i>Epilepsia</i> , 2012, 53, 1526-1538.	2.6	148
90	ESPERA study: Applicability of the new ILAE criteria for antiepileptic drug resistance of focal epilepsies in current clinical practice. <i>Epilepsy and Behavior</i> , 2012, 25, 166-169.	0.9	3

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91	Timing of antiepileptic drug withdrawal and long-term seizure outcome after paediatric epilepsy surgery (TimeToStop): a retrospective observational study. <i>Lancet Neurology</i> , The, 2012, 11, 784-791.	4.9	115
92	De novo mutations in ATP1A3 cause alternating hemiplegia of childhood. <i>Nature Genetics</i> , 2012, 44, 1030-1034.	9.4	345
93	Alternating hemiplegia of childhood: Metabolic studies in the largest European series of patients. <i>European Journal of Paediatric Neurology</i> , 2012, 16, 10-14.	0.7	20
94	Diagnosing and treating epileptic drop attacks, atypical absences and episodes of nonconvulsive status epilepticus. <i>Epileptic Disorders</i> , 2011, 13, 1-2.	0.7	29
95	Rufinamide from clinical trials to clinical practice in the United States and Europe. <i>Epileptic Disorders</i> , 2011, 13, 27-43.	0.7	7
96	All children who experience epileptic falls do not necessarily have Lennox-Gastaut syndrome... but many do. <i>Epileptic Disorders</i> , 2011, 13, 3-13.	0.7	57
97	Mutations and deletions in PCDH19 account for various familial or isolated epilepsies in females. <i>Human Mutation</i> , 2011, 32, E1959-E1975.	1.1	109
98	Definition of drug resistant epilepsy: Consensus proposal by the ad hoc Task Force of the ILAE Commission on Therapeutic Strategies. <i>Epilepsia</i> , 2010, 51, 1069-1077.	2.6	3,400
99	The evolution of antiepileptic drug development and regulation. <i>Epileptic Disorders</i> , 2010, 12, 3-15.	0.7	60
100	When the past challenges the present: are older antiepileptic drugs still the best choice in childhood absence epilepsy?. <i>Lancet Neurology</i> , The, 2010, 9, 457-459.	4.9	6
101	Outcome and Prognosis of Status Epilepticus in Children. <i>Seminars in Pediatric Neurology</i> , 2010, 17, 195-200.	1.0	18
102	Absence of Mutation in the <i>SLC2A1</i> Gene in a Cohort of Patients with Alternating Hemiplegia of Childhood (AHC). <i>Neuropediatrics</i> , 2010, 41, 267-269.	0.3	13
103	Evidence of a non-progressive course of alternating hemiplegia of childhood: study of a large cohort of children and adults. <i>Brain</i> , 2010, 133, 3598-3610.	3.7	126
104	Alternating Hemiplegia of Childhood: Early Characteristics and Evolution of a Neurodevelopmental Syndrome. <i>Pediatrics</i> , 2009, 123, e534-e541.	1.0	129
105	Lennox-Gastaut syndrome: a consensus approach on diagnosis, assessment, management, and trial methodology. <i>Lancet Neurology</i> , The, 2009, 8, 82-93.	4.9	412
106	Dravet syndrome: From electroclinical characteristics to molecular biology. <i>Epilepsia</i> , 2009, 50, 3-9.	2.6	31
107	A novel three base-pair <i>LG11</i> deletion leading to loss of function in a family with autosomal dominant lateral temporal epilepsy and migraine-like episodes. <i>Epilepsy Research</i> , 2009, 85, 118-122.	0.8	19
108	Perceived impact of epilepsy in teenagers and young adults: An international survey. <i>Epilepsy and Behavior</i> , 2008, 12, 395-401.	0.9	75

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109	Spectrum of SCN1A gene mutations associated with Dravet syndrome: analysis of 333 patients. <i>Journal of Medical Genetics</i> , 2008, 46, 183-191.	1.5	302
110	Greater Response to Placebo in Children Than in Adults: A Systematic Review and Meta-Analysis in Drug-Resistant Partial Epilepsy. <i>PLoS Medicine</i> , 2008, 5, e166.	3.9	124
111	Key clinical features to identify girls with CDKL5 mutations. <i>Brain</i> , 2008, 131, 2647-2661.	3.7	242
112	Surgical treatment of epilepsy in Sturge-Weber syndrome in children. <i>Journal of Neurosurgery: Pediatrics</i> , 2007, 106, 20-28.	0.8	45
113	Surgical Outcome in Tuberous Sclerosis Complex: A Multicenter Survey. <i>Epilepsia</i> , 2007, 48, 1625-1628.	2.6	78
114	Outcome of status epilepticus in children. <i>Epilepsia</i> , 2007, 48, 91-93.	2.6	19
115	Children with epilepsy: are they the same on both sides of the Atlantic, and do the same treatments work?. <i>Epileptic Disorders</i> , 2007, 9, 351-352.	0.7	4
116	Treatment of pediatric epilepsy: European expert opinion, 2007. <i>Epileptic Disorders</i> , 2007, 9, 353-412.	0.7	220
117	Nocturnal Hypermotor Seizures, Suggesting Frontal Lobe Epilepsy, Can Originate in the Insula. <i>Epilepsia</i> , 2006, 47, 755-765.	2.6	233
118	Role of valproate across the ages. Treatment of epilepsy in children. <i>Acta Neurologica Scandinavica</i> , 2006, 114, 1-13.	1.0	54
119	Landau-Kleffner syndrome is not an eponymic badge of ignorance. <i>Epilepsy Research</i> , 2006, 70, 239-247.	0.8	31
120	Parental mosaicism can cause recurrent transmission of SCN1A mutations associated with severe myoclonic epilepsy of infancy. <i>Human Mutation</i> , 2006, 27, 389-389.	1.1	93
121	Zonisamide for the treatment of epilepsy. <i>Expert Review of Neurotherapeutics</i> , 2006, 6, 1283-1292.	1.4	22
122	Optimizing therapy of seizures in children and adolescents with ADHD. <i>Neurology</i> , 2006, 67, S49-51.	1.5	32
123	Isolated paroxysmal arousals as focal epilepsy. <i>Epileptic Disorders</i> , 2006, 8, 45-52.	0.7	4
124	Leptomeningeal Enhancement and Enlarged Choroid Plexus Simulating the Appearance of Sturge-Weber Disease in a Child with Tuberous Sclerosis. <i>Epilepsia</i> , 2005, 46, 595-596.	2.6	8
125	From eponyms to acronyms. <i>Brain and Development</i> , 2005, 27, 163.	0.6	1
126	Topiramate prevents excitotoxic damage in the newborn rodent brain. <i>Neurobiology of Disease</i> , 2005, 20, 837-848.	2.1	80

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127	Les Ã©pilepsies partielles pharmaco-rÃ©sistantes Quels sont les critÃ©res d'Ã©ligibilitÃ© Ã un traitement chirurgical chez lâ€™enfant ?. Revue Neurologique, 2004, 160, 210-219.	0.6	10
128	Topiramate: efficacy and tolerability in children according to epilepsy syndromes. Epilepsy Research, 2003, 53, 225-232.	0.8	81
129	Hypothalamic Hamartoma and Seizures: A Treatable Epileptic Encephalopathy. Epilepsia, 2003, 44, 969-973.	2.6	153
130	Sleep Organization in Children With Partial Refractory Epilepsy. Journal of Child Neurology, 2003, 18, 763-766.	0.7	68
131	Benign idiopathic occipital epilepsy: report of a case of the late (Gastaut) type [corrected]. Epileptic Disorders, 2003, 5, 57-9.	0.7	10
132	Hypothalamic hamartoma and epilepsy: the pathway of discovery. Epileptic Disorders, 2003, 5, 173-5.	0.7	16
133	Positron emission tomography in epileptogenic hypothalamic hamartomas. Epileptic Disorders, 2003, 5, 219-27.	0.7	19
134	Heart rate variability during sleep in children with partial epilepsy. Journal of Sleep Research, 2002, 11, 153-160.	1.7	61
135	Evaluation of the positional candidate gene CHRNA7 at the juvenile myoclonic epilepsy locus (EJM2) on chromosome 15q13â€“14. Epilepsy Research, 2002, 49, 157-172.	0.8	50
136	Epilepsy and neuroprotection: an illustrated review. Epileptic Disorders, 2002, 4, 173-82.	0.7	24
137	Treatment options in pediatric epilepsy syndromes. Epileptic Disorders, 2002, 4, 217-25.	0.7	4
138	Polymorphism analysis of JRK/JH8, the human homologue of mouse jerky, and description of a rare mutation in a case of CAE evolving to JME. Epilepsy Research, 2001, 46, 157-167.	0.8	32
139	MECP2 mutations account for most cases of typical forms of Rett syndrome. Human Molecular Genetics, 2000, 9, 1377-1384.	1.4	254
140	Genetic mapping of a major susceptibility locus for juvenile myoclonic epilepsy on chromosome 15q. Human Molecular Genetics, 1997, 6, 1329-1334.	1.4	220
141	The epilepsy of Sturge-Weber syndrome: Clinical features and treatment in 23 patients. Acta Neurologica Scandinavica, 1992, 86, 18-22.	1.0	79
142	Spontaneous calcific cerebral embolus from a calcific aortic stenosis in a middle cerebral artery infarct.. Stroke, 1989, 20, 691-693.	1.0	51
143	Sturge-Weber syndrome. , 0, , 189-195.		0