Alexis Arzimanoglou

List of Publications by Year
in descending order

143 papers	15,416 citations	47006 47 h-index	18130 120 g-index
153 all docs	153 docs citations	153 times ranked	13797 citing authors

#	Article	IF	Citations
1	CNTNAP1-encephalopathy: Six novel patients surviving the neonatal period. European Journal of Paediatric Neurology, 2022, 37, 98-104.	1.6	4
2	The p.Glu787Lys variant in the GRIA3 gene causes developmental and epileptic encephalopathy mimicking structural epilepsy in a female patient. European Journal of Medical Genetics, 2022, 65, 104442.	1.3	4
3	Sleep disorders and ADHD symptoms in children and adolescents with typical absence seizures: An observational study. Epilepsy and Behavior, 2022, 128, 108513.	1.7	2
4	Electroclinical Features and Long-term Seizure Outcome in Patients With Eyelid Myoclonia With Absences. Neurology, 2022, 98, .	1.1	15
5	Gain-of-function and loss-of-function GABRB3 variants lead to distinct clinical phenotypes in patients with developmental and epileptic encephalopathies. Nature Communications, 2022, 13, 1822.	12.8	32
6	The gain of function SCN1A disorder spectrum: novel epilepsy phenotypes and therapeutic implications. Brain, 2022, 145, 3816-3831.	7.6	43
7	Epileptic spasms are associated with increased stereoâ€electroencephalography derived functional connectivity in tuberous sclerosis complex. Epilepsia, 2022, 63, 2359-2370.	5.1	4
8	Basal Ganglia Dysmorphism in Patients With Aicardi Syndrome. Neurology, 2021, 96, e1319-e1333.	1.1	6
9	A survey of the European Reference Network EpiCARE on clinical practice for selected rare epilepsies. Epilepsia Open, 2021, 6, 160-170.	2.4	3
10	The aetiologies of epilepsy. Epileptic Disorders, 2021, 23, 1-16.	1.3	35
11	Clinical spectrum of MTOR-related hypomelanosis of Ito with neurodevelopmental abnormalities. Genetics in Medicine, 2021, 23, 1484-1491.	2.4	14
12	Revision of the diagnostic criteria of alternating hemiplegia of childhood. European Journal of Paediatric Neurology, 2021, 32, A4-A5.	1.6	16
13	Alternating hemiplegia of childhood: evolution over time and mouse model corroboration. Brain Communications, 2021, 3, fcab128.	3.3	8
14	Safety and efficacy of rufinamide in children and adults with Lennox-Gastaut syndrome: A post hoc analysis from Study 022. Epilepsy and Behavior, 2021, 124, 108275.	1.7	2
15	Hypothalamic Hamartomas. Neurology, 2021, 97, 864-873.	1.1	12
16	Trends in pediatric epilepsy surgery in Europe between 2008 and 2015: Countryâ€, centerâ€, and ageâ€specific variation. Epilepsia, 2020, 61, 216-227.	5.1	44
17	Epileptogenicity in tuberous sclerosis complex: A stereoelectroencephalographic study. Epilepsia, 2020, 61, 81-95.	5.1	31
18	Neural correlates of verbal working memory in children with epilepsy with centro-temporal spikes. NeuroImage: Clinical, 2020, 28, 102392.	2.7	4

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19	Corticosteroids versus clobazam in epileptic encephalopathy with ESES: a European multicentre randomised controlled clinical trial (RESCUE ESES*). Trials, 2020, 21, 957.	1.6	8
20	Establishing criteria for pediatric epilepsy surgery center levels of care: Report from the ILAE Pediatric Epilepsy Surgery Task Force. Epilepsia, 2020, 61, 2629-2642.	5.1	19
21	An accelerated shift in the use of remote systems in epilepsy due to the COVID-19 pandemic. Epilepsy and Behavior, 2020, 112, 107376.	1.7	29
22	Interrater agreement of classification of photoparoxysmal electroencephalographic response. Epilepsia, 2020, 61, e124-e128.	5.1	6
23	Did the COVIDâ€19 pandemic silence the needs of people with epilepsy?. Epileptic Disorders, 2020, 22, 439-442.	1.3	46
24	Cardiac phenotype in ATP1A3 -related syndromes. Neurology, 2020, 95, e2866-e2879.	1.1	19
25	Seizure outcome and use of antiepileptic drugs after epilepsy surgery according to histopathological diagnosis: a retrospective multicentre cohort study. Lancet Neurology, The, 2020, 19, 748-757.	10.2	177
26	Exome sequencing in 57 patients with self-limited focal epilepsies of childhood with typical or atypical presentations suggests novel candidate genes. European Journal of Paediatric Neurology, 2020, 27, 104-110.	1.6	17
27	Movement disorders in patients with alternating hemiplegia. Neurology, 2020, 94, e1378-e1385.	1.1	14
28	Brain volumetrics in alternating hemiplegia of childhood. European Journal of Paediatric Neurology, 2020, 26, 1.	1.6	1
29	Epilepsy in LAMA2â€related muscular dystrophy: An electroâ€clinicoâ€radiological characterization. Epilepsia, 2020, 61, 971-983.	5.1	12
30	Meta-analysis of drug efficacy in adult vs pediatric trials of patients with PGTC seizures. Neurology, 2020, 94, e1845-e1852.	1.1	2
31	Earlyâ€onset epileptic encephalopathy with migrating focal seizures associated with a FARS2 homozygous nonsense variant. Epileptic Disorders, 2020, 22, 327-335.	1.3	6
32	The COVIDâ€19 outbreak and approaches to performing EEG in Europe. Epileptic Disorders, 2020, 22, 548-554.	1.3	12
33	BLAST paradigm: A new test to assess brief attentional fluctuations in children with epilepsy, ADHD, and normally developing children. Epilepsy and Behavior, 2019, 99, 106470.	1.7	3
34	Novel study design to assess the efficacy and tolerability of antiseizure medications for focalâ€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). Epilepsia Open, 2019, 4, 537-543.	2.4	20
35	The evaluation and costs of transition programs for youth with epilepsy. Epilepsy and Behavior, 2019, 93, 133-137.	1.7	12
36	Resective surgery in tuberous Sclerosis complex, from Penfield to 2018: A critical review. Revue Neurologique, 2019, 175, 163-182.	1.5	19

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37	Role of observational studies in supporting extrapolation of efficacy data from adults to children with epilepsy â€" A systematic review of the literature using lacosamide as an example. European Journal of Paediatric Neurology, 2019, 23, 589-603.	1.6	5
38	Disappearance of symptomatic generalized 3-Hz discharges after focal surgery in a patient with tuberous sclerosis. Seizure: the Journal of the British Epilepsy Association, 2019, 67, 71-72.	2.0	0
39	GRIN2A-related disorders: genotype and functional consequence predict phenotype. Brain, 2019, 142, 80-92.	7.6	143
40	Evaluation of long-term safety, tolerability, and behavioral outcomes with adjunctive rufinamide in pediatric patients (â‰¥1 to <4 years old) with Lennox-Gastaut syndrome: Final results from randomized study 303. European Journal of Paediatric Neurology, 2019, 23, 126-135.	1.6	18
41	Classification of paroxysmal events and the fourâ€dimensional epilepsy classification system. Epileptic Disorders, 2019, 21, 1-29.	1.3	20
42	Roadmap for a competencyâ€based educational curriculum in epileptology: report of the Epilepsy Education Task Force of the International League Against Epilepsy. Epileptic Disorders, 2019, 21, 129-140.	1.3	50
43	Predictive factors and prognostic value for status epilepticus in newborns. European Journal of Paediatric Neurology, 2019, 23, 270-279.	1.6	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. Epileptic Disorders, 2019, 21, 71-75.	1.3	9
46	A Review of the New Antiepileptic Drugs for Focal-Onset Seizures in Pediatrics: Role of Extrapolation. Paediatric Drugs, 2018, 20, 249-264.	3.1	35
47	Movement disorders in children: The need to observe, describe in detail and integrate your findings to the global clinical picture. European Journal of Paediatric Neurology, 2018, 22, 217-218.	1.6	1
48	Individualized prediction of seizure relapse and outcomes following antiepileptic drug withdrawal after pediatric epilepsy surgery. Epilepsia, 2018, 59, e28-e33.	5.1	23
49	Neonatal tremor episodes and hyperekplexia-like presentation at onset in a child with SCN8A developmental and epileptic encephalopathy. Epileptic Disorders, 2018, 20, 289-294.	1.3	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. Epileptic Disorders, 2018, 20, 239-256.	1.3	7
51	Why the TimeToStop trial failed to recruit: a survey on antiepileptic drug withdrawal after paediatric epilepsy surgery. Epileptic Disorders, 2018, 20, 374-385.	1.3	12
52	Frameless robot-assisted stereoelectroencephalography for refractory epilepsy in pediatric patients: accuracy, usefulness, and technical issues. Acta Neurochirurgica, 2018, 160, 2489-2500.	1.7	20
53	Epilepsy surgery near or in eloquent cortex in childrenâ€"Practice patterns and recommendations for minimizing and reporting deficits. Epilepsia, 2018, 59, 1484-1491.	5.1	18
54	Electrical status epilepticus in sleep, a constitutive feature of Christianson syndrome?. European Journal of Paediatric Neurology, 2018, 22, 1124-1132.	1.6	11

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55	European trends in epilepsy surgery. Neurology, 2018, 91, e96-e106.	1.1	108
56	SEEG in ... Family. Neuropediatrics, 2018, 49, S1-S12.	0.6	1
57	Treatment issues for children with epilepsy transitioning to adult care. Epilepsy and Behavior, 2017, 69, 153-160.	1.7	33
58	Histopathological Findings in Brain Tissue Obtained during Epilepsy Surgery. New England Journal of Medicine, 2017, 377, 1648-1656.	27.0	621
59	Extrapolating evidence of antiepileptic drug efficacy in adults to children â‰¥2Âyears of age with focal seizures: The case for disease similarity. Epilepsia, 2017, 58, 1686-1696.	5.1	49
60	Behavioral and <scp>fMRI</scp> responses to fearful faces are altered in benign childhood epilepsy with centrotemporal spikes (BCECTS). Epilepsia, 2017, 58, 1716-1727.	5.1	19
61	How long for epilepsy remission in the <scp>ILAE</scp> definition?. Epilepsia, 2017, 58, 1486-1487.	5.1	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). Epileptic Disorders, 2017, 19, 233-298.	1.3	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). Epileptic Disorders, 2017, 19, 385-437.	1.3	48
64	Expert Opinion on the Management of Lennoxâ€“Gastaut Syndrome: Treatment Algorithms and Practical Considerations. Frontiers in Neurology, 2017, 8, 505.	2.4	129
65	ADHD in childhood epilepsy: Clinical determinants of severity and of the response to methylphenidate. Epilepsia, 2016, 57, 1069-1077.	5.1	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. European Journal of Paediatric Neurology, 2016, 20, 393-402.	1.6	29
67	Costâ€effectiveness analysis of epilepsy surgery in a controlled cohort of adult patients with intractable partial epilepsy: A 5â€year followâ€up study. Epilepsia, 2016, 57, 1669-1679.	5.1	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. Epilepsy and Behavior, 2016, 58, 1-6.	1.7	18
69	Safety of levetiracetam among infants younger than 12 months â€“ Results from a European multicenter observational study. European Journal of Paediatric Neurology, 2016, 20, 368-375.	1.6	11
70	Clinical profile of patients with ATP1A3 mutations in Alternating Hemiplegia of Childhoodâ€”a study of 155 patients. Orphanet Journal of Rare Diseases, 2015, 10, 123.	2.7	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. Trials, 2015, 16, 482.	1.6	18
72	Faulty cardiac repolarization reserve in alternating hemiplegia of childhood broadens the phenotype. Brain, 2015, 138, 2859-2874.	7.6	30

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73	Intelligence quotient improves after antiepileptic drug withdrawal following pediatric epilepsy surgery. Annals of Neurology, 2015, 78, 104-114.	5.3	97
74	Vagus nerve stimulation for drugâ€resistant epilepsy: A European longâ€term study up to 24Âmonths in 347Âchildren. Epilepsia, 2014, 55, 1576-1584.	5.1	185
75	Time to relapse after epilepsy surgery in children: AED withdrawal policies are a contributing factor. Epileptic Disorders, 2014, 16, 305-311.	1.3	5
76	A subset of genomic alterations detected in rolandic epilepsies contains candidate or known epilepsy genes including <scp>GRIN</scp>2A and <scp>PRRT</scp>2. Epilepsia, 2014, 55, 370-378.	5.1	69
77	Intrainsular functional connectivity in human. Human Brain Mapping, 2014, 35, 2779-2788.	3.6	46
78	The administration of rescue medication to children with prolonged acute convulsive seizures in a non-hospital setting: an exploratory survey of healthcare professionalsâ€™ perspectives. European Journal of Pediatrics, 2014, 173, 773-779.	2.7	8
79	ILAE Official Report: A practical clinical definition of epilepsy. Epilepsia, 2014, 55, 475-482.	5.1	3,770
80	Functional connectivity of insular efferences. Human Brain Mapping, 2014, 35, 5279-5294.	3.6	66
81	Distinct neurological disorders with ATP1A3 mutations. Lancet Neurology, The, 2014, 13, 503-514.	10.2	206
82	Visual and auditory socio-cognitive perception in unilateral temporal lobe epilepsy in children and adolescents: a prospective controlled study. Epileptic Disorders, 2014, 16, 456-470.	1.3	19
83	The outcome of childhood epilepsy: what improvements are needed?. Epileptic Disorders, 2013, 15, 101-104.	1.3	9
84	GRIN2A mutations in acquired epileptic aphasia and related childhood focal epilepsies and encephalopathies with speech and language dysfunction. Nature Genetics, 2013, 45, 1061-1066.	21.4	380
85	The administration of rescue medication to children with prolonged acute convulsive seizures in the community: What happens in practice?. European Journal of Paediatric Neurology, 2013, 17, 14-23.	1.6	39
86	Caveats and pitfalls of â€œenduring valueâ€•publications: Is only the first author responsible?. Epilepsy and Behavior, 2013, 28, 533-534.	1.7	0
87	Therapeutic approach to epileptic encephalopathies. Epilepsia, 2013, 54, 45-50.	5.1	74
88	Are we failing to provide adequate rescue medication to children at risk of prolonged convulsive seizures in schools?. Archives of Disease in Childhood, 2013, 98, 777-780.	1.9	21
89	Epileptic encephalopathies of the Landauâ€Kleffner and continuous spike and waves during slowâ€wave sleep types: Genomic dissection makes the link with autism. Epilepsia, 2012, 53, 1526-1538.	5.1	148
90	ESPERA study: Applicability of the new ILAE criteria for antiepileptic drug resistance of focal epilepsies in current clinical practice. Epilepsy and Behavior, 2012, 25, 166-169.	1.7	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. Lancet Neurology, The, 2012, 11, 784-791.	10.2	115
92	De novo mutations in ATP1A3 cause alternating hemiplegia of childhood. Nature Genetics, 2012, 44, 1030-1034.	21.4	345
93	Alternating hemiplegia of childhood: Metabolic studies in the largest European series of patients. European Journal of Paediatric Neurology, 2012, 16, 10-14.	1.6	20
94	Diagnosing and treating epileptic drop attacks, atypical absences and episodes of nonconvulsive status epilepticus. Epileptic Disorders, 2011, 13, 1-2.	1.3	29
95	Rufinamide from clinical trials to clinical practice in the United States and Europe. Epileptic Disorders, 2011, 13, 27-43.	1.3	7
96	All children who experience epileptic falls do not necessarily have Lennox-Gastaut syndrome... but many do. Epileptic Disorders, 2011, 13, 3-13.	1.3	57
97	Mutations and deletions in PCDH19 account for various familial or isolated epilepsies in females. Human Mutation, 2011, 32, E1959-E1975.	2.5	109
98	Definition of drug resistant epilepsy: Consensus proposal by the ad hoc Task Force of the ILAE Commission on Therapeutic Strategies. Epilepsia, 2010, 51, 1069-1077.	5.1	3,400
99	The evolution of antiepileptic drug development and regulation. Epileptic Disorders, 2010, 12, 3-15.	1.3	60
100	When the past challenges the present: are older antiepileptic drugs still the best choice in childhood absence epilepsy?. Lancet Neurology, The, 2010, 9, 457-459.	10.2	6
101	Outcome and Prognosis of Status Epilepticus in Children. Seminars in Pediatric Neurology, 2010, 17, 195-200.	2.0	18
102	Absence of Mutation in theSLC2A1Gene in a Cohort of Patients with Alternating Hemiplegia of Childhood (AHC). Neuropediatrics, 2010, 41, 267-269.	0.6	13
103	Evidence of a non-progressive course of alternating hemiplegia of childhood: study of a large cohort of children and adults. Brain, 2010, 133, 3598-3610.	7.6	126
104	Alternating Hemiplegia of Childhood: Early Characteristics and Evolution of a Neurodevelopmental Syndrome. Pediatrics, 2009, 123, e534-e541.	2.1	129
105	Lennox-Gastaut syndrome: a consensus approach on diagnosis, assessment, management, and trial methodology. Lancet Neurology, The, 2009, 8, 82-93.	10.2	412
106	Dravet syndrome: From electroclinical characteristics to molecular biology. Epilepsia, 2009, 50, 3-9.	5.1	31
107	A novel three base-pair LGI1 deletion leading to loss of function in a family with autosomal dominant lateral temporal epilepsy and migraine-like episodes. Epilepsy Research, 2009, 85, 118-122.	1.6	19
108	Perceived impact of epilepsy in teenagers and young adults: An international survey. Epilepsy and Behavior, 2008, 12, 395-401.	1.7	75

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109	Spectrum of SCN1A gene mutations associated with Dravet syndrome: analysis of 333 patients. Journal of Medical Genetics, 2008, 46, 183-191.	3.2	302
110	Greater Response to Placebo in Children Than in Adults: A Systematic Review and Meta-Analysis in Drug-Resistant Partial Epilepsy. PLoS Medicine, 2008, 5, e166.	8.4	124
111	Key clinical features to identify girls with CDKL5 mutations. Brain, 2008, 131, 2647-2661.	7.6	242
112	Surgical treatment of epilepsy in Sturgeâ€“Weber syndrome in children. Journal of Neurosurgery: Pediatrics, 2007, 106, 20-28.	1.3	45
113	Surgical Outcome in Tuberous Sclerosis Complex: A Multicenter Survey. Epilepsia, 2007, 48, 1625-1628.	5.1	78
114	Outcome of status epilepticus in children. Epilepsia, 2007, 48, 91-93.	5.1	19
115	Children with epilepsy: are they the same on both sides of the Atlantic, and do the same treatments work?. Epileptic Disorders, 2007, 9, 351-352.	1.3	4
116	Treatment of pediatric epilepsy: European expert opinion, 2007. Epileptic Disorders, 2007, 9, 353-412.	1.3	220
117	Nocturnal Hypermotor Seizures, Suggesting Frontal Lobe Epilepsy, Can Originate in the Insula. Epilepsia, 2006, 47, 755-765.	5.1	233
118	Role of valproate across the ages. Treatment of epilepsy in children. Acta Neurologica Scandinavica, 2006, 114, 1-13.	2.1	54
119	Landauâ€“Kleffner syndrome is not an eponymic badge of ignorance. Epilepsy Research, 2006, 70, 239-247.	1.6	31
120	Parental mosaicism can cause recurrent transmission ofSCN1A mutations associated with severe myoclonic epilepsy of infancy. Human Mutation, 2006, 27, 389-389.	2.5	93
121	Zonisamide for the treatment of epilepsy. Expert Review of Neurotherapeutics, 2006, 6, 1283-1292.	2.8	22
122	Optimizing therapy of seizures in children and adolescents with ADHD. Neurology, 2006, 67, S49-51.	1.1	32
123	Isolated paroxysmal arousals as focal epilepsy. Epileptic Disorders, 2006, 8, 45-52.	1.3	4
124	Leptomeningeal Enhancement and Enlarged Choroid Plexus Simulating the Appearance of Sturge-Weber Disease in a Child with Tuberous Sclerosis. Epilepsia, 2005, 46, 595-596.	5.1	8
125	From eponyms to acronyms. Brain and Development, 2005, 27, 163.	1.1	1
126	Topiramate prevents excitotoxic damage in the newborn rodent brain. Neurobiology of Disease, 2005, 20, 837-848.	4.4	80

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127	Les Ã©pilepsies partielles pharmaco-rÃ©sistantes Quels sont les critÃ¨res dâ€™Ã©ligibilitÃ© Ã un traitement chirurgical chez lâ€™enfant ?. Revue Neurologique, 2004, 160, 210-219.	1.5	10
128	Topiramate: efficacy and tolerability in children according to epilepsy syndromes. Epilepsy Research, 2003, 53, 225-232.	1.6	81
129	Hypothalamic Hamartoma and Seizures: A Treatable Epileptic Encephalopathy. Epilepsia, 2003, 44, 969-973.	5.1	153
130	Sleep Organization in Children With Partial Refractory Epilepsy. Journal of Child Neurology, 2003, 18, 763-766.	1.4	68
131	Benign idiopathic occipital epilepsy: report of a case of the late (Gastaut) type [corrected]. Epileptic Disorders, 2003, 5, 57-9.	1.3	10
132	Hypothalamic hamartoma and epilepsy: the pathway of discovery. Epileptic Disorders, 2003, 5, 173-5.	1.3	16
133	Positron emission tomography in epileptogenic hypothalamic hamartomas. Epileptic Disorders, 2003, 5, 219-27.	1.3	19
134	Heart rate variability during sleep in children with partial epilepsy. Journal of Sleep Research, 2002, 11, 153-160.	3.2	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.	1.6	50
136	Epilepsy and neuroprotection: an illustrated review. Epileptic Disorders, 2002, 4, 173-82.	1.3	24
137	Treatment options in pediatric epilepsy syndromes. Epileptic Disorders, 2002, 4, 217-25.	1.3	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.	1.6	32
139	MECP2 mutations account for most cases of typical forms of Rett syndrome. Human Molecular Genetics, 2000, 9, 1377-1384.	2.9	254
140	Genetic mapping of a major susceptibility locus for juvenile myoclonic epilepsy on chromosome 15q. Human Molecular Genetics, 1997, 6, 1329-1334.	2.9	220
141	The epilepsy of Sturge-Weber syndrome: Clinical features and treatment in 23 patients. Acta Neurologica Scandinavica, 1992, 86, 18-22.	2.1	79
142	Spontaneous calcific cerebral embolus from a calcific aortic stenosis in a middle cerebral artery infarct.. Stroke, 1989, 20, 691-693.	2.0	51
143	Sturgeâ€“Weber syndrome. , 0, , 189-195.		0