Alexis Arzimanoglou

List of Publications by Year
in descending order

143 papers	15,416 citations	46984 47 h-index	18115 120 g-index
153 all docs	153 docs citations	153 times ranked	13797 citing authors

#	Article	IF	Citations
1	ILAE Official Report: A practical clinical definition of epilepsy. Epilepsia, 2014, 55, 475-482.	2.6	3,770
2	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.	2.6	3,400
3	Histopathological Findings in Brain Tissue Obtained during Epilepsy Surgery. New England Journal of Medicine, 2017, 377, 1648-1656.	13.9	621
4	Lennox-Gastaut syndrome: a consensus approach on diagnosis, assessment, management, and trial methodology. Lancet Neurology, The, 2009, 8, 82-93.	4.9	412
5	GRIN2A mutations in acquired epileptic aphasia and related childhood focal epilepsies and encephalopathies with speech and language dysfunction. Nature Genetics, 2013, 45, 1061-1066.	9.4	380
6	De novo mutations in ATP1A3 cause alternating hemiplegia of childhood. Nature Genetics, 2012, 44, 1030-1034.	9.4	345
7	Spectrum of SCN1A gene mutations associated with Dravet syndrome: analysis of 333 patients. Journal of Medical Genetics, 2008, 46, 183-191.	1.5	302
8	MECP2 mutations account for most cases of typical forms of Rett syndrome. Human Molecular Genetics, 2000, 9, 1377-1384.	1.4	254
9	Key clinical features to identify girls with CDKL5 mutations. Brain, 2008, 131, 2647-2661.	3.7	242
10	Nocturnal Hypermotor Seizures, Suggesting Frontal Lobe Epilepsy, Can Originate in the Insula. Epilepsia, 2006, 47, 755-765.	2.6	233
11	Genetic mapping of a major susceptibility locus for juvenile myoclonic epilepsy on chromosome 15q. Human Molecular Genetics, 1997, 6, 1329-1334.	1.4	220
12	Treatment of pediatric epilepsy: European expert opinion, 2007. Epileptic Disorders, 2007, 9, 353-412.	0.7	220
13	Distinct neurological disorders with ATP1A3 mutations. Lancet Neurology, The, 2014, 13, 503-514.	4.9	206
14	Vagus nerve stimulation for drugâ€resistant epilepsy: A European longâ€term study up to 24Âmonths in 347Âchildren. Epilepsia, 2014, 55, 1576-1584.	2.6	185
15	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.	4.9	177
16	Hypothalamic Hamartoma and Seizures: A Treatable Epileptic Encephalopathy. Epilepsia, 2003, 44, 969-973.	2.6	153
17	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.	2.6	148
18	GRIN2A-related disorders: genotype and functional consequence predict phenotype. Brain, 2019, 142, 80-92.	3.7	143

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19	Alternating Hemiplegia of Childhood: Early Characteristics and Evolution of a Neurodevelopmental Syndrome. Pediatrics, 2009, 123, e534-e541.	1.0	129
20	Expert Opinion on the Management of Lennoxâ€“Gastaut Syndrome: Treatment Algorithms and Practical Considerations. Frontiers in Neurology, 2017, 8, 505.	1.1	129
21	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.	3.7	126
22	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.	3.9	124
23	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.	1.2	117
24	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.	4.9	115
25	Mutations and deletions in PCDH19 account for various familial or isolated epilepsies in females. Human Mutation, 2011, 32, E1959-E1975.	1.1	109
26	European trends in epilepsy surgery. Neurology, 2018, 91, e96-e106.	1.5	108
27	Intelligence quotient improves after antiepileptic drug withdrawal following pediatric epilepsy surgery. Annals of Neurology, 2015, 78, 104-114.	2.8	97
28	Parental mosaicism can cause recurrent transmission ofSCN1A mutations associated with severe myoclonic epilepsy of infancy. Human Mutation, 2006, 27, 389-389.	1.1	93
29	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.	2.6	90
30	Topiramate: efficacy and tolerability in children according to epilepsy syndromes. Epilepsy Research, 2003, 53, 225-232.	0.8	81
31	Topiramate prevents excitotoxic damage in the newborn rodent brain. Neurobiology of Disease, 2005, 20, 837-848.	2.1	80
32	The epilepsy of Sturge-Weber syndrome: Clinical features and treatment in 23 patients. Acta Neurologica Scandinavica, 1992, 86, 18-22.	1.0	79
33	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.	0.7	79
34	Surgical Outcome in Tuberous Sclerosis Complex: A Multicenter Survey. Epilepsia, 2007, 48, 1625-1628.	2.6	78
35	Perceived impact of epilepsy in teenagers and young adults: An international survey. Epilepsy and Behavior, 2008, 12, 395-401.	0.9	75
36	Therapeutic approach to epileptic encephalopathies. Epilepsia, 2013, 54, 45-50.	2.6	74

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37	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.	2.6	69
38	Sleep Organization in Children With Partial Refractory Epilepsy. Journal of Child Neurology, 2003, 18, 763-766.	0.7	68
39	Functional connectivity of insular efferences. Human Brain Mapping, 2014, 35, 5279-5294.	1.9	66
40	Heart rate variability during sleep in children with partial epilepsy. Journal of Sleep Research, 2002, 11, 153-160.	1.7	61
41	The evolution of antiepileptic drug development and regulation. Epileptic Disorders, 2010, 12, 3-15.	0.7	60
42	All children who experience epileptic falls do not necessarily have Lennox-Gastaut syndrome... but many do. Epileptic Disorders, 2011, 13, 3-13.	0.7	57
43	Role of valproate across the ages. Treatment of epilepsy in children. Acta Neurologica Scandinavica, 2006, 114, 1-13.	1.0	54
44	Spontaneous calcific cerebral embolus from a calcific aortic stenosis in a middle cerebral artery infarct.. Stroke, 1989, 20, 691-693.	1.0	51
45	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
46	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.	0.7	50
47	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.	2.6	49
48	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.	0.7	48
49	Intrainsular functional connectivity in human. Human Brain Mapping, 2014, 35, 2779-2788.	1.9	46
50	Did the COVIDâ€19 pandemic silence the needs of people with epilepsy?. Epileptic Disorders, 2020, 22, 439-442.	0.7	46
51	Surgical treatment of epilepsy in Sturgeâ€“Weber syndrome in children. Journal of Neurosurgery: Pediatrics, 2007, 106, 20-28.	0.8	45
52	Trends in pediatric epilepsy surgery in Europe between 2008 and 2015: Countryâ€, centerâ€, and ageâ€specific variation. Epilepsia, 2020, 61, 216-227.	2.6	44
53	The gain of function SCN1A disorder spectrum: novel epilepsy phenotypes and therapeutic implications. Brain, 2022, 145, 3816-3831.	3.7	43
54	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.	0.7	39

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55	A Review of the New Antiepileptic Drugs for Focal-Onset Seizures in Pediatrics: Role of Extrapolation. Paediatric Drugs, 2018, 20, 249-264.	1.3	35
56	The aetiologies of epilepsy. Epileptic Disorders, 2021, 23, 1-16.	0.7	35
57	Treatment issues for children with epilepsy transitioning to adult care. Epilepsy and Behavior, 2017, 69, 153-160.	0.9	33
58	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
59	Optimizing therapy of seizures in children and adolescents with ADHD. Neurology, 2006, 67, S49-51.	1.5	32
60	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.	5.8	32
61	Landauâ€“Kleffner syndrome is not an eponymic badge of ignorance. Epilepsy Research, 2006, 70, 239-247.	0.8	31
62	Dravet syndrome: From electroclinical characteristics to molecular biology. Epilepsia, 2009, 50, 3-9.	2.6	31
63	ADHD in childhood epilepsy: Clinical determinants of severity and of the response to methylphenidate. Epilepsia, 2016, 57, 1069-1077.	2.6	31
64	Epileptogenicity in tuberous sclerosis complex: A stereoelectroencephalographic study. Epilepsia, 2020, 61, 81-95.	2.6	31
65	Faulty cardiac repolarization reserve in alternating hemiplegia of childhood broadens the phenotype. Brain, 2015, 138, 2859-2874.	3.7	30
66	Diagnosing and treating epileptic drop attacks, atypical absences and episodes of nonconvulsive status epilepticus. Epileptic Disorders, 2011, 13, 1-2.	0.7	29
67	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.	0.7	29
68	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
69	Epilepsy and neuroprotection: an illustrated review. Epileptic Disorders, 2002, 4, 173-82.	0.7	24
70	Individualized prediction of seizure relapse and outcomes following antiepileptic drug withdrawal after pediatric epilepsy surgery. Epilepsia, 2018, 59, e28-e33.	2.6	23
71	Zonisamide for the treatment of epilepsy. Expert Review of Neurotherapeutics, 2006, 6, 1283-1292.	1.4	22
72	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.	1.0	21

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73	Alternating hemiplegia of childhood: Metabolic studies in the largest European series of patients. European Journal of Paediatric Neurology, 2012, 16, 10-14.	0.7	20
74	Frameless robot-assisted stereoelectroencephalography for refractory epilepsy in pediatric patients: accuracy, usefulness, and technical issues. Acta Neurochirurgica, 2018, 160, 2489-2500.	0.9	20
75	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.	1.3	20
76	Classification of paroxysmal events and the fourâ€dimensional epilepsy classification system. Epileptic Disorders, 2019, 21, 1-29.	0.7	20
77	Outcome of status epilepticus in children. Epilepsia, 2007, 48, 91-93.	2.6	19
78	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.	0.8	19
79	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.	0.7	19
80	Behavioral and <scp>fMRI</scp> responses to fearful faces are altered in benign childhood epilepsy with centrotemporal spikes (BCECTS). Epilepsia, 2017, 58, 1716-1727.	2.6	19
81	Resective surgery in tuberous Sclerosis complex, from Penfield to 2018: A critical review. Revue Neurologique, 2019, 175, 163-182.	0.6	19
82	Establishing criteria for pediatric epilepsy surgery center levels of care: Report from the ILAE Pediatric Epilepsy Surgery Task Force. Epilepsia, 2020, 61, 2629-2642.	2.6	19
83	Cardiac phenotype in ATP1A3 -related syndromes. Neurology, 2020, 95, e2866-e2879.	1.5	19
84	Positron emission tomography in epileptogenic hypothalamic hamartomas. Epileptic Disorders, 2003, 5, 219-27.	0.7	19
85	Outcome and Prognosis of Status Epilepticus in Children. Seminars in Pediatric Neurology, 2010, 17, 195-200.	1.0	18
86	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.	0.7	18
87	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.	0.9	18
88	Epilepsy surgery near or in eloquent cortex in childrenâ€”Practice patterns and recommendations for minimizing and reporting deficits. Epilepsia, 2018, 59, 1484-1491.	2.6	18
89	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.	0.7	18
90	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.	0.7	17

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91	Revision of the diagnostic criteria of alternating hemiplegia of childhood. European Journal of Paediatric Neurology, 2021, 32, A4-A5.	0.7	16
92	Hypothalamic hamartoma and epilepsy: the pathway of discovery. Epileptic Disorders, 2003, 5, 173-5.	0.7	16
93	Electroclinical Features and Long-term Seizure Outcome in Patients With Eyelid Myoclonia With Absences. Neurology, 2022, 98, .	1.5	15
94	Movement disorders in patients with alternating hemiplegia. Neurology, 2020, 94, e1378-e1385.	1.5	14
95	Clinical spectrum of MTOR-related hypomelanosis of Ito with neurodevelopmental abnormalities. Genetics in Medicine, 2021, 23, 1484-1491.	1.1	14
96	Absence of Mutation in theSLC2A1Gene in a Cohort of Patients with Alternating Hemiplegia of Childhood (AHC). Neuropediatrics, 2010, 41, 267-269.	0.3	13
97	Neonatal tremor episodes and hyperekplexia-like presentation at onset in a child with SCN8A developmental and epileptic encephalopathy. Epileptic Disorders, 2018, 20, 289-294.	0.7	13
98	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
99	The evaluation and costs of transition programs for youth with epilepsy. Epilepsy and Behavior, 2019, 93, 133-137.	0.9	12
100	Epilepsy in LAMA2â€related muscular dystrophy: An electroâ€clinicoâ€radiological characterization. Epilepsia, 2020, 61, 971-983.	2.6	12
101	The COVIDâ€19 outbreak and approaches to performing EEG in Europe. Epileptic Disorders, 2020, 22, 548-554.	0.7	12
102	Hypothalamic Hamartomas. Neurology, 2021, 97, 864-873.	1.5	12
103	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.	0.7	11
104	Electrical status epilepticus in sleep, a constitutive feature of Christianson syndrome?. European Journal of Paediatric Neurology, 2018, 22, 1124-1132.	0.7	11
105	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.	0.6	10
106	Benign idiopathic occipital epilepsy: report of a case of the late (Gastaut) type [corrected]. Epileptic Disorders, 2003, 5, 57-9.	0.7	10
107	The outcome of childhood epilepsy: what improvements are needed?. Epileptic Disorders, 2013, 15, 101-104.	0.7	9
108	Cognitive impairment and behavioral disorders in Encephalopathy related to Status Epilepticus during slow Sleep: diagnostic assessment and outcome. Epileptic Disorders, 2019, 21, 71-75.	0.7	9

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109	Leptomeningeal Enhancement and Enlarged Choroid Plexus Simulating the Appearance of Sturge-Weber Disease in a Child with Tuberous Sclerosis. Epilepsia, 2005, 46, 595-596.	2.6	8
110	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.	1.3	8
111	Corticosteroids versus clobazam in epileptic encephalopathy with ESES: a European multicentre randomised controlled clinical trial (RESCUE ESES*). Trials, 2020, 21, 957.	0.7	8
112	Alternating hemiplegia of childhood: evolution over time and mouse model corroboration. Brain Communications, 2021, 3, fcab128.	1.5	8
113	Rufinamide from clinical trials to clinical practice in the United States and Europe. Epileptic Disorders, 2011, 13, 27-43.	0.7	7
114	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.	0.7	7
115	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.	4.9	6
116	Interrater agreement of classification of photoparoxysmal electroencephalographic response. Epilepsia, 2020, 61, e124-e128.	2.6	6
117	Basal Ganglia Dysmorphism in Patients With Aicardi Syndrome. Neurology, 2021, 96, e1319-e1333.	1.5	6
118	Earlyâ€onset epileptic encephalopathy with migrating focal seizures associated with a FARS2 homozygous nonsense variant. Epileptic Disorders, 2020, 22, 327-335.	0.7	6
119	Time to relapse after epilepsy surgery in children: AED withdrawal policies are a contributing factor. Epileptic Disorders, 2014, 16, 305-311.	0.7	5
120	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.	0.7	5
121	How long for epilepsy remission in the <scp>ILAE</scp> definition?. Epilepsia, 2017, 58, 1486-1487.	2.6	4
122	Neural correlates of verbal working memory in children with epilepsy with centro-temporal spikes. NeuroImage: Clinical, 2020, 28, 102392.	1.4	4
123	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.	0.7	4
124	CNTNAP1-encephalopathy: Six novel patients surviving the neonatal period. European Journal of Paediatric Neurology, 2022, 37, 98-104.	0.7	4
125	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.	0.7	4
126	Treatment options in pediatric epilepsy syndromes. Epileptic Disorders, 2002, 4, 217-25.	0.7	4

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127	Isolated paroxysmal arousals as focal epilepsy. Epileptic Disorders, 2006, 8, 45-52.	0.7	4
128	Epileptic spasms are associated with increased stereoâ€electroencephalography derived functional connectivity in tuberous sclerosis complex. Epilepsia, 2022, 63, 2359-2370.	2.6	4
129	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.	0.9	3
130	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.	0.9	3
131	A survey of the European Reference Network EpiCARE on clinical practice for selected rare epilepsies. Epilepsia Open, 2021, 6, 160-170.	1.3	3
132	Meta-analysis of drug efficacy in adult vs pediatric trials of patients with PGTC seizures. Neurology, 2020, 94, e1845-e1852.	1.5	2
133	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.	0.9	2
134	Predictive factors and prognostic value for status epilepticus in newborns. European Journal of Paediatric Neurology, 2019, 23, 270-279.	0.7	2
135	Sleep disorders and ADHD symptoms in children and adolescents with typical absence seizures: An observational study. Epilepsy and Behavior, 2022, 128, 108513.	0.9	2
136	From eponyms to acronyms. Brain and Development, 2005, 27, 163.	0.6	1
137	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.	0.7	1
138	Brain volumetrics in alternating hemiplegia of childhood. European Journal of Paediatric Neurology, 2020, 26, 1.	0.7	1
139	SEEG in ... Family. Neuropediatrics, 2018, 49, S1-S12.	0.3	1
140	Sturgeâ€“Weber syndrome. , 0, , 189-195.		0
141	Caveats and pitfalls of â€œenduring valueâ€•publications: Is only the first author responsible?. Epilepsy and Behavior, 2013, 28, 533-534.	0.9	0
142	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.	0.9	0
143	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