Giangennaro G Coppola

List of Publications by Year
in descending order

199 papers	6,290 citations	53660 45 h-index	110170 64 g-index
204 all docs	204 docs citations	204 times ranked	5727 citing authors

#	Article	IF	Citations
1	Migrating Partial Seizures in Infancy: A Malignant Disorder with Developmental Arrest. Epilepsia, 1995, 36, 1017-1024.	2.6	251
2	Bone and calcium metabolism and antiepileptic drugs. Clinical Neurology and Neurosurgery, 2010, 112, 1-10.	0.6	206
3	The ketogenic diet in children, adolescents and young adults with refractory epilepsy: an Italian multicentric experience. Epilepsy Research, 2002, 48, 221-227.	0.8	134
4	Benign Familial Neonatal Convulsions Caused by Altered Gating of KCNQ2/KCNQ3 Potassium Channels. Journal of Neuroscience, 2002, 22, RC199-RC199.	1.7	120
5	Topiramate as add-on drug in severe myoclonic epilepsy in infancy: an Italian multicenter open trial. Epilepsy Research, 2002, 49, 45-48.	0.8	116
6	Lamotrigine versus Valproic Acid as First-line Monotherapy in Newly Diagnosed Typical Absence Seizures: An Open-label, Randomized, Parallel-group Study. Epilepsia, 2004, 45, 1049-1053.	2.6	110
7	Melatonin in wakeâ€“sleep disorders in children, adolescents and young adults with mental retardation with or without epilepsy: a double-blind, cross-over, placebo-controlled trial. Brain and Development, 2004, 26, 373-376.	0.6	110
8	The metabolic syndrome in overweight epileptic patients treated with valproic acid. Epilepsia, 2010, 51, 268-273.	2.6	109
9	Weight gain following treatment with valproic acid: pathogenetic mechanisms and clinical implications. Obesity Reviews, 2011, 12, e32-43.	3.1	109
10	Malignant migrating partial seizures in infancy: An epilepsy syndrome of unknown etiology. Epilepsia, 2009, 50, 49-51.	2.6	101
11	Impact of Dietary Fats on Brain Functions. Current Neuropharmacology, 2018, 16, 1059-1085.	1.4	95
12	Heterozygous Reelin Mutations Cause Autosomal-Dominant Lateral Temporal Epilepsy. American Journal of Human Genetics, 2015, 96, 992-1000.	2.6	94
13	Levetiracetam or oxcarbazepine as monotherapy in newly diagnosed benign epilepsy of childhood with centrotemporal spikes (BECTS): An open-label, parallel group trial. Brain and Development, 2007, 29, 281-284.	0.6	80
14	Bone mineral density in children, adolescents, and young adults with epilepsy. Epilepsia, 2009, 50, 2140-2146.	2.6	80
15	Antiepileptic drugs, sex hormones, and PCOS. Epilepsia, 2011, 52, 199-211.	2.6	78
16	Characterization of two de novo KCNT1 mutations in children with malignant migrating partial seizures in infancy. Molecular and Cellular Neurosciences, 2016, 72, 54-63.	1.0	77
17	Mutational scanning of potassium, sodium and chloride ion channels in malignant migrating partial seizures in infancy. Brain and Development, 2006, 28, 76-79.	0.6	70
18	Efficacy and safety of levetiracetam: An add-on trial in children with refractory epilepsy. Seizure: the Journal of the British Epilepsy Association, 2005, 14, 248-253.	0.9	68

#	Article	IF	Citations
19	Low glycemic index diet in children and young adults with refractory epilepsy: First Italian experience. Seizure: the Journal of the British Epilepsy Association, 2011, 20, 526-528.	0.9	68
20	A multicenter, randomized, placeboâ€controlled trial of levetiracetam in children and adolescents with newly diagnosed absence epilepsy. Epilepsia, 2011, 52, 802-809.	2.6	67
21	Should â€œmigralepsyâ€•be considered an obsolete concept? A multicenter retrospective clinical/EEG study and review of the literature. Epilepsy and Behavior, 2011, 21, 52-59.	0.9	65
22	Improving molecular diagnosis in epilepsy by a dedicated high-throughput sequencing platform. European Journal of Human Genetics, 2015, 23, 354-362.	1.4	64
23	Management of psychogenic nonâ€epileptic seizures: a multidisciplinary approach. European Journal of Neurology, 2019, 26, 205.	1.7	64
24	Anticonvulsant drugs and hematological disease. Neurological Sciences, 2014, 35, 983-993.	0.9	63
25	Effectiveness and tolerability of perampanel in children and adolescents with refractory epilepsiesâ€”An Italian observational multicenter study. Epilepsy Research, 2016, 127, 93-100.	0.8	62
26	Lacosamide in pediatric and adult patients: Comparison of efficacy and safety. Seizure: the Journal of the British Epilepsy Association, 2013, 22, 210-216.	0.9	60
27	Plasma free carnitine in epilepsy children, adolescents and young adults treated with old and new antiepileptic drugs with or without ketogenic diet. Brain and Development, 2006, 28, 358-365.	0.6	59
28	Neuropsychological impairment in childhood absence epilepsy: Review of the literature. Journal of the Neurological Sciences, 2015, 359, 59-66.	0.3	59
29	Benign convulsions associated with mild gastroenteritis: A multicenter clinical study. Epilepsy Research, 2011, 93, 107-114.	0.8	57
30	Treatment of malignant migrating partial epilepsy of infancy with rufinamide: report of five cases. Epileptic Disorders, 2011, 13, 18-21.	0.7	55
31	Conduct disorders and psychopathy in children and adolescents: aetiology, clinical presentation and treatment strategies of callous-unemotional traits. Italian Journal of Pediatrics, 2017, 43, 84.	1.0	54
32	Epilepsy and genetic in Rett syndrome: A review. Brain and Behavior, 2019, 9, e01250.	1.0	53
33	Levetiracetam monotherapy for children and adolescents with benign rolandic seizures. Seizure: the Journal of the British Epilepsy Association, 2007, 16, 271-275.	0.9	52
34	Rufinamide in children and adults with Lennoxâ€“Gastaut syndrome: First Italian multicenter experience. Seizure: the Journal of the British Epilepsy Association, 2010, 19, 587-591.	0.9	52
35	The pharmacological management of Lennox-Gastaut syndrome and critical literature review. Seizure: the Journal of the British Epilepsy Association, 2018, 63, 17-25.	0.9	52
36	Benign familial neonatal convulsions (BFNC) resulting from mutation of the KCNQ2 voltage sensor. European Journal of Human Genetics, 2000, 8, 994-997.	1.4	51

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37	Efficacy and safety of levetiracetam in infants and young children with refractory epilepsy. Seizure: the Journal of the British Epilepsy Association, 2007, 16, 345-350.	0.9	51
38	Levetiracetam in absence epilepsy. Developmental Medicine and Child Neurology, 2008, 50, 850-853.	1.1	51
39	Peri-ictal and inter-ictal headache in children and adolescents with idiopathic epilepsy: a multicenter cross-sectional study. Child's Nervous System, 2011, 27, 1419-1423.	0.6	50
40	A Novel Hyperekplexia-causing Mutation in the Pre-transmembrane Segment 1 of the Human Glycine Receptor Î±1 Subunit Reduces Membrane Expression and Impairs Gating by Agonists. Journal of Biological Chemistry, 2004, 279, 25598-25604.	1.6	49
41	Ketogenic diet for the treatment of catastrophic epileptic encephalopathies in childhood. European Journal of Paediatric Neurology, 2010, 14, 229-234.	0.7	49
42	West syndrome associated with 14q12 duplications harboring FOXG1. Neurology, 2011, 76, 1600-1602.	1.5	49
43	The effects of the ketogenic diet in refractory partial seizures with reference to tuberous sclerosis. European Journal of Paediatric Neurology, 2006, 10, 148-151.	0.7	48
44	The impact of the ketogenic diet on arterial morphology and endothelial function in children and young adults with epilepsy: A caseâ€“control study. Seizure: the Journal of the British Epilepsy Association, 2014, 23, 260-265.	0.9	48
45	Topiramate as add-on drug in children, adolescents and young adults with Lennox-Gastaut syndrome: an Italian multicentric study. Epilepsy Research, 2002, 51, 147-153.	0.8	46
46	Levetiracetam during 1-year follow-up in children, adolescents, and young adults with refractory epilepsy. Epilepsy Research, 2004, 59, 35-42.	0.8	46
47	Levetiracetam in juvenile myoclonic epilepsy: longâ€term efficacy in newly diagnosed adolescents. Developmental Medicine and Child Neurology, 2008, 50, 29-32.	1.1	45
48	Clinical Guidelines for Management of Bone Health in Rett Syndrome Based on Expert Consensus and Available Evidence. PLoS ONE, 2016, 11, e0146824.	1.1	45
49	Topiramate in children and adolescents with epilepsy and mental retardation: A prospective study on behavior and cognitive effects. Epilepsy and Behavior, 2008, 12, 253-256.	0.9	43
50	Low long-term efficacy and tolerability of add-on rufinamide in patients with Dravet syndrome. Epilepsy and Behavior, 2011, 21, 282-284.	0.9	42
51	Melatonin in wake-sleep disorders in children, adolescents and young adults with mental retardation with or without epilepsy: a double-blind, cross-over, placebo-controlled trial. Brain and Development, 2004, 26, 373-6.	0.6	42
52	Mozart's music in children with drug-refractory epileptic encephalopathies. Epilepsy and Behavior, 2015, 50, 18-22.	0.9	41
53	Update on the safety of second generation antipsychotics in youths: a call for collaboration among paediatricians and child psychiatrists. Italian Journal of Pediatrics, 2016, 42, 51.	1.0	41
54	Rufinamide in refractory childhood epileptic encephalopathies other than Lennox-Gastaut syndrome. European Journal of Neurology, 2011, 18, 246-251.	1.7	40

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55	Lack of pathogenic mutations in six patients with MMPSI. Epilepsy Research, 2014, 108, 340-344.	0.8	40
56	Lamotrigine as first-line drug in childhood absence epilepsy: a clinical and neurophysiological study. Brain and Development, 2004, 26, 26-29.	0.6	38
57	Clinical features of psychogenic non-epileptic seizures in prepubertal and pubertal patients with idiopathic epilepsy. Neurological Sciences, 2009, 30, 319-323.	0.9	38
58	Antiepileptic drug withdrawal in childhood epilepsy: What are the risk factors associated with seizure relapse?. European Journal of Paediatric Neurology, 2012, 16, 599-604.	0.7	38
59	A 12â€month longitudinal study of calcium metabolism and bone turnover during valproate monotherapy. European Journal of Neurology, 2010, 17, 232-237.	1.7	37
60	The ketogenic diet for Dravet syndrome and other epileptic encephalopathies: An Italian consensus. Epilepsia, 2011, 52, 83-89.	2.6	37
61	Hormonal and reproductive disturbances in epileptic male patients: Emerging issues. Reproductive Toxicology, 2011, 31, 519-527.	1.3	37
62	Neuropsychological impairment in children with Rolandic epilepsy and in their siblings. Epilepsy and Behavior, 2013, 28, 108-112.	0.9	36
63	Ketogenic diet and childhood neurological disorders other than epilepsy: an overview. Expert Review of Neurotherapeutics, 2017, 17, 461-473.	1.4	36
64	Nonalcoholic fatty liver disease in adolescents receiving valproic acid. Epilepsy and Behavior, 2011, 20, 382-385.	0.9	35
65	Topiramate in refractory partial-onset seizures in children, adolescents and young adults: a multicentric open trial. Epilepsy Research, 2001, 43, 255-260.	0.8	33
66	Low penetrance of autosomal dominant lateral temporal epilepsy in Italian families without <scp>LGI</scp>1 mutations. Epilepsia, 2013, 54, 1288-1297.	2.6	32
67	Current role of rufinamide in the treatment of childhood epilepsy: Literature review and treatment guidelines. European Journal of Paediatric Neurology, 2014, 18, 685-690.	0.7	32
68	New developments in the management of partial-onset epilepsy: role of brivaracetam. Drug Design, Development and Therapy, 2017, Volume11, 643-657.	2.0	32
69	Perampanel tolerability in children and adolescents with focal epilepsy: Effects on behavior and executive functions. Epilepsy and Behavior, 2020, 103, 106879.	0.9	32
70	A mutation (V260M) in the middle of the M2 pore-lining domain of the glycine receptor causes hereditary hyperekplexia. European Journal of Human Genetics, 2001, 9, 873-876.	1.4	31
71	Valproic acid and phenobarbital blood levels during the first month of treatment with the ketogenic diet. Acta Neurologica Scandinavica, 2010, 122, 303-307.	1.0	31
72	Bone mineral density in a population of children and adolescents with cerebral palsy and mental retardation with or without epilepsy. Epilepsia, 2012, 53, 2172-2177.	2.6	31

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73	Malignant migrating partial seizures in infancy. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2013, 111, 605-609.	1.0	31
74	Efficacy and tolerability of add-on lacosamide in children with Lennox-Gastaut syndrome. Acta Neurologica Scandinavica, 2014, 129, 420-424.	1.0	31
75	The clinical phenotype of autosomal dominant lateral temporal lobe epilepsy related to reelin mutations. Epilepsy and Behavior, 2017, 68, 103-107.	0.9	31
76	Lamotrigine as add-on drug in children and adolescents with refractory epilepsy and mental delay: an open trial. Brain and Development, 1997, 19, 398-402.	0.6	30
77	Cognitive and linguistic abnormalities in benign childhood epilepsy with centrotemporal spikes. Acta Paediatrica, International Journal of Paediatrics, 2011, 100, 768-772.	0.7	30
78	Pharmacotherapy for children and adolescents with epilepsy. Expert Opinion on Pharmacotherapy, 2011, 12, 175-194.	0.9	30
79	Long-term follow-up in children with benign convulsions associated with gastroenteritis. European Journal of Paediatric Neurology, 2014, 18, 572-577.	0.7	30
80	Oral-facial-digital syndrome type VI: is C5orf42 really the major gene?. Human Genetics, 2015, 134, 123-126.	1.8	30
81	Neonatal sporadic hyperekplexia: a rare and often unrecognized entity. Brain and Development, 1997, 19, 226-228.	0.6	29
82	Correlating the Clinical and Genetic Features of Benign Familial Neonatal Seizures (BFNS) with the Functional Consequences of Underlying Mutations. Channels, 2007, 1, 228-233.	1.5	29
83	First long-term experience with the orphan drug rufinamide in children with myoclonic-astatic epilepsy (Doose syndrome). European Journal of Paediatric Neurology, 2012, 16, 459-463.	0.7	29
84	Nutritional problems in children with neuromotor disabilities: an Italian case series. Italian Journal of Pediatrics, 2014, 40, 61.	1.0	29
85	Mozart's music in children with drug-refractory epileptic encephalopathies: Comparison of two protocols. Epilepsy and Behavior, 2018, 78, 100-103.	0.9	29
86	Non-resective surgery and radiosurgery for treatment of drug-resistant epilepsy. Epilepsy Research, 2012, 99, 193-201.	0.8	28
87	Obese children suffer more often from migraine. Acta Paediatrica, International Journal of Paediatrics, 2012, 101, e416-21.	0.7	28
88	Social Cognition in Neurodevelopmental Disorders and Epilepsy. Frontiers in Neurology, 2021, 12, 658823.	1.1	28
89	Autosomal dominant lateral temporal epilepsy: Absence of mutations in ADAM22 and Kv1 channel genes encoding LGI1-associated proteins. Epilepsy Research, 2008, 80, 1-8.	0.8	26
90	Epilepsy, sex hormones and antiepileptic drugs in female patients. Expert Review of Neurotherapeutics, 2009, 9, 1803-1814.	1.4	26

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91	Low penetrance and effect on protein secretion of LGI1 mutations causing autosomal dominant lateral temporal epilepsy. Epilepsia, 2011, 52, 1258-1264.	2.6	26
92	Zonisamide in children and young adults with refractory epilepsy: An open label, multicenter Italian study. Epilepsy Research, 2009, 83, 112-116.	0.8	25
93	Electroclinical Features and Long-Term Outcome of Cryptogenic Epilepsy in Children with Down Syndrome. Journal of Pediatrics, 2013, 163, 1754-1758.	0.9	25
94	Epilepsy in Children With Menkes Disease. Journal of Child Neurology, 2014, 29, 1757-1764.	0.7	25
95	Anticonvulsant properties of an oral ketone ester in a pentylenetetrazole-model of seizure. Brain Research, 2015, 1618, 50-54.	1.1	25
96	Current role of perampanel in pediatric epilepsy. Italian Journal of Pediatrics, 2017, 43, 51.	1.0	25
97	Association between SCN1A gene polymorphisms and drug resistant epilepsy in pediatric patients. Seizure: the Journal of the British Epilepsy Association, 2018, 55, 30-35.	0.9	25
98	The Ketogenic Diet for the Treatment of Mood Disorders in Comorbidity With Epilepsy in Children and Adolescents. Frontiers in Pharmacology, 2020, 11, 578396.	1.6	25
99	Neonatal Hyperekplexia: A Case Report. Epilepsia, 1992, 33, 817-820.	2.6	24
100	Short-term Nonhormonal and Nonsteroid Treatment in West Syndrome. Epilepsia, 2003, 44, 1085-1088.	2.6	24
101	Levetiracetam in submaximal subcutaneous pentylentetrazol-induced seizures in rats. Seizure: the Journal of the British Epilepsy Association, 2010, 19, 296-299.	0.9	24
102	Epilepsy in the setting of full trisomy 18: A multicenter study on 18 affected children with and without structural brain abnormalities. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2016, 172, 288-295.	0.7	24
103	Treatment of Partial Seizures in Childhood. CNS Drugs, 2004, 18, 133-156.	2.7	23
104	Catamenial epilepsy: hormonal aspects. Gynecological Endocrinology, 2010, 26, 783-790.	0.7	23
105	Rufinamide for refractory focal seizures: An open-label, multicenter European study. Seizure: the Journal of the British Epilepsy Association, 2013, 22, 33-36.	0.9	23
106	Epilepsy in patients with Cornelia de Lange syndrome: A clinical series. Seizure: the Journal of the British Epilepsy Association, 2013, 22, 356-359.	0.9	23
107	Psychogenic nonepileptic seizures in pediatric population: A review. Brain and Behavior, 2019, 9, e01406.	1.0	23
108	Putative Mechanisms of Action and Clinical Use of Lithium in Children and Adolescents: A Critical Review. Current Neuropharmacology, 2019, 17, 318-341.	1.4	23

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109	Long term outcome in children affected by absence epilepsy with onset before the age of three years. Epilepsy and Behavior, 2011, 20, 366-369.	0.9	22
110	The Effect of Plasma Protein Binding on the Therapeutic Monitoring of Antiseizure Medications. Pharmaceutics, 2021, 13, 1208.	2.0	21
111	Role of folic acid depletion on homocysteine serum level in children and adolescents with epilepsy and different MTHFR C677T genotypes. Seizure: the Journal of the British Epilepsy Association, 2012, 21, 340-343.	0.9	20
112	Update on rufinamide in childhood epilepsy. Neuropsychiatric Disease and Treatment, 2011, 7, 399.	1.0	19
113	Efficacy and safety of rufinamide in children under four years of age with drug-resistant epilepsies. European Journal of Paediatric Neurology, 2014, 18, 641-645.	0.7	19
114	Investigational small molecules in phase II clinical trials for the treatment of epilepsy. Expert Opinion on Investigational Drugs, 2018, 27, 971-979.	1.9	19
115	Rufinamide for the treatment of refractory epilepsy secondary to neuronal migration disorders. Epilepsy Research, 2014, 108, 542-546.	0.8	18
116	Teachers of various school grades and representations of epilepsy: problems, relational aspects and perspectives of life quality. Italian Journal of Pediatrics, 2015, 41, 70.	1.0	18
117	Analysis of LGI1 promoter sequence, PDYN and GABBR1 polymorphisms in sporadic and familial lateral temporal lobe epilepsy. Neuroscience Letters, 2008, 436, 23-26.	1.0	17
118	Refractory absence seizures: An Italian multicenter retrospective study. European Journal of Paediatric Neurology, 2015, 19, 660-664.	0.7	17
119	Facial Emotion Recognition in Children and Adolescents with Specific Learning Disorder. Brain Sciences, 2020, 10, 473.	1.1	17
120	Temporal lobe dual pathology in malignant migrating partial seizures in infancy. Epileptic Disorders, 2007, 9, 145-148.	0.7	17
121	Vaccination and Occurrence of Seizures in SCN1A Mutationâ€“Positive Patients: A Multicenter Italian Study. Pediatric Neurology, 2014, 50, 228-232.	1.0	16
122	Anticonvulsant drugs for generalized tonic-clonic epilepsy. Expert Opinion on Pharmacotherapy, 2017, 18, 925-936.	0.9	16
123	Digital Devices Use and Language Skills in Children between 8 and 36 Month. Brain Sciences, 2020, 10, 656.	1.1	15
124	Cerebellar Vermis Defect, Oligophrenia, Congenital Ataxia, and Hepatic Fibrocirrhosis without Coloboma and Renal Abnormalities: Report of Three Cases. Neuropediatrics, 2002, 33, 180-185.	0.3	14
125	Low-dose lamotrigine in West syndrome. Epilepsy Research, 2002, 51, 199-200.	0.8	14
126	Neuropsychological profiles and outcomes in children with new onset frontal lobe epilepsy. Epilepsy and Behavior, 2016, 55, 79-83.	0.9	14

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127	Adaptive Behavior, Emotional/Behavioral Problems and Parental Stress in Children With Autism Spectrum Disorder. Frontiers in Neuroscience, 2021, 15, 751465.	1.4	14
128	Pontocerebellar hypoplasia type 2 (PCH2): report of two siblings. Brain and Development, 2000, 22, 188-192.	0.6	13
129	Benign idiopathic partial seizures in the velocardiofacial syndrome: Report of two cases. American Journal of Medical Genetics Part A, 2001, 103, 172-175.	2.4	13
130	Epilepsy and electroencephalographic anomalies in chromosome 2 aberrations. Epilepsy Research, 2008, 79, 63-70.	0.8	13
131	Efficacy and safety of felbamate in children under 4â€ƒyears of age: a retrospective chart review. European Journal of Neurology, 2008, 15, 940-946.	1.7	13
132	Effects of the abrupt switch from solution to modified-release granule formulation of valproate. Acta Neurologica Scandinavica, 2012, 125, e14-e18.	1.0	13
133	Satisfaction with antiepileptic drugs in children and adolescents with newly diagnosed and chronic epilepsy. Epilepsy Research, 2012, 100, 142-151.	0.8	13
134	Clinical dissection of childhood occipital epilepsy of Gastaut and prognostic implication. European Journal of Neurology, 2016, 23, 241-246.	1.7	13
135	Ketogenic diet prevents neuronal firing increase within the substantia nigra during pentylenetetrazole-induced seizure in rats. Brain Research Bulletin, 2016, 125, 168-172.	1.4	13
136	Monitoring And Managing Depression In Adolescents With Epilepsy: Current Perspectives . Neuropsychiatric Disease and Treatment, 2019, Volume 15, 2773-2780.	1.0	13
137	Parental stress in a sample of children with epilepsy. Acta Neurologica Scandinavica, 2019, 140, 87-92.	1.0	13
138	Impact of COVID-19 Pandemic on Children and Adolescents with Neuropsychiatric Disorders: Emotional/Behavioral Symptoms and Parental Stress. International Journal of Environmental Research and Public Health, 2022, 19, 3795.	1.2	13
139	Early onset absence epilepsy with onset in the first year of life: A multicenter cohort study. Epilepsia, 2013, 54, 66-69.	2.6	12
140	Weight Regain after Discontinuation of Topiramate Treatment in Patients with Migraine: a Prospective Observational Study. CNS Drugs, 2015, 29, 163-169.	2.7	12
141	Long-term outcome of autistic spectrum disorder: a retrospective case study in a southern italian region. Italian Journal of Pediatrics, 2017, 43, 83.	1.0	12
142	Cognitive, adaptive, and behavioral effects of adjunctive rufinamide in Lennoxâ€“Gastaut syndrome: A prospective observational clinical study. Epilepsy and Behavior, 2020, 112, 107445.	0.9	12
143	Bone Mineral Density in Angelman Syndrome. Pediatric Neurology, 2007, 37, 411-416.	1.0	11
144	Epilepsy in Menkes disease: An electroclinical long-term study of 28 patients. Epilepsy Research, 2014, 108, 1597-1603.	0.8	11

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145	Update on the role of eslicarbazepine acetate in the treatment of partial-onset epilepsy. Neuropsychiatric Disease and Treatment, 2016, 12, 1251.	1.0	11
146	Gelastic seizures and low-grade hypothalamic astrocytoma: a case report. Brain and Development, 2002, 24, 183-186.	0.6	10
147	Long-term outcome of epilepsy in Kabuki syndrome. Seizure: the Journal of the British Epilepsy Association, 2011, 20, 650-654.	0.9	10
148	Genetic heterogeneity in malignant migrating partial seizures of infancy. Annals of Neurology, 2014, 75, 324-326.	2.8	10
149	Brain anatomical substrates of mirror movements in Kallmann syndrome. NeuroImage, 2015, 104, 52-58.	2.1	10
150	Epilepsy and Electroencephalographic Abnormalities in SATB2-Associated Syndrome. Pediatric Neurology, 2020, 112, 94-100.	1.0	10
151	Psychiatric Symptoms and Parental Stress in Children and Adolescents With Epilepsy. Frontiers in Neurology, 2021, 12, 778410.	1.1	10
152	Association of intronic variants of the KCNAB1 gene with lateral temporal epilepsy. Epilepsy Research, 2011, 94, 110-116.	0.8	9
153	Different Immune Signature in Youths Experiencing Antipsychotic-Induced Weight Gain Compared to Untreated Obese Patients. Journal of Child and Adolescent Psychopharmacology, 2017, 27, 844-848.	0.7	9
154	Differences in Metabolic Factors Between Antipsychotic-Induced Weight Gain and Non-pharmacological ObesityÂin Youths. Clinical Drug Investigation, 2018, 38, 457-462.	1.1	9
155	Parental stress in pediatric epilepsy after therapy withdrawal. Epilepsy and Behavior, 2019, 94, 239-242.	0.9	9
156	Focal polymicrogyria, continuous spike-and-wave discharges during slow-wave sleep and Cohen syndrome: a case report. Brain and Development, 2003, 25, 446-449.	0.6	8
157	Reflex myoclonic epilepsy in infancy: A multicenter clinical study. Epilepsy Research, 2013, 103, 237-244.	0.8	8
158	Different calorie restriction treatments have similar anti-seizure efficacy. Seizure: the Journal of the British Epilepsy Association, 2016, 35, 45-49.	0.9	8
159	Electroclinical features of epilepsy monosomy 1p36 syndrome and their implications. Acta Neurologica Scandinavica, 2018, 138, 523-530.	1.0	8
160	Gelastic seizures not associated with hypothalamic hamartoma: A long-term follow-up study. Epilepsy and Behavior, 2020, 103, 106578.	0.9	8
161	What are the challenges with the pharmacological management of epilepsy in patients with Attention Deficit Hyperactivity Disorder (ADHD)?. Expert Opinion on Pharmacotherapy, 2020, 21, 737-739.	0.9	8
162	Perampanel as first add-on antiseizure medication: Italian consensus clinical practice statements. BMC Neurology, 2021, 21, 410.	0.8	8

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163	Phenytoin neurotoxicity in a child carrying new STXBP1 and CYP2C9 gene mutations. Seizure: the Journal of the British Epilepsy Association, 2016, 34, 26-28.	0.9	7
164	Cognitive Profile, Emotional-Behavioral Features, and Parental Stress in Boys With 47,XYY Syndrome. Cognitive and Behavioral Neurology, 2019, 32, 87-94.	0.5	7
165	Familial Hemiplegic Migraine with an ATP1A4 Mutation: Clinical Spectrum and Carbamazepine Efficacy. Brain Sciences, 2020, 10, 372.	1.1	7
166	Double-blind, placebo-controlled, cross-over trial of allopurinol as add-on therapy in childhood refractory epilepsy. Brain and Development, 1996, 18, 50-52.	0.6	6
167	Vigabatrin as add-on therapy in children and adolescents with refractory epilepsy: an open trial. Brain and Development, 1997, 19, 459-463.	0.6	6
168	Simultaneous Onset of Infantile Spasms in Monozygotic Twins. Pediatric Neurology, 2010, 43, 127-130.	1.0	6
169	Gastaut type-idiopathic childhood occipital epilepsy and childhood absence epilepsy: A clinically significant association?. Seizure: the Journal of the British Epilepsy Association, 2010, 19, 368-372.	0.9	6
170	Selection of antiseizure medications for first add-on use: A consensus paper. Epilepsy and Behavior, 2021, 122, 108087.	0.9	6
171	Topiramate in frontal lobe epilepsy. Acta Neurologica Scandinavica, 2007, 115, 132-135.	1.0	5
172	Effects of antiseizure monotherapy on visuospatial memory in pediatric age. European Journal of Paediatric Neurology, 2021, 32, 106-114.	0.7	5
173	Perampanel dosage in plasma samples: development and validation of a novel HPLC method with combined UV-Fluorescence detection. Journal of Pharmaceutical and Biomedical Analysis, 2021, 204, 114252.	1.4	5
174	Epilepsy and Cognitive Impairment in Childhood and Adolescence: A Mini-Review. Current Neuropharmacology, 2023, 21, 1646-1665.	1.4	5
175	Epilepsy and occipital calcifications with or without celiac disease: Report of four cases. Journal of Epilepsy, 1994, 7, 130-136.	0.4	4
176	Unusual compulsive motor activity during treatment with clothiapine in a mentally retarded adolescent. Brain and Development, 2004, 26, 409-411.	0.6	4
177	Successful Treatment of Refractory Seizures With Rufinamide in Children With Schizencephaly. Journal of Child Neurology, 2015, 30, 1079-1083.	0.7	4
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