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	Epilepsy and Cognitive Impairment in Childhood and Adolescence: A Mini-Review. Current Neuropharmacology, 2023, 21, 1646-1665.	1.4	5
2	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
3	Social Cognition in Neurodevelopmental Disorders and Epilepsy. Frontiers in Neurology, 2021, 12, 658823.	1.1	28
4	Effects of antiseizure monotherapy on visuospatial memory in pediatric age. European Journal of Paediatric Neurology, 2021, 32, 106-114.	0.7	5
5	The Effect of Plasma Protein Binding on the Therapeutic Monitoring of Antiseizure Medications. Pharmaceutics, 2021, 13, 1208.	2.0	21
6	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
7	1.Neuropsychiatric disorders and parental stress during the covid-19 pandemic: an Italian retrospective longitudinal study. Archiv Euromedica, 2021, 11, 5-9.	0.1	0
8	Selection of antiseizure medications for first add-on use: A consensus paper. Epilepsy and Behavior, 2021, 122, 108087.	0.9	6
9	DIGITALTOOLS AND LANGUAGE IN CHILDREN AGED BETWEEN 8 TO 36 MONTHS. Archiv Euromedica, 2021, 11, 20-23.	0.1	0
10	VISUOSPATIALSKILLS AND ANTISEIZURE MEDICATIONSIN CHI. Archiv Euromedica, 2021, 11, 24-27.	0.1	0
11	Perampanel as first add-on antiseizure medication: Italian consensus clinical practice statements. BMC Neurology, 2021, 21, 410.	0.8	8
12	Adaptive Behavior, Emotional/Behavioral Problems and Parental Stress in Children With Autism Spectrum Disorder. Frontiers in Neuroscience, 2021, 15, 751465.	1.4	14
13	Psychiatric Symptoms and Parental Stress in Children and Adolescents With Epilepsy. Frontiers in Neurology, 2021, 12, 778410.	1.1	10
14	Gelastic seizures not associated with hypothalamic hamartoma: A long-term follow-up study. Epilepsy and Behavior, 2020, 103, 106578.	0.9	8
15	Infantile spasms followed by childhood absence epilepsy: A case series. Seizure: the Journal of the British Epilepsy Association, 2020, 74, 77-80.	0.9	3
16	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
17	Psychogenic Non-Epileptic Status as Refractory, Generalized Hypertonic Posturing: Report of Two Adolescents. Medicina (Lithuania), 2020, 56, 508.	0.8	3
18	Digital Devices Use and Language Skills in Children between 8 and 36 Month. Brain Sciences, 2020, 10, 656.	1.1	15

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19	Facial Emotion Recognition in Children and Adolescents with Specific Learning Disorder. Brain Sciences, 2020, 10, 473.	1.1	17
20	Epilepsy and Electroencephalographic Abnormalities in SATB2-Associated Syndrome. Pediatric Neurology, 2020, 112, 94-100.	1.0	10
21	Withdrawal seizures: possible risk factors. Expert Review of Neurotherapeutics, 2020, 20, 667-672.	1.4	4
22	Familial Hemiplegic Migraine with an ATP1A4 Mutation: Clinical Spectrum and Carbamazepine Efficacy. Brain Sciences, 2020, 10, 372.	1.1	7
23	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
24	Perampanel tolerability in children and adolescents with focal epilepsy: Effects on behavior and executive functions. Epilepsy and Behavior, 2020, 103, 106879.	0.9	32
25	Reading and writing difficultiesin third and sixth-grade students: a cross-sectional survey. Minerva Pediatrics, 2020, , .	0.2	1
26	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
27	Psychogenic nonepileptic seizures in pediatric population: A review. Brain and Behavior, 2019, 9, e01406.	1.0	23
28	A Calorie-Restricted Ketogenic Diet Reduces Cerebral Cortex Vascularization in Prepubertal Rats. Nutrients, 2019, 11, 2681.	1.7	3
29	Monitoring And Managing Depression In Adolescents With Epilepsy: Current Perspectives . Neuropsychiatric Disease and Treatment, 2019, Volume 15, 2773-2780.	1.0	13
30	Parental stress in a sample of children with epilepsy. Acta Neurologica Scandinavica, 2019, 140, 87-92.	1.0	13
31	Parental stress in pediatric epilepsy after therapy withdrawal. Epilepsy and Behavior, 2019, 94, 239-242.	0.9	9
32	Epilepsy and genetic in Rett syndrome: A review. Brain and Behavior, 2019, 9, e01250.	1.0	53
33	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
34	Management of psychogenic nonâ€epileptic seizures: a multidisciplinary approach. European Journal of Neurology, 2019, 26, 205.	1.7	64
35	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
36	Familial Hemiplegic Migraine: A New Gene in an Italian Family. Archives of Clinical and Medical Case Reports, 2019, 03, .	0.0	2

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37	Attention deficit hyperactivity disorder in genetically-determined intellectual disability. Minerva Pediatrica, 2019, 71, 310-312.	2.6	0
38	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
39	Differences in Metabolic Factors Between Antipsychotic-Induced Weight Gain and Non-pharmacological ObesityÂin Youths. Clinical Drug Investigation, 2018, 38, 457-462.	1.1	9
40	Mozart's music in children with drug-refractory epileptic encephalopathies: Comparison of two protocols. Epilepsy and Behavior, 2018, 78, 100-103.	0.9	29
41	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
42	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
43	Electroclinical features of epilepsy monosomy 1p36 syndrome and their implications. Acta Neurologica Scandinavica, 2018, 138, 523-530.	1.0	8
44	Impact of Dietary Fats on Brain Functions. Current Neuropharmacology, 2018, 16, 1059-1085.	1.4	95
45	Qualitative and quantitative revaluation of specific learning disabilities: a multicentric study. Minerva Pediatrics, 2018, , .	0.2	1
46	Infantile spasms in early-onset Niemannâ€“Pick disease with a novel compound heterozygous mutations in SMPD1 gene. European Journal of Molecular and Clinical Medicine, 2017, 2, 155.	0.5	1
47	The clinical phenotype of autosomal dominant lateral temporal lobe epilepsy related to reelin mutations. Epilepsy and Behavior, 2017, 68, 103-107.	0.9	31
48	Anticonvulsant drugs for generalized tonic-clonic epilepsy. Expert Opinion on Pharmacotherapy, 2017, 18, 925-936.	0.9	16
49	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
50	Current role of perampanel in pediatric epilepsy. Italian Journal of Pediatrics, 2017, 43, 51.	1.0	25
51	Ketogenic diet and childhood neurological disorders other than epilepsy: an overview. Expert Review of Neurotherapeutics, 2017, 17, 461-473.	1.4	36
52	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
53	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
54	New developments in the management of partial-onset epilepsy: role of brivaracetam. Drug Design, Development and Therapy, 2017, Volume11, 643-657.	2.0	32

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55	Update on the role of eslicarbazepine acetate in the treatment of partial-onset epilepsy. Neuropsychiatric Disease and Treatment, 2016, 12, 1251.	1.0	11
56	Clinical dissection of childhood occipital epilepsy of Gastaut and prognostic implication. European Journal of Neurology, 2016, 23, 241-246.	1.7	13
57	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
58	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
59	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
60	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
61	Psychiatric Manifestation of EAST Syndrome. Journal of Clinical Psychopharmacology, 2016, 36, 185-187.	0.7	0
62	Kleefstra-variant syndrome with heterozygous mutations in EHMT1 and KCNQ2 genes: a case report. Neurological Sciences, 2016, 37, 829-831.	0.9	4
63	Different calorie restriction treatments have similar anti-seizure efficacy. Seizure: the Journal of the British Epilepsy Association, 2016, 35, 45-49.	0.9	8
64	Neuropsychological profiles and outcomes in children with new onset frontal lobe epilepsy. Epilepsy and Behavior, 2016, 55, 79-83.	0.9	14
65	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
66	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
67	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
68	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
69	Heterozygous Reelin Mutations Cause Autosomal-Dominant Lateral Temporal Epilepsy. American Journal of Human Genetics, 2015, 96, 992-1000.	2.6	94
70	Neuropsychological impairment in childhood absence epilepsy: Review of the literature. Journal of the Neurological Sciences, 2015, 359, 59-66.	0.3	59
71	Weight Regain after Discontinuation of Topiramate Treatment in Patients with Migraine: a Prospective Observational Study. CNS Drugs, 2015, 29, 163-169.	2.7	12
72	Oral-facial-digital syndrome type VI: is C5orf42 really the major gene?. Human Genetics, 2015, 134, 123-126.	1.8	30

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73	Refractory absence seizures: An Italian multicenter retrospective study. European Journal of Paediatric Neurology, 2015, 19, 660-664.	0.7	17
74	Corrigendum to â€˜efficacy and safety of rufinamide in children under four years of age with drug-resistant epilepsiesâ€™ [YEJPN 18 (5) (2015) 641â€“645]. European Journal of Paediatric Neurology, 2015, 19, 388.	0.7	0
75	Mozart's music in children with drug-refractory epileptic encephalopathies. Epilepsy and Behavior, 2015, 50, 18-22.	0.9	41
76	Successful Treatment of Refractory Seizures With Rufinamide in Children With Schizencephaly. Journal of Child Neurology, 2015, 30, 1079-1083.	0.7	4
77	Anticonvulsant properties of an oral ketone ester in a pentylenetetrazole-model of seizure. Brain Research, 2015, 1618, 50-54.	1.1	25
78	Brain anatomical substrates of mirror movements in Kallmann syndrome. NeuroImage, 2015, 104, 52-58.	2.1	10
79	Improving molecular diagnosis in epilepsy by a dedicated high-throughput sequencing platform. European Journal of Human Genetics, 2015, 23, 354-362.	1.4	64
80	Epilepsy in Children With Menkes Disease. Journal of Child Neurology, 2014, 29, 1757-1764.	0.7	25
81	Efficacy and tolerability of add-on lacosamide in children with Lennox-Gastaut syndrome. Acta Neurologica Scandinavica, 2014, 129, 420-424.	1.0	31
82	Genetic heterogeneity in malignant migrating partial seizures of infancy. Annals of Neurology, 2014, 75, 324-326.	2.8	10
83	Nutritional problems in children with neuromotor disabilities: an Italian case series. Italian Journal of Pediatrics, 2014, 40, 61.	1.0	29
84	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
85	Lack of pathogenic mutations in six patients with MMPSI. Epilepsy Research, 2014, 108, 340-344.	0.8	40
86	Long-term follow-up in children with benign convulsions associated with gastroenteritis. European Journal of Paediatric Neurology, 2014, 18, 572-577.	0.7	30
87	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
88	Anticonvulsant drugs and hematological disease. Neurological Sciences, 2014, 35, 983-993.	0.9	63
89	Vaccination and Occurrence of Seizures in SCN1A Mutationâ€“Positive Patients: A Multicenter Italian Study. Pediatric Neurology, 2014, 50, 228-232.	1.0	16
90	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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91	Identification of malnutrition in children with severe neuromotor disabilities: A still overlooked aspect in our country. Digestive and Liver Disease, 2014, 46, e99.	0.4	0
92	Brainstem arteriovenous malformation presenting with dyspraxic handwriting in a young girl. Brain and Development, 2014, 36, 541-544.	0.6	0
93	Rufinamide for the treatment of refractory epilepsy secondary to neuronal migration disorders. Epilepsy Research, 2014, 108, 542-546.	0.8	18
94	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
95	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
96	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
97	Neuropsychological impairment in children with Rolandic epilepsy and in their siblings. Epilepsy and Behavior, 2013, 28, 108-112.	0.9	36
98	Reflex myoclonic epilepsy in infancy: A multicenter clinical study. Epilepsy Research, 2013, 103, 237-244.	0.8	8
99	Electroclinical Features and Long-Term Outcome of Cryptogenic Epilepsy in Children with Down Syndrome. Journal of Pediatrics, 2013, 163, 1754-1758.	0.9	25
100	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
101	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
102	Epileptic Encephalopathies in Children. Epilepsy Research & Treatment, 2013, 2013, 1-1.	1.4	0
103	Early onset absence epilepsy with onset in the first year of life: A multicenter cohort study. Epilepsia, 2013, 54, 66-69.	2.6	12
104	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
105	Commentary on â€œBenign Afebrile Convulsions in the Course of Mild Acute Gastroenteritisâ€• Pediatric Emergency Care, 2012, 28, 830.	0.5	0
106	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
107	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
108	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

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109	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
110	Effects of the abrupt switch from solution to modified-release granule formulation of valproate. Acta Neurologica Scandinavica, 2012, 125, e14-e18.	1.0	13
111	Non-resective surgery and radiosurgery for treatment of drug-resistant epilepsy. Epilepsy Research, 2012, 99, 193-201.	0.8	28
112	Satisfaction with antiepileptic drugs in children and adolescents with newly diagnosed and chronic epilepsy. Epilepsy Research, 2012, 100, 142-151.	0.8	13
113	Obese children suffer more often from migraine. Acta Paediatrica, International Journal of Paediatrics, 2012, 101, e416-21.	0.7	28
114	Antiepileptic drugs, sex hormones, and PCOS. Epilepsia, 2011, 52, 199-211.	2.6	78
115	Nonalcoholic fatty liver disease in adolescents receiving valproic acid. Epilepsy and Behavior, 2011, 20, 382-385.	0.9	35
116	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
117	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
118	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
119	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
120	Long-term outcome of epilepsy in Kabuki syndrome. Seizure: the Journal of the British Epilepsy Association, 2011, 20, 650-654.	0.9	10
121	Update on rufinamide in childhood epilepsy. Neuropsychiatric Disease and Treatment, 2011, 7, 399.	1.0	19
122	Weight gain following treatment with valproic acid: pathogenetic mechanisms and clinical implications. Obesity Reviews, 2011, 12, e32-43.	3.1	109
123	Rufinamide in refractory childhood epileptic encephalopathies other than Lennox-Gastaut syndrome. European Journal of Neurology, 2011, 18, 246-251.	1.7	40
124	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
125	The ketogenic diet for Dravet syndrome and other epileptic encephalopathies: An Italian consensus. Epilepsia, 2011, 52, 83-89.	2.6	37
126	Low penetrance and effect on protein secretion of LGI1 mutations causing autosomal dominant lateral temporal epilepsy. Epilepsia, 2011, 52, 1258-1264.	2.6	26

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127	Cognitive and linguistic abnormalities in benign childhood epilepsy with centrotemporal spikes. Acta Paediatrica, International Journal of Paediatrics, 2011, 100, 768-772.	0.7	30
128	Hormonal and reproductive disturbances in epileptic male patients: Emerging issues. Reproductive Toxicology, 2011, 31, 519-527.	1.3	37
129	Treatment of malignant migrating partial epilepsy of infancy with rufinamide: report of five cases. Epileptic Disorders, 2011, 13, 18-21.	0.7	55
130	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
131	Benign convulsions associated with mild gastroenteritis: A multicenter clinical study. Epilepsy Research, 2011, 93, 107-114.	0.8	57
132	Association of intronic variants of the KCNAB1 gene with lateral temporal epilepsy. Epilepsy Research, 2011, 94, 110-116.	0.8	9
133	Pharmacotherapy for children and adolescents with epilepsy. Expert Opinion on Pharmacotherapy, 2011, 12, 175-194.	0.9	30
134	West syndrome associated with 14q12 duplications harboring FOXG1. Neurology, 2011, 76, 1600-1602.	1.5	49
135	Ketogenic diet for the treatment of catastrophic epileptic encephalopathies in childhood. European Journal of Paediatric Neurology, 2010, 14, 229-234.	0.7	49
136	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
137	The metabolic syndrome in overweight epileptic patients treated with valproic acid. Epilepsia, 2010, 51, 268-273.	2.6	109
138	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
139	Bone and calcium metabolism and antiepileptic drugs. Clinical Neurology and Neurosurgery, 2010, 112, 1-10.	0.6	206
140	Simultaneous Onset of Infantile Spasms in Monozygotic Twins. Pediatric Neurology, 2010, 43, 127-130.	1.0	6
141	Levetiracetam in submaximal subcutaneous pentylentetrazol-induced seizures in rats. Seizure: the Journal of the British Epilepsy Association, 2010, 19, 296-299.	0.9	24
142	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
143	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
144	Catamenial epilepsy: hormonal aspects. Gynecological Endocrinology, 2010, 26, 783-790.	0.7	23

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145	Startle Epilepsy Associated with Infantile Hemiplegia (SEIH): Video-Polygraphic Features and Long-Term Outcome. Neuropediatrics, 2009, 40, 97-100.	0.3	2
146	Clinical features of psychogenic non-epileptic seizures in prepubertal and pubertal patients with idiopathic epilepsy. Neurological Sciences, 2009, 30, 319-323.	0.9	38
147	Bone mineral density in children, adolescents, and young adults with epilepsy. Epilepsia, 2009, 50, 2140-2146.	2.6	80
148	Malignant migrating partial seizures in infancy: An epilepsy syndrome of unknown etiology. Epilepsia, 2009, 50, 49-51.	2.6	101
149	Zonisamide in children and young adults with refractory epilepsy: An open label, multicenter Italian study. Epilepsy Research, 2009, 83, 112-116.	0.8	25
150	Epilepsy, sex hormones and antiepileptic drugs in female patients. Expert Review of Neurotherapeutics, 2009, 9, 1803-1814.	1.4	26
151	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
152	Epilepsy and electroencephalographic anomalies in chromosome 2 aberrations. Epilepsy Research, 2008, 79, 63-70.	0.8	13
153	Levetiracetam in juvenile myoclonic epilepsy: longâ€term efficacy in newly diagnosed adolescents. Developmental Medicine and Child Neurology, 2008, 50, 29-32.	1.1	45
154	Levetiracetam in absence epilepsy. Developmental Medicine and Child Neurology, 2008, 50, 850-853.	1.1	51
155	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
156	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
157	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
158	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
159	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
160	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
161	Bone Mineral Density in Angelman Syndrome. Pediatric Neurology, 2007, 37, 411-416.	1.0	11
162	Topiramate in frontal lobe epilepsy. Acta Neurologica Scandinavica, 2007, 115, 132-135.	1.0	5

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163	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
164	Temporal lobe dual pathology in malignant migrating partial seizures in infancy. Epileptic Disorders, 2007, 9, 145-148.	0.7	17
165	Tuberous sclerosis complex and hydrosyringomielia: Report of two cases. European Journal of Paediatric Neurology, 2006, 10, 37-40.	0.7	1
166	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
167	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
168	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
169	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
170	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
171	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
172	Short-term Nonhormonal and Nonsteroid Treatment in West Syndrome. Epilepsia, 2004, 45, 887-887.	2.6	0
173	Lamotrigine as first-line drug in childhood absence epilepsy: a clinical and neurophysiological study. Brain and Development, 2004, 26, 26-29.	0.6	38
174	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
175	Unusual compulsive motor activity during treatment with clothiapine in a mentally retarded adolescent. Brain and Development, 2004, 26, 409-411.	0.6	4
176	Levetiracetam during 1-year follow-up in children, adolescents, and young adults with refractory epilepsy. Epilepsy Research, 2004, 59, 35-42.	0.8	46
177	Treatment of Partial Seizures in Childhood. CNS Drugs, 2004, 18, 133-156.	2.7	23
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