

Tiziana Granata

List of Publications by Year in descending order

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Version: 2024-02-01

174
papers

10,753
citations

41627

51
h-index

40945

97
g-index

177
all docs

177
docs citations

177
times ranked

10446
citing authors

#	ARTICLE	IF	CITATIONS
1	Brain Inflammation in Epilepsy: Experimental and Clinical Evidence. <i>Epilepsia</i> , 2005, 46, 1724-1743.	2.6	921
2	Pathogenesis, diagnosis and treatment of Rasmussen encephalitis: A European consensus statement. <i>Brain</i> , 2005, 128, 454-471.	3.7	490
3	Seizure-Promoting Effect of Blood-Brain Barrier Disruption. <i>Epilepsia</i> , 2007, 48, 732-742.	2.6	442
4	Congenital malformations due to antiepileptic drugs. <i>Epilepsy Research</i> , 1999, 33, 145-158.	0.8	351
5	De novo mutations in ATP1A3 cause alternating hemiplegia of childhood. <i>Nature Genetics</i> , 2012, 44, 1030-1034.	9.4	345
6	Antagonism of peripheral inflammation reduces the severity of status epilepticus. <i>Neurobiology of Disease</i> , 2009, 33, 171-181.	2.1	270
7	Spectrum of <i>SCN1A</i> mutations in severe myoclonic epilepsy of infancy. <i>Neurology</i> , 2003, 60, 1961-1967.	1.5	241
8	Ultra-Rare Genetic Variation in the Epilepsies: A Whole-Exome Sequencing Study of 17,606 Individuals. <i>American Journal of Human Genetics</i> , 2019, 105, 267-282.	2.6	237
9	Fenfluramine hydrochloride for the treatment of seizures in Dravet syndrome: a randomised, double-blind, placebo-controlled trial. <i>Lancet</i> , The, 2019, 394, 2243-2254.	6.3	227
10	Band heterotopia: Correlation of outcome with magnetic resonance imaging parameters. <i>Annals of Neurology</i> , 1994, 36, 609-617.	2.8	206
11	Blood-brain barrier dysfunction and epilepsy: Pathophysiologic role and therapeutic approaches. <i>Epilepsia</i> , 2012, 53, 1877-1886.	2.6	199
12	Inflammatory pathways of seizure disorders. <i>Trends in Neurosciences</i> , 2014, 37, 55-65.	4.2	196
13	Malformations in Offspring of Women with Epilepsy: A Prospective Study. <i>Epilepsia</i> , 1999, 40, 1231-1236.	2.6	181
14	Experience with immunomodulatory treatments in Rasmussen's encephalitis. <i>Neurology</i> , 2003, 61, 1807-1810.	1.5	161
15	Rasmussen encephalitis. <i>Handbook of Clinical Neurology</i> / Edited By P J Vinken and G W Bruyn, 2013, 111, 511-519.	1.0	158
16	Antibodies against GluR3 peptides are not specific for Rasmussen's encephalitis but are also present in epilepsy patients with severe, early onset disease and intractable seizures. <i>Journal of Neuroimmunology</i> , 2002, 131, 179-185.	1.1	151
17	Periventricular Nodular Heterotopia: Classification, Epileptic History, and Genesis of Epileptic Discharges. <i>Epilepsia</i> , 2006, 47, 86-97.	2.6	150
18	Familial schizencephaly associated with <i>EMX2</i> mutation. <i>Neurology</i> , 1997, 48, 1403-1406.	1.5	145

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19	Epileptic phenotypes associated with mitochondrial disorders. <i>Neurology</i> , 2001, 56, 1340-1346.	1.5	143
20	Rasmussen's encephalitis. <i>Neurology</i> , 2003, 60, 422-425.	1.5	137
21	Efficacy of Anti-Inflammatory Therapy in a Model of Acute Seizures and in a Population of Pediatric Drug Resistant Epileptics. <i>PLoS ONE</i> , 2011, 6, e18200.	1.1	130
22	Polymicrogyria and deletion 22q11.2 syndrome: Window to the etiology of a common cortical malformation. <i>American Journal of Medical Genetics, Part A</i> , 2006, 140A, 2416-2425.	0.7	125
23	Clinical profile of patients with ATP1A3 mutations in Alternating Hemiplegia of Childhood—a study of 155 patients. <i>Orphanet Journal of Rare Diseases</i> , 2015, 10, 123.	1.2	117
24	Diagnostic imaging in 13 cases of Rasmussen's encephalitis: can early MRI suggest the diagnosis?. <i>Neuroradiology</i> , 2003, 45, 171-183.	1.1	116
25	An open-label trial of levetiracetam in severe myoclonic epilepsy of infancy. <i>Neurology</i> , 2007, 69, 250-254.	1.5	115
26	Hemispherotomy and functional hemispherectomy: Indications and outcome. <i>Epilepsy Research</i> , 2010, 89, 104-112.	0.8	115
27	Intellectual and language findings and their relationship to EEG characteristics in benign childhood epilepsy with centrotemporal spikes. <i>Epilepsy and Behavior</i> , 2007, 10, 278-285.	0.9	113
28	Adult-Onset Rasmussen's Encephalitis: Anatomical-Electrographic-Clinical Features of 7 Italian Cases. <i>Epilepsia</i> , 2006, 47, 41-46.	2.6	108
29	Long-term selective IgG immunoadsorption improves Rasmussen's encephalitis. <i>Neurology</i> , 1998, 51, 302-305.	1.5	106
30	Somatic and germline mosaicisms in Severe Myoclonic Epilepsy of Infancy. <i>Biochemical and Biophysical Research Communications</i> , 2006, 341, 489-493.	1.0	102
31	Cognitive development in Dravet syndrome: A retrospective, multicenter study of 26 patients. <i>Epilepsia</i> , 2011, 52, 386-392.	2.6	99
32	Suppressive Efficacy by a Commercially Available Blue Lens on PPR in 610 Photosensitive Epilepsy Patients. <i>Epilepsia</i> , 2006, 47, 529-533.	2.6	96
33	<i>HCN1</i> mutation spectrum: from neonatal epileptic encephalopathy to benign generalized epilepsy and beyond. <i>Brain</i> , 2018, 141, 3160-3178.	3.7	96
34	Dravet syndrome: Early clinical manifestations and cognitive outcome in 37 Italian patients. <i>Brain and Development</i> , 2010, 32, 71-77.	0.6	94
35	Clinical and genetic factors predicting Dravet syndrome in infants with <i>SCN1A</i> mutations. <i>Neurology</i> , 2017, 88, 1037-1044.	1.5	93
36	Topical Review: Schizencephaly: Clinical Spectrum, Epilepsy, and Pathogenesis. <i>Journal of Child Neurology</i> , 2005, 20, 313-318.	0.7	92

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37	Polygenic burden in focal and generalized epilepsies. <i>Brain</i> , 2019, 142, 3473-3481.	3.7	90
38	Brain MRI Findings in Severe Myoclonic Epilepsy in Infancy and Genotype?Phenotype Correlations. <i>Epilepsia</i> , 2007, 48, 1092-1096.	2.6	89
39	<i>VAR2</i> and <i>TARS2</i> Mutations in Patients with Mitochondrial Encephalomyopathies. <i>Human Mutation</i> , 2014, 35, 983-989.	1.1	86
40	Anakinra usage in febrile infection related epilepsy syndrome: an international cohort. <i>Annals of Clinical and Translational Neurology</i> , 2020, 7, 2467-2474.	1.7	80
41	Immune-mediated epilepsies. <i>Epilepsia</i> , 2011, 52, 5-11.	2.6	76
42	Positive response to immunomodulatory therapy in an adult patient with Rasmussen's encephalitis. <i>Neurology</i> , 2001, 56, 248-250.	1.5	74
43	Management of the patient with medically refractory epilepsy. <i>Expert Review of Neurotherapeutics</i> , 2009, 9, 1791-1802.	1.4	72
44	The Movement disorder associated with NMDAR antibody-encephalitis is complex and characteristic: an expert video-rating study. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2019, 90, 724-726.	0.9	71
45	Periventricular Nodular Heterotopia: Epileptogenic Findings. <i>Epilepsia</i> , 1997, 38, 1173-1182.	2.6	66
46	A Number of Schizencephaly Patients Including 2 Brothers Are Heterozygous for Germline Mutations in the Homeobox Gene <i>EMX2</i> . <i>European Journal of Human Genetics</i> , 1997, 5, 186-190.	1.4	66
47	Schizencephaly: Neuroradiologic and Epileptologic Findings. <i>Epilepsia</i> , 1996, 37, 1185-1193.	2.6	63
48	Malformations in offspring of 305 epileptic women: a prospective study. <i>Acta Neurologica Scandinavica</i> , 1992, 85, 204-207.	1.0	63
49	Focal seizures with affective symptoms are a major feature of <i>PCDH19</i> gene-related epilepsy. <i>Epilepsia</i> , 2012, 53, 2111-2119.	2.6	63
50	Xp22.3 genomic deletions involving the <i>CDKL5</i> gene in girls with early onset epileptic encephalopathy. <i>Epilepsia</i> , 2010, 51, 647-654.	2.6	60
51	The Etiological Role of Blood-Brain Barrier Dysfunction in Seizure Disorders. <i>Cardiovascular Psychiatry and Neurology</i> , 2011, 2011, 1-9.	0.8	58
52	Neuronal migration disorders and epilepsy: a morphological analysis of three surgically treated patients. <i>Epilepsy Research</i> , 1996, 26, 49-58.	0.8	51
53	A role for inflammation in status epilepticus is revealed by a review of current therapeutic approaches. <i>Epilepsia</i> , 2013, 54, 30-32.	2.6	51
54	Perisylvian, including insular, childhood epilepsy: Presurgical workup and surgical outcome. <i>Epilepsia</i> , 2017, 58, 1360-1369.	2.6	51

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55	Clinical course and variability of non-Rasmussen, nonstroke motor and sensory epilepsy partialis continua: A European survey and analysis of 65 cases. <i>Epilepsia</i> , 2011, 52, 1168-1176.	2.6	50
56	ILAE Classification of Epilepsies: Its Applicability and Practical Value of Different Diagnostic Categories. <i>Epilepsia</i> , 1996, 37, 1051-1059.	2.6	49
57	West syndrome associated with 14q12 duplications harboring FOXP1. <i>Neurology</i> , 2011, 76, 1600-1602.	1.5	49
58	Familial Occurrence of Febrile Seizures and Epilepsy in Severe Myoclonic Epilepsy of Infancy (SMEI) Patients with SCN1A Mutations. <i>Epilepsia</i> , 2006, 47, 1629-1635.	2.6	48
59	A novel de novo HCN1 loss-of-function mutation in genetic generalized epilepsy causing increased neuronal excitability. <i>Neurobiology of Disease</i> , 2018, 118, 55-63.	2.1	47
60	Epilepsy subtype-specific copy number burden observed in a genome-wide study of 17,458 subjects. <i>Brain</i> , 2020, 143, 2106-2118.	3.7	47
61	Did the COVID-19 pandemic silence the needs of people with epilepsy?. <i>Epileptic Disorders</i> , 2020, 22, 439-442.	0.7	46
62	Epileptic and imaging findings in perinatal hypoxic-ischemic encephalopathy with ulegyria. <i>Epilepsy Research</i> , 2003, 55, 235-243.	0.8	45
63	Variant late infantile ceroid lipofuscinoses associated with novel mutations in CLN6. <i>Biochemical and Biophysical Research Communications</i> , 2009, 379, 892-897.	1.0	45
64	Overview of presurgical assessment and surgical treatment of epilepsy from the Italian League Against Epilepsy. <i>Epilepsia</i> , 2013, 54, 35-48.	2.6	45
65	Trends in pediatric epilepsy surgery in Europe between 2008 and 2015: Country-, center-, and age-specific variation. <i>Epilepsia</i> , 2020, 61, 216-227.	2.6	44
66	Rasmussen's syndrome. <i>Neurological Sciences</i> , 2003, 24, s239-s243.	0.9	43
67	Changes in Unbound and Total Valproic Acid Concentrations After Replacement of Carbamazepine with Oxcarbazepine. <i>Therapeutic Drug Monitoring</i> , 1992, 14, 376-379.	1.0	41
68	Partial Seizures Associated with Antiphospholipid Antibodies in Childhood. <i>Neuropediatrics</i> , 1998, 29, 249-254.	0.3	41
69	Revelation of a Novel CLN5 Mutation in Early Juvenile Neuronal Ceroid Lipofuscinosis. <i>Neuropediatrics</i> , 2007, 38, 46-49.	0.3	41
70	Hemispherotomy in Rasmussen encephalitis: Long-term outcome in an Italian series of 16 patients. <i>Epilepsy Research</i> , 2014, 108, 1106-1119.	0.8	41
71	Outcome of childhood-onset epilepsy from adolescence to adulthood: Transition issues. <i>Epilepsy and Behavior</i> , 2017, 69, 161-169.	0.9	41
72	Electroclinical spectrum of the neuronal ceroid lipofuscinoses associated with CLN6 mutations. <i>Neurology</i> , 2015, 85, 316-324.	1.5	40

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73	Relapse risk factors in anti-N-methyl-D-aspartate receptor encephalitis. <i>Developmental Medicine and Child Neurology</i> , 2019, 61, 1101-1107.	1.1	40
74	Lamotrigine Plasma Concentrations in Children and Adults. <i>Therapeutic Drug Monitoring</i> , 1997, 19, 620-627.	1.0	40
75	Defining the electroclinical phenotype and outcome of PCDH19-related epilepsy: A multicenter study. <i>Epilepsia</i> , 2018, 59, 2260-2271.	2.6	39
76	Cognitive and neuropsychological evolution in children with anti-NMDAR encephalitis. <i>Journal of Neurology</i> , 2016, 263, 765-771.	1.8	38
77	Efficacy and safety of Fenfluramine hydrochloride for the treatment of seizures in Dravet syndrome: A real-world study. <i>Epilepsia</i> , 2020, 61, 2405-2414.	2.6	37
78	Burst Suppression and Impairment of Neocortical Ontogenesis: Electroclinical and Neuropathologic Findings in Two Infants with Early Myoclonic Encephalopathy. <i>Epilepsia</i> , 1993, 34, 800-808.	2.6	36
79	Small 6q16.1 Deletions Encompassing POU3F2 Cause Susceptibility to Obesity and Variable Developmental Delay with Intellectual Disability. <i>American Journal of Human Genetics</i> , 2016, 98, 363-372.	2.6	36
80	Lamotrigine in Resistant Childhood Epilepsy. <i>Neuropediatrics</i> , 1993, 24, 332-336.	0.3	35
81	Transporters in Drug-Refractory Epilepsy: Clinical Significance. <i>Clinical Pharmacology and Therapeutics</i> , 2010, 87, 13-15.	2.3	35
82	Sub-genic intolerance, ClinVar, and the epilepsies: A whole-exome sequencing study of 29,165 individuals. <i>American Journal of Human Genetics</i> , 2021, 108, 965-982.	2.6	35
83	Intrauterine growth in the offspring of epileptic mothers. <i>Acta Neurologica Scandinavica</i> , 1992, 86, 555-557.	1.0	34
84	Spectral properties of EEG fast activity ictal discharges associated with infantile spasms. <i>Clinical Neurophysiology</i> , 1999, 110, 593-603.	0.7	34
85	Pediatric NMDAR encephalitis: A single center observation study with a closer look at movement disorders. <i>European Journal of Paediatric Neurology</i> , 2018, 22, 301-307.	0.7	34
86	Transition into adulthood: Tuberous sclerosis complex, Sturge-Weber syndrome, and Rasmussen encephalitis. <i>Epilepsia</i> , 2014, 55, 29-33.	2.6	32
87	HCN ion channels and accessory proteins in epilepsy: genetic analysis of a large cohort of patients and review of the literature. <i>Epilepsy Research</i> , 2019, 153, 49-58.	0.8	32
88	Single-Dose Pharmacokinetics of Lamotrigine in Children: Influence of Age and Antiepileptic Comedication. <i>Therapeutic Drug Monitoring</i> , 2001, 23, 217-222.	1.0	31
89	Neonatal developmental and epileptic encephalopathy due to autosomal recessive variants in <i>SLC13A5</i> gene. <i>Epilepsia</i> , 2020, 61, 2474-2485.	2.6	31
90	Faulty cardiac repolarization reserve in alternating hemiplegia of childhood broadens the phenotype. <i>Brain</i> , 2015, 138, 2859-2874.	3.7	30

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91	Paroxysmal dyskinesias in childhood. <i>Pediatric Neurology</i> , 2003, 28, 168-172.	1.0	27
92	Impaired surface $\alpha 2 \beta 3$ GABAA receptor expression in familial epilepsy due to a GABRG2 frameshift mutation. <i>Neurobiology of Disease</i> , 2013, 50, 135-141.	2.1	27
93	Labeling of rat neurons by anti-GluR3 IgG from patients with Rasmussen encephalitis. <i>Neurology</i> , 2001, 57, 324-327.	1.5	25
94	Similar binding to glutamate receptors by Rasmussen and partial epilepsy patients' sera. <i>Neurology</i> , 2002, 59, 1998-2001.	1.5	25
95	Electroclinical Features and Long-Term Outcome of Cryptogenic Epilepsy in Children with Down Syndrome. <i>Journal of Pediatrics</i> , 2013, 163, 1754-1758.	0.9	25
96	A Loss-of-Function HCN4 Mutation Associated With Familial Benign Myoclonic Epilepsy in Infancy Causes Increased Neuronal Excitability. <i>Frontiers in Molecular Neuroscience</i> , 2018, 11, 269.	1.4	25
97	Electroencephalographic Recordings of Focal Seizures in Patients Affected by Periventricular Nodular Heterotopia: Role of the Heterotopic Nodules in the Genesis of Epileptic Discharges. <i>Journal of Child Neurology</i> , 2005, 20, 369-377.	0.7	24
98	A de novo heterozygous mutation in KCNC2 gene implicated in severe developmental and epileptic encephalopathy. <i>European Journal of Medical Genetics</i> , 2020, 63, 103848.	0.7	24
99	Joubert syndrome with bilateral polymicrogyria: Clinical and neuropathological findings in two brothers. <i>American Journal of Medical Genetics, Part A</i> , 2009, 149A, 1511-1515.	0.7	22
100	The blood-brain barrier hypothesis in drug resistant epilepsy. <i>Brain</i> , 2012, 135, e211-e211.	3.7	22
101	Refractory Absence Epilepsy and Glut1 Deficiency Syndrome: A New Case Report and Literature Review. <i>Neuropediatrics</i> , 2014, 45, 328-332.	0.3	22
102	Epilepsy in type 1 Chiari malformation. <i>Neurological Sciences</i> , 2011, 32, 303-306.	0.9	21
103	FCD Type II and mTOR pathway: Evidence for different mechanisms involved in the pathogenesis of dysmorphic neurons. <i>Epilepsy Research</i> , 2017, 129, 146-156.	0.8	21
104	Kv7.3 Compound Heterozygous Variants in Early Onset Encephalopathy Reveal Additive Contribution of C-Terminal Residues to PIP2-Dependent K ⁺ Channel Gating. <i>Molecular Neurobiology</i> , 2018, 55, 7009-7024.	1.9	21
105	No evidence of GABRG2 mutations in severe myoclonic epilepsy of infancy. <i>Epilepsy Research</i> , 2003, 53, 196-200.	0.8	20
106	ICTAL EEG Fast Activity in West Syndrome: From Onset to Outcome. <i>Epilepsia</i> , 2007, 48, 2101-2110.	2.6	20
107	Vagus nerve stimulation for drug-resistant Epilepsia Partialis Continua: Report of four cases. <i>Epilepsy Research</i> , 2013, 107, 163-171.	0.8	20
108	Dravet syndrome: Early electroclinical findings and long-term outcome in adolescents and adults. <i>Epilepsia</i> , 2019, 60, S49-S58.	2.6	20

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109	Progressive Myoclonus Epilepsies. <i>Neurology: Genetics</i> , 2021, 7, e641.	0.9	20
110	A causality algorithm to guide diagnosis and treatment of catatonia due to autoimmune conditions in children and adolescents. <i>Schizophrenia Research</i> , 2018, 200, 68-76.	1.1	19
111	An Italian multicentre study of perampanel in progressive myoclonus epilepsies. <i>Epilepsy Research</i> , 2019, 156, 106191.	0.8	19
112	Cardiac phenotype in <i>ATP1A3</i> -related syndromes. <i>Neurology</i> , 2020, 95, e2866-e2879.	1.5	19
113	Linguistic Development in a Patient with Landau-Kleffner Syndrome: A Nine-Year Follow-Up. <i>Neuropediatrics</i> , 1995, 26, 19-25.	0.3	18
114	Long-term outcome after limited cortical resections in two cases of adult-onset Rasmussen encephalitis. <i>Epilepsia</i> , 2014, 55, e38-e43.	2.6	18
115	Variable course of Unverricht-Lundborg disease. <i>Neurology</i> , 2017, 89, 1691-1697.	1.5	18
116	Screening of SLC2A1 in a large cohort of patients suspected for Glut1 deficiency syndrome: identification of novel variants and associated phenotypes. <i>Journal of Neurology</i> , 2019, 266, 1439-1448.	1.8	18
117	Encephalopathies with intracranial calcification in children: clinical and genetic characterization. <i>Orphanet Journal of Rare Diseases</i> , 2018, 13, 135.	1.2	17
118	Comparative Pharmacokinetic Study of Chewable and Conventional Carbamazepine in Children. <i>Epilepsia</i> , 1993, 34, 158-160.	2.6	16
119	Scurvy hidden behind neuropsychiatric symptoms. <i>Neurological Sciences</i> , 2011, 32, 1091-1093.	0.9	16
120	Alternating Hemiplegia and Epilepsia Partialis Continua: A new phenotype for a novel compound TBC1D24 mutation. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2017, 47, 71-73.	0.9	16
121	Gait abnormalities in people with Dravet syndrome: A cross-sectional multi-center study. <i>European Journal of Paediatric Neurology</i> , 2019, 23, 808-818.	0.7	16
122	Progressive myoclonus epilepsy caused by a gain-of-function KCNA2 mutation. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2019, 65, 106-108.	0.9	16
123	Neuromuscular syndrome associated with the 3291T>C mutation of mitochondrial DNA: a second case. <i>Neuromuscular Disorders</i> , 2000, 10, 415-418.	0.3	15
124	Subcortical nodular heterotopia: a functional MRI and somatosensory evoked potentials study. <i>Neurological Sciences</i> , 2004, 25, 225-229.	0.9	15
125	Mitochondrial Complex III Deficiency Caused by TTC19 Defects: Report of a Novel Mutation and Review of Literature. <i>JIMD Reports</i> , 2015, 22, 115-120.	0.7	15
126	Epileptic phenotypes in children with early-onset mitochondrial diseases. <i>Acta Neurologica Scandinavica</i> , 2019, 140, 184-193.	1.0	15

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127	The Contribution of Tertiary Centers to the Quality of the Diagnosis and Treatment of Epilepsy. <i>Epilepsia</i> , 1997, 38, 1338-1343.	2.6	14
128	Clinical and genetic features of paroxysmal kinesigenic dyskinesia in Italian patients. <i>European Journal of Paediatric Neurology</i> , 2016, 20, 152-157.	0.7	14
129	Focal seizure, focal dyskinesia, or both? A complex motor phenomenon reveals anti-NMDAR encephalitis. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2015, 27, 16-18.	0.9	13
130	Topical Review: Schizencephaly: Clinical Spectrum, Epilepsy, and Pathogenesis. <i>Journal of Child Neurology</i> , 2004, 19, 313-318.	0.7	12
131	Comprehensive care of children with Dravet syndrome. <i>Epilepsia</i> , 2011, 52, 90-94.	2.6	12
132	Epilepsy and NREM-parasomnia: A complex and reciprocal relationship. <i>Sleep Medicine</i> , 2012, 13, 442-444.	0.8	12
133	Electroencephalographic (EEG) Photoparoxysmal Responses Under 5 Years of Age. <i>Journal of Child Neurology</i> , 2015, 30, 1824-1830.	0.7	12
134	Movement-activated cortical myoclonus in Dravet syndrome. <i>Epilepsy Research</i> , 2017, 130, 47-52.	0.8	12
135	Early clinical and EEG findings associated with the outcome in childhood absence epilepsy. <i>Epilepsy and Behavior</i> , 2019, 98, 273-278.	0.9	12
136	Double cortex syndrome: Electroclinical study of three cases. <i>Italian Journal of Neurological Sciences</i> , 1994, 15, 15-23.	0.1	11
137	Electroclinical findings and long-term outcomes in epileptic patients with inv dup (15). <i>Acta Neurologica Scandinavica</i> , 2018, 137, 575-581.	1.0	11
138	Psychiatric autoimmune conditions in children and adolescents: Is catatonia a severity marker?. <i>Progress in Neuro-Psychopharmacology and Biological Psychiatry</i> , 2021, 104, 110028.	2.5	11
139	Do the functional properties of HCN1 mutants correlate with the clinical features in epileptic patients?. <i>Progress in Biophysics and Molecular Biology</i> , 2021, 166, 147-155.	1.4	11
140	White matter and cerebellar involvement in alternating hemiplegia of childhood. <i>Journal of Neurology</i> , 2020, 267, 1300-1311.	1.8	10
141	Periventricular nodular heterotopia. <i>Handbook of Clinical Neurology</i> / Edited By P J Vinken and G W Bruyn, 2007, 87, 177-189.	1.0	9
142	Association of intronic variants of the KCNAB1 gene with lateral temporal epilepsy. <i>Epilepsy Research</i> , 2011, 94, 110-116.	0.8	9
143	Symptomatic and presumed symptomatic focal epilepsies in childhood: An observational, prospective multicentre study. <i>Epilepsia</i> , 2016, 57, 1808-1816.	2.6	9
144	Reflex myoclonic epilepsy in infancy: A multicenter clinical study. <i>Epilepsy Research</i> , 2013, 103, 237-244.	0.8	8

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145	Focal seizures versus epileptic spasms in children with focal cortical dysplasia and epilepsy onset in the first year. <i>Epilepsy Research</i> , 2015, 109, 203-209.	0.8	8
146	Electroclinical features of epilepsy monosomy 1p36 syndrome and their implications. <i>Acta Neurologica Scandinavica</i> , 2018, 138, 523-530.	1.0	8
147	Early add-on immunoglobulin administration in Rasmussen encephalitis: The hypothesis of neuroimmunomodulation. <i>Medical Hypotheses</i> , 2011, 77, 917-920.	0.8	7
148	Gabapentin treatment in a patient with KCNQ2 developmental epileptic encephalopathy. <i>Pharmacological Research</i> , 2020, 160, 105200.	3.1	7
149	Multicenter prospective longitudinal study in 34 patients with Dravet syndrome: Neuropsychological development in the first six years of life. <i>Brain and Development</i> , 2021, 43, 419-430.	0.6	7
150	Basal Ganglia Dymorphism in Patients With Aicardi Syndrome. <i>Neurology</i> , 2021, 96, e1319-e1333.	1.5	6
151	Inflammation in pediatric epilepsies: Update on clinical features and treatment options. <i>Epilepsy and Behavior</i> , 2022, 131, 107959.	0.9	6
152	Cortico-muscular and cortico-cortical coherence changes resulting from Perampanel treatment in patients with cortical myoclonus. <i>Clinical Neurophysiology</i> , 2021, 132, 1057-1063.	0.7	6
153	Serial electrophysiological studies of the visual pathway in patients treated with vigabatrin. <i>International Congress Series</i> , 2005, 1278, 41-44.	0.2	5
154	Metabolic and degenerative disorders. <i>Handbook of Clinical Neurology</i> / Edited By P J Vinken and G W Bruyn, 2012, 108, 485-511.	1.0	5
155	Focal status and acute encephalopathy in a 13-year-old boy with de novo DNM1L mutation: Video-polygraphic pattern and clues for differential diagnosis. <i>Brain and Development</i> , 2021, 43, 644-651.	0.6	5
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