Roberto Michelucci

List of Publications by Year
in descending order

153	7,211	46918 47	71532 76
papers	citations	h-index	g-index

156	156	156	6771
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Risk of hospitalization and death for <scp>COVID</scp>â€19 in persons with epilepsy over a 20â€month period: The <scp>EpiLink</scp> Bologna cohort, Italy. Epilepsia, 2022, 63, 2279-2289.	2.6	8
2	Intravenous immunoglobulin therapy in COVID-19-related encephalopathy. Journal of Neurology, 2021, 268, 2671-2675.	1.8	35
3	Progressive myoclonus epilepsiesâ€”Residual unsolved cases have marked genetic heterogeneity including dolichol-dependent protein glycosylation pathway genes. American Journal of Human Genetics, 2021, 108, 722-738.	2.6	41
4	Italian cohort of Lafora disease: Clinical features, disease evolution, and genotype-phenotype correlations. Journal of the Neurological Sciences, 2021, 424, 117409.	0.3	11
5	fMRI-Based Effective Connectivity in Surgical Remediable Epilepsies: A Pilot Study. Brain Topography, 2021, 34, 632-650.	0.8	6
6	Progressive Myoclonus Epilepsies. Neurology: Genetics, 2021, 7, e641.	0.9	20
7	EEG findings in COVID-19 related encephalopathy. Clinical Neurophysiology, 2020, 131, 2265-2267.	0.7	31
8	Autosomal dominant lateral temporal lobe epilepsy associated with a novel reelin mutation. Epileptic Disorders, 2020, 22, 443-448.	0.7	8
9	Early neurological manifestations of hospitalized COVID-19 patients. Neurological Sciences, 2020, 41, 2029-2031.	0.9	72
10	Advances in genetic testing and optimization of clinical management in children and adults with epilepsy. Expert Review of Neurotherapeutics, 2020, 20, 251-269.	1.4	45
11	Treatment with metformin in twelve patients with Lafora disease. Orphanet Journal of Rare Diseases, 2019, 14, 149.	1.2	34
12	The best evidence for progressive myoclonic epilepsy: A pathway to precision therapy. Seizure: the Journal of the British Epilepsy Association, 2019, 71, 247-257.	0.9	57
13	An Italian multicentre study of perampanel in progressive myoclonus epilepsies. Epilepsy Research, 2019, 156, 106191.	0.8	19
14	Hypertension, seizures, and epilepsy: a review on pathophysiology and management. Neurological Sciences, 2019, 40, 1775-1783.	0.9	51
15	LGI1 tumor tissue expression and serum autoantibodies in patients with primary malignant glioma. Clinical Neurology and Neurosurgery, 2018, 170, 27-33.	0.6	3
16	Mutations in MICALâ€1cause autosomalâ€dominant lateral temporal epilepsy. Annals of Neurology, 2018, 83, 483-493.	2.8	25
17	CNTNAP2 mutations and autosomal dominant epilepsy with auditory features. Epilepsy Research, 2018, 139, 51-53.	0.8	3
18	Is autopsy tissue a valid control for epilepsy surgery tissue in micro<scp>RNA</scp> studies?. Epilepsia Open, 2017, 2, 90-95.	1.3	11

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19	The clinical phenotype of autosomal dominant lateral temporal lobe epilepsy related to reelin mutations. Epilepsy and Behavior, 2017, 68, 103-107.	0.9	31
20	Coexistence of meningoencephalocele and hippocampal sclerosis: a new type of dual pathology. Acta Neurochirurgica, 2017, 159, 767-769.	0.9	4
21	Myoclonus epilepsy and ataxia due to <scp>KCNC</scp>1 mutation: Analysis of 20 cases and <scp>K</scp>⁺ channel properties. Annals of Neurology, 2017, 81, 677-689.	2.8	69
22	Variable course of Unverricht-Lundborg disease. Neurology, 2017, 89, 1691-1697.	1.5	18
23	Relationship among clinical, pathological and bio-molecular features in low-grade epilepsy-associated neuroepithelial tumors. Journal of Clinical Neuroscience, 2017, 44, 158-163.	0.8	15
24	Myoclonus and seizures in progressive myoclonus epilepsies: pharmacology and therapeutic trials. Epileptic Disorders, 2016, 18, 145-153.	0.7	48
25	In response: <scp>DEPDC</scp>5 mutations in epilepsy with auditory features. Epilepsia, 2016, 57, 336-336.	2.6	1
26	Cerebrospinal fluid amounts of HLA-G in dimeric form are strongly associated to patients with MRI inactive multiple sclerosis. Multiple Sclerosis Journal, 2016, 22, 245-249.	1.4	11
27	Pathology-Based Approach to Seizure Outcome After Surgery for Pharmacoresistant Medial Temporal Lobe Epilepsy. World Neurosurgery, 2016, 90, 448-453.	0.7	8
28	Spinal muscular atrophy associated with progressive myoclonic epilepsy: A rare condition caused by mutations in <scp>ASAH</scp>1. Epilepsia, 2015, 56, 692-698.	2.6	33
29	MicroRNA profiles in hippocampal granule cells and plasma of rats with pilocarpine-induced epilepsy â€“ comparison with human epileptic samples. Scientific Reports, 2015, 5, 14143.	1.6	101
30	<scp>DEPDC</scp>5 mutations are not a frequent cause of familial temporal lobe epilepsy. Epilepsia, 2015, 56, e168-71.	2.6	37
31	Heterozygous Reelin Mutations Cause Autosomal-Dominant Lateral Temporal Epilepsy. American Journal of Human Genetics, 2015, 96, 992-1000.	2.6	94
32	Focal cortical dysplasias in temporal lobe epilepsy surgery: Challenge in defining unusual variants according to the last ILAE classification. Epilepsy and Behavior, 2015, 45, 212-216.	0.9	11
33	BRAF V600E mutation in neocortical posterior temporal epileptogenic gangliogliomas. Journal of Clinical Neuroscience, 2015, 22, 1250-1253.	0.8	16
34	No evidence of a role for cystatin <scp>B</scp> gene in juvenile myoclonic epilepsy. Epilepsia, 2015, 56, e40-3.	2.6	7
35	Intraoperative monitoring of somatosensory (SSEPs) and transcranial electric motor-evoked potentials (tce-MEPs) during surgical correction of neuromuscular scoliosis in patients with central or peripheral nervous system diseases. European Spine Journal, 2015, 24, 931-936.	1.0	24
36	Teaching Neuro Images : Diffusion tensor tractography of cortico-ponto-cerebellar pathways in Rasmussen encephalitis. Neurology, 2015, 85, e15-6.	1.5	2

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37	An educational campaign about epilepsy among Italian primary school teachers. 2. The results of a focused training program. Epilepsy and Behavior, 2015, 42, 93-97.	0.9	33
38	A recurrent de novo mutation in KCNC1 causes progressive myoclonus epilepsy. Nature Genetics, 2015, 47, 39-46.	9.4	245
39	Epilepsy associated tumors: Review article. World Journal of Clinical Cases, 2014, 2, 623.	0.3	58
40	Mild <scp>L</scp>afora disease: Clinical, neurophysiologic, and genetic findings. Epilepsia, 2014, 55, e129-33.	2.6	43
41	Pattern of care and effectiveness of treatment for glioblastoma patients in the real world: Results from a prospective population-based registry. Could survival differ in a high-volume center?. Neuro-Oncology Practice, 2014, 1, 166-171.	1.0	23
42	Copy number variations and susceptibility to lateral temporal epilepsy: A study of 21 pedigrees. Epilepsia, 2014, 55, 1651-1658.	2.6	10
43	Mutant <scp>BRAF</scp> in lowâ€grade epilepsyâ€associated tumors and focal cortical dysplasia. Annals of Clinical and Translational Neurology, 2014, 1, 130-134.	1.7	33
44	An educational campaign toward epilepsy among Italian primary school teachers. Epilepsy and Behavior, 2014, 32, 84-91.	0.9	22
45	Progressive myoclonic epilepsies. Neurology, 2014, 82, 405-411.	1.5	87
46	Identification of miRNAs Differentially Expressed in Human Epilepsy with or without Granule Cell Pathology. PLoS ONE, 2014, 9, e105521.	1.1	36
47	Seizure outcome in surgically treated drug-resistant mesial temporal lobe epilepsy based on the recent histopathological classifications. Journal of Neurosurgery, 2013, 119, 37-47.	0.9	59
48	Withdrawal of antiepileptic drugs: Guidelines of the <scp>I</scp>talian <scp>L</scp>eague <scp>A</scp>gainst <scp>E</scp>pilepsy. Epilepsia, 2013, 54, 2-12.	2.6	112
49	Autosomal dominant essential tremor: a novel family with anticipation. Neurological Sciences, 2013, 34, 761-763.	0.9	7
50	An Overlooked Cause of Acute Symptomatic Seizures: Psychogenic Polydipsia. American Journal of Medicine, 2013, 126, e1-e2.	0.6	2
51	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
52	Acute rhabdomyolysis induced by tonicâ€“clonic epileptic seizures in a patient with glucose-6-phosphate dehydrogenase deficiency. Journal of Neurology, 2013, 260, 2669-2671.	1.8	6
53	Seizure outcome of surgical treatment of focal epilepsy associated with low-grade tumors in children. Journal of Neurosurgery: Pediatrics, 2013, 11, 214-223.	0.8	50
54	Epilepsy in primary cerebral tumors: The characteristics of epilepsy at the onset (results from the) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 6	2.6	33

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55	LGI1 microdeletion in autosomal dominant lateral temporal epilepsy. Neurology, 2012, 78, 1299-1303.	1.5	28
56	Electroclinical presentation and genotypeâ€“phenotype relationships in patients with Unverrichtâ€Lundborg disease carrying compound heterozygous CSTB point and indel mutations. Epilepsia, 2012, 53, 2120-2127.	2.6	38
57	Genetics of Epilepsy and Relevance to Current Practice. Current Neurology and Neuroscience Reports, 2012, 12, 445-455.	2.0	18
58	Clinical course and variability of non-Rasmussen, nonstroke motor and sensory epilepsia partialis continua: A European survey and analysis of 65 cases. Epilepsia, 2011, 52, 1168-1176.	2.6	50
59	Low penetrance and effect on protein secretion of LGI1 mutations causing autosomal dominant lateral temporal epilepsy. Epilepsia, 2011, 52, 1258-1264.	2.6	26
60	A functional polymorphism in the SCN1A gene does not influence antiepileptic drug responsiveness in Italian patients with focal epilepsy. Epilepsia, 2011, 52, e40-e44.	2.6	50
61	Clinical and neurophysiologic features of progressive myoclonus epilepsy without renal failure caused bySCARB2mutations. Epilepsia, 2011, 52, 2356-2363.	2.6	63
62	Rud syndrome with focal cortical dysplasia: A case report. Brain and Development, 2011, 33, 683-686.	0.6	4
63	Oligodendroglial hamartoma: a potential source of misdiagnosis for oligodendroglioma. Journal of Neuro-Oncology, 2011, 101, 325-328.	1.4	9
64	The prevention of neural complications in the surgical treatment of scoliosis: the role of the neurophysiological intraoperative monitoring. European Spine Journal, 2011, 20, 105-114.	1.0	161
65	Association of intronic variants of the KCNAB1 gene with lateral temporal epilepsy. Epilepsy Research, 2011, 94, 110-116.	0.8	9
66	Familial temporal lobe epilepsy with psychic auras associated with a novel LGI1 mutation. Neurology, 2011, 76, 1173-1176.	1.5	49
67	ADAM23, a Gene Related to LGI1, Is Not Linked to Autosomal Dominant Lateral Temporal Epilepsy. Epilepsy Research & Treatment, 2011, 2011, 1-6.	1.4	2
68	Tubular proteinuria in mice and humans lacking the intrinsic lysosomal protein SCARB2/Limp-2. American Journal of Physiology - Renal Physiology, 2011, 300, F1437-F1447.	1.3	23
69	A PTG Variant Contributes to a Milder Phenotype in Lafora Disease. PLoS ONE, 2011, 6, e21294.	1.1	93
70	p-ANCA pachymeningitis presenting with isolated â€œoptic neuropathyâ€. Neurological Sciences, 2010, 31, 639-641.	0.9	10
71	Postictal hyperfamiliarity for unknown faces. Epilepsy and Behavior, 2010, 19, 518-521.	0.9	5
72	Familial Lateral Temporal Lobe Epilepsy. , 2010, , 1139-1145.		0

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73	Management of Myoclonus. , 2010, , 1271-1278.		0
74	Other Possible Familial Focal Epilepsies Not Yet Recognized by the ILAE. , 2010, , 1153-1157.		0
75	Seizure outcome of epilepsy surgery in focal epilepsies associated with temporomesial glioneuronal tumors: lesionectomy compared with tailored resection. Journal of Neurosurgery, 2009, 111, 1275-1282.	0.9	101
76	SCARB2 mutations in progressive myoclonus epilepsy (PME) without renal failure. Annals of Neurology, 2009, 66, 532-536.	2.8	90
77	LGI1mutations in autosomal dominant and sporadic lateral temporal epilepsy. Human Mutation, 2009, 30, 530-536.	1.1	155
78	Lateral temporal lobe epilepsies: Clinical and genetic features. Epilepsia, 2009, 50, 52-54.	2.6	52
79	Familial mesial temporal lobe epilepsy (FMTLE). Journal of Neurology, 2008, 255, 16-23.	1.8	60
80	Impact of treatment on the short-term prognosis of status epilepticus in two population-based cohorts. Journal of Neurology, 2008, 255, 197-204.	1.8	23
81	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
82	Familial epilepsy and developmental dysphasia: Description of an Italian pedigree with autosomal dominant inheritance and screening of candidate loci. Epilepsy Research, 2008, 80, 9-17.	0.8	9
83	Effects of levetiracetam on EEG abnormalities in juvenile myoclonic epilepsy. Epilepsia, 2008, 49, 663-669.	2.6	54
84	Partial monosomy Xq(Xq23â†'qter) and trisomy 4p(4p15.33â†'pter) in a woman with intractable focal epilepsy, borderline intellectual functioning, and dysmorphic features. Brain and Development, 2008, 30, 425-429.	0.6	10
85	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
86	A Novel Loss-of-Function LGI1 Mutation Linked to Autosomal Dominant Lateral Temporal Epilepsy. Archives of Neurology, 2008, 65, 939-42.	4.9	33
87	SUDDEN FALLS DUE TO SEIZURE-INDUCED CARDIAC ASYSTOLE IN DRUG-RESISTANT FOCAL EPILEPSY. Neurology, 2008, 70, 1933-1935.	1.5	86
88	STRUCTURAL ANOMALY OF LEFT LATERAL TEMPORAL LOBE IN EPILEPSY DUE TO MUTATED LGI1. Neurology, 2007, 69, 1298-1300.	1.5	34
89	A DE NOVO LGI1 MUTATION CAUSING IDIOPATHIC PARTIAL EPILEPSY WITH TELEPHONE-INDUCED SEIZURES. Neurology, 2007, 68, 2150-2151.	1.5	59
90	Neuropathy in multiple myeloma treated with thalidomide. Neurology, 2007, 69, 573-581.	1.5	121

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91	Idiopathic Epilepsies with Seizures Precipitated by Fever and SCN1A Abnormalities. Epilepsia, 2007, 48, 1678-1685.	2.6	154
92	Mutational Analysis of EFHC1 Gene in Italian Families with Juvenile Myoclonic Epilepsy. Epilepsia, 2007, 48, 1686-1690.	2.6	44
93	Charles Bonnet syndrome in hemianopia, following anteroâ€mesial temporal lobectomy for drugâ€resistant epilepsy. Epileptic Disorders, 2007, 9, 271-275.	0.7	15
94	The LGI1/Epitempin gene encodes two protein isoforms differentially expressed in human brain. Journal of Neurochemistry, 2006, 98, 985-991.	2.1	24
95	Recommendations of the Italian League Against Epilepsy Working Group on Generic Products of Antiepileptic Drugs. Epilepsia, 2006, 47, 16-20.	2.6	75
96	Adult-Onset Rasmussen's Encephalitis: Anatomical-Electrographic-Clinical Features of 7 Italian Cases. Epilepsia, 2006, 47, 41-46.	2.6	108
97	Lateralizing Value of the Auditory Aura in Partial Seizures. Epilepsia, 2006, 47, 68-72.	2.6	57
98	Genetic analysis of the LGI/Epitempin gene family in sporadic and familial lateral temporal lobe epilepsy. Epilepsy Research, 2006, 70, 118-126.	0.8	6
99	Open label, long-term, pragmatic study on levetiracetam in the treatment of juvenile myoclonic epilepsy. Epilepsy Research, 2006, 71, 32-39.	0.8	72
100	Increased expression ofLGI1 gene triggers growth inhibition and apoptosis of neuroblastoma cells. Journal of Cellular Physiology, 2006, 207, 711-721.	2.0	22
101	Optimizing therapy of seizures in neurosurgery. Neurology, 2006, 67, S14-8.	1.5	38
102	Biting Behavior, Aggression, and Seizures. Epilepsia, 2005, 46, 654-663.	2.6	46
103	Lafora disease due to EPM2B mutations. Neurology, 2005, 64, 982-986.	1.5	98
104	Clinical features and long term outcome of epilepsy in periventricular nodular heterotopia. Simple compared with plus forms. Journal of Neurology, Neurosurgery and Psychiatry, 2004, 75, 873-878.	0.9	54
105	Idiopathic partial epilepsy with auditory features (IPEAF): a clinical and genetic study of 53 sporadic cases. Brain, 2004, 127, 1343-1352.	3.7	82
106	Telephoneâ€induced Seizures: A New Type of Reflex Epilepsy. Epilepsia, 2004, 45, 280-283.	2.6	36
107	A de novo LGI1 mutation in sporadic partial epilepsy with auditory features. Annals of Neurology, 2004, 56, 455-456.	2.8	54
108	Autosomal Dominant Lateral Temporal Epilepsy: Clinical Spectrum, New Epitempin Mutations, and Genetic Heterogeneity in Seven European Families. Epilepsia, 2003, 44, 1289-1297.	2.6	134

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109	The Provision of Epilepsy Care acrossâ€ƒEurope. Epilepsia, 2003, 44, 727-731.	2.6	26
110	Early ictal speech and motor inhibition in fronto-mesial epileptic seizures: a polygraphic study in one patient. Clinical Neurophysiology, 2003, 114, 56-62.	0.7	12
111	Transcranial magnetic stimulation and epilepsy. Clinical Neurophysiology, 2003, 114, 777-798.	0.7	178
112	Brainstem involvement in Unverrichtâ€“Lundborg disease (EPM1): An MRI and ¹ H MRS study. Neurology, 2002, 58, 1686-1689.	1.5	55
113	Chapter 62 Transcranial magnetic stimulation in epilepsy and Parkinson's disease: drug induced changes in motor excitability. Supplements To Clinical Neurophysiology, 2002, , 416-421.	2.1	1
114	Mutations in the LGI1/Epitempin gene on 10q24 cause autosomal dominant lateral temporal epilepsy. Human Molecular Genetics, 2002, 11, 1119-1128.	1.4	289
115	Identification and characterization of a novel human brain-specific gene, homologous to S. scrofa tmp83.5 , in the chromosome 10q24 critical region for temporal lobe epilepsy and spastic paraplegia. Gene, 2002, 282, 87-94.	1.0	8
116	Guidelines for the use of EEG methodology in the diagnosis of epilepsy. Acta Neurologica Scandinavica, 2002, 106, 1-7.	1.0	150
117	Autosomal Dominant Partial Epilepsy with Auditory Features: Description of a New Family. Epilepsia, 2000, 41, 967-970.	2.6	40
118	Encephalopathy with electrical status epilepticus during slow sleep or ESES syndrome including the acquired aphasia. Clinical Neurophysiology, 2000, 111, S94-S102.	0.7	262
119	A novel protein tyrosine phosphatase gene is mutated in progressive myoclonus epilepsy of the Lafora type (EPM2). Human Molecular Genetics, 1999, 8, 345-352.	1.4	196
120	The Preclinical and Therapeutic Activity of the Novel Anticonvulsant Topiramate. CNS Neuroscience & Therapeutics, 1998, 4, 165-186.	4.0	4
121	The Use of Diazepam and Clonazepam in Epilepsy. Epilepsia, 1998, 39, S7.	2.6	19
122	Familial Cortical Tremor, Epilepsy, and Mental Retardation. Archives of Neurology, 1998, 55, 1569.	4.9	48
123	Self-induction of visually-induced seizures. Advances in Neurology, 1998, 75, 179-92.	0.8	8
124	Unstable minisatellite expansion causing recessively inherited myoclonus epilepsy, EPM1. Nature Genetics, 1997, 15, 393-396.	9.4	207
125	Transcranial magnetic stimulation in partial epilepsy: drug-induced changes of motor excitability. Acta Neurologica Scandinavica, 1996, 94, 24-30.	1.0	44
126	Double-Blind, Placebo-Controlled Trial of Topiramate (600 mg Daily) for the Treatment of Refractory Partial Epilepsy. Epilepsia, 1996, 37, 763-768.	2.6	178

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127	Electroclinical features of idiopathic generalised epilepsy with persisting absences in adult life.. Journal of Neurology, Neurosurgery and Psychiatry, 1996, 61, 471-477.	0.9	38
128	Epileptic negative myoclonus. Advances in Neurology, 1995, 67, 181-97.	0.8	18
129	PME of Unverricht-Lundborg type in the Mediterranean region: linkage and linkage disequilibrium confirm the assignment to the EPM1 locus. Human Genetics, 1994, 93, 668-674.	1.8	15
130	Long-term follow-up study of vigabatrin in the treatment of refractory epilepsy. Journal of Epilepsy, 1994, 7, 88-93.	0.4	9
131	Rapidâ€rate transcranial magnetic stimulation and hemispheric language dominance. Neurology, 1994, 44, 1697-1697.	1.5	87
132	Vigabatrinâ€induced decrease in serum phenytoin concentration does not involve a change in phenytoin bioavailability. British Journal of Clinical Pharmacology, 1993, 36, 603-606.	1.1	34
133	Transthyretin-Related TTR Hereditary Amyloidosis of the Vitreous Body: Clinical and Molecular Characterization in two Italian Families. Ophthalmic Paediatrics and Genetics, 1993, 14, 9-16.	0.4	8
134	Single-blind, placebo-controlled dose-modification study of vigabatrin in refractory epileptic patients. Journal of Epilepsy, 1992, 5, 248-252.	0.4	6
135	Identical genetic locus for Baltic and Mediterranean myoclonus. Lancet, The, 1992, 339, 1080-1081.	6.3	50
136	Slowly progressive familial dementia with recurrent strokes and white matter hypodensities on CT scan. Italian Journal of Neurological Sciences, 1992, 13, 135-140.	0.1	39
137	Single-blind, placebo-controlled multicenter trial of vigabatrin in the treatment of epilepsy. Italian Journal of Neurological Sciences, 1992, 13, 741-747.	0.1	20
138	The electrical status epilepticus syndrome. Epilepsy Research Supplement, 1992, 6, 111-5.	0.0	23
139	Transient Global Amnesia as a Postictal State from Recurrent Partial Seizures. Epilepsia, 1991, 32, 882-885.	2.6	46
140	Vigabatrin. Epilepsy Research Supplement, 1991, 3, 193-6.	0.0	0
141	The Ramsay Hunt syndrome revisited: Mediterranean myoclonus versus mitochondrial encephalomyopathy with ragged-red fibers and Baltic myoclonus. Acta Neurologica Scandinavica, 1990, 81, 8-15.	1.0	49
142	Transcranial magnetic stimulation in epileptic patients. Neurology, 1990, 40, 1132-1132.	1.5	87
143	Trigeminal neuralgia associated with contralateral intracranial tumour: a false localising sign caused by vascular compression? Report of two cases.. Journal of Neurology, Neurosurgery and Psychiatry, 1989, 52, 1202-1203.	0.9	10
144	Dyssynergia cerebellaris myoclonica (Ramsay Hunt syndrome): a condition unrelated to mitochondrial encephalomyopathies.. Journal of Neurology, Neurosurgery and Psychiatry, 1989, 52, 262-265.	0.9	22

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145	Response to vigabatrin in relation to seizure type.. British Journal of Clinical Pharmacology, 1989, 27, 119S-124S.	1.1	44
146	Extreme somatosensory evoked potentials (ESEPs) elicited by tapping of hands or feet in children: A somatosensory cerebral evoked potentials study. Neurophysiologie Clinique, 1988, 18, 123-128.	1.0	29
147	Double-blind Study of Vigabatrin in the Treatment of Drug-Resistant Epilepsy. Archives of Neurology, 1987, 44, 907-910.	4.9	141
148	Intracranial microvascular decompression for â€œcryptogenicâ€•hemifacial spasm, trigeminal and glossopharyngeal neuralgia, paroxysmal vertigo and tinnitus: I. Surgical technique and results. Italian Journal of Neurological Sciences, 1986, 7, 359-366.	0.1	10
149	Intracranial microvascular decompression for â€œcryptogenicâ€•hemifacial spasm, tirgeminal and glossopharyngeal neuralgia, paroxysmal vertigo and tinnitus: II. Clinical study and long-term follow up. Italian Journal of Neurological Sciences, 1986, 7, 367-374.	0.1	9
150	CT Findings in Progressive Supranuclear Palsy. Journal of Computer Assisted Tomography, 1984, 8, 406-409.	0.5	15
151	Sturge-Weber Syndrome without Port-Wine Facial Nevus. Pediatric Neurosurgery, 1983, 10, 387-392.	0.4	12
152	Benzodiazepines: efficacy in status epilepticus. Advances in Neurology, 1983, 34, 465-75.	0.8	15
153	Locked-in syndrome in multiple sclerosis with sparing of the ventral portion of the pons. Annals of Neurology, 1982, 12, 393-394.	2.8	21