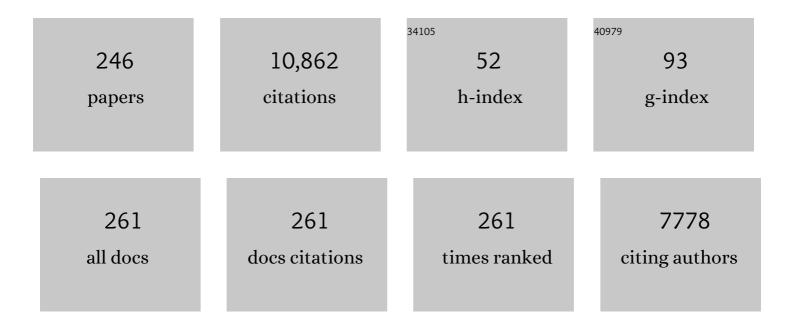
## **Paolo Tinuper**

List of Publications by Year in descending order

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PAOLO TINUDED

#	Article	IF	CITATIONS
1	Fatal Familial Insomnia and Dysautonomia with Selective Degeneration of Thalamic Nuclei. New England Journal of Medicine, 1986, 315, 997-1003.	27.0	688
2	Fatal Familial Insomnia, a Prion Disease with a Mutation at Codon 178 of the Prion Protein Gene. New England Journal of Medicine, 1992, 326, 444-449.	27.0	578
3	Nocturnal frontal lobe epilepsy: A clinical and polygraphic overview of 100 consecutive cases. Brain, 1999, 122, 1017-1031.	7.6	453
4	EFNS guideline on the management of status epilepticus in adults. European Journal of Neurology, 2010, 17, 348-355.	3.3	415
5	A recurrent de novo mutation in KCNC1 causes progressive myoclonus epilepsy. Nature Genetics, 2015, 47, 39-46.	21.4	245
6	Definition and diagnostic criteria of sleep-related hypermotor epilepsy. Neurology, 2016, 86, 1834-1842.	1.1	245
7	Fatal familial insomnia. Neurology, 1992, 42, 312-312.	1.1	211
8	Nocturnal Paroxysmal Dystonia with Short‣asting Attacks: Three Cases with Evidence for an Epileptic Frontal Lobe Origin of Seizures. Epilepsia, 1990, 31, 549-556.	5.1	195
9	Mutations in mammalian target of rapamycin regulator <i>DEPDC5</i> cause focal epilepsy with brain malformations. Annals of Neurology, 2014, 75, 782-787.	5.3	193
10	Mutations in the mammalian target of rapamycin pathway regulators <i>NPRL2</i> and <i>NPRL3</i> cause focal epilepsy. Annals of Neurology, 2016, 79, 120-131.	5.3	190
11	Movement disorders in sleep: Guidelines for differentiating epileptic from non-epileptic motor phenomena arising from sleep. Sleep Medicine Reviews, 2007, 11, 255-267.	8.5	172
12	EFNS guideline on the management of status epilepticus. European Journal of Neurology, 2006, 13, 445-450.	3.3	157
13	The longâ€ŧerm effect of vagus nerve stimulation on quality of life in patients with pharmacoresistant focal epilepsy: The PuLsE (Open Prospective Randomized Longâ€ŧerm Effectiveness) trial. Epilepsia, 2014, 55, 893-900.	5.1	149
14	ILAE definition of the Idiopathic Generalized Epilepsy Syndromes: Position statement by the ILAE Task Force on Nosology and Definitions. Epilepsia, 2022, 63, 1475-1499.	5.1	148
15	The landscape of epilepsy-related GATOR1 variants. Genetics in Medicine, 2019, 21, 398-408.	2.4	137
16	Autosomal Dominant Lateral Temporal Epilepsy: Clinical Spectrum, New Epitempin Mutations, and Genetic Heterogeneity in Seven European Families. Epilepsia, 2003, 44, 1289-1297.	5.1	134
17	Genome-wide association analysis of genetic generalized epilepsies implicates susceptibility loci at 1q43, 2p16.1, 2q22.3 and 17q21.32. Human Molecular Genetics, 2012, 21, 5359-5372.	2.9	134
18	Epileptic Nocturnal Wanderings. Sleep, 1995, 18, 749-756.	1.1	123

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19	Treatment of Status Epilepticus in Adults: Guidelines of the Italian League Against Epilepsy. Epilepsia, 2006, 47, 9-15.	5.1	117
20	Clinical Features of Fatal Familial Insomnia: Phenotypic Variability in Relation to a Polymorphism at Codon 129 of the Prion Protein Gene. Brain Pathology, 1998, 8, 515-520.	4.1	110
21	Non-paraneoplastic limbic encephalitis associated with anti-glutamic acid decarboxylase antibodies. Journal of Neuroimmunology, 2008, 199, 155-159.	2.3	110
22	Increased frequency of arousal parasomnias in families with nocturnal frontal lobe epilepsy: A common mechanism?. Epilepsia, 2010, 51, 1852-1860.	5.1	110
23	Pre-symptomatic diagnosis in fatal familial insomnia: serial neurophysiological and 18FDG-PET studies. Brain, 2006, 129, 668-675.	7.6	109
24	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.	1.6	101
25	Intronic ATTTC repeat expansions in STARD7 in familial adult myoclonic epilepsy linked to chromosome 2. Nature Communications, 2019, 10, 4920.	12.8	99
26	Suppressive Efficacy by a Commercially Available Blue Lens on PPR in 610 Photosensitive Epilepsy Patients. Epilepsia, 2006, 47, 529-533.	5.1	96
27	Cerebral metabolism in fatal familial insomnia: Relation to duration, neuropathology, and distribution of protease-resistent prion protein. Neurology, 1997, 49, 126-133.	1.1	93
28	[18F]FDG PET in fatal familial insomnia. Neurology, 1993, 43, 2565-2565.	1.1	91
29	Sleep-wake cycle abnormalities in fatal familial insomnia. Evidence of the role of the thalamus in sleep regulation. Electroencephalography and Clinical Neurophysiology, 1995, 94, 398-405.	0.3	88
30	Progressive myoclonic epilepsies. Neurology, 2014, 82, 405-411.	1.1	87
31	SUDDEN FALLS DUE TO SEIZURE-INDUCED CARDIAC ASYSTOLE IN DRUG-RESISTANT FOCAL EPILEPSY. Neurology, 2008, 70, 1933-1935.	1.1	86
32	Variation in Lamotrigine Plasma Concentrations with Hormonal Contraceptive Monthly Cycles in Patients with Epilepsy. Epilepsia, 2006, 47, 1573-1575.	5.1	85
33	Idiopathic partial epilepsy with auditory features (IPEAF): a clinical and genetic study of 53 sporadic cases. Brain, 2004, 127, 1343-1352.	7.6	82
34	International League Against Epilepsy classification and definition of epilepsy syndromes with onset at a variable age: position statement by the ILAE Task Force on Nosology and Definitions. Epilepsia, 2022, 63, 1443-1474.	5.1	81
35	The thalamus participates in the regulation of the sleep-waking cycle. A clinico-pathological study in fatal familial thalamic degeneration. Electroencephalography and Clinical Neurophysiology, 1989, 73, 117-123.	0.3	75
36	Startleâ€Induced Epileptic Seizures. Epilepsia, 1984, 25, 712-720.	5.1	71

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37	Paroxysmal arousals during sleep. Neurology, 1990, 40, 1063-1063.	1.1	70
38	Myoclonus epilepsy and ataxia due to <scp><i>KCNC</i></scp> <i>1</i> mutation: Analysis of 20 cases and <scp>K</scp> <sup>+</sup> channel properties. Annals of Neurology, 2017, 81, 677-689.	5.3	69
39	Videopolygraphic and functional MRI study of musicogenic epilepsy. A case report and literature review. Epilepsy and Behavior, 2008, 13, 685-692.	1.7	68
40	Nocturnal Frontal Lobe Epilepsy. Current Neurology and Neuroscience Reports, 2014, 14, 424.	4.2	68
41	Paroxysmal periodic motor attacks during sleep: clinical and polygraphic features. Electroencephalography and Clinical Neurophysiology, 1993, 86, 161-166.	0.3	63
42	Excessive Daytime Sleepiness and Subjective Sleep Quality in Patients with Nocturnal Frontal Lobe Epilepsy: A Case-Control Study. Epilepsia, 2006, 47, 73-77.	5.1	62
43	From nocturnal frontal lobe epilepsy to Sleep-Related Hypermotor Epilepsy: A 35-year diagnostic challenge. Seizure: the Journal of the British Epilepsy Association, 2017, 44, 87-92.	2.0	62
44	Complex Segmental Duplications Mediate a Recurrent dup(X)(p11.22-p11.23) Associated with Mental Retardation, Speech Delay, and EEG Anomalies in Males and Females. American Journal of Human Genetics, 2009, 85, 394-400.	6.2	60
45	A DE NOVO LGI1 MUTATION CAUSING IDIOPATHIC PARTIAL EPILEPSY WITH TELEPHONE-INDUCED SEIZURES. Neurology, 2007, 68, 2150-2151.	1.1	59
46	Seizure outcome in surgically treated drug-resistant mesial temporal lobe epilepsy based on the recent histopathological classifications. Journal of Neurosurgery, 2013, 119, 37-47.	1.6	59
47	Cardiovascular dysautonomia in fatal familial insomnia. Clinical Autonomic Research, 1991, 1, 15-21.	2.5	58
48	Epilepsy associated tumors: Review article. World Journal of Clinical Cases, 2014, 2, 623.	0.8	58
49	GATOR1 complex: the common genetic actor in focal epilepsies. Journal of Medical Genetics, 2016, 53, 503-510.	3.2	58
50	Lateralizing Value of the Auditory Aura in Partial Seizures. Epilepsia, 2006, 47, 68-72.	5.1	57
51	Parasomnias and nocturnal frontal lobe epilepsy (NFLE): Lights and shadows – Controversial points in the differential diagnosis. Sleep Medicine, 2011, 12, S27-S32.	1.6	57
52	Interobserver Reliability of Video Recording in the Diagnosis of Nocturnal Frontal Lobe Seizures. Epilepsia, 2007, 48, 1506-1511.	5.1	55
53	Epilepsy with auditory features. Neurology: Genetics, 2015, 1, e5.	1.9	55
54	Visual Ictal Phenomena in a Case of Lafora Disease Proven by Skin Biopsy. Epilepsia, 1983, 24, 214-218.	5.1	54

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55	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.	1.9	54
56	A de novo LGI1 mutation in sporadic partial epilepsy with auditory features. Annals of Neurology, 2004, 56, 455-456.	5.3	54
57	Endogenous benzodiazepine receptor ligands in idiopathic recurring stupor. Lancet, The, 1992, 340, 1002-1004.	13.7	52
58	Idiopathic recurring stupor: A case with possible involvement of the gamma-aminobutyric acid (GABA)ergic system. Annals of Neurology, 1992, 31, 503-506.	5.3	52
59	Past and present public knowledge and attitudes toward epilepsy in Italy. Epilepsy and Behavior, 2010, 18, 110-115.	1.7	52
60	Prognostic factors in patients with mesial temporal lobe epilepsy. Epilepsia, 2009, 50, 41-44.	5.1	51
61	Patterns of prescription of antiepileptic drugs in patients with refractory epilepsy at tertiary referral centres in Italy. Epilepsy Research, 2010, 91, 273-282.	1.6	50
62	Knowledge and attitudes toward epilepsy among primary and secondary schoolteachers in Italy. Epilepsy and Behavior, 2011, 22, 285-292.	1.7	50
63	The prognostic value of the electroencephalogram in antiepileptic drug withdrawal in partial epilepsies. Neurology, 1996, 47, 76-78.	1.1	49
64	Physiologic autonomic arousal heralds motor manifestations of seizures in nocturnal frontal lobe epilepsy: Implications for pathophysiology. Sleep Medicine, 2012, 13, 252-262.	1.6	49
65	Cost-of-Illness of Epilepsy in Italy. Pharmacoeconomics, 2000, 17, 197-208.	3.3	48
66	Clinical features of sleepâ€related hypermotor epilepsy in relation to the seizureâ€onset zone: A review of 135 surgically treated cases. Epilepsia, 2019, 60, 707-717.	5.1	48
67	Partial Epilepsy of Long Duration: Changing Semiology with Age. Epilepsia, 1996, 37, 162-164.	5.1	47
68	Hyperkinetic manifestations in nocturnal frontal lobe epilepsy. Semeiological features and physiopathological hypothesis. Neurological Sciences, 2005, 26, s210-s214.	1.9	46
69	lctal characteristics of psychogenic nonepileptic seizures: What we have learned from video/EEG recordings—A literature review. Epilepsy and Behavior, 2011, 22, 144-153.	1.7	46
70	Sudden unexpected death in epilepsy (SUDEP): a pilot study on truth telling among Italian epileptologists. Neurological Sciences, 2011, 32, 331-335.	1.9	46
71	Diagnostic accuracy of a structured interview for nocturnal frontal lobe epilepsy (SINFLE): A proposal for developing diagnostic criteria. Sleep Medicine, 2012, 13, 81-87.	1.6	45
72	Overview of presurgical assessment and surgical treatment of epilepsy from the Italian League Against Epilepsy. Epilepsia, 2013, 54, 35-48.	5.1	45

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73	Sleep-related hypermotor epilepsy: prevalence, impact and management strategies. Nature and Science of Sleep, 2018, Volume 10, 317-326.	2.7	45
74	The parasomnias: Mechanisms and treatment. Epilepsia, 2012, 53, 12-19.	5.1	44
75	A prospective study of direct medical costs in a large cohort of consecutively enrolled patients with refractory epilepsy in Italy. Epilepsia, 2015, 56, 1162-1173.	5.1	44
76	Sleep-related hypermotor epilepsy. Neurology, 2017, 88, 70-77.	1.1	43
77	Headache in epilepsy: prevalence and clinical features. Journal of Headache and Pain, 2015, 16, 556.	6.0	42
78	Epilepsy surgery of "low grade epilepsy associated neuroepithelial tumors― A retrospective nationwide Italian study. Epilepsia, 2017, 58, 1832-1841.	5.1	41
79	Specific motor patterns of arousal disorders in adults: aÂvideo-polysomnographic analysis of 184 episodes. Sleep Medicine, 2018, 41, 102-109.	1.6	41
80	Idiopathic recurring stupor. Neurology, 1994, 44, 621-621.	1.1	40
81	Interobserver reliability of ICSD–R minimal diagnostic criteria for the parasomnias. Journal of Neurology, 2005, 252, 712-717.	3.6	39
82	Arousal disorders. Sleep Medicine, 2011, 12, S22-S26.	1.6	39
83	Motor Overactivity and Loss of Motor Circadian Rhythm in Fatal Familial Insomnia: An Actigraphic Study. Sleep, 1997, 20, 739-742.	1.1	38
84	"I heard voices…â€! From semiology, a historical review, and a new hypothesis on the presumed epilepsy of Joan of Arc. Epilepsy and Behavior, 2006, 9, 152-157.	1.7	38
85	Oneiric stupor: The peculiar behaviour of agrypnia excitata. Sleep Medicine, 2011, 12, S64-S67.	1.6	38
86	Epileptic drop attacks in partial epilepsy: clinical features, evolution, and prognosis. Journal of Neurology, Neurosurgery and Psychiatry, 1998, 64, 231-237.	1.9	37
87	Telephoneâ€induced Seizures: A New Type of Reflex Epilepsy. Epilepsia, 2004, 45, 280-283.	5.1	36
88	Endozepine stupor. Recurring stupor linked to endozepine-4 accumulation. Brain, 1998, 121, 127-133.	7.6	35
89	Clinical Features and Pathophysiology of Disorders of Arousal in Adults: A Window Into the Sleeping Brain. Frontiers in Neurology, 2019, 10, 526.	2.4	35
90	Intravenous immunoglobulin therapy in COVID-19-related encephalopathy. Journal of Neurology, 2021, 268, 2671-2675.	3.6	35

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91	Sub-genic intolerance, ClinVar, and the epilepsies: A whole-exome sequencing study of 29,165 individuals. American Journal of Human Genetics, 2021, 108, 965-982.	6.2	35
92	Use of Clobazam in Certain Forms of Status Epilepticus and in Startleâ€Induced Epileptic Seizures. Epilepsia, 1986, 27, S18-26.	5.1	34
93	Autonomic and hormonal ictal changes in gelastic seizures from hypothalamic hamartomas. Electroencephalography and Clinical Neurophysiology, 1998, 107, 317-322.	0.3	34
94	STRUCTURAL ANOMALY OF LEFT LATERAL TEMPORAL LOBE IN EPILEPSY DUE TO MUTATED LGI1. Neurology, 2007, 69, 1298-1300.	1.1	34
95	Benign versive or circling epilepsy with bilateral 3-cps spike-and-wave discharges in late childhood. Annals of Neurology, 1986, 19, 301-303.	5.3	33
96	Low penetrance of autosomal dominant lateral temporal epilepsy in Italian families without <i><scp>LGI</scp>1</i> mutations. Epilepsia, 2013, 54, 1288-1297.	5.1	32
97	Management of status epilepticus in adults. Position paper of the Italian League against Epilepsy. Epilepsy and Behavior, 2020, 102, 106675.	1.7	32
98	Electroencephalogram and HIV Infection: A Prospective Study in 100 Patients. Clinical EEG (electroencephalography), 1990, 21, 145-150.	0.9	31
99	Epilepsy in ring chromosome 20 syndrome. Epilepsy Research, 2016, 128, 83-93.	1.6	30
100	LACK OF VEGETATIVE AND ENDOCRINE ORCADIAN RHYTHMS IN FATAL FAMILIAL THALAMIC DEGENERATION. Clinical Endocrinology, 1987, 26, 573-580.	2.4	29
101	ldentity by descent fine mapping of familial adult myoclonus epilepsy (FAME) to 2p11.2–2q11.2. Human Genetics, 2016, 135, 1117-1125.	3.8	29
102	Tailored surgery for drug-resistant epilepsy due to temporal pole encephalocele and microdysgenesis. Seizure: the Journal of the British Epilepsy Association, 2014, 23, 164-166.	2.0	28
103	Morvan chorea and agrypnia excitata: When video-polysomnographic recording guides the diagnosis. Sleep Medicine, 2011, 12, 1041-1043.	1.6	27
104	Prevalence of Nocturnal Frontal Lobe Epilepsy in the Adult Population of Bologna and Modena, Emilia-Romagna Region, Italy. Sleep, 2015, 38, 479-485.	1.1	27
105	Increasing volume and complexity of pediatric epilepsy surgery with stable seizure outcome between 2008 and 2014: A nationwide multicenter study. Epilepsy and Behavior, 2017, 75, 151-157.	1.7	27
106	Polysomnographic features differentiating disorder of arousals from sleep-related hypermotor epilepsy. Sleep, 2019, 42, .	1.1	27
107	Absence of sleep EEG markers in fatal familial insomnia healthy carriers: a spectral analysis study. Clinical Neurophysiology, 2001, 112, 1888-1892.	1.5	26
108	Endozepines in recurrent stupor. Sleep Medicine Reviews, 2005, 9, 477-487.	8.5	26

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109	Autosomal Dominant Early-onset Cortical Myoclonus, Photic-induced Myoclonus, and Epilepsy in a Large Pedigree. Epilepsia, 2006, 47, 1643-1649.	5.1	26
110	Autosomal dominant lateral temporal epilepsy: Absence of mutations in ADAM22 and Kv1 channel genes encoding LGI1-associated proteins. Epilepsy Research, 2008, 80, 1-8.	1.6	26
111	Incidence of sudden unexpected death in nocturnal frontal lobe epilepsy: a cohort study. Sleep Medicine, 2015, 16, 232-236.	1.6	26
112	Eating Seizures. European Neurology, 1983, 22, 227-231.	1.4	25
113	Sleep-related hypermotor epilepsy (SHE): Contribution of known genes in 103 patients. Seizure: the Journal of the British Epilepsy Association, 2020, 74, 60-64.	2.0	25
114	Seizures with paroxysmal arousals in sleepâ€related hypermotor epilepsy (SHE): Dissecting epilepsy from NREM parasomnias. Epilepsia, 2020, 61, 2194-2202.	5.1	24
115	Impact of treatment on the short-term prognosis of status epilepticus in two population-based cohorts. Journal of Neurology, 2008, 255, 197-204.	3.6	23
116	A novel pedigree with familial cortical myoclonic tremor and epilepsy ( <scp>FCMTE</scp> ): Clinical characterization, refinement of the <scp>FCMTE</scp> 2 locus, and confirmation of a founder haplotype. Epilepsia, 2013, 54, 1298-1306.	5.1	23
117	Off-Label Prescribing of Antiepileptic Drugs in Pharmacoresistant Epilepsy: A Cross-Sectional Drug Utilization Study of Tertiary Care Centers in Italy. CNS Drugs, 2014, 28, 939-949.	5.9	23
118	Effect of valproic acid on perampanel pharmacokinetics in patients with epilepsy. Epilepsia, 2018, 59, e103-e108.	5.1	23
119	Accelerated long-term forgetting in temporal lobe epilepsy: Evidence of improvement after left temporal pole lobectomy. Epilepsy and Behavior, 2011, 22, 793-795.	1.7	22
120	Epilepsy in coeliac disease: not just a matter of calcifications. Neurological Sciences, 2011, 32, 1069-1074.	1.9	22
121	Nocturnal epileptic seizures versus the arousal parasomnias. Somnologie, 2008, 12, 25-37.	1.5	21
122	<i>PRIMA1</i> mutation: a new cause of nocturnal frontal lobe epilepsy. Annals of Clinical and Translational Neurology, 2015, 2, 821-830.	3.7	21
123	Psychiatric comorbidities in patients from seven families with autosomal dominant cortical tremor, myoclonus, and epilepsy. Epilepsy and Behavior, 2016, 56, 38-43.	1.7	21
124	Adverse pregnancy outcomes in women exposed to gabapentin and pregabalin: data from a population-based study. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 223-224.	1.9	21
125	The Impact of the COVID-19 Pandemic on People With Epilepsy. An Italian Survey and a Global Perspective. Frontiers in Neurology, 2020, 11, 613719.	2.4	21
126	Therapy in Sleep-Related Hypermotor Epilepsy (SHE). Current Treatment Options in Neurology, 2020, 22, 1.	1.8	21

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127	Progressive Myoclonus Epilepsies. Neurology: Genetics, 2021, 7, e641.	1.9	20
128	Gait disorders in fatal familial insomnia. Movement Disorders, 2014, 29, 420-424.	3.9	19
129	An Italian multicentre study of perampanel in progressive myoclonus epilepsies. Epilepsy Research, 2019, 156, 106191.	1.6	19
130	Occipital Lobe Epilepsy: A Chronic Condition Related to Transient Occipital Lobe Involvement in Eclampsia. Epilepsia, 1994, 35, 644-647.	5.1	18
131	Analysis of LGI1 promoter sequence, PDYN and GABBR1 polymorphisms in sporadic and familial lateral temporal lobe epilepsy. Neuroscience Letters, 2008, 436, 23-26.	2.1	17
132	Estrogen-related seizure exacerbation following hormone therapy for assisted reproduction in women with epilepsy. Seizure: the Journal of the British Epilepsy Association, 2018, 61, 200-202.	2.0	17
133	Standard procedures for the diagnostic pathway of sleepâ€related epilepsies and comorbid sleep disorders: an EAN, ESRS and ILAEâ€Europe consensus review. European Journal of Neurology, 2021, 28, 15-32.	3.3	17
134	Unexpected gamma glutamyltransferase rise increase during levetiracetam monotherapy. Epileptic Disorders, 2010, 12, 81-82.	1.3	16
135	BRAF V600E mutation in neocortical posterior temporal epileptogenic gangliogliomas. Journal of Clinical Neuroscience, 2015, 22, 1250-1253.	1.5	16
136	Genetic heterogeneity in autosomal dominant nocturnal frontal lobe epilepsy. Italian Journal of Neurological Sciences, 1997, 18, 183-183.	0.1	15
137	Endozepine Stupor in Children. Cephalalgia, 1997, 17, 658-661.	3.9	15
138	Nocturnal Frontal Lobe Epilepsy: New pathophysiological interpretations. Sleep Medicine, 2011, 12, S39-S42.	1.6	15
139	A clinical and genetic study of 33 new cases with early-onset absence epilepsy. Epilepsy Research, 2011, 95, 221-226.	1.6	15
140	Relationship among clinical, pathological and bio-molecular features in low-grade epilepsy-associated neuroepithelial tumors. Journal of Clinical Neuroscience, 2017, 44, 158-163.	1.5	15
141	Brain functional connectivity in sleep-related hypermotor epilepsy. NeuroImage: Clinical, 2018, 17, 873-881.	2.7	15
142	Contribution of ultrarare variants in mTOR pathway genes to sporadic focal epilepsies. Annals of Clinical and Translational Neurology, 2019, 6, 475-485.	3.7	15
143	Occipital Seizures in Lafora Disease: A Further Case Documented by EEG. Clinical EEG (electroencephalography), 1985, 16, 167-170.	0.9	14
144	Partial epilepsy with prominent auditory symptoms not linked to chromosome 10q. Epileptic Disorders, 2002, 4, 183-7.	1.3	14

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145	Suspected covert lorazepam administration misdiagnosed as recurrent endozepine stupor. Brain, 1998, 121, 2201-2201.	7.6	13
146	Does the prion protein gene 129 codon polymorphism influence sleep? Evidence from a fatal familial insomnia kindred. Clinical Neurophysiology, 2002, 113, 1948-1953.	1.5	13
147	Parasomnias Versus Epilepsy: Common Grounds and a Need to Change the Approach to the Problem. Epilepsia, 2007, 48, 1033-1034.	5.1	13
148	Health Technology Assessment report on the presurgical evaluation and surgical treatment of drugâ€resistant epilepsy. Epilepsia, 2013, 54, 49-58.	5.1	13
149	Profile of neuropsychological impairment in Sleep-related Hypermotor Epilepsy. Sleep Medicine, 2018, 48, 8-15.	1.6	13
150	Standard procedures for the diagnostic pathway of sleepâ€related epilepsies and comorbid sleep disorders: A European Academy of Neurology, European Sleep Research Society and International League against Epilepsyâ€Europe consensus review. Journal of Sleep Research, 2020, 29, e13184.	3.2	13
151	Split-screen synchronized display. A useful video-EEG technique for studying paroxysmal phenomena. Epileptic Disorders, 2004, 6, 27-30.	1.3	13
152	Auditory aura in nocturnal frontal lobe epilepsy: a red flag to suspect an extra-frontal epileptogenic zone. Sleep Medicine, 2014, 15, 1417-1423.	1.6	12
153	Migraine with Aura and Photosensitive Epileptic Seizures: A Case Report. Cephalalgia, 1991, 11, 151-153.	3.9	11
154	Clinical and polygraphic study of familial paroxysmal kinesigenic dyskinesia with <i>PRRT2</i> mutation. Epileptic Disorders, 2013, 15, 123-127.	1.3	11
155	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.	1.7	11
156	Intermittent head drops: the differential spectrum. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, 414-419.	1.9	11
157	Paraneoplastic cerebellar degeneration and lambertâ€eaton myasthenia in a patient with merkel cell carcinoma and voltageâ€gated calcium channel antibodies. Muscle and Nerve, 2017, 56, 998-1000.	2.2	11
158	A stereo EEG study in a patient with sleep-related hypermotor epilepsy due to DEPDC5 mutation. Seizure: the Journal of the British Epilepsy Association, 2017, 53, 51-54.	2.0	11
159	Antidepressant effect of vagal nerve stimulation in epilepsy patients: a systematic review. Neurological Sciences, 2020, 41, 3075-3084.	1.9	11
160	The Arousal Disorders Questionnaire: a new and effective screening tool for confusional arousals, Sleepwalking and Sleep Terrors in epilepsy and sleep disorders units. Sleep Medicine, 2021, 80, 279-285.	1.6	11
161	Italian cohort of Lafora disease: Clinical features, disease evolution, and genotype-phenotype correlations. Journal of the Neurological Sciences, 2021, 424, 117409.	0.6	11
162	Classic Migraine Attack Complicated by Confusional State: EEG and CT Study. Cephalalgia, 1985, 5, 63-68.	3.9	10

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163	Facial Asymmetry in Partial Epilepsies. Epilepsia, 1992, 33, 1097-1100.	5.1	10
164	Tobacco habits in nocturnal frontal lobe epilepsy. Epilepsy and Behavior, 2013, 26, 114-117.	1.7	10
165	Autosomal dominant partial epilepsy with auditory features: A new locus on chromosome 19q13.11–q13.31. Epilepsia, 2014, 55, 841-848.	5.1	10
166	Epilepsy in <i>MT</i> â€ <i>ATP6</i> ―related mils/NARP: correlation of elettroclinical features with heteroplasmy. Annals of Clinical and Translational Neurology, 2021, 8, 704-710.	3.7	10
167	Defective lipid signalling caused by mutations in <i>PIK3C2B</i> underlies focal epilepsy. Brain, 2022, 145, 2313-2331.	7.6	10
168	Electroencephalographic and Anatomo-Clinical Evidences of Posterior Cerebral Damage in Hypertensive Encephalopathy. Clinical EEG (electroencephalography), 1984, 15, 53-60.	0.9	9
169	Brainstem Auditory Evoked Responses in Lafora Disease. Clinical EEG (electroencephalography), 1985, 16, 202-207.	0.9	9
170	Familial frontal lobe epilepsy and its relationship with other nocturnal paroxysmal events. Epilepsia, 2010, 51, 51-53.	5.1	9
171	Association of intronic variants of the KCNAB1 gene with lateral temporal epilepsy. Epilepsy Research, 2011, 94, 110-116.	1.6	9
172	Psychogenic nonepileptic seizures. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2012, 107, 277-295.	1.8	9
173	A novel mutation of CLN3 associated with delayed-classic juvenile ceroid lipofuscinois and autophagic vacuolar myopathy. European Journal of Medical Genetics, 2015, 58, 540-544.	1.3	9
174	ELECTROENCEPHALOGRAPHY AND HIV INFECTION. Lancet, The, 1989, 333, 554.	13.7	8
175	Abnormal sympathetic skin responses in thalamic lesions. Electroencephalography and Clinical Neurophysiology - Evoked Potentials, 1992, 85, 225-227.	2.0	8
176	Definition of the neurological phenotype associated with dup (X)(p11.22-p11.23). Epileptic Disorders, 2011, 13, 240-251.	1.3	8
177	Narcolepsy Type 1 and Idiopathic Generalized Epilepsy: Diagnostic and Therapeutic Challenges in Dual Cases. Journal of Clinical Sleep Medicine, 2015, 11, 1257-1262.	2.6	8
178	Pathology-Based Approach to Seizure Outcome After Surgery for Pharmacoresistant Medial Temporal Lobe Epilepsy. World Neurosurgery, 2016, 90, 448-453.	1.3	8
179	Epilepsy with auditory features: Longâ€ŧerm outcome and predictors of terminal remission. Epilepsia, 2018, 59, 834-843.	5.1	8
180	Phenotype variability of GLUT1 deficiency syndrome: Description of a case series with novel SLC2A1 gene mutations. Epilepsy and Behavior, 2018, 79, 169-173.	1.7	8

#	Article	IF	CITATIONS
181	Emilia-Romagna Study on Pregnancy and Exposure to Antiepileptic drugs (ESPEA): a population-based study on prescription patterns, pregnancy outcomes and fetal health. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 983-988.	1.9	8
182	Relationship between plasma concentrations and clinical effects of perampanel: A prospective observational study. Epilepsy and Behavior, 2020, 112, 107385.	1.7	8
183	Perampanel as first add-on antiseizure medication: Italian consensus clinical practice statements. BMC Neurology, 2021, 21, 410.	1.8	8
184	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.	5.1	8
185	Posttraumatic Epilepsy Prediction and Prophylaxis: Open Problems. Archives of Neurology, 1983, 40, 831-831.	4.5	7
186	Plasma endogenous benzodiazepine-like activity in sleep disorders with excessive daytime sleepiness. Neurology, 1995, 45, 1783-1783.	1.1	7
187	Clinical and EEG features of partial epilepsy with secondary bilateral synchrony. Journal of Epilepsy, 1995, 8, 210-214.	0.4	7
188	A seizure response dog: video recording ofÂreacting behaviour during repetitive prolonged seizures. Epileptic Disorders, 2010, 12, 142-145.	1.3	7
189	Successful removal and reimplant of vagal nerve stimulator device after 10 years. Annals of Indian Academy of Neurology, 2012, 15, 128.	0.5	7
190	Colpocephaly in two siblings: further evidence of a genetic transmission. Developmental Medicine and Child Neurology, 2000, 42, 280-282.	2.1	7
191	Impairment of Consciousness and Memory in Migraine: A Review. Headache, 1987, 27, 30-33.	3.9	6
192	Paroxysmal Blinking Provoked by Head Movements. European Neurology, 1989, 29, 298-300.	1.4	6
193	Genetic analysis of the LGI/Epitempin gene family in sporadic and familial lateral temporal lobe epilepsy. Epilepsy Research, 2006, 70, 118-126.	1.6	6
194	<i><scp>DEPDC</scp>5</i> mutations in epilepsy with auditory features. Epilepsia, 2016, 57, 335-335.	5.1	6
195	Sleepâ€related hypermotor epilepsy: A prediction cohort study on sleep/awake patterns of seizures. Epilepsia, 2019, 60, e115-e120.	5.1	6
196	Does screening for adverse effects improve health outcomes in epilepsy?. Neurology, 2020, 95, e239-e246.	1.1	6
197	Epilepsy with auditory features: Contribution of known genes in 112 patients. Seizure: the Journal of the British Epilepsy Association, 2021, 85, 115-118.	2.0	6
198	fMRI-Based Effective Connectivity in Surgical Remediable Epilepsies: A Pilot Study. Brain Topography, 2021, 34, 632-650.	1.8	6

#	Article	IF	CITATIONS
199	Selection of antiseizure medications for first add-on use: A consensus paper. Epilepsy and Behavior, 2021, 122, 108087.	1.7	6
200	Seizure worsening in pregnancy in women with sleep-related hypermotor epilepsy (SHE): A historical cohort study. Seizure: the Journal of the British Epilepsy Association, 2021, 91, 258-262.	2.0	6
201	Prescription patterns of antiepileptic drugs in young women: development of a tool to distinguish between epilepsy and psychiatric disorders. Pharmacoepidemiology and Drug Safety, 2016, 25, 763-769.	1.9	5
202	Prevalence of Sleep-Related Hypermotor Epilepsy—Formerly Named Nocturnal Frontal Lobe Epilepsy—in the Adult Population of the Emilia-Romagna Region, Italy. Sleep, 2017, 40, .	1.1	5
203	Sleep related hyper motor epilepsy (SHE): a unique syndrome with heterogeneous genetic etiologies. Sleep Science and Practice, 2019, 3, .	1.3	5
204	Accurate Detection of Hot-Spot MTOR Somatic Mutations in Archival Surgical Specimens of Focal Cortical Dysplasia by Molecular Inversion Probes. Molecular Diagnosis and Therapy, 2020, 24, 571-577.	3.8	5
205	CT scan in a case of progressive generalized dystonia with amyotrophic paraplegia. Italian Journal of Neurological Sciences, 1983, 4, 335-337.	0.1	4
206	Praxisâ€induced seizures misdiagnosed as cataplexy: A case report. Movement Disorders, 2008, 23, 2105-2107.	3.9	4
207	LGI1 microdeletions are not a frequent cause of partial epilepsy with auditory features (PEAF). Epilepsy Research, 2014, 108, 972-977.	1.6	4
208	The "voices―of Joan of Arc and epilepsy with auditory features. Epilepsy and Behavior, 2016, 61, 281.	1.7	4
209	Coexistence of meningoencephalocele and hippocampal sclerosis: a new type of dual pathology. Acta Neurochirurgica, 2017, 159, 767-769.	1.7	4
210	Incidence of sudden unexpected death in epilepsy in sleep-related hypermotor epilepsy, formerly named nocturnal frontal lobe epilepsy. Sleep Medicine, 2017, 29, 98.	1.6	4
211	Cortical myoclonic tremor induced by fixation-off sensitivity. Neurology, 2018, 91, 1061-1063.	1.1	4
212	Absence Status Appearing on Eye-Closure. Clinical EEG (electroencephalography), 1985, 16, 111-118.	0.9	3
213	Marfan's Syndrome, Recurrent Complex Partial Status Epilepticus and Myoclonus: A Case Report. Clinical EEG (electroencephalography), 1988, 19, 33-36.	0.9	3
214	Transient Unresponsiveness in the Elderly: Possible Episodes of Idiopathic Recurring Stupor. Archives of Neurology, 1995, 52, 232-232.	4.5	3
215	Celiac Disease, Epilepsy, and Occipital Calcifications: Histopathological Study and Clinical Outcome. Journal of Epilepsy, 1996, 9, 206-209.	0.4	3
216	Semiological study of ictal affective behaviour in epilepsy and mental retardation limited to females (EFMR). Epileptic Disorders, 2012, 14, 304-309.	1.3	3

#	Article	IF	CITATIONS
217	Limbic encephalitis with anti-GAD antibodies and Thomsen myotonia: a casual or causal association?. Epileptic Disorders, 2014, 16, 362-365.	1.3	3
218	A survey of the European Reference Network EpiCARE on clinical practice for selected rare epilepsies. Epilepsia Open, 2021, 6, 160-170.	2.4	3
219	Sudden arousals from slow-wave sleep and panic disorder. Sleep, 1998, 21, 548, 551.	1.1	3
220	Complex Segmental Duplications Mediate a Recurrent dup(X)(p11.22-p11.23) Associated with Mental Retardation, Speech Delay, and EEG Anomalies in Males and Females. American Journal of Human Genetics, 2009, 85, 419.	6.2	2
221	Proton MR Spectroscopy in Patients With Sleep-Related Hypermotor Epilepsy (SHE): Evidence of Altered Cingulate Cortex Metabolism. Sleep, 2017, 40, .	1.1	2
222	Insight into epileptic and physiological déjà vu : from a multicentric cohort study. European Journal of Neurology, 2019, 26, 407-414.	3.3	2
223	Ictal vasodepressive syncope in temporal lobe epilepsy. Clinical Neurophysiology, 2020, 131, 155-157.	1.5	2
224	Epilepsy With Auditory Features: From Etiology to Treatment. Frontiers in Neurology, 2021, 12, 807939.	2.4	2
225	Benign unilateral seizures or epilepsy Journal of Neurology, Neurosurgery and Psychiatry, 1983, 46, 871-873.	1.9	1
226	Video-polygraphic recording in "Pure―pilomotor seizures. Journal of Epilepsy, 1991, 4, 103-106.	0.4	1
227	Letters to the Editor. Epilepsia, 1997, 38, 1363-1364.	5.1	1
228	Autosomal dominant nocturnal frontal lobe epilepsy. , 0, , 70-73.		1
229	Nocturnal Frontal Epilepsies: Diagnostic and Therapeutic Challenges for Sleep Specialists. Sleep Medicine Clinics, 2012, 7, 105-112.	2.6	1
230	Validation Study of Italian Version of Inventory for Déjà Vu Experiences Assessment (I-IDEA): A Screening Tool to Detect Déjà Vu Phenomenon in Italian Healthy Individuals. Behavioral Sciences (Basel, Switzerland), 2017, 7, 50.	2.1	1
231	Juvenile absence epilepsy relapsing as recurrent absence status, mimicking transient global amnesia, in an elderly patient. Epileptic Disorders, 2018, 20, 557-561.	1.3	1
232	Autosomal Dominant Nocturnal Frontal Lobe Epilepsy. , 2010, , 1125-1134.		1
233	Elio Lugaresi, MD (1926–2015). Neurology, 2016, 86, 2124-2125.	1.1	1

The Syndrome of Nocturnal Frontal Lobe Epilepsy. , 1997, , 125-135.

1

IF # ARTICLE CITATIONS Letters to the Editor. Epilepsia, 1997, 38, 738-740. 5.1 Nocturnal frontal lobe epilepsy., 2001, , 97-110. 236 0 Introduction. Epilepsia, 2009, 50, 1-1. 5.1 Response to the letter "New avenues to prevent sudden unexpected death in nocturnal frontal lobe epilepsy: follow the route established by omega-3 polyunsaturated fatty acids― Sleep Medicine, 2015, 16, 238 1.6 0 1022-1023. Epilepsy and Sleep: Close Connections and Reciprocal Influences. Neuropsychiatric Symptoms of Neurológical Disease, 2016, , 117-139. Advanced morphological neuroimaging study in lateral temporal lobe epilepsy: A multicentric study. Epilepsy and Behavior, 2017, 74, 69-72. 240 1.7 0 Nocturnal motor behaviors with unexpected EEG and brain MRI findings. Sleep Medicine, 2018, 52, 241 116-117. Reply to Dr. Capovilla on "Reply to the article "Management of status epilepticus in adults. Position 242 1.7 0 paper of the Italian League Against Epilepsyâ€â€• Epilepsy and Behavior, 2020, 107, 107048. Prognostic and Electroclinical Features of Grand Mal Epilepsies. European Neurology, 1986, 25, 339-345. 1.4 244 Fatal Familial Insomnia: A Human Model of Prion Disease., 1998, 33-35. 0 Predictors of hyperkinetic seizures. Epilepsy and Behavior, 2022, 129, 108629. 1.7 The 50th anniversary of the Italian League Against Epilepsy (Lega Italiana Contro l'Epilessia). Epilepsy 246 1.0 0 and Behavior Reports, 2022, 19, 100553.