Francesca Bisulli

List of Publications by Year in Descending Order

172 papers	4,255 citations	37 h-index	59 g-index
196 ext. papers	5,135 ext. citations	4.6 avg, IF	5.03 L-index

#	Paper	IF	Citations
172	Predictors of hyperkinetic seizures.. *Epilepsy and Behavior*, **2022**, 129, 108629	3.2	
171	Questionnaire-based assessment of sleep disorders in an adult population of Tuberous Sclerosis Complex.. *Sleep Medicine*, **2022**, 92, 81-87	4.6	
170	Mild neurological phenotype in a family carrying a novel N-terminal null GRIN2A variant.. *European Journal of Medical Genetics*, **2022**, 65, 104500	2.6	
169	duplication syndrome: The electroclinical features of a case with long-term evolution.. *Epilepsy and Behavior Reports*, **2022**, 19, 100541	1.3	
168	FDG-PET findings and alcohol-responsive myoclonus in a patient with Unverricht-Lundborg disease. *Epilepsy and Behavior Reports*, **2022**, 100551	1.3	
167	If seizures left speechless: CA-P-S C-A-R-E, a proposal of a new ictal language evaluation protocol. *Neurological Sciences*, **2021**, 42, 3249-3255	3.5	
166	Epilepsy With Auditory Features: From Etiology to Treatment.. *Frontiers in Neurology*, **2021**, 12, 807939	4.1	0
165	TELEmedicine for EPIlepsy Care (TELE-EPIC): protocol of a randomised, open controlled non-inferiority clinical trial. *BMJ Open*, **2021**, 11, e053980	3	0
164	Intravenous immunoglobulin therapy in COVID-19-related encephalopathy. *Journal of Neurology*, **2021**, 268, 2671-2675	5.5	13
163	Brain dysfunction in COVID-19 and CAR-T therapy: cytokine storm-associated encephalopathy. *Annals of Clinical and Translational Neurology*, **2021**, 8, 968-979	5.3	16
162	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	4.6	4
161	Italian cohort of Lafora disease: Clinical features, disease evolution, and genotype-phenotype correlations. *Journal of the Neurological Sciences*, **2021**, 424, 117409	3.2	3
160	Results From an Italian Expanded Access Program on Cannabidiol Treatment in Highly Refractory Dravet Syndrome and Lennox-Gastaut Syndrome. *Frontiers in Neurology*, **2021**, 12, 673135	4.1	5
159	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	11	6
158	fMRI-Based Effective Connectivity in Surgical Remediable Epilepsies: A Pilot Study. *Brain Topography*, **2021**, 34, 632-650	4.3	4
157	Is Focal Cortical Dysplasia/Epilepsy Caused by Somatic Mutations Always a Unilateral Disorder?. *Neurology: Genetics*, **2021**, 7, e540	3.8	9
156	A survey of the European Reference Network EpiCARE on clinical practice for selected rare epilepsies. *Epilepsia Open*, **2021**, 6, 160-170	4	0

155	Epilepsy with auditory features: Contribution of known genes in 112 patients. *Seizure: the Journal of the British Epilepsy Association*, **2021**, 85, 115-118	3.2	4
154	Risk of SUDEP during infancy. *Epilepsy and Behavior*, **2021**, 107896	3.2	0
153	Akinetic mutism in COVID-19-related encephalopathy: A cytokine-mediated maladaptive sickness behavioral response?. *Brain, Behavior, & Immunity - Health*, **2021**, 15, 100272	5.1	1
152	Natural history of Lafora disease: a prognostic systematic review and individual participant data meta-analysis. *Orphanet Journal of Rare Diseases*, **2021**, 16, 362	4.2	4
151	Postictal Psychosis in Epilepsy: A Clinicogenetic Study. *Annals of Neurology*, **2021**, 90, 464-476	9.4	1
150	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	3.2	1
149	Epilepsy in MT-ATP6 - related mils/NARP: correlation of elettroclinical features with heteroplasmy. *Annals of Clinical and Translational Neurology*, **2021**, 8, 704-710	5.3	1
148	The Impact of the COVID-19 Pandemic on People With Epilepsy. An Italian Survey and a Global Perspective. *Frontiers in Neurology*, **2020**, 11, 613719	4.1	12
147	Autosomal dominant lateral temporal lobe epilepsy associated with a novel reelin mutation. *Epileptic Disorders*, **2020**, 22, 443-448	1.9	3
146	Epilepsy subtype-specific copy number burden observed in a genome-wide study of 17 458 subjects. *Brain*, **2020**, 143, 2106-2118	11.2	14
145	Antidepressant effect of vagal nerve stimulation in epilepsy patients: a systematic review. *Neurological Sciences*, **2020**, 41, 3075-3084	3.5	8
144	Low CSF hypocretin-1 levels in an adult patient with hypothalamic hamartoma. *Neurology*, **2020**, 94, 670-672	6.5	2
143	Advances in genetic testing and optimization of clinical management in children and adults with epilepsy. *Expert Review of Neurotherapeutics*, **2020**, 20, 251-269	4.3	17
142	Therapy in Sleep-Related Hypermotor Epilepsy (SHE). *Current Treatment Options in Neurology*, **2020**, 22, 1	4.4	13
141	FDG-PET assessment and metabolic patterns in Lafora disease. *European Journal of Nuclear Medicine and Molecular Imaging*, **2020**, 47, 1576-1584	8.8	5
140	Ictal vasodepressive syncope in temporal lobe epilepsy. *Clinical Neurophysiology*, **2020**, 131, 155-157	4.3	2
139	Sleep-related hypermotor epilepsy (SHE): Contribution of known genes in 103 patients. *Seizure: the Journal of the British Epilepsy Association*, **2020**, 74, 60-64	3.2	17
138	Relationship between plasma concentrations and clinical effects of perampanel: A prospective observational study. *Epilepsy and Behavior*, **2020**, 112, 107385	3.2	1

137	EEG findings in COVID-19 related encephalopathy. *Clinical Neurophysiology*, **2020**, 131, 2265-2267	4.3	21
136	Epilepsy with eyelid myoclonias and Sotos syndrome features in a patient with compound heterozygous missense variants in APC2 gene. *Seizure: the Journal of the British Epilepsy Association* , **2020**, 83, 169-171	3.2	2
135	Encephalopathy in COVID-19 Presenting With Acute Aphasia Mimicking Stroke. *Frontiers in Neurology*, **2020**, 11, 587226	4.1	15
134	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	4.5	3
133	Whole-exome sequencing in adult patients with developmental and epileptic encephalopathy: It is never too late. *Clinical Genetics*, **2020**, 98, 477-485	4	9
132	COVID-19-Associated Encephalopathy and Cytokine-Mediated Neuroinflammation. *Annals of Neurology*, **2020**, 88, 860-861	9.4	32
131	COVID-19-related encephalopathy presenting with aphasia resolving following tocilizumab treatment. *Journal of Neuroimmunology*, **2020**, 349, 577400	3.5	24
130	Interrater agreement of classification of photoparoxysmal electroencephalographic response. *Epilepsia*, **2020**, 61, e124-e128	6.4	4
129	Seizures with paroxysmal arousals in sleep-related hypermotor epilepsy (SHE): Dissecting epilepsy from NREM parasomnias. *Epilepsia*, **2020**, 61, 2194-2202	6.4	4
128	Did the COVID-19 pandemic silence the needs of people with epilepsy?. *Epileptic Disorders*, **2020**, 22, 439-442	1.9	25
127	An Italian multicentre study of perampanel in progressive myoclonus epilepsies. *Epilepsy Research*, **2019**, 156, 106191	3	12
126	Polygraphic Techniques **2019**, 259-279		
125	Clinical Features and Pathophysiology of Disorders of Arousal in Adults: A Window Into the Sleeping Brain. *Frontiers in Neurology*, **2019**, 10, 526	4.1	15
124	Contribution of ultrarare variants in mTOR pathway genes to sporadic focal epilepsies. *Annals of Clinical and Translational Neurology*, **2019**, 6, 475-485	5.3	9
123	The landscape of epilepsy-related GATOR1 variants. *Genetics in Medicine*, **2019**, 21, 398-408	8.1	75
122	Ultra-Rare Genetic Variation in the Epilepsies: A Whole-Exome Sequencing Study of 17,606 Individuals. *American Journal of Human Genetics*, **2019**, 105, 267-282	11	104
121	Sleep related hyper motor epilepsy (SHE): a unique syndrome with heterogeneous genetic etiologies. *Sleep Science and Practice*, **2019**, 3,	1.2	3
120	Treatment with metformin in twelve patients with Lafora disease. *Orphanet Journal of Rare Diseases*, **2019**, 14, 149	4.2	13

119	Sleep-related hypermotor epilepsy: A prediction cohort study on sleep/awake patterns of seizures. *Epilepsia*, **2019**, 60, e115-e120	6.4	5
118	Polysomnographic features differentiating disorder of arousals from sleep-related hypermotor epilepsy. *Sleep*, **2019**, 42,	1.1	15
117	Intronic ATTTC repeat expansions in STARD7 in familial adult myoclonic epilepsy linked to chromosome 2. *Nature Communications*, **2019**, 10, 4920	17.4	48
116	SCN1A mutations in focal epilepsy with auditory features: widening the spectrum of GEFS plus **2019**, 21, 185-191		2
115	Super refractory status epilepticus in Lafora disease interrupted by vagus nerve stimulation: A case report. *Brain Stimulation*, **2019**, 12, 1605-1607	5.1	3
114	Insight into epileptic and physiological dj vu: from a multicentric cohort study. *European Journal of Neurology*, **2019**, 26, 407-414	6	1
113	Epilepsy with auditory features: Long-term outcome and predictors of terminal remission. *Epilepsia*, **2018**, 59, 834-843	6.4	7
112	Brain functional connectivity in sleep-related hypermotor epilepsy. *NeuroImage: Clinical*, **2018**, 17, 873-881	3.5	10
111	Phenotype variability of GLUT1 deficiency syndrome: Description of a case series with novel SLC2A1 gene mutations. *Epilepsy and Behavior*, **2018**, 79, 169-173	3.2	5
110	Profile of neuropsychological impairment in Sleep-related Hypermotor Epilepsy. *Sleep Medicine*, **2018**, 48, 8-15	4.6	7
109	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	5.5	7
108	Specific motor patterns of arousal disorders in adults: a video-polysomnographic analysis of 184 episodes. *Sleep Medicine*, **2018**, 41, 102-109	4.6	24
107	Rare coding variants in genes encoding GABA receptors in genetic generalised epilepsies: an exome-based case-control study. *Lancet Neurology, The*, **2018**, 17, 699-708	24.1	44
106	Effect of valproic acid on perampanel pharmacokinetics in patients with epilepsy. *Epilepsia*, **2018**, 59, e103-e108	6.4	15
105	Cortical myoclonic tremor induced by fixation-off sensitivity: An unusual cause of insomnia. *Neurology*, **2018**, 91, 1061-1063	6.5	0
104	Juvenile absence epilepsy relapsing as recurrent absence status, mimicking transient global amnesia, in an elderly patient. *Epileptic Disorders*, **2018**, 20, 557-561	1.9	
103	Sleep-related hypermotor epilepsy: prevalence, impact and management strategies. *Nature and Science of Sleep*, **2018**, 10, 317-326	3.6	28
102	Nocturnal motor behaviors with unexpected EEG and brain MRI findings. *Sleep Medicine*, **2018**, 52, 116-117	4.6	

101	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	3.2	9
100	Incidence of sudden unexpected death in epilepsy in sleep-related hypermotor epilepsy, formerly named nocturnal frontal lobe epilepsy. *Sleep Medicine*, **2017**, 29, 98	4.6	3
99	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	3
98	Myoclonus epilepsy and ataxia due to KCNC1 mutation: Analysis of 20 cases and K channel properties. *Annals of Neurology*, **2017**, 81, 677-689	9.4	39
97	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	3.2	47
96	Validation Study of Italian Version of Inventory for D□Vu Experiences Assessment (I-IDEA): A Screening Tool to Detect D□Vu Phenomenon in Italian Healthy Individuals. *Behavioral Sciences (Basel, Switzerland)*, **2017**, 7,	2.3	1
95	Alterations in the □ligand, thrombospondin-1, in a rat model of spontaneous absence epilepsy and in patients with idiopathic/genetic generalized epilepsies. *Epilepsia*, **2017**, 58, 1993-2001	6.4	4
94	Advanced morphological neuroimaging study in lateral temporal lobe epilepsy: A multicentric study. *Epilepsy and Behavior*, **2017**, 74, 69-72	3.2	
93	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	3.2	9
92	Sleep-related hypermotor epilepsy: Long-term outcome in a large cohort. *Neurology*, **2017**, 88, 70-77	6.5	34
91	Epilepsy in ring chromosome 20 syndrome. *Epilepsy Research*, **2016**, 128, 83-93	3	22
90	Mutations in the mammalian target of rapamycin pathway regulators NPRL2 and NPRL3 cause focal epilepsy. *Annals of Neurology*, **2016**, 79, 120-31	9.4	136
89	Psychiatric comorbidities in patients from seven families with autosomal dominant cortical tremor, myoclonus, and epilepsy. *Epilepsy and Behavior*, **2016**, 56, 38-43	3.2	13
88	Pathology-Based Approach to Seizure Outcome After Surgery for Pharmacoresistant Medial Temporal Lobe Epilepsy. *World Neurosurgery*, **2016**, 90, 448-453	2.1	4
87	Epilepsy and Sleep: Close Connections and Reciprocal Influences. *Neuropsychiatric Symptoms of Neurological Disease*, **2016**, 117-139		
86	Identity by descent fine mapping of familial adult myoclonus epilepsy (FAME) to 2p11.2-2q11.2. *Human Genetics*, **2016**, 135, 1117-25	6.3	16
85	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, 763-9	2.6	5
84	DEPDC5 mutations in epilepsy with auditory features. *Epilepsia*, **2016**, 57, 335	6.4	4

83	GATOR1 complex: the common genetic actor in focal epilepsies. *Journal of Medical Genetics*, **2016**, 53, 503-10	5.8	39
82	Definition and diagnostic criteria of sleep-related hypermotor epilepsy. *Neurology*, **2016**, 86, 1834-42	6.5	182
81	BRAF V600E mutation in neocortical posterior temporal epileptogenic gangliogliomas. *Journal of Clinical Neuroscience*, **2015**, 22, 1250-3	2.2	12
80	Incidence of sudden unexpected death in nocturnal frontal lobe epilepsy: a cohort study. *Sleep Medicine*, **2015**, 16, 232-6	4.6	20
79	A novel mutation af Cln3 associated with delayed-classic juvenile ceroid lipofuscinois and autophagic vacuolar myopathy. *European Journal of Medical Genetics*, **2015**, 58, 540-4	2.6	8
78	Headache in epilepsy: prevalence and clinical features. *Journal of Headache and Pain*, **2015**, 16, 556	8.8	28
77	PRIMA1 mutation: a new cause of nocturnal frontal lobe epilepsy. *Annals of Clinical and Translational Neurology*, **2015**, 2, 821-30	5.3	19
76	Epilepsy with auditory features: A heterogeneous clinico-molecular disease. *Neurology: Genetics*, **2015**, 1, e5	3.8	38
75	Prevalence of nocturnal frontal lobe epilepsy in the adult population of Bologna and Modena, Emilia-Romagna region, Italy. *Sleep*, **2015**, 38, 479-85	1.1	22
74	Focal cortical dysplasias in temporal lobe epilepsy surgery: Challenge in defining unusual variants according to the last ILAE classification. *Epilepsy and Behavior*, **2015**, 45, 212-6	3.2	9
73	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, 1022-3	4.6	
72	Nocturnal frontal lobe epilepsy. *Current Neurology and Neuroscience Reports*, **2014**, 14, 424	6.6	51
71	LGI1 microdeletions are not a frequent cause of partial epilepsy with auditory features (PEAF). *Epilepsy Research*, **2014**, 108, 972-7	3	4
70	Epilepsy associated tumors: Review article. *World Journal of Clinical Cases*, **2014**, 2, 623-41	1.6	40
69	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	2.2	19
68	Autosomal dominant partial epilepsy with auditory features: a new locus on chromosome 19q13.11-q13.31. *Epilepsia*, **2014**, 55, 841-8	6.4	9
67	Limbic encephalitis with anti-GAD antibodies and Thomsen myotonia: a casual or causal association?. *Epileptic Disorders*, **2014**, 16, 362-5	1.9	2
66	Mutations in mammalian target of rapamycin regulator DEPDC5 cause focal epilepsy with brain malformations. *Annals of Neurology*, **2014**, 75, 782-7	9.4	153

65	Auditory aura in nocturnal frontal lobe epilepsy: a red flag to suspect an extra-frontal epileptogenic zone. *Sleep Medicine*, **2014**, 15, 1417-23	4.6	6
64	Tailored surgery for drug-resistant epilepsy due to temporal pole encephalocele and microdysgenesis. *Seizure: the Journal of the British Epilepsy Association*, **2014**, 23, 164-6	3.2	19
63	Clinical and polygraphic study of familial paroxysmal kinesigenic dyskinesia with PRRT2 mutation. *Epileptic Disorders*, **2013**, 15, 123-7	1.9	10
62	Seizure outcome in surgically treated drug-resistant mesial temporal lobe epilepsy based on the recent histopathological classifications. *Journal of Neurosurgery*, **2013**, 119, 37-47	3.2	54
61	A novel pedigree with familial cortical myoclonic tremor and epilepsy (FCMTE): clinical characterization, refinement of the FCMTE2 locus, and confirmation of a founder haplotype. *Epilepsia*, **2013**, 54, 1298-306	6.4	16
60	Tobacco habits in nocturnal frontal lobe epilepsy. *Epilepsy and Behavior*, **2013**, 26, 114-7	3.2	9
59	Mutations in DEPDC5 cause familial focal epilepsy with variable foci. *Nature Genetics*, **2013**, 45, 546-51	36.3	238
58	Low penetrance of autosomal dominant lateral temporal epilepsy in Italian families without LGI1 mutations. *Epilepsia*, **2013**, 54, 1288-97	6.4	28
57	Diagnostic accuracy of a structured interview for nocturnal frontal lobe epilepsy (SINFLE): a proposal for developing diagnostic criteria. *Sleep Medicine*, **2012**, 13, 81-7	4.6	41
56	Physiologic autonomic arousal heralds motor manifestations of seizures in nocturnal frontal lobe epilepsy: implications for pathophysiology. *Sleep Medicine*, **2012**, 13, 252-62	4.6	42
55	The parasomnias: mechanisms and treatment. *Epilepsia*, **2012**, 53 Suppl 7, 12-9	6.4	26
54	Nocturnal Frontal Epilepsies: Diagnostic and Therapeutic Challenges for Sleep Specialists. *Sleep Medicine Clinics*, **2012**, 7, 105-112	3.6	1
53	Semiological study of ictal affective behaviour in epilepsy and mental retardation limited to females (EFMR). *Epileptic Disorders*, **2012**, 14, 304-9	1.9	2
52	Genome-wide linkage meta-analysis identifies susceptibility loci at 2q34 and 13q31.3 for genetic generalized epilepsies. *Epilepsia*, **2012**, 53, 308-18	6.4	28
51	Successful removal and reimplant of vagal nerve stimulator device after 10 years. *Annals of Indian Academy of Neurology*, **2012**, 15, 128-9	0.9	6
50	Ictal characteristics of psychogenic nonepileptic seizures: what we have learned from video/EEG recordings--a literature review. *Epilepsy and Behavior*, **2011**, 22, 144-53	3.2	36
49	Arousal disorders. *Sleep Medicine*, **2011**, 12 Suppl 2, S22-6	4.6	32
48	Parasomnias and nocturnal frontal lobe epilepsy (NFLE): lights and shadows--controversial points in the differential diagnosis. *Sleep Medicine*, **2011**, 12 Suppl 2, S27-32	4.6	46

47	Nocturnal frontal lobe epilepsy: new pathophysiological interpretations. *Sleep Medicine*, **2011**, 12 Suppl 2, S39-42	4.6	14
46	Determinants of health-related quality of life in pharmacoresistant epilepsy: results from a large multicenter study of consecutively enrolled patients using validated quantitative assessments. *Epilepsia*, **2011**, 52, 2181-91	6.4	182
45	Definition of the neurological phenotype associated with dup (X)(p11.22-p11.23). *Epileptic Disorders*, **2011**, 13, 240-51	1.9	6
44	Epilepsy in coeliac disease: not just a matter of calcifications. *Neurological Sciences*, **2011**, 32, 1069-74	3.5	21
43	Association of intronic variants of the KCNAB1 gene with lateral temporal epilepsy. *Epilepsy Research*, **2011**, 94, 110-6	3	8
42	A clinical and genetic study of 33 new cases with early-onset absence epilepsy. *Epilepsy Research*, **2011**, 95, 221-6	3	15
41	Unexpected gamma glutamyltransferase rise increase during levetiracetam monotherapy. *Epileptic Disorders*, **2010**, 12, 81-2	1.9	10
40	A seizure response dog: video recording of reacting behaviour during repetitive prolonged seizures. *Epileptic Disorders*, **2010**, 12, 142-5	1.9	5
39	Familial frontal lobe epilepsy and its relationship with other nocturnal paroxysmal events. *Epilepsia*, **2010**, 51 Suppl 1, 51-3	6.4	8
38	Increased frequency of arousal parasomnias in families with nocturnal frontal lobe epilepsy: a common mechanism?. *Epilepsia*, **2010**, 51, 1852-60	6.4	93
37	Autosomal Dominant Nocturnal Frontal Lobe Epilepsy **2010**, 1125-1134		1
36	Seizure outcome of epilepsy surgery in focal epilepsies associated with temporomesial glioneuronal tumors: lesionectomy compared with tailored resection. *Journal of Neurosurgery*, **2009** , 111, 1275-82	3.2	87
35	Prognostic factors in patients with mesial temporal lobe epilepsy. *Epilepsia*, **2009**, 50 Suppl 1, 41-4	6.4	43
34	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	11	56
33	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	11	2
32	Non-paraneoplastic limbic encephalitis associated with anti-glutamic acid decarboxylase antibodies. *Journal of Neuroimmunology*, **2008**, 199, 155-9	3.5	99
31	Analysis of LGI1 promoter sequence, PDYN and GABBR1 polymorphisms in sporadic and familial lateral temporal lobe epilepsy. *Neuroscience Letters*, **2008**, 436, 23-6	3.3	16
30	Videopolygraphic and functional MRI study of musicogenic epilepsy. A case report and literature review. *Epilepsy and Behavior*, **2008**, 13, 685-92	3.2	56

29	Sudden falls due to seizure-induced cardiac asystole in drug-resistant focal epilepsy. *Neurology*, **2008**, 70, 1933-5	6.5	20
28	Nocturnal epileptic seizures versus the arousal parasomnias. *Somnologie*, **2008**, 12, 25-37	2	20
27	Autosomal dominant lateral temporal epilepsy: absence of mutations in ADAM22 and Kv1 channel genes encoding LGI1-associated proteins. *Epilepsy Research*, **2008**, 80, 1-8	3	24
26	Parasomnias versus epilepsy: common grounds and a need to change the approach to the problem. *Epilepsia*, **2007**, 48, 1033-4; author reply 1034	6.4	13
25	Interobserver reliability of video recording in the diagnosis of nocturnal frontal lobe seizures. *Epilepsia*, **2007**, 48, 1506-11	6.4	49
24	Structural anomaly of left lateral temporal lobe in epilepsy due to mutated LGI1. *Neurology*, **2007**, 69, 1298-300	6.5	32
23	A de novo LGI1 mutation causing idiopathic partial epilepsy with telephone-induced seizures. *Neurology*, **2007**, 68, 2150-1	6.5	49
22	Movement disorders in sleep: guidelines for differentiating epileptic from non-epileptic motor phenomena arising from sleep. *Sleep Medicine Reviews*, **2007**, 11, 255-67	10.2	144
21	Multicentre observational study evaluating immediate and progressive switching from carbamazepine to oxcarbazepine in patients with epilepsy. *Functional Neurology*, **2007**, 22, 111-5	2.2	6
20	Epilepsy as a consequence of cerebral malaria in area in which malaria is endemic in Mali, West Africa. *Epilepsia*, **2006**, 47, 873-9	6.4	59
19	Autosomal dominant early-onset cortical myoclonus, photic-induced myoclonus, and epilepsy in a large pedigree. *Epilepsia*, **2006**, 47, 1643-9	6.4	19
18	Lateralizing value of the auditory aura in partial seizures. *Epilepsia*, **2006**, 47 Suppl 5, 68-72	6.4	48
17	Excessive daytime sleepiness and subjective sleep quality in patients with nocturnal frontal lobe epilepsy: a case-control study. *Epilepsia*, **2006**, 47 Suppl 5, 73-7	6.4	54
16	Genetic analysis of the LGI/Epitempin gene family in sporadic and familial lateral temporal lobe epilepsy. *Epilepsy Research*, **2006**, 70, 118-26	3	6
15	Interobserver reliability of ICSD-R minimal diagnostic criteria for the parasomnias. *Journal of Neurology*, **2005**, 252, 712-7	5.5	34
14	Hyperkinetic manifestations in nocturnal frontal lobe epilepsy. Semeiological features and physiopathological hypothesis. *Neurological Sciences*, **2005**, 26 Suppl 3, s210-4	3.5	42
13	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, 873-8	5.5	41
12	Idiopathic partial epilepsy with auditory features (IPEAF): a clinical and genetic study of 53 sporadic cases. *Brain*, **2004**, 127, 1343-52	11.2	73

11	Telephone-induced seizures: a new type of reflex epilepsy. *Epilepsia*, **2004**, 45, 280-3	6.4	29
10	A de novo LGI1 mutation in sporadic partial epilepsy with auditory features. *Annals of Neurology*, **2004**, 56, 455-6	9.4	47
9	Split-screen synchronized display. A useful video-EEG technique for studying paroxysmal phenomena. *Epileptic Disorders*, **2004**, 6, 27-30	1.9	13
8	Autosomal dominant lateral temporal epilepsy: clinical spectrum, new epitempin mutations, and genetic heterogeneity in seven European families. *Epilepsia*, **2003**, 44, 1289-97	6.4	119
7	Ictal pattern of EEG and muscular activation in symptomatic infantile spasms: a videopolygraphic and computer analysis. *Epilepsia*, **2002**, 43, 1559-63	6.4	18
6	Partial epilepsy with prominent auditory symptoms not linked to chromosome 10q. *Epileptic Disorders*, **2002**, 4, 183-7	1.9	13
5	Ictal bradycardia in partial epileptic seizures: Autonomic investigation in three cases and literature review. *Brain*, **2001**, 124, 2361-71	11.2	135
4	Efficacy of lamotrigine add-on therapy in severe partial epilepsy in adults with drop seizures and secondary bilateral synchrony on EEG. *Epileptic Disorders*, **2001**, 3, 151-6	1.9	4
3	Epileptic negative myoclonus and brief asymmetric tonic seizures. A supplementary sensorimotor area involvement for both negative and positive motor phenomena. *Epileptic Disorders*, **2000**, 2, 163-8	1.9	10
2	Gaze-evoked amaurosis heralding orbital angiomyoma. Case report. *Italian Journal of Neurological Sciences*, **1997**, 18, 31-4		4
1	Autosomal dominant nocturnal frontal lobe epilepsy70-73		1