Angelo Labate

List of Publications by Year
in descending order

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200	6,088	66343	98798
papers	citations	42 h-index	67 g-index

210	210	210	7797
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The <scp>ENIGMAâ€Epilepsy</scp> working group: Mapping disease from large data sets. Human Brain Mapping, 2022, 43, 113-128.	3.6	47
2	A systemsâ€level analysis highlights microglial activation as a modifying factor in common epilepsies. Neuropathology and Applied Neurobiology, 2022, 48, .	3.2	22
3	Topographic divergence of atypical cortical asymmetry and atrophy patterns in temporal lobe epilepsy. Brain, 2022, 145, 1285-1298.	7.6	18
4	Atlas of lesion locations and postsurgical seizure freedom in focal cortical dysplasia: A MELD study. Epilepsia, 2022, 63, 61-74.	5.1	36
5	The impact of one-year COVID-19 containment measures in patients with mesial temporal lobe epilepsy: A longitudinal survey-based study. Epilepsy and Behavior, 2022, 128, 108600.	1.7	1
6	Clinical and Instrumental Characterization of Patients With Late-Onset Epilepsy. Frontiers in Neurology, 2022, 13, 851897.	2.4	4
7	Psychiatric and Behavioural Side Effects Associated With Perampanel in Patients With Temporal Lobe Epilepsy. A Real-World Experience. Frontiers in Neurology, 2022, 13, 839985.	2.4	7
8	Electroclinical Features and Long-term Seizure Outcome in Patients With Eyelid Myoclonia With Absences. Neurology, 2022, 98, .	1.1	15
9	Non-Coding RNAs: New Biomarkers and Therapeutic Targets for Temporal Lobe Epilepsy. International Journal of Molecular Sciences, 2022, 23, 3063.	4.1	8
10	Brivaracetam as add-on treatment in patients with post-stroke epilepsy: real-world data from the BRIVAracetam add-on First Italian netwoRk Study (BRIVAFIRST). Seizure: the Journal of the British Epilepsy Association, 2022, 97, 37-42.	2.0	4
11	Epileptic belly dancing: a videoâ€polygraphic recording. Epileptic Disorders, 2022, 24, 442-444.	1.3	0
12	Late epileptic seizures following cerebral venous thrombosis: a systematic review and meta-analysis. Neurological Sciences, 2022, 43, 5229-5236.	1.9	8
13	Eventâ€based modeling in temporal lobe epilepsy demonstrates progressive atrophy from crossâ€sectional data. Epilepsia, 2022, 63, 2081-2095.	5.1	11
14	Networks Underlie Temporal Onset of Dysplasiaâ€Related Epilepsy: A <scp>MELD</scp> Study. Annals of Neurology, 2022, 92, 503-511.	5.3	7
15	Epilepsy, Immunity and Neuropsychiatric Disorders. Current Neuropharmacology, 2022, 20, .	2.9	4
16	The impact of sexual abuse on psychopathology of patients with psychogenic nonepileptic seizures. Neurological Sciences, 2021, 42, 1423-1428.	1.9	3
17	Perampanel as first add-on choice on the treatment of mesial temporal lobe epilepsy: an observational real-life study. Neurological Sciences, 2021, 42, 1389-1394.	1.9	13
18	Circulating microRNAs as Potential Novel Diagnostic Biomarkers to Predict Drug Resistance in Temporal Lobe Epilepsy: A Pilot Study. International Journal of Molecular Sciences, 2021, 22, 702.	4.1	30

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19	Artificial intelligence for classification of temporal lobe epilepsy with ROI-level MRI data: A worldwide ENIGMA-Epilepsy study. NeuroImage: Clinical, 2021, 31, 102765.	2.7	25
20	Facemask headache: a new nosographic entity among healthcare providers in COVID-19 era. Neurological Sciences, 2021, 42, 1267-1276.	1.9	22
21	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.	6.2	41
22	Diagnostic and therapeutic approach to drug-resistant juvenile myoclonic epilepsy. Expert Review of Neurotherapeutics, 2021, 21, 1265-1273.	2.8	12
23	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
24	Orbito-frontal thinning together with a somatoform dissociation might be the fingerprint of PNES. Epilepsy and Behavior, 2021, 121, 108044.	1.7	9
25	Predictive factors of Status Epilepticus and its recurrence in patients with adultâ€"onset seizures: A multicenter, long followâ€"up cohort study. Seizure: the Journal of the British Epilepsy Association, 2021, 91, 397-401.	2.0	7
26	A case of epiletic â€œbelly dancingâ€• Journal of the Neurological Sciences, 2021, 429, 119088.	0.6	0
27	Facemask headache: A new nosographic entity among healthcare providers in COVID-19 era. Journal of the Neurological Sciences, 2021, 429, 119801.	0.6	0
28	Usefulness of 24-hour ambulatory EEG monitoring in the diagnosis of typical absences. Journal of the Neurological Sciences, 2021, 429, 117684.	0.6	1
29	Abnormal cortical and subcortical structure in juvenile myoclonic epilepsy demonstrated with advanced MRI analysis. Journal of the Neurological Sciences, 2021, 429, 118300.	0.6	0
30	Random-forest classification of psychogenic non-epileptic seizures and temporal lobe epilepsy. Journal of the Neurological Sciences, 2021, 429, 117781.	0.6	0
31	Impaired embodied cognition in patients with mesial temporal lobe epilepsy and hippocampal sclerosis. Journal of the Neurological Sciences, 2021, 429, 117841.	0.6	0
32	A multimodal neuroimaging approach to non lesional frontal lobe epilepsy. Journal of the Neurological Sciences, 2021, 429, 117689.	0.6	0
33	Status epilepticus amauroticus in a patient with familial photosensitive occipital epilepsy. Journal of the Neurological Sciences, 2021, 429, 119142.	0.6	1
34	Mild case of Unverricht-Lundborg disease presenting as Juvenile myoclonic epilepsy. Journal of the Neurological Sciences, 2021, 429, 117839.	0.6	0
35	A brainstem hypermetabolism in a patient with essential palatal tremor: A simultaneous 18F-FDG-PET/3â€T-MRI study. Journal of the Neurological Sciences, 2021, 429, 119603.	0.6	0
36	Enlarging the clinical spectrum of chorea-acanthocytosis. Neurological Sciences, 2021, 43, 1453.	1.9	1

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37	Late-Onset Ictal Asystole and Falls Related to Severe Coronary Artery Stenosis: A Case Report. Frontiers in Neurology, 2021, 12, 780564.	2.4	0
38	Brainstem Posterior Reversible Encephalopathy Syndrome in an Asymptomatic Patient. Canadian Journal of Neurological Sciences, 2020, 47, 267-269.	0.5	0
39	Network-based atrophy modeling in the common epilepsies: A worldwide ENIGMA study. Science Advances, 2020, 6, .	10.3	97
40	Modulation of GABAergic dysfunction due to SCN1A mutation linked to Hippocampal Sclerosis. Annals of Clinical and Translational Neurology, 2020, 7, 1726-1731.	3.7	4
41	Looking for indicative magnetic resonance imaging signs of hippocampal developmental abnormalities in patients with mesial temporal lobe epilepsy and healthy controls. Epilepsia, 2020, 61, 1714-1722.	5.1	5
42	White matter abnormalities across different epilepsy syndromes in adults: an ENIGMA-Epilepsy study. Brain, 2020, 143, 2454-2473.	7.6	123
43	Management of status epilepticus in patients with liver or kidney disease: a narrative review. Expert Review of Neurotherapeutics, 2020, 21, 1-14.	2.8	4
44	The efficacy of perampanel as adjunctive therapy in drug-resistant focal epilepsy in a â€œreal worldâ€• context: focus on temporal lobe epilepsy. Journal of the Neurological Sciences, 2020, 415, 116903.	0.6	18
45	Epilepsy subtype-specific copy number burden observed in a genome-wide study of 17â€‰458 subjects. Brain, 2020, 143, 2106-2118.	7.6	47
46	Antidepressant effect of vagal nerve stimulation in epilepsy patients: a systematic review. Neurological Sciences, 2020, 41, 3075-3084.	1.9	11
47	A familial t(4;8) translocation segregates with epilepsy and migraine with aura. Annals of Clinical and Translational Neurology, 2020, 7, 855-859.	3.7	7
48	Late drugâ€resistance in mild MTLE: Can it be influenced by preexisting white matter alterations?. Epilepsia, 2020, 61, 924-934.	5.1	7
49	Terminology for psychogenic nonepileptic seizures: The contribution of neuroimaging. Epilepsy and Behavior, 2020, 109, 107063.	1.7	1
50	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.	6.2	237
51	Response to commentary on recommendations for the use of structural <scp>MRI</scp> in the care of patients with epilepsy: A consensus report from the <scp>ILAE</scp> Neuroimaging Task Force. Epilepsia, 2019, 60, 2143-2144.	5.1	74
52	Value of Multimodal Imaging Approach to Diagnosis of Neurosarcoidosis. Brain Sciences, 2019, 9, 243.	2.3	13
53	Imaging Genetics for Benign Mesial Temporal Lobe Epilepsy. , 2019, , 48-54.		2
54	Functional activity changes in memory and emotional systems of healthy subjects with dÃ©jÃ vu. Epilepsy and Behavior, 2019, 97, 8-14.	1.7	3

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55	Recommendations for the use of structural magnetic resonance imaging in the care of patients with epilepsy: A consensus report from the International League Against Epilepsy Neuroimaging Task Force. Epilepsia, 2019, 60, 1054-1068.	5.1	184
56	Hypertension, seizures, and epilepsy: a review on pathophysiology and management. Neurological Sciences, 2019, 40, 1775-1783.	1.9	51
57	HCN ion channels and accessory proteins in epilepsy: genetic analysis of a large cohort of patients and review of the literature. Epilepsy Research, 2019, 153, 49-58.	1.6	32
58	Psychiatric Assessment in Patients with Mild Temporal Lobe Epilepsy. Behavioural Neurology, 2019, 2019, 1-9.	2.1	8
59	Insight into epileptic and physiological dÃ©jÃ vu : from a multicentric cohort study. European Journal of Neurology, 2019, 26, 407-414.	3.3	2
60	Kufs disease due to mutation ofCLN6: clinical, pathological and molecular genetic features. Brain, 2019, 142, 59-69.	7.6	28
61	Management of psychogenic nonâ€epileptic seizures: a multidisciplinary approach. European Journal of Neurology, 2019, 26, 205.	3.3	64
62	Comparative risk of major congenital malformations with eight different antiepileptic drugs: a prospective cohort study of the EURAP registry. Lancet Neurology, The, 2018, 17, 530-538.	10.2	348
63	Midbrain meningioma causing subacute parkinsonism. Neurology: Clinical Practice, 2018, 8, 166-168.	1.6	0
64	Structural brain abnormalities in the common epilepsies assessed in a worldwide ENIGMA study. Brain, 2018, 141, 391-408.	7.6	352
65	Value of clinical features to differentiate refractory epilepsy from mimics: a prospective longitudinal cohort study. European Journal of Neurology, 2018, 25, 711-717.	3.3	5
66	Brandâ€toâ€generic levetiracetam switching: a 4â€year prospective observational realâ€life study. European Journal of Neurology, 2018, 25, 666-671.	3.3	17
67	Psychopathological constellation in patients with PNES: A new hypothesis. Epilepsy and Behavior, 2018, 78, 297-301.	1.7	21
68	The meaning of anxiety in patients with PNES. Epilepsy and Behavior, 2018, 87, 248.	1.7	3
69	The application of artificial intelligence to understand the pathophysiological basis of psychogenic nonepileptic seizures. Epilepsy and Behavior, 2018, 87, 167-172.	1.7	29
70	A novel de novo HCN1 loss-of-function mutation in genetic generalized epilepsy causing increased neuronal excitability. Neurobiology of Disease, 2018, 118, 55-63.	4.4	47
71	Neuropsychological profile of mild temporal lobe epilepsy. Epilepsy and Behavior, 2018, 85, 222-226.	1.7	10
72	A Loss-of-Function HCN4 Mutation Associated With Familial Benign Myoclonic Epilepsy in Infancy Causes Increased Neuronal Excitability. Frontiers in Molecular Neuroscience, 2018, 11, 269.	2.9	25

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73	Role of Pharmacogenomics in Antiepileptic Drug Therapy: Current Status and Future Perspectives. Current Pharmaceutical Design, 2018, 23, 5760-5765.	1.9	12
74	Diagnostic Biomarkers of Epilepsy. Current Pharmaceutical Biotechnology, 2018, 19, 440-450.	1.6	20
75	The mystery of unexplained traumatic sudden falls. A clinical case that adds a new feasible cause. Neurological Sciences, 2017, 38, 1115-1117.	1.9	7
76	AMPA receptors and perampanel behind selected epilepsies: current evidence and future perspectives. Expert Opinion on Pharmacotherapy, 2017, 18, 1751-1764.	1.8	54
77	Epilepsy and sleep disorders improve in adolescents and adults with Angelman syndrome: A multicenter study on 46 patients. Epilepsy and Behavior, 2017, 75, 225-229.	1.7	20
78	Advanced morphological neuroimaging study in lateral temporal lobe epilepsy: A multicentric study. Epilepsy and Behavior, 2017, 74, 69-72.	1.7	0
79	Hypersomnia hiding a bipolar disorder. Neurological Sciences, 2017, 38, 2057-2058.	1.9	0
80	Ictal 18F-FDG PET/MRI in a Patient With Cortical Heterotopia and Focal Epilepsy. Clinical Nuclear Medicine, 2017, 42, 768-769.	1.3	8
81	Gerstmannâ€“Straussler–Scheinker disease with <scp>PRNP</scp> P102L heterozygous mutation presenting as progressive myoclonus epilepsy. European Journal of Neurology, 2017, 24, e87-e88.	3.3	5
82	Acute bulbar palsy without ophtalmoplegia associated with anti-GD3 IgM antibodies. Neurological Sciences, 2017, 38, 521-523.	1.9	1
83	Deep Learning Representation from Electroencephalography of Early-Stage Creutzfeldt-Jakob Disease and Features for Differentiation from Rapidly Progressive Dementia. International Journal of Neural Systems, 2017, 27, 1650039.	5.2	104
84	Validation Study of Italian Version of Inventory for DÃ©jÃ Vu Experiences Assessment (I-IDEA): A Screening Tool to Detect DÃ©jÃ Vu Phenomenon in Italian Healthy Individuals. Behavioral Sciences (Basel, Switzerland), 2017, 7, 50.	2.1	1
85	The Natural History of Epilepsy in 163 Untreated Patients: Looking for â€œOligoepilepsyâ€• PLoS ONE, 2016, 11, e0161722.	2.5	2
86	Integrity of the corpus callosum in patients with benign temporal lobe epilepsy. Epilepsia, 2016, 57, 590-596.	5.1	17
87	An SNP site in pri-miR-124, a brain expressed miRNA gene, no contribution to mesial temporal lobe epilepsy in an Italian sample. Neurological Sciences, 2016, 37, 1335-1339.	1.9	15
88	Long-term outcome of mild mesial temporal lobe epilepsy. Neurology, 2016, 86, 1904-1910.	1.1	40
89	Lacosamide in patients with temporal lobe epilepsy: An observational multicentric open-label study. Epilepsy and Behavior, 2016, 58, 111-114.	1.7	12
90	Relevance of clinical context in the diagnosticâ€therapeutic approach to status epilepticus. Epilepsia, 2016, 57, 1527-1529.	5.1	4

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91	Pharmacological modulation in mesial temporal lobe epilepsy: Current status and future perspectives. Pharmacological Research, 2016, 113, 421-425.	7.1	20
92	Epilepsy in cerebrovascular diseases: Review of experimental and clinical data with metaâ€analysis of risk factors. Epilepsia, 2016, 57, 1205-1214.	5.1	122
93	Genetics of reflex seizures and epilepsies in humans and animals. Epilepsy Research, 2016, 121, 47-54.	1.6	60
94	Tremor pattern differentiates drug-induced resting tremor from Parkinson disease. Parkinsonism and Related Disorders, 2016, 25, 100-103.	2.2	17
95	PRIMA1 mutation: a new cause of nocturnal frontal lobe epilepsy. Annals of Clinical and Translational Neurology, 2015, 2, 821-830.	3.7	21
96	<scp>DEPDC</scp>5 mutations are not a frequent cause of familial temporal lobe epilepsy. Epilepsia, 2015, 56, e168-71.	5.1	37
97	Why should we change the term psychogenic nonepileptic seizures?. Epilepsia, 2015, 56, 1178-1179.	5.1	5
98	Letter: Beyond and within <scp>CA</scp>1 subfield in magnetic resonance imaging negative temporal lobe epilepsy. Epilepsia, 2015, 56, 1471-1471.	5.1	0
99	White matter abnormalities differentiate severe from benign temporal lobe epilepsy. Epilepsia, 2015, 56, 1109-1116.	5.1	38
100	Brivaracetam: review of its pharmacology and potential use as adjunctive therapy in patients with partial onset seizures. Drug Design, Development and Therapy, 2015, 9, 5719.	4.3	11
101	Profile of brivaracetam and its potential in the treatment of epilepsy. Neuropsychiatric Disease and Treatment, 2015, 11, 2967.	2.2	14
102	3â€T magnetic resonance imaging simultaneous automated multimodal approach improves detection of ambiguous visual hippocampal sclerosis. European Journal of Neurology, 2015, 22, 725.	3.3	13
103	Polymorphism of the multidrug resistance 1 gene MDR1/ABCB1 C3435T and response to antiepileptic drug treatment in temporal lobe epilepsy. Seizure: the Journal of the British Epilepsy Association, 2015, 24, 124-126.	2.0	50
104	PCDH19 mutations in female patients from Southern Italy. Seizure: the Journal of the British Epilepsy Association, 2015, 24, 118-120.	2.0	7
105	Epilepsy and Immune System: A Tour Around the Current Literature. , 2015, , 163-182.		0
106	No evidence of a role for cystatin <scp>B</scp> gene in juvenile myoclonic epilepsy. Epilepsia, 2015, 56, e40-3.	5.1	7
107	A puzzling case without solution: isolated late-onset epileptic seizure. Neurological Sciences, 2015, 36, 2303-2304.	1.9	1
108	Epilepsy associated with Leukoaraiosis mainly affects temporal lobe: a casual or causal relationship?. Epilepsy Research, 2015, 109, 1-8.	1.6	39

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109	Neuro-anatomical differences among epileptic and non-epileptic dÃ©jÃ-vu. Cortex, 2015, 64, 1-7.	2.4	14
110	Positivity to p-ANCA in patients with status epilepticus. BMC Neurology, 2014, 14, 148.	1.8	5
111	Mild <scp>L</scp>afora disease: Clinical, neurophysiologic, and genetic findings. Epilepsia, 2014, 55, e129-33.	5.1	43
112	Generalized versus partial reflex seizures: A review. Seizure: the Journal of the British Epilepsy Association, 2014, 23, 512-520.	2.0	70
113	Rosacea-like facial rash related to metformin administration in a young woman. BMC Pharmacology & Toxicology, 2014, 15, 3.	2.4	12
114	Reversible symmetrical external capsule hyperintensity as an early finding of autoimmune encephalitis. Neurological Sciences, 2014, 35, 1147-1149.	1.9	6
115	Thalamotemporal impairment in benign temporal lobe epilepsy: Same hypotheses?. Epilepsia, 2014, 55, 944-944.	5.1	0
116	The role of calcium channel mutations in human epilepsy. Progress in Brain Research, 2014, 213, 87-96.	1.4	18
117	Permutation entropy of scalp EEG: A tool to investigate epilepsies. Clinical Neurophysiology, 2014, 125, 13-20.	1.5	59
118	Autosomal dominant lateral temporal epilepsy (ADLTE): Absence of chromosomal rearrangements in LGI1 gene. Epilepsy Research, 2014, 108, 597-599.	1.6	2
119	Teaching NeuroImages: Pseudohypertrophic cerebral cortex in end-stage Creutzfeldt-Jakob disease. Neurology, 2013, 80, e21-e21.	1.1	5
120	Hippocampal sclerosis worsens autosomal dominant nocturnal frontal lobe epilepsy (ADNFLE) phenotype related to CHRNB2 mutation. European Journal of Neurology, 2013, 20, 591-593.	3.3	7
121	Mutations in PRRT2 result in familial infantile seizures with heterogeneous phenotypes including febrile convulsions and probable SUDEP. Epilepsy Research, 2013, 104, 280-284.	1.6	29
122	Failure to confirm association of a polymorphism in KCNMB4 gene with mesial temporal lobe epilepsy. Epilepsy Research, 2013, 106, 284-287.	1.6	4
123	Family history and frontal lobe seizures predict long-term remission in newly diagnosed cryptogenic focal epilepsy. Epilepsy Research, 2013, 107, 101-108.	1.6	19
124	Comment on BrÃ¡zdil (2012) â€œUnveiling the mystery of dÃ¨jÃ-vÃ¹: The structural anatomy of dÃ¨jÃ-vÃ¹â€•. Cortex, 2013, 49, 1162.	2.4	3
125	Relationship between genetic variant in pre-microRNA-146a and genetic predisposition to temporal lobe epilepsy: A caseâ€“control study. Gene, 2013, 516, 181-183.	2.2	22
126	Pharmacodynamic potentiation of antiepileptic drugsâ€™ effects by some HMG-CoA reductase inhibitors against audiogenic seizures in DBA/2 mice. Pharmacological Research, 2013, 70, 1-12.	7.1	49

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127	No evidence for a role of the coding variant of the Toll-like receptor 4 gene in temporal lobe epilepsy. Seizure: the Journal of the British Epilepsy Association, 2013, 22, 791-793.	2.0	2
128	Detection of hippocampal atrophy in patients with temporal lobe epilepsy: A 3-Tesla MRI shape. Epilepsy and Behavior, 2013, 28, 489-493.	1.7	25
129	Lamotrigine positively affects the development of psychiatric comorbidity in epileptic animals, while psychiatric comorbidity aggravates seizures. Epilepsy and Behavior, 2013, 28, 232-240.	1.7	30
130	Septo-optic dysplasia plus bilateral perisylvian polymicrogyria: a case report. Neurological Sciences, 2013, 34, 1479-1480.	1.9	3
131	A possible case of natalizumab-dependent suicide attempt: A brief review about drugs and suicide. Journal of Pharmacology and Pharmacotherapeutics, 2013, 4, 90.	0.4	5
132	Advanced MRI Morphologic Study Shows No Atrophy in Healthy Individuals with Hippocampal Hyperintensity. American Journal of Neuroradiology, 2013, 34, 1585-1588.	2.4	5
133	Divergent effects of the <scp>T</scp>1174S <scp>SCN1A</scp> mutation associated with seizures and hemiplegic migraine. Epilepsia, 2013, 54, 927-935.	5.1	63
134	Antiâ€“Nâ€methylâ€Dâ€aspartateâ€glutamicâ€receptor encephalitis presenting as paroxysmal exerciseâ€induced foot weakness. Movement Disorders, 2013, 28, 820-822.	3.9	11
135	Post-stroke Movement Disorders: Clinical Manifestations and Pharmacological Management. Current Neuropharmacology, 2012, 10, 254-262.	2.9	49
136	Homozygous c.649dupC mutation in PRRT2 worsens the BFIS/PKD phenotype with mental retardation, episodic ataxia, and absences. Epilepsia, 2012, 53, e196-9.	5.1	78
137	A Functional Genetic Variation of the 5â€HTR2A Receptor Affects Age at Onset in Patients with Temporal Lobe Epilepsy. Annals of Human Genetics, 2012, 76, 277-282.	0.8	5
138	Topiramate and temporal lobe epilepsy: an open-label study. Epileptic Disorders, 2012, 14, 163-166.	1.3	5
139	Cardiac denervation precedes nigrostriatal damage in idiopathic rapid eye movement sleep behavior disorder. Movement Disorders, 2012, 27, 1068-1069.	3.9	15
140	Migraine attack triggering a generalised seizure: is this a case of migralepsy or ictal epileptic headache?. Neurological Sciences, 2012, 33, 957-959.	1.9	7
141	Neuroanatomic correlates of psychogenic nonepileptic seizures: A cortical thickness and VBM study. Epilepsia, 2012, 53, 377-385.	5.1	140
142	Usefulness of rectal biopsy for the diagnosis of Kufs disease: a controlled study and review of the literature. European Journal of Neurology, 2012, 19, 1331-1336.	3.3	5
143	Unilateral basal ganglia atrophy in a patient with tuberous sclerosis complex and hemichorea. Movement Disorders, 2012, 27, 458-460.	3.9	5
144	Benign mesial temporal lobe epilepsy. Nature Reviews Neurology, 2011, 7, 237-240.	10.1	76

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145	Rating scale for psychogenic nonepileptic seizures: Scale development and clinimetric testing. Epilepsy and Behavior, 2011, 21, 128-131.	1.7	21
146	Blocking out the real diagnosis. Lancet, The, 2011, 377, 690.	13.7	5
147	Blocking out the real diagnosis â€“ Authors' reply. Lancet, The, 2011, 378, 316.	13.7	0
148	Neocortical thinning in â€œbenignâ€•mesial temporal lobe epilepsy. Epilepsia, 2011, 52, 712-717.	5.1	51
149	A functional polymorphism in the SCN1A gene does not influence antiepileptic drug responsiveness in Italian patients with focal epilepsy. Epilepsia, 2011, 52, e40-e44.	5.1	50
150	COMBINED USE OF CARDIAC M-I123-IODOBENZYLGUANIDINE SCINTIGRAPHY AND 123I-FP-CIT SINGLE PHOTON EMISSION COMPUTED TOMOGRAPHY IN OLDER ADULTS WITH RAPID EYE MOVEMENT SLEEP BEHAVIOR DISORDER. Journal of the American Geriatrics Society, 2011, 59, 928-929.	2.6	11
151	Bellâ€™s palsy: a manifestation of prediabetes?. Acta Neurologica Scandinavica, 2011, 123, 68-72.	2.1	20
152	Age at onset predicts good seizure outcome in sporadic non-lesional and mesial temporal sclerosis based temporal lobe epilepsy. Journal of Neurology, Neurosurgery and Psychiatry, 2011, 82, 555-559.	1.9	36
153	Tacrolimus-Induced Polyneuropathy After Heart Transplantation. Clinical Neuropharmacology, 2010, 33, 161-162.	0.7	18
154	Voxel-based morphometry of adulthood patients with temporal lobe epilepsy. BMC Geriatrics, 2010, 10, .	2.7	0
155	Voxelâ€based morphometry of sporadic epileptic patients with mesiotemporal sclerosis. Epilepsia, 2010, 51, 506-510.	5.1	43
156	Temporal lobe abnormalities on brain MRI in healthy volunteers. Neurology, 2010, 74, 553-557.	1.1	47
157	TEMPORAL LOBE ABNORMALITIES ON BRAIN MRI IN HEALTHY VOLUNTEERS: A PROSPECTIVE CASE-CONTROL STUDY. Neurology, 2010, 75, 377-378.	1.1	15
158	Non-paraneoplastic limbic encephalitis characterized by mesio-temporal seizures and extratemporal lesions: A case report. Seizure: the Journal of the British Epilepsy Association, 2010, 19, 446-449.	2.0	17
159	Anti-GM1 antibodies are not associated with cerebral atrophy in patients with multiple sclerosis. Multiple Sclerosis Journal, 2009, 15, 114-115.	3.0	7
160	The DRD2 TaqIA polymorphism associated with changed midbrain volumes in healthy individuals. Genes, Brain and Behavior, 2009, 8, 459-463.	2.2	28
161	Prognostic factors in patients with mesial temporal lobe epilepsy. Epilepsia, 2009, 50, 41-44.	5.1	51
162	Familial mesial temporal lobe epilepsies: Clinical and genetic features. Epilepsia, 2009, 50, 55-57.	5.1	30

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163	Involvement of cardiac sympathetic nerve endings in a patient with idiopathic RBD and intact nigrostriatal pathway. Parkinsonism and Related Disorders, 2009, 15, 789-791.	2.2	13
164	Unexpected Detection of Melanoma Brain Metastasis by PET With Iodine-124 Î²CIT. Clinical Nuclear Medicine, 2009, 34, 698-699.	1.3	13
165	Anti-NMDA receptor encephalitis: aÂvideo case report. Epileptic Disorders, 2009, 11, 267-269.	1.3	11
166	Familial mesial temporal lobe epilepsy (FMTLE). Journal of Neurology, 2008, 255, 16-23.	3.6	60
167	Orolingual tremor as unusual presentation of antiâ€Hu paraneoplastic syndrome. Movement Disorders, 2008, 23, 1791-1792.	3.9	6
168	Spontaneous obliteration of MRI-silent cerebral angiomatosis revealed by CT angiography in a patient with Sturgeâ€“Weber syndrome. Journal of the Neurological Sciences, 2008, 264, 168-172.	0.6	6
169	Hippocampal and thalamic atrophy in mild temporal lobe epilepsy. Neurology, 2008, 71, 1094-1101.	1.1	91
170	Impact of catechol-O-methyltransferase Val108/158 Met genotype on hippocampal and prefrontal gray matter volume. NeuroReport, 2008, 19, 405-408.	1.2	66
171	MAO A VNTR polymorphism and variation in human morphology: a VBM study. NeuroReport, 2008, 19, 1107-1110.	1.2	24
172	Monotherapy for partial epilepsy: focus on levetiracetam. Neuropsychiatric Disease and Treatment, 2008, 4, 33.	2.2	40
173	Hepatonecrosis and cholangitis related to long-term phenobarbital therapy: An autopsy report of two patients. Seizure: the Journal of the British Epilepsy Association, 2007, 16, 653-656.	2.0	10
174	Association between the M129V variant allele of PRNP gene and mild temporal lobe epilepsy in women. Neuroscience Letters, 2007, 421, 1-4.	2.1	13
175	Serotonin transporter gene (5-Htt): Association analysis with temporal lobe epilepsy. Neuroscience Letters, 2007, 421, 52-56.	2.1	32
176	Further evidence of genetic heterogeneity in families with autosomal dominant nocturnal frontal lobe epilepsy. Epilepsy Research, 2007, 74, 70-73.	1.6	29
177	Usefulness of a morning routine EEG recording in patients with juvenile myoclonic epilepsy. Epilepsy Research, 2007, 77, 17-21.	1.6	48
178	Electroclinical Features of a Family with Simple Febrile Seizures and Temporal Lobe Epilepsy Associated with SCN1A Loss-of-Function Mutation. Epilepsia, 2007, 48, 1691-1696.	5.1	44
179	Mutational Analysis of EFHC1 Gene in Italian Families with Juvenile Myoclonic Epilepsy. Epilepsia, 2007, 48, 1686-1690.	5.1	44
180	SUNCT and high nocturnal prolactin levels: some new unusual characteristics. Journal of Headache and Pain, 2007, 8, 114-118.	6.0	12

#	Article	IF	Citations
181	Levetiracetam in patients with generalised epilepsy and myoclonic seizures: An open label study. Seizure: the Journal of the British Epilepsy Association, 2006, 15, 214-218.	2.0	67
182	Is Language Lateralization in Temporal Lobe Epilepsy Patients Related to the Nature of the Epileptogenic Lesion?. Epilepsia, 2006, 47, 916-920.	5.1	38
183	Reflex periodic spasms induced by eating. Brain and Development, 2006, 28, 170-174.	1.1	28
184	MRI evidence of mesial temporal sclerosis in sporadic "benign" temporal lobe epilepsy. Neurology, 2006, 66, 562-565.	1.1	91
185	ApoE Epsilon4 Allele and Disease Duration Affect Verbal Learning in Mild Temporal Lobe Epilepsy. Epilepsia, 2005, 46, 110-117.	5.1	53
186	Amygdala dysplasia with temporal lobe epilepsy and obsessive-compulsive disorder: An fMRI/EEG study. Neurology, 2005, 64, 1309-1310.	1.1	4
187	Identification of an Nav1.1 sodium channel (SCN1A) loss-of-function mutation associated with familial simple febrile seizures. Proceedings of the National Academy of Sciences of the United States of America, 2005, 102, 18177-18182.	7.1	193
188	How reliable are fMRIâ€“EEG studies of epilepsy? A nonparametric approach to analysis validation and optimization. NeuroImage, 2005, 24, 192-199.	4.2	41
189	Startle epilepsy complicating aspartylglucosaminuria. Brain and Development, 2004, 26, 130-133.	1.1	12
190	Gene conversion events in adult-onset spinal muscular atrophy. Acta Neurologica Scandinavica, 2004, 109, 151-154.	2.1	12
191	Prodynorphin Gene Promoter Polymorphism and Temporal Lobe Epilepsy. Epilepsia, 2003, 44, 1255-1256.	5.1	27
192	Two Novel SCN1A Missense Mutations in Generalized Epilepsy with Febrile Seizures Plus. Epilepsia, 2003, 44, 1257-1258.	5.1	56
193	GABA(B) receptor 1 polymorphism (G1465A) is associated with temporal lobe epilepsy. Neurology, 2003, 60, 560-563.	1.1	127
194	Temporal Lobe Epilepsy as a Unique Manifestation of Multiple Sclerosis. Canadian Journal of Neurological Sciences, 2003, 30, 228-232.	0.5	36
195	Silent Celiac Disease in Patients with Childhood Localization-Related Epilepsies. Epilepsia, 2002, 42, 1153-1155.	5.1	54
196	A Novel Mutation in the Notch3 Gene in an Italian Family With Cerebral Autosomal Dominant Arteriopathy With Subcortical Infarcts and Leukoencephalopathy. Archives of Neurology, 2001, 58, 1418.	4.5	29
197	Benign Partial Epilepsies of Adolescence: A Report of 37 New Cases. Epilepsia, 2001, 42, 1549-1552.	5.1	10
198	The parkin gene is not involved in late-onset Parkinsonâ€™s disease. Neurology, 2001, 57, 359-362.	1.1	48

#	Article	IF	Citations
199	Cerebral venous thrombosis and isolated intracranial hypertension without papilledema in CDH. Neurology, 2001, 57, 31-36.	1.1	114
200	Juvenile Huntingtonâ€™s disease presenting as progressive myoclonic epilepsy. Neurology, 2001, 57, 708-711.	1.1	62