Antonio Gambardella

List of Publications by Year
in descending order

480 papers	20,911 citations	13099 68 h-index	17592 121 g-index
496 all docs	496 docs citations	496 times ranked	20650 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	Hyper-religiosity and visual hallucinations in a patient with frontotemporal dementia carrying a double variant in GRN gene. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2022, 23, 87-90.	1.7	0
3	Face atopic dermatitis resistant to dupilumab: a case series of three patients successfully treated with upadacitinib. Journal of the European Academy of Dermatology and Venereology, 2022, 36, .	2.4	23
4	A systemsâ€level analysis highlights microglial activation as a modifying factor in common epilepsies. Neuropathology and Applied Neurobiology, 2022, 48, .	3.2	22
5	Topographic divergence of atypical cortical asymmetry and atrophy patterns in temporal lobe epilepsy. Brain, 2022, 145, 1285-1298.	7.6	18
6	Atlas of lesion locations and postsurgical seizure freedom in focal cortical dysplasia: A MELD study. Epilepsia, 2022, 63, 61-74.	5.1	36
7	Cognitive impairment and type 2 diabetes mellitus: Focus of SGLT2 inhibitors treatment. Pharmacological Research, 2022, 176, 106062.	7.1	44
8	Association of ultraâ€rare coding variants with genetic generalized epilepsy: A caseâ€“control whole exome sequencing study. Epilepsia, 2022, 63, 723-735.	5.1	8
9	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
10	Cerebellar Agenesis and Bilateral Polimicrogyria Associated with Rare Variants of CUB and Sushi Multiple Domains 1 Gene (CSMD1): A Longitudinal Neuropsychological and Neuroradiological Case Study. International Journal of Environmental Research and Public Health, 2022, 19, 1224.	2.6	0
11	Loss of Neuron Navigator 2 Impairs Brain and Cerebellar Development. Cerebellum, 2022, , 1.	2.5	5
12	Electroclinical Features and Long-term Seizure Outcome in Patients With Eyelid Myoclonia With Absences. Neurology, 2022, 98, .	1.1	15
13	Non-Coding RNAs: New Biomarkers and Therapeutic Targets for Temporal Lobe Epilepsy. International Journal of Molecular Sciences, 2022, 23, 3063.	4.1	8
14	Epileptic belly dancing: a videoâ€polygraphic recording. Epileptic Disorders, 2022, 24, 442-444.	1.3	0
15	Human iPSC Modeling of Genetic Febrile Seizure Reveals Aberrant Molecular and Physiological Features Underlying an Impaired Neuronal Activity. Biomedicines, 2022, 10, 1075.	3.2	10
16	Late epileptic seizures following cerebral venous thrombosis: a systematic review and meta-analysis. Neurological Sciences, 2022, 43, 5229-5236.	1.9	8
17	Eventâ€based modeling in temporal lobe epilepsy demonstrates progressive atrophy from crossâ€sectional data. Epilepsia, 2022, 63, 2081-2095.	5.1	11
18	Resting-State EEG Classification forÂPNES Diagnosis. Lecture Notes in Computer Science, 2022, , 526-538.	1.3	2

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19	Networks Underlie Temporal Onset of Dysplasiaâ€Related Epilepsy: A <scp>MELD</scp> Study. Annals of Neurology, 2022, 92, 503-511.	5.3	7
20	Epilepsy, Immunity and Neuropsychiatric Disorders. Current Neuropharmacology, 2022, 20, .	2.9	4
21	The impact of sexual abuse on psychopathology of patients with psychogenic nonepileptic seizures. Neurological Sciences, 2021, 42, 1423-1428.	1.9	3
22	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
23	Coverage of the requirements of first and second level stroke unit in Italy. Neurological Sciences, 2021, 42, 1073-1079.	1.9	1
24	Varicella zoster immunity loss in multiple sclerosis patient treated with ocrelizumab. Clinical Immunology, 2021, 223, 108554.	3.2	6
25	Opicapone-induced reversible myopathy in a patient with advanced Parkinsonâ€™s disease and familial hyperCKemia. Neurological Sciences, 2021, 42, 2583-2585.	1.9	1
26	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
27	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
28	Facemask headache: a new nosographic entity among healthcare providers in COVID-19 era. Neurological Sciences, 2021, 42, 1267-1276.	1.9	22
29	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
30	Incidental evidence of hypointensity in brain grey nuclei on routine MR imaging: when to suspect a neurodegenerative disorder?. Neurological Sciences, 2021, , 1.	1.9	1
31	Diagnostic and therapeutic approach to drug-resistant juvenile myoclonic epilepsy. Expert Review of Neurotherapeutics, 2021, 21, 1265-1273.	2.8	12
32	Assessing the role of rare genetic variants in drugâ€resistant, nonâ€lesional focal epilepsy. Annals of Clinical and Translational Neurology, 2021, 8, 1376-1387.	3.7	16
33	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
34	Dupilumab and conjunctivitis: a case series of twenty patients. Journal of the European Academy of Dermatology and Venereology, 2021, 35, e612-e614.	2.4	9
35	Rare manifestations and malignancies in tuberous sclerosis complex: findings from the TuberOus SClerosis registry to increAse disease awareness (TOSCA). Orphanet Journal of Rare Diseases, 2021, 16, 301.	2.7	15
36	GuillainÂ-BarrÃ© syndrome following BNT162b2 COVID-19 vaccine. Neurological Sciences, 2021, 42, 4401-4402.	1.9	51

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37	Orbito-frontal thinning together with a somatoform dissociation might be the fingerprint of PNES. Epilepsy and Behavior, 2021, 121, 108044.	1.7	9
38	Postictal Psychosis in Epilepsy: A Clinicogenetic Study. Annals of Neurology, 2021, 90, 464-476.	5.3	11
39	Selection of antiseizure medications for first add-on use: A consensus paper. Epilepsy and Behavior, 2021, 122, 108087.	1.7	6
40	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
41	Two cases of unilateral wasting and weakness of distal upper limb: Similar onset and different diagnosis in young patients. Journal of the Neurological Sciences, 2021, 429, 119405.	0.6	0
42	A case of epiletic â€œbelly dancingâ€• Journal of the Neurological Sciences, 2021, 429, 119088.	0.6	0
43	Clinical importance of mimics. Journal of the Neurological Sciences, 2021, 429, 117933.	0.6	0
44	End of dose interval (EDI) symptoms in patients undergoing treatment with natalizumab. Journal of the Neurological Sciences, 2021, 429, 118177.	0.6	0
45	Facemask headache: A new nosographic entity among healthcare providers in COVID-19 era. Journal of the Neurological Sciences, 2021, 429, 119801.	0.6	0
46	Usefulness of 24-hour ambulatory EEG monitoring in the diagnosis of typical absences. Journal of the Neurological Sciences, 2021, 429, 117684.	0.6	1
47	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
48	Random-forest classification of psychogenic non-epileptic seizures and temporal lobe epilepsy. Journal of the Neurological Sciences, 2021, 429, 117781.	0.6	0
49	Parkinsonism and corticospinal tracts involvement in hepatic encephalopathy. Journal of the Neurological Sciences, 2021, 429, 118611.	0.6	0
50	Sex differences in cortical hemodynamic response to levodopa in Parkinson's disease patients: A functional NIRS study. Journal of the Neurological Sciences, 2021, 429, 119550.	0.6	0
51	Impaired embodied cognition in patients with mesial temporal lobe epilepsy and hippocampal sclerosis. Journal of the Neurological Sciences, 2021, 429, 117841.	0.6	0
52	A multimodal neuroimaging approach to non lesional frontal lobe epilepsy. Journal of the Neurological Sciences, 2021, 429, 117689.	0.6	0
53	Headache and sleep-related breathing disorders among patients with sclerosteosis and disease carriers. Journal of the Neurological Sciences, 2021, 429, 119265.	0.6	0
54	PRECLINICAL MARKERS OF NEUROPATHIC DAMAGE IN TYPE 1 DIABETES MELLITUS. Journal of the Neurological Sciences, 2021, 429, 118542.	0.6	0

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55	Opicapone-induced reversible myopathy in a patient with advanced Parkinson's disease and familial hyperckemia. Journal of the Neurological Sciences, 2021, 429, 119555.	0.6	0
56	Myositis associated to COVID-19 mimics an acute exacerbation in myasthenia gravis patient. Journal of the Neurological Sciences, 2021, 429, 119865.	0.6	0
57	Status epilepticus amauroticus in a patient with familial photosensitive occipital epilepsy. Journal of the Neurological Sciences, 2021, 429, 119142.	0.6	1
58	A case of neuromyelitis optica spectrum disorder (NMOSD) and acute myositis following SARS-CoV-2 infection. Journal of the Neurological Sciences, 2021, 429, 119862.	0.6	5
59	Perampanel as first add-on antiseizure medication: Italian consensus clinical practice statements. BMC Neurology, 2021, 21, 410.	1.8	8
60	Mild case of Unverricht-Lundborg disease presenting as Juvenile myoclonic epilepsy. Journal of the Neurological Sciences, 2021, 429, 117839.	0.6	0
61	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
62	Epilepsy in â€œSunflower syndromeâ€• electroclinical features, therapeutic response, and long-term follow-up. Seizure: the Journal of the British Epilepsy Association, 2021, 93, 8-12.	2.0	7
63	Neuroma of the Inferior Alveolar Nerve. Neurology: Clinical Practice, 2021, 11, e582-e584.	1.6	0
64	Correlation EEG and Clinic. , 2021, , 75-94.		0
65	Progressive Myoclonus Epilepsies. Neurology: Genetics, 2021, 7, e641.	1.9	20
66	Late-Onset Ictal Asystole and Falls Related to Severe Coronary Artery Stenosis: A Case Report. Frontiers in Neurology, 2021, 12, 780564.	2.4	0
67	Usefulness of cardiac parasympathetic index in CPAP â€treated patients with obstructive sleep apnea: A preliminary study. Journal of Sleep Research, 2020, 29, e12893.	3.2	2
68	Brainstem Posterior Reversible Encephalopathy Syndrome in an Asymptomatic Patient. Canadian Journal of Neurological Sciences, 2020, 47, 267-269.	0.5	0
69	Homozygous STXBP1 variant causes encephalopathy and gain-of-function in synaptic transmission. Brain, 2020, 143, 441-451.	7.6	46
70	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
71	Objective bilateral tinnitus from palatal nystagmus. Audio and video features of a rare case of palatal myoclonus. American Journal of Otolaryngology - Head and Neck Medicine and Surgery, 2020, 41, 102739.	1.3	2
72	Network-based atrophy modeling in the common epilepsies: A worldwide ENIGMA study. Science Advances, 2020, 6, .	10.3	97

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73	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
74	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
75	Natural clusters of tuberous sclerosis complex (TSC)-associated neuropsychiatric disorders (TAND): new findings from the TOSCA TAND research project. Journal of Neurodevelopmental Disorders, 2020, 12, 24.	3.1	16
76	Burden of Illness and Quality of Life in Tuberous Sclerosis Complex: Findings From the TOSCA Study. Frontiers in Neurology, 2020, 11, 904.	2.4	20
77	White matter abnormalities across different epilepsy syndromes in adults: an ENIGMA-Epilepsy study. Brain, 2020, 143, 2454-2473.	7.6	123
78	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
79	An invited commentary on â€œPrognostic significance of the controlling nutritional status (CONUT) score in patients with colorectal cancer: A systematic review and meta-analysisâ€•[Int. J. Surg. 78 (2020) 91â€“96]. International Journal of Surgery, 2020, 79, 38-39.	2.7	0
80	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
81	Epilepsy subtype-specific copy number burden observed in a genome-wide study of 17â€‰458 subjects. Brain, 2020, 143, 2106-2118.	7.6	47
82	Focal neuromyotonia associated with a C9ORF72 expansion mutation. Muscle and Nerve, 2020, 62, E63-E65.	2.2	1
83	COVID-19 risk contagion: Organization and procedures in a South Italy geriatric oncology ward. Journal of Geriatric Oncology, 2020, 11, 1187-1188.	1.0	27
84	Autonomic Nervous System and Cognitive Impairment in Older Patients: Evidence From Long-Term Heart Rate Variability in Real-Life Setting. Frontiers in Aging Neuroscience, 2020, 12, 40.	3.4	27
85	Testing association of rare genetic variants with resistance to three common antiseizure medications. Epilepsia, 2020, 61, 657-666.	5.1	22
86	Antiepileptic Drug Teratogenicity and De Novo Genetic Variation Load. Annals of Neurology, 2020, 87, 897-906.	5.3	9
87	A Family With a Complex Phenotype Caused by Two Different Rare Metabolic Disorders: GLUT1 and Very-Long-Chain Fatty Acid Dehydrogenase (VLCAD) Deficiencies. Frontiers in Neurology, 2020, 11, 514.	2.4	2
88	A case of Atopic dermatitis and Hidradenitis Suppurativa successfully treated with Dupilumab. Journal of the European Academy of Dermatology and Venereology, 2020, 34, e284-e286.	2.4	12
89	Advances in genetic testing and optimization of clinical management in children and adults with epilepsy. Expert Review of Neurotherapeutics, 2020, 20, 251-269.	2.8	45
90	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

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91	Late drugâ€resistance in mild MTLE: Can it be influenced by preexisting white matter alterations?. Epilepsia, 2020, 61, 924-934.	5.1	7
92	Terminology for psychogenic nonepileptic seizures: The contribution of neuroimaging. Epilepsy and Behavior, 2020, 109, 107063.	1.7	1
93	Listeria infection after treatment with alemtuzumab: a case report and literature review. Would antibiotic prophylaxis be considered?. Infezioni in Medicina, 2020, 28, 258-262.	1.1	3
94	The landscape of epilepsy-related GATOR1 variants. Genetics in Medicine, 2019, 21, 398-408.	2.4	137
95	Newly Diagnosed and Growing Subependymal Giant Cell Astrocytoma in Adults With Tuberous Sclerosis Complex: Results From the International TOSCA Study. Frontiers in Neurology, 2019, 10, 821.	2.4	18
96	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
97	Clinical Characteristics of Subependymal Giant Cell Astrocytoma in Tuberous Sclerosis Complex. Frontiers in Neurology, 2019, 10, 705.	2.4	22
98	Polygenic burden in focal and generalized epilepsies. Brain, 2019, 142, 3473-3481.	7.6	90
99	Treatment Patterns and Use of Resources in Patients With Tuberous Sclerosis Complex: Insights From the TOSCA Registry. Frontiers in Neurology, 2019, 10, 1144.	2.4	11
100	Declining malformation rates with changed antiepileptic drug prescribing. Neurology, 2019, 93, e831-e840.	1.1	69
101	Value of Multimodal Imaging Approach to Diagnosis of Neurosarcoidosis. Brain Sciences, 2019, 9, 243.	2.3	13
102	An Italian multicentre study of perampanel in progressive myoclonus epilepsies. Epilepsy Research, 2019, 156, 106191.	1.6	19
103	Imaging Genetics for Benign Mesial Temporal Lobe Epilepsy. , 2019, , 48-54.		2
104	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
105	REM-Sleep Behavior Disorder in Patients With Essential Tremor: What Is Its Clinical Significance?. Frontiers in Neurology, 2019, 10, 315.	2.4	12
106	Hypertension, seizures, and epilepsy: a review on pathophysiology and management. Neurological Sciences, 2019, 40, 1775-1783.	1.9	51
107	Intestinal-Cell Kinase and Juvenile Myoclonic Epilepsy. New England Journal of Medicine, 2019, 380, e24.	27.0	4
108	Abdominal Fat SIRT6 Expression and Its Relationship with Inflammatory and Metabolic Pathways in Pre-Diabetic Overweight Patients. International Journal of Molecular Sciences, 2019, 20, 1153.	4.1	27

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109	Validated outcome of treatment changes according to International League Against Epilepsy criteria in adults with drugâ€resistant focal epilepsy. Epilepsia, 2019, 60, 1114-1123.	5.1	23
110	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
111	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
112	Psychiatric Assessment in Patients with Mild Temporal Lobe Epilepsy. Behavioural Neurology, 2019, 2019, 1-9.	2.1	8
113	0590 Usefulness Of Cardiac Parasympathetic Index In Cpap-treated Patients With Obstructive Sleep Apnea: A Preliminary Study. Sleep, 2019, 42, A235-A236.	1.1	0
114	Insight into epileptic and physiological dÃ©jÃ vu : from a multicentric cohort study. European Journal of Neurology, 2019, 26, 407-414.	3.3	2
115	A genomeâ€wide association study of sodium levels and drug metabolism in an epilepsy cohort treated with carbamazepine and oxcarbazepine. Epilepsia Open, 2019, 4, 102-109.	2.4	9
116	Kufs disease due to mutation ofCLN6: clinical, pathological and molecular genetic features. Brain, 2019, 142, 59-69.	7.6	28
117	Synovial sarcoma diagnosis on fine needle cytology sample confirmed by fluorescence inÂsitu hybridisation. Cytopathology, 2019, 30, 314-317.	0.7	3
118	Epilepsy in tuberous sclerosis complex: Findings from the <scp>TOSCA</scp> Study. Epilepsia Open, 2019, 4, 73-84.	2.4	125
119	Management of psychogenic nonâ€epileptic seizures: a multidisciplinary approach. European Journal of Neurology, 2019, 26, 205.	3.3	64
120	Adjuvant anastrozole versus exemestane versus letrozole, upfront or after 2 years of tamoxifen, in endocrine-sensitive breast cancer (FATA-GIM3): a randomised, phase 3 trial. Lancet Oncology, The, 2018, 19, 474-485.	10.7	59
121	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
122	Structural brain abnormalities in the common epilepsies assessed in a worldwide ENIGMA study. Brain, 2018, 141, 391-408.	7.6	352
123	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
124	Brandâ€toâ€generic levetiracetam switching: a 4â€year prospective observational realâ€life study. European Journal of Neurology, 2018, 25, 666-671.	3.3	17
125	PRRT2 controls neuronal excitability by negatively modulating Na+ channel 1.2/1.6 activity. Brain, 2018, 141, 1000-1016.	7.6	99
126	Lacosamide monotherapy in clinical practice: A retrospective chart review. Acta Neurologica Scandinavica, 2018, 138, 186-194.	2.1	24

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127	Psychopathological constellation in patients with PNES: A new hypothesis. Epilepsy and Behavior, 2018, 78, 297-301.	1.7	21
128	On the Classification of EEG Signal by Using an SVM Based Algorithm. Smart Innovation, Systems and Technologies, 2018, , 271-278.	0.6	15
129	Sexual function and sex hormones in breast cancer patients. Endocrine, 2018, 60, 510-515.	2.3	13
130	The application of artificial intelligence to understand the pathophysiological basis of psychogenic nonepileptic seizures. Epilepsy and Behavior, 2018, 87, 167-172.	1.7	29
131	Potential Role of miRNAs as Theranostic Biomarkers of Epilepsy. Molecular Therapy - Nucleic Acids, 2018, 13, 275-290.	5.1	37
132	Inflammatory Cytokines and SIRT1 Levels in Subcutaneous Abdominal Fat: Relationship With Cardiac Performance in Overweight Pre-diabetics Patients. Frontiers in Physiology, 2018, 9, 1030.	2.8	41
133	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
134	Neuropsychological profile of mild temporal lobe epilepsy. Epilepsy and Behavior, 2018, 85, 222-226.	1.7	10
135	Comparison between Electrocardiographic and Earlobe Pulse Photoplethysmographic Detection for Evaluating Heart Rate Variability in Healthy Subjects in Short- and Long-Term Recordings. Sensors, 2018, 18, 844.	3.8	55
136	Rare coding variants in genes encoding GABAA receptors in genetic generalised epilepsies: an exome-based case-control study. Lancet Neurology, The, 2018, 17, 699-708.	10.2	67
137	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
138	Relationship between severity of migraine and vitamin D deficiency: a case-control study. Neurological Sciences, 2018, 39, 167-168.	1.9	19
139	Cardiac parasympathetic index identifies subjects with adult obstructive sleep apnea: A simultaneous polysomnographic-heart rate variability study. PLoS ONE, 2018, 13, e0193879.	2.5	11
140	Role of Pharmacogenomics in Antiepileptic Drug Therapy: Current Status and Future Perspectives. Current Pharmaceutical Design, 2018, 23, 5760-5765.	1.9	12
141	Diagnostic Biomarkers of Epilepsy. Current Pharmaceutical Biotechnology, 2018, 19, 440-450.	1.6	20
142	Risk factors for unprovoked epileptic seizures in multiple sclerosis: a systematic review and meta-analysis. Neurological Sciences, 2017, 38, 399-406.	1.9	35
143	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
144	Hyperkinetic psychogenic movement disorders remain a diagnosis at first sight. Neurology, 2017, 88, 114-115.	1.1	1

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145	Carbamazepineâ€•and oxcarbazepineâ€•induced hyponatremia in people with epilepsy. Epilepsia, 2017, 58, 1227-1233.	5.1	54
146	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
147	Alterations in the Î±₂Î´ ligand, thrombospondinâ€1, in a rat model of spontaneous absence epilepsy and in patients with idiopathic/genetic generalized epilepsies. Epilepsia, 2017, 58, 1993-2001.	5.1	8
148	Variable course of Unverricht-Lundborg disease. Neurology, 2017, 89, 1691-1697.	1.1	18
149	Advanced morphological neuroimaging study in lateral temporal lobe epilepsy: A multicentric study. Epilepsy and Behavior, 2017, 74, 69-72.	1.7	0
150	Ictal 18F-FDG PET/MRI in a Patient With Cortical Heterotopia and Focal Epilepsy. Clinical Nuclear Medicine, 2017, 42, 768-769.	1.3	8
151	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
152	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
153	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
154	The Natural History of Epilepsy in 163 Untreated Patients: Looking for â€œOligoepilepsyâ€• PLoS ONE, 2016, 11, e0161722.	2.5	2
155	Integrity of the corpus callosum in patients with benign temporal lobe epilepsy. Epilepsia, 2016, 57, 590-596.	5.1	17
156	Cardiac sympathetic index identifies patients with Parkinson's disease and REM behavior disorder. Parkinsonism and Related Disorders, 2016, 26, 62-66.	2.2	20
157	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
158	Definition and diagnostic criteria of sleep-related hypermotor epilepsy. Neurology, 2016, 86, 1834-1842.	1.1	245
159	Long-term outcome of mild mesial temporal lobe epilepsy. Neurology, 2016, 86, 1904-1910.	1.1	40
160	Lacosamide in patients with temporal lobe epilepsy: An observational multicentric open-label study. Epilepsy and Behavior, 2016, 58, 111-114.	1.7	12
161	Relevance of clinical context in the diagnosticâ€therapeutic approach to status epilepticus. Epilepsia, 2016, 57, 1527-1529.	5.1	4
162	Pharmacological modulation in mesial temporal lobe epilepsy: Current status and future perspectives. Pharmacological Research, 2016, 113, 421-425.	7.1	20

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163	Epilepsy in cerebrovascular diseases: Review of experimental and clinical data with metaâ€analysis of risk factors. Epilepsia, 2016, 57, 1205-1214.	5.1	122
164	Genetics of reflex seizures and epilepsies in humans and animals. Epilepsy Research, 2016, 121, 47-54.	1.6	60
165	Multidisciplinary approach to breast angiosarcoma in an elderly patient: Repeated local relapses and significant objective responses. International Journal of Immunopathology and Pharmacology, 2016, 29, 537-542.	2.1	10
166	Tremor pattern differentiates drug-induced resting tremor from Parkinson disease. Parkinsonism and Related Disorders, 2016, 25, 100-103.	2.2	17
167	PRIMA1 mutation: a new cause of nocturnal frontal lobe epilepsy. Annals of Clinical and Translational Neurology, 2015, 2, 821-830.	3.7	21
168	<scp>DEPDC</scp>5 mutations are not a frequent cause of familial temporal lobe epilepsy. Epilepsia, 2015, 56, e168-71.	5.1	37
169	Why should we change the term psychogenic nonepileptic seizures?. Epilepsia, 2015, 56, 1178-1179.	5.1	5
170	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
171	Clinical profile of patients with ATP1A3 mutations in Alternating Hemiplegia of Childhoodâ€"a study of 155 patients. Orphanet Journal of Rare Diseases, 2015, 10, 123.	2.7	117
172	White matter abnormalities differentiate severe from benign temporal lobe epilepsy. Epilepsia, 2015, 56, 1109-1116.	5.1	38
173	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
174	Profile of brivaracetam and its potential in the treatment of epilepsy. Neuropsychiatric Disease and Treatment, 2015, 11, 2967.	2.2	14
175	Professional Acquisition ofM. bovisin Calabria Region (Southern Italy): A Challenging Case of Osteomyelitis in a Migrant Patient from Bulgaria. Case Reports in Infectious Diseases, 2015, 2015, 1-4.	0.5	0
176	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
177	Angiosarcoma of the breast: a new therapeutic approach?. International Journal of Surgery Case Reports, 2015, 13, 30-32.	0.6	8
178	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
179	C19orf12 gene mutations in patients with neurodegeneration with brain iron accumulation. Parkinsonism and Related Disorders, 2015, 21, 813-816.	2.2	21
180	PCDH19 mutations in female patients from Southern Italy. Seizure: the Journal of the British Epilepsy Association, 2015, 24, 118-120.	2.0	7

#	Article	IF	Citations
181	Epilepsy and Immune System: A Tour Around the Current Literature. , 2015, , 163-182.		0
182	A new SLC20A2 mutation identified in southern Italy family with primary familial brain calcification. Gene, 2015, 568, 109-111.	2.2	16
183	No evidence of a role for cystatin <scp>B</scp> gene in juvenile myoclonic epilepsy. Epilepsia, 2015, 56, e40-3.	5.1	7
184	A puzzling case without solution: isolated late-onset epileptic seizure. Neurological Sciences, 2015, 36, 2303-2304.	1.9	1
185	A hyaluronic acid-based compound inhibits fibroblast senescence induced by oxidative stress in vitro and prevents oral mucositis in vivo. Journal of Cellular Physiology, 2015, 230, 1421-1429.	4.1	32
186	A recurrent de novo mutation in KCNC1 causes progressive myoclonus epilepsy. Nature Genetics, 2015, 47, 39-46.	21.4	245
187	Epilepsy associated with Leukoaraiosis mainly affects temporal lobe: a casual or causal relationship?. Epilepsy Research, 2015, 109, 1-8.	1.6	39
188	Neuro-anatomical differences among epileptic and non-epileptic dÃ©jÃ-vu. Cortex, 2015, 64, 1-7.	2.4	14
189	Positivity to p-ANCA in patients with status epilepticus. BMC Neurology, 2014, 14, 148.	1.8	5
190	Mild <scp>L</scp>afora disease: Clinical, neurophysiologic, and genetic findings. Epilepsia, 2014, 55, e129-33.	5.1	43
191	DEPDC5 mutations in families presenting as autosomal dominant nocturnal frontal lobe epilepsy. Neurology, 2014, 82, 2101-2106.	1.1	126
192	Generalized versus partial reflex seizures: A review. Seizure: the Journal of the British Epilepsy Association, 2014, 23, 512-520.	2.0	70
193	Rosacea-like facial rash related to metformin administration in a young woman. BMC Pharmacology & Toxicology, 2014, 15, 3.	2.4	12
194	Mutations in the Matrin 3 gene cause familial amyotrophic lateral sclerosis. Nature Neuroscience, 2014, 17, 664-666.	14.8	398
195	Genetic counselling in ALS: facts, uncertainties and clinical suggestions. Journal of Neurology, Neurosurgery and Psychiatry, 2014, 85, 478-485.	1.9	99
196	Reversible symmetrical external capsule hyperintensity as an early finding of autoimmune encephalitis. Neurological Sciences, 2014, 35, 1147-1149.	1.9	6
197	Thalamotemporal impairment in benign temporal lobe epilepsy: Same hypotheses?. Epilepsia, 2014, 55, 944-944.	5.1	0
198	The role of calcium channel mutations in human epilepsy. Progress in Brain Research, 2014, 213, 87-96.	1.4	18

#	Article	IF	Citations
199	Reduced thalamic volume in Parkinson disease with REM sleep behavior disorder: Volumetric study. Parkinsonism and Related Disorders, 2014, 20, 1004-1008.	2.2	54
200	Permutation entropy of scalp EEG: A tool to investigate epilepsies. Clinical Neurophysiology, 2014, 125, 13-20.	1.5	59
201	Autosomal dominant lateral temporal epilepsy (ADLTE): Absence of chromosomal rearrangements in LGI1 gene. Epilepsy Research, 2014, 108, 597-599.	1.6	2
202	Progressive myoclonic epilepsies. Neurology, 2014, 82, 405-411.	1.1	87
203	Chronic neuroborreliosis by B. garinii: an unusual case presenting with epilepsy and multifocal brain MRI lesions. New Microbiologica, 2014, 37, 393-7.	0.1	5
204	Sacsin-Related Spastic Ataxia Caused by a Novel Missense Mutation p.Arg272His in a Patient from Sicily, Southern Italy. Cerebellum, 2013, 12, 589-592.	2.5	3
205	Potential involvement of GRIN2B encoding the NMDA receptor subunit NR2B in the spectrum of Alzheimerâ€™s disease. Journal of Neural Transmission, 2013, 121, 533-42.	2.8	35
206	Amyotrophic lateral sclerosis: a new missense mutation in the SOD1 gene. Neurobiology of Aging, 2013, 34, 1709.e3-1709.e5.	3.1	19
207	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
208	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
209	Failure to confirm association of a polymorphism in KCNMB4 gene with mesial temporal lobe epilepsy. Epilepsy Research, 2013, 106, 284-287.	1.6	4
210	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
211	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
212	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
213	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
214	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
215	Consensus on diagnosis and management of JME: From founder's observations to current trends. Epilepsy and Behavior, 2013, 28, S87-S90.	1.7	142
216	Low penetrance of autosomal dominant lateral temporal epilepsy in Italian families without <scp>LGI</scp>1 mutations. Epilepsia, 2013, 54, 1288-1297.	5.1	32

#	Article	IF	Citations
217	Septo-optic dysplasia plus bilateral perisylvian polymicrogyria: a case report. Neurological Sciences, 2013, 34, 1479-1480.	1.9	3
218	Diffusivity of Cerebellar Hemispheres Enables Discrimination of Cerebellar or Parkinsonian Multiple System Atrophy from Progressive Supranuclear Palsyâ€“Richardson Syndrome and Parkinson Disease. Radiology, 2013, 267, 843-850.	7.3	31
219	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
220	Advanced MRI Morphologic Study Shows No Atrophy in Healthy Individuals with Hippocampal Hyperintensity. American Journal of Neuroradiology, 2013, 34, 1585-1588.	2.4	5
221	Exonâ€disrupting deletions of <scp>NRXN1</scp> in idiopathic generalized epilepsy. Epilepsia, 2013, 54, 256-264.	5.1	59
222	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
223	Antiâ€“Nâ€methylâ€Dâ€aspartateâ€glutamicâ€receptor encephalitis presenting as paroxysmal exerciseâ€induced foot weakness. Movement Disorders, 2013, 28, 820-822.	3.9	11
224	Contribution of Cerebrospinal Fluid Thymosin Î²4 Levels to the Clinical Differentiation of Creutzfeldt-Jakob Disease. Archives of Neurology, 2012, 69, 868-72.	4.5	11
225	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
226	Ataxin-1 and ataxin-2 intermediate-length PolyQ expansions in amyotrophic lateral sclerosis. Neurology, 2012, 79, 2315-2320.	1.1	70
227	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
228	FUS mutations in sporadic amyotrophic lateral sclerosis: Clinical and genetic analysis. Neurobiology of Aging, 2012, 33, 837.e1-837.e5.	3.1	32
229	C9ORF72 hexanucleotide repeat expansions in the Italian sporadic ALS population. Neurobiology of Aging, 2012, 33, 1848.e15-1848.e20.	3.1	76
230	CAV3 T78M mutation as polymorphic variant in South Italy. Neuromuscular Disorders, 2012, 22, 669-670.	0.6	8
231	Mutation analysis of the PINK1 gene in Southern Italian patients with early- and late-onset parkinsonism. Parkinsonism and Related Disorders, 2012, 18, 651-653.	2.2	7
232	Clinical characteristics of patients with familial amyotrophic lateral sclerosis carrying the pathogenic GGGGCC hexanucleotide repeat expansion of C9ORF72. Brain, 2012, 135, 784-793.	7.6	182
233	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
234	Topiramate and temporal lobe epilepsy: an open-label study. Epileptic Disorders, 2012, 14, 163-166.	1.3	5

#	Article	IF	Citations
235	The p.Arg416Cys mutation in SPG3a gene associated with a pure form of spastic paraplegia. Muscle and Nerve, 2012, 45, 919-920.	2.2	5
236	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
237	Neuroanatomic correlates of psychogenic nonepileptic seizures: A cortical thickness and VBM study. Epilepsia, 2012, 53, 377-385.	5.1	140
238	Genomeâ€wide linkage metaâ€analysis identifies susceptibility loci at 2q34 and 13q31.3 for genetic generalized epilepsies. Epilepsia, 2012, 53, 308-318.	5.1	32
239	Frequency of the C9orf72 hexanucleotide repeat expansion in patients with amyotrophic lateral sclerosis and frontotemporal dementia: a cross-sectional study. Lancet Neurology, The, 2012, 11, 323-330.	10.2	1,039
240	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
241	Unilateral basal ganglia atrophy in a patient with tuberous sclerosis complex and hemichorea. Movement Disorders, 2012, 27, 458-460.	3.9	5
242	Dentatorubral-pallidoluysian atrophy: Haplotype of Asian origin in 2 Italian families. Movement Disorders, 2012, 27, 460-461.	3.9	4
243	Generalised epilepsy with febrile seizures plus (GEFS+): molecular analysis in a restricted area. Child's Nervous System, 2012, 28, 141-145.	1.1	7
244	Benign mesial temporal lobe epilepsy. Nature Reviews Neurology, 2011, 7, 237-240.	10.1	76
245	NOS2A as a candidate gene in Relapsingâ€“Remitting Multiple Sclerosis: A haplotype study using selected subsets of single nucleotide polymorphisms. Journal of the Neurological Sciences, 2011, 304, 75-77.	0.6	2
246	Rating scale for psychogenic nonepileptic seizures: Scale development and clinimetric testing. Epilepsy and Behavior, 2011, 21, 128-131.	1.7	21
247	A new Italian instrument for the assessment of irritability in patients with epilepsy. Epilepsy and Behavior, 2011, 21, 275-281.	1.7	15
248	Presenilin enhancerâ€2 gene: Identification of a novel promoter mutation in a patient with earlyâ€onset familial Alzheimer's disease. Alzheimer's and Dementia, 2011, 7, 574-578.	0.8	16
249	Blocking out the real diagnosis. Lancet, The, 2011, 377, 690.	13.7	5
250	Blocking out the real diagnosis â€“ Authors' reply. Lancet, The, 2011, 378, 316.	13.7	0
251	Neocortical thinning in â€œbenignâ€•mesial temporal lobe epilepsy. Epilepsia, 2011, 52, 712-717.	5.1	51
252	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

#	Article	IF	Citations
253	Clinical and neurophysiologic features of progressive myoclonus epilepsy without renal failure caused bySCARB2mutations. Epilepsia, 2011, 52, 2356-2363.	5.1	63
254	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
255	Dose-dependent risk of malformations with antiepileptic drugs: an analysis of data from the EURAP epilepsy and pregnancy registry. Lancet Neurology, The, 2011, 10, 609-617.	10.2	654
256	A novel NF1 gene mutation in an Italian family with neurofibromatosis type 1. Child's Nervous System, 2011, 27, 635-638.	1.1	4
257	Presenilin-2 gene mutation presenting as Lewy Body dementia?. Neurological Sciences, 2011, 32, 533-534.	1.9	9
258	Association study between four polymorphisms in the HFE, TF and TFR genes and Parkinsonâ€™s disease in Southern Italy. Neurological Sciences, 2011, 32, 525-527.	1.9	13
259	A System for the Analysis of Snore Signals. Procedia Computer Science, 2011, 4, 1101-1108.	2.0	5
260	Sporadic motor neuron disease in a familial novel SOD1 mutation: Incomplete penetrance or chance association?. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2011, 12, 220-222.	2.1	7
261	Mutations of the orexin system, a regulator of sleep arousal, are not a common cause of ADNFLE. Neurology, 2011, 76, 1272-1273.	1.1	1
262	ADAM23, a Gene Related to LGI1, Is Not Linked to Autosomal Dominant Lateral Temporal Epilepsy. Epilepsy Research & Treatment, 2011, 2011, 1-6.	1.4	2
263	TARDBP gene mutations in south Italian patients with amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2011, 82, 587-588.	1.9	25
264	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
265	Leberâ€™s hereditary optic neuropathy associated with a multiple-sclerosis-like picture in a man. Multiple Sclerosis Journal, 2011, 17, 763-766.	3.0	15
266	Ile587Val Polymorphism of the eIF2B5 Gene as Susceptibility Factor for Multiple Sclerosis. Neuroscience and Medicine, 2011, 02, 117-119.	0.2	2
267	Single nucleotide polymorphism in the MMP-9 gene is associated with susceptibility to develop multiple sclerosis in an Italian case-control study. Journal of Neuroimmunology, 2010, 225, 175-179.	2.3	25
268	Early involvement of sympathetic cardiac nerve endings in a patient with rem sleep behaviour disorders. BMC Geriatrics, 2010, 10, .	2.7	0
269	Voxel-based morphometry of adulthood patients with temporal lobe epilepsy. BMC Geriatrics, 2010, 10, .	2.7	0
270	Voxelâ€based morphometry of sporadic epileptic patients with mesiotemporal sclerosis. Epilepsia, 2010, 51, 506-510.	5.1	43

#	Article	IF	Citations
271	Benign temporal lobe epilepsy. Epilepsia, 2010, 51, 45-46.	5.1	4
272	Characteristics of a large population of patients with refractory epilepsy attending tertiary referral centers in Italy. Epilepsia, 2010, 51, 921-925.	5.1	35
273	Relationship between adverse effects of antiepileptic drugs, number of coprescribed drugs, and drug load in a large cohort of consecutive patients with drugâ€refractory epilepsy. Epilepsia, 2010, 51, 797-804.	5.1	160
274	Alexithymia and Cancer-Related Fatigue: A Controlled Cross-Sectional Study. Tumori, 2010, 96, 131-137.	1.1	10
275	Temporal lobe abnormalities on brain MRI in healthy volunteers. Neurology, 2010, 74, 553-557.	1.1	47
276	TEMPORAL LOBE ABNORMALITIES ON BRAIN MRI IN HEALTHY VOLUNTEERS: A PROSPECTIVE CASE-CONTROL STUDY. Neurology, 2010, 75, 377-378.	1.1	15
277	An Embedded System for EEG Acquisition and Processing for Brain Computer Interface Applications. Lecture Notes in Electrical Engineering, 2010, , 137-154.	0.4	9
278	Proteomic profiling of cerebrospinal fluid in Creutzfeldtâ€"Jakob disease. Expert Review of Proteomics, 2010, 7, 907-917.	3.0	14
279	Mutation analysis of the SPG4 gene in Italian patients with pure and complicated forms of spastic paraplegia. Journal of the Neurological Sciences, 2010, 288, 96-100.	0.6	27
280	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
281	Design of an electronic device for brain computer interface applications. , 2009, , .		3
282	CONVENTIONAL MRI AND NOTCH3 GENE SCREENING IN SPORADIC CADASIL. Neurology, 2009, 72, 469-471.	1.1	3
283	SCARB2 mutations in progressive myoclonus epilepsy (PME) without renal failure. Annals of Neurology, 2009, 66, 532-536.	5.3	90
284	Prognostic factors in patients with mesial temporal lobe epilepsy. Epilepsia, 2009, 50, 41-44.	5.1	51
285	Eyelid fluttering, typical EEG pattern, and impaired intellectual function: A homogeneous epileptic condition among the patients presenting with eyelid myoclonia. Epilepsia, 2009, 50, 1536-1541.	5.1	48
286	Clinical spectrum of SCN1A mutations. Epilepsia, 2009, 50, 20-23.	5.1	87
287	Familial mesial temporal lobe epilepsies: Clinical and genetic features. Epilepsia, 2009, 50, 55-57.	5.1	30
288	A novel ICA-based hardware system for reconfigurable and portable BCI. , 2009, , .		12

#	Article	IF	Citations
289	Anti-NMDA receptor encephalitis: aÂvideo case report. Epileptic Disorders, 2009, 11, 267-269.	1.3	11
290	Familial mesial temporal lobe epilepsy (FMTLE). Journal of Neurology, 2008, 255, 16-23.	3.6	60
291	Comparison of the safety and efficacy of paclitaxel plus gemcitabine combination in young and elderly patients with locally advanced or metastatic non-small cell lung cancer. Critical Reviews in Oncology/Hematology, 2008, 65, 164-171.	4.4	9
292	Ictal impending danger â€“ â€œsixth sense seizuresâ€•â€“ in patients with benign focal epileptic seizures of adolescence. Epilepsy Research, 2008, 79, 90-96.	1.6	4
293	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
294	Effects of levetiracetam on EEG abnormalities in juvenile myoclonic epilepsy. Epilepsia, 2008, 49, 663-669.	5.1	54
295	Metabolic aspects of the extreme longevity. Experimental Gerontology, 2008, 43, 74-78.	2.8	34
296	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
297	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
298	Health-related quality of life in epilepsy: Findings obtained with a new Italian instrument. Epilepsy and Behavior, 2008, 13, 119-126.	1.7	29
299	Limited chronic focal encephalitis. Neurology, 2008, 70, 374-377.	1.1	37
300	Hippocampal and thalamic atrophy in mild temporal lobe epilepsy. Neurology, 2008, 71, 1094-1101.	1.1	91
301	Monotherapy for partial epilepsy: focus on levetiracetam. Neuropsychiatric Disease and Treatment, 2008, 4, 33.	2.2	40
302	Efficacy of the combination of cisplatin with either gemcitabine and vinorelbine or gemcitabine and paclitaxel in the treatment of locally advanced or metastatic non-small-cell lung cancer: a phase III randomised trial of the Southern Italy Cooperative Oncology Group (SICOG 0101). Annals of Oncology, 2007, 18, 324-330.	1.2	45
303	Quality of life outcomes of immediate or delayed treatment of early epilepsy and single seizures. Neurology, 2007, 68, 1188-1196.	1.1	105
304	Modulatory Proteins Can Rescue a Trafficking Defective Epileptogenic Na_v1.1 Na⁺Channel Mutant. Journal of Neuroscience, 2007, 27, 11037-11046.	3.6	169
305	Screening of Depressive Symptoms in Youngâ€“Old Hemodialysis Patients: Relationship between Beck Depression Inventory and 15-Item Geriatric Depression Scale. Nephron Clinical Practice, 2007, 106, c187-c192.	2.3	18
306	Psychiatric events in epilepsy. Seizure: the Journal of the British Epilepsy Association, 2007, 16, 586-592.	2.0	9

#	Article	IF	Citations
307	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
308	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
309	Serotonin transporter gene (5-Htt): Association analysis with temporal lobe epilepsy. Neuroscience Letters, 2007, 421, 52-56.	2.1	32
310	Intra-patient alternated dose escalation of paclitaxel and gemcitabine versus paclitaxel followed by fixed dose rate infusion of gemcitabine in fit elderly non-small cell lung cancer patients. Lung Cancer, 2007, 56, 263-271.	2.0	5
311	Further evidence of genetic heterogeneity in families with autosomal dominant nocturnal frontal lobe epilepsy. Epilepsy Research, 2007, 74, 70-73.	1.6	29
312	Usefulness of a morning routine EEG recording in patients with juvenile myoclonic epilepsy. Epilepsy Research, 2007, 77, 17-21.	1.6	48
313	Photosensitive Benign Myoclonic Epilepsy in Infancy. Epilepsia, 2007, 48, 96-100.	5.1	29
314	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
315	Mutational Analysis of EFHC1 Gene in Italian Families with Juvenile Myoclonic Epilepsy. Epilepsia, 2007, 48, 1686-1690.	5.1	44
316	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
317	Novel spastin (SPG4) mutations in Italian patients with hereditary spastic paraplegia. Neuromuscular Disorders, 2006, 16, 387-390.	0.6	15
318	Suppressive Efficacy by a Commercially Available Blue Lens on PPR in 610 Photosensitive Epilepsy Patients. Epilepsia, 2006, 47, 529-533.	5.1	96
319	Clinical and Genetic Findings in 26 Italian Patients with Lafora Disease. Epilepsia, 2006, 47, 640-643.	5.1	71
320	Linkage Analysis and Disease Models in Benign Familial Infantile Seizures: A Study of 16 Families. Epilepsia, 2006, 47, 1029-1034.	5.1	23
321	Open label, long-term, pragmatic study on levetiracetam in the treatment of juvenile myoclonic epilepsy. Epilepsy Research, 2006, 71, 32-39.	1.6	72
322	Reflex periodic spasms induced by eating. Brain and Development, 2006, 28, 170-174.	1.1	28
323	Genetic heterogeneity in patients with pantothenate kinaseâ€“associated neurodegeneration and classic magnetic resonance imaging eye-of-the-tiger pattern. Movement Disorders, 2006, 21, 252-254.	3.9	26
324	Alteration of the in vivo nicotinic receptor density in ADNFLE patients: a PET study. Brain, 2006, 129, 2047-2060.	7.6	165

#	Article	IF	Citations
325	MRI evidence of mesial temporal sclerosis in sporadic "benign" temporal lobe epilepsy. Neurology, 2006, 66, 562-565.	1.1	91
326	ApoE Epsilon4 Allele and Disease Duration Affect Verbal Learning in Mild Temporal Lobe Epilepsy. Epilepsia, 2005, 46, 110-117.	5.1	53
327	Capecitabine plus oxaliplatin for the first-line treatment of elderly patients with metastatic colorectal carcinoma. Cancer, 2005, 104, 282-289.	4.1	91
328	CNS findings in three cases of septo-optic dysplasia, including one with semilobar holoprosencephaly. American Journal of Medical Genetics, Part A, 2005, 136A, 357-357.	1.2	5
329	A tailored regimen including capecitabine and oxaliplatin for treating elderly patients with metastatic colorectal carcinoma. Critical Reviews in Oncology/Hematology, 2005, 53, 133-139.	4.4	10
330	Sex Differences in Clinical and Genetic Determinants of Levodopa Peak-Dose Dyskinesias in Parkinson Disease. Archives of Neurology, 2005, 62, 601.	4.5	195
331	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
332	Immediate versus deferred antiepileptic drug treatment for early epilepsy and single seizures: a randomised controlled trial. Lancet, The, 2005, 365, 2007-2013.	13.7	369
333	Movement time and aging: a normative study in healthy subjects with the â€œMovement Time AnalyzerÂ®â€. Aging Clinical and Experimental Research, 2005, 17, 207-210.	2.9	13
334	Brachial amyotrophic diplegia associated with a novel SOD1 mutation (L106P). Neurology, 2005, 64, 1477-1478.	1.1	16
335	Capecitabine (XEL) plus oxaliplatin (OX) tailored regimen for the first-line treatment of elderly patients (pts) with metastatic colorectal cancer (MCC). Final results of the Southern Italy Cooperative Oncology Group (SICOG) trial 0108. Journal of Clinical Oncology, 2005, 23, 3582-3582.	1.6	1
336	Increased Risk for Alzheimer Disease With the Interaction of MPO and A2M Polymorphisms. Archives of Neurology, 2004, 61, 341.	4.5	45
337	Mutations and polymorphisms of the CLCN2 gene in idiopathic epilepsy. Neurology, 2004, 63, 1500-1502.	1.1	63
338	Suggestive evidence for linkage to chromosome 13qter for autosomal dominant type 1 porencephaly. Neurology, 2004, 62, 1613-1615.	1.1	29
339	The fragile X premutation presenting as postprandial hypotension. Neurology, 2004, 63, 2188-2189.	1.1	21
340	Direct Cost of Medical Management of Epilepsy among Adults in Italy: A Prospective Cost-of-Illness Study (EPICOS). Epilepsia, 2004, 45, 171-178.	5.1	64
341	A Novel Exon 1 Mutation in a Patient with Atypical Lafora Progressive Myoclonus Epilepsy Seen as Childhoodâ€onset Cognitive Deficit. Epilepsia, 2004, 45, 294-295.	5.1	27
342	Morbidity in Patients with Epilepsy: Type and Complications: A European Cohort Study. Epilepsia, 2004, 45, 71-76.	5.1	27

#	Article	IF	Citations
343	Accidents in Patients with Epilepsy: Types, Circumstances, and Complications: A European Cohort Study. Epilepsia, 2004, 45, 667-672.	5.1	89
344	EURAP: An International Registry of Antiepileptic Drugs and Pregnancy. Epilepsia, 2004, 45, 1463-1464.	5.1	70
345	Startle epilepsy complicating aspartylglucosaminuria. Brain and Development, 2004, 26, 130-133.	1.1	12
346	Gene conversion events in adult-onset spinal muscular atrophy. Acta Neurologica Scandinavica, 2004, 109, 151-154.	2.1	12
347	Benign familial neonatal-infantile seizures: Characterization of a new sodium channelopathy. Annals of Neurology, 2004, 55, 550-557.	5.3	250
348	Prodynorphin Gene Promoter Polymorphism and Temporal Lobe Epilepsy. Epilepsia, 2003, 44, 1255-1256.	5.1	27
349	Two Novel SCN1A Missense Mutations in Generalized Epilepsy with Febrile Seizures Plus. Epilepsia, 2003, 44, 1257-1258.	5.1	56
350	Irinotecan plus leucovorin-modulated 5-fluorouracil I.V. bolus every other week may be a suitable therapeutic option also for elderly patients with metastatic colorectal carcinoma. British Journal of Cancer, 2003, 89, 992-996.	6.4	19
351	GABA(B) receptor 1 polymorphism (G1465A) is associated with temporal lobe epilepsy. Neurology, 2003, 60, 560-563.	1.1	127
352	Temporal Lobe Epilepsy as a Unique Manifestation of Multiple Sclerosis. Canadian Journal of Neurological Sciences, 2003, 30, 228-232.	0.5	36
353	Genetic association of Î± ₂ -macroglobulin polymorphisms with AD in southern Italy. Neurology, 2002, 59, 756-758.	1.1	31
354	Body Weight Influences Pharmacokinetics of Levodopa in Parkinson's Disease. Clinical Neuropharmacology, 2002, 25, 79-82.	0.7	71
355	A large family with pure autosomal dominant hereditary spastic paraplegia from southern Italy mapping to chromosome 14q11.2-q24.3. Journal of Neurology, 2002, 249, 1413-1416.	3.6	4
356	Anti-GM1 ganglioside antibodies in Parkinson's disease. Acta Neurologica Scandinavica, 2002, 106, 54-57.	2.1	57
357	Silent Celiac Disease in Patients with Childhood Localization-Related Epilepsies. Epilepsia, 2002, 42, 1153-1155.	5.1	54
358	Morbidity and Accidents in Patients with Epilepsy: Results of a European Cohort Study. Epilepsia, 2002, 43, 1076-1083.	5.1	159
359	Benign Familial Infantile Convulsions: Mapping of a Novel Locus on Chromosome 2q24 and Evidence for Genetic Heterogeneity. American Journal of Human Genetics, 2001, 68, 1521-1526.	6.2	66
360	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

#	Article	IF	Citations
361	Benign Partial Epilepsies of Adolescence: A Report of 37 New Cases. Epilepsia, 2001, 42, 1549-1552.	5.1	10
362	Clinical and genetic study of a large Charcotâ€“Marieâ€“Tooth type 2A family from southern Italy. Neurology, 2001, 56, 100-103.	1.1	28
363	The parkin gene is not involved in late-onset Parkinsonâ€™s disease. Neurology, 2001, 57, 359-362.	1.1	48
364	Cerebral venous thrombosis and isolated intracranial hypertension without papilledema in CDH. Neurology, 2001, 57, 31-36.	1.1	114
365	Juvenile Huntingtonâ€™s disease presenting as progressive myoclonic epilepsy. Neurology, 2001, 57, 708-711.	1.1	62
366	Role of Free Fatty Acids on Cardiac Autonomic Nervous System in Noninsulin-Dependent Diabetic Patients: Effects of Metabolic Control. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 2769-2774.	3.6	24
367	A clinical spectrum of the myoclonic manifestations associated with typical absences in childhood absence epilepsy. A video-polygraphic study. Epileptic Disorders, 2001, 3, 57-62.	1.3	21
368	A new locus for autosomal dominant nocturnal frontal lobe epilepsy maps to chromosome 1. Neurology, 2000, 55, 1467-1471.	1.1	82
369	Effects of insulin on the cardiac autonomic nervous system in insulin-resistant states. Clinical Science, 2000, 98, 129.	4.3	19
370	Identification of a new polymorphism (c134G>A) in the exon 2 of the myelin protein zero gene. Human Mutation, 2000, 15, 299-299.	2.5	1
371	Vitamin E deficiency due to chylomicron retention disease in Marinesco-Sjï¿½gren syndrome. Annals of Neurology, 2000, 47, 260-264.	5.3	40
372	The dopamine D2 receptor gene is a susceptibility locus for Parkinson's disease. Movement Disorders, 2000, 15, 120-126.	3.9	108
373	Effects of glucose ingestion on cardiac autonomic nervous system in healthy centenarians: differences with aged subjects. European Journal of Clinical Investigation, 2000, 30, 277-284.	3.4	20
374	Familial temporal lobe epilepsy. Epilepsy Research, 2000, 38, 127-132.	1.6	58
375	Interhemispheric threshold differences in idiopathic generalized epilepsies with versive or circling seizures determined with focal magnetic transcranial stimulation. Epilepsy Research, 2000, 40, 1-6.	1.6	24
376	Charcot-Marie-Tooth type 4B is caused by mutations in the gene encoding myotubularin-related protein-2. Nature Genetics, 2000, 25, 17-19.	21.4	462
377	The nicotinic receptor Î²2 subunit is mutant in nocturnal frontal lobe epilepsy. Nature Genetics, 2000, 26, 275-276.	21.4	433
378	Social Aspects of Epilepsy in the Adult in Seven European Countries. Epilepsia, 2000, 41, 998-1004.	5.1	59

#	Article	IF	Citations
379	The long-duration response to <scp>l</scp> -dopa in the treatment of early PD. Neurology, 2000, 54, 1910-1915.	1.1	39
380	Plasma Leptin Concentrations and Cardiac Autonomic Nervous System in Healthy Subjects with Different Body Weights. Journal of Clinical Endocrinology and Metabolism, 2000, 85, 1810-1814.	3.6	81
381	Genetic Refinement and Physical Mapping of the CMT4B Gene on Chromosome 11q22. Genomics, 2000, 63, 271-278.	2.9	18
382	Chronic Vitamin E Administration Improves Brachial Reactivity and Increases Intracellular Magnesium Concentration in Type II Diabetic Patients. Journal of Clinical Endocrinology and Metabolism, 2000, 85, 109-115.	3.6	72
383	Chronic Vitamin E Administration Improves Brachial Reactivity and Increases Intracellular Magnesium Concentration in Type II Diabetic Patients. Journal of Clinical Endocrinology and Metabolism, 2000, 85, 109-115.	3.6	28
384	Plasma Leptin Concentrations and Cardiac Autonomic Nervous System in Healthy Subjects with Different Body Weights. Journal of Clinical Endocrinology and Metabolism, 2000, 85, 1810-1814.	3.6	28
385	Genetic Refinement And Physical Mapping Of The CMT4B Gene On Chromosome 11Q22. Journal of the Peripheral Nervous System, 2000, 5, 116-116.	3.1	0
386	CORRELATION BETWEEN HYPERTENSION AND OVERWEIGHT IN ELDERLY OUT-PATIENTS. Journal of Hypertension, 2000, 18, S25.	0.5	0
387	Identification of a new polymorphism (c134G>A) in the exon 2 of the myelin protein zero gene Communicated by: Mark H. Paalman Online Citation: Human Mutation, Mutation and Polymorphism Report #100 (1999) Online http://journals.wiley.com/1059-7794/pdf/mutation/mpr100.pdf. Human Mutation, 2000, 15, 299.	2.5	3
388	Pharmacodynamics of the long-duration response to levodopa in PD. Neurology, 1999, 53, 557-557.	1.1	32
389	Low Plasma Insulinâ€Like Growth Factorâ€1 Concentrations Predict Worsening of Insulinâ€Mediated Glucose Uptake in Older People. Journal of the American Geriatrics Society, 1999, 47, 1312-1318.	2.6	29
390	Loss of long-duration response to levodopa over time in PD. Neurology, 1999, 52, 763-763.	1.1	60
391	Dopamine D ₂ receptor gene polymorphism and the risk of levodopa-induced dyskinesias in PD. Neurology, 1999, 53, 1425-1425.	1.1	116
392	Emotion-Induced Myoclonic Absence-Like Seizures in a Patient with Inv-Dup(15) Syndrome: A Clinical, EEG, and Molecular Genetic Study. Epilepsia, 1999, 40, 1316-1319.	5.1	15
393	Apolipoprotein E Polymorphisms and the Risk of Nonlesional Temporal Lobe Epilepsy. Epilepsia, 1999, 40, 1804-1807.	5.1	36
394	Idiopathic generalized epilepsies with versive or circling seizures. Acta Neurologica Scandinavica, 1999, 99, 219-224.	2.1	26
395	Hyperekplexia in a patient with a brainstem vascular anomaly. Acta Neurologica Scandinavica, 1999, 99, 255-259.	2.1	14
396	Autosomal Recessive Hereditary Motor and Sensory Neuropathy with Focally Folded Myelin Sheaths (CMT4B). Annals of the New York Academy of Sciences, 1999, 883, 47-55.	3.8	16

#	Article	IF	Citations
397	A hypofibrinolytic state in overweight patients with cerebral venous thrombosis and isolated intracranial hypertension. Journal of Neurology, 1999, 246, 1086-1089.	3.6	30
398	Prognostic importance of insulin-mediated glucose uptake in aged patients with congestive heart failure secondary to mitral and/or aortic valve disease. American Journal of Cardiology, 1999, 83, 1338-1344.	1.6	75
399	Apolipoprotein E polymorphisms and Parkinson's disease. Neuroscience Letters, 1999, 277, 83-86.	2.1	18
400	Intralipid infusion combined with propranolol administration has favorable metabolic effects in elderly malnourished cancer patients. Metabolism: Clinical and Experimental, 1999, 48, 291-297.	3.4	19
401	Effects of different insulin infusion rates on heart rate variability in lean and obese subjects. Metabolism: Clinical and Experimental, 1999, 48, 755-762.	3.4	39
402	Baseline heart rate variability in healthy centenarians: differences compared with aged subjects (>75 years old). Clinical Science, 1999, 97, 579.	4.3	8
403	APOE and risk of cognitive impairment in multiple sclerosis. Acta Neurologica Scandinavica, 1999, 100, 290-295.	2.1	51
404	Autosomal recessive hereditary motor and sensory neuropathy with focally folded myelin sheaths (CMT4B). Annals of the New York Academy of Sciences, 1999, 883, 47-55.	3.8	2
405	Myoclonic Absence-Like Seizures and Chromosome Abnormality Syndromes. Epilepsia, 1998, 39, 660-663.	5.1	52
406	Usefulness of latero-orbital electrodes in detecting interictal epileptiform activity â€“ a study of 60 patients with complex partial seizures. Electroencephalography and Clinical Neurophysiology, 1998, 107, 174-176.	0.3	3
407	Spinal muscular atrophy due to an isolated deletion of exon 8 of the telomeric survival motor neuron gene. Annals of Neurology, 1998, 44, 836-839.	5.3	32
408	Treatment of palatal myoclonus with sumatriptan. Movement Disorders, 1998, 13, 195-195.	3.9	5
409	CAG repeat length and clinical features in three Italian families with spinocerebellar ataxia type 2 (SCA2): early impairment of Wisconsin Card Sorting Test and saccade velocity. Journal of Neurology, 1998, 245, 647-652.	3.6	59
410	Lowering fatty acids potentiates acute insulin response in first degree relatives of people with Type II diabetes. Diabetologia, 1998, 41, 1127-1132.	6.3	83
411	Plasma sex hormones are significantly associated with plasma leptin concentration in healthy subjects. Clinical Endocrinology, 1998, 48, 291-297.	2.4	59
412	Effect of metformin on food intake in obese subjects. European Journal of Clinical Investigation, 1998, 28, 441-446.	3.4	115
413	Assessment of sense of taste in Italian centenarians,. Archives of Gerontology and Geriatrics, 1998, 26, 177-183.	3.0	6
414	Laboratory parameters of Italian centenarians. Archives of Gerontology and Geriatrics, 1998, 27, 67-74.	3.0	9

#	Article	IF	Citations
415	Genetic heterogeneity in autosomal recessive hereditary motor and sensory neuropathy with focally folded myelin sheaths (CMT4B). Neurology, 1998, 50, 799-801.	1.1	46
416	Kufs' disease presenting as late-onset epilepsia partialis continua. Neurology, 1998, 51, 1180-1182.	1.1	12
417	Randomized trial comparing two different high doses of methylprednisolone in MS A clinical and MRI study. Neurology, 1998, 50, 1833-1836.	1.1	133
418	Oxidative Stress and Advancing Age: Results in Healthy Centenarians. Journal of the American Geriatrics Society, 1998, 46, 833-838.	2.6	105
419	Mild Non-lesional Temporal Lobe Epilepsy: A Common, Unrecognized Disorder with Onset in Adulthood. Canadian Journal of Neurological Sciences, 1998, 25, 282-286.	0.5	55
420	MRI of skeletal muscles in autosomal recessive hereditary motor and sensory neuropathy with focally folded myelin sheaths. European Neurology, 1998, 39, 191-2.	1.4	4
421	The significance of electrocorticographic findings in focal cortical dysplasia: a review of their clinical, electrophysiological and neurochemical characteristics. Electroencephalography and Clinical Neurophysiology Supplement, 1998, 48, 77-96.	0.0	14
422	Serum Levels of Insulin-Like Growth Factor-I (IGF-I) and IGF-Binding Protein-3 in Healthy Centenarians: Relationship with Plasma Leptin and Lipid Concentrations, Insulin Action, and Cognitive Function. Journal of Clinical Endocrinology and Metabolism, 1997, 82, 2204-2209.	3.6	166
423	Hereditary demyelinating neuropathy of infancy: a genetically complex syndrome. Brain, 1997, 120, 2113-2115.	7.6	11
424	Preserved Antilipolytic Insulin Action Is Associated with a Less Atherogenic Plasma Lipid Profile in Healthy Centenarians. Journal of the American Geriatrics Society, 1997, 45, 1504-1509.	2.6	18
425	Disappearance of periodic sharp wave complexes in Creutzfeldt-Jakob disease. Neurophysiologie Clinique, 1997, 27, 277-282.	2.2	8
426	Insulin resistance and advancing age: What role for dehydroepiandrosterone sulfate?. Metabolism: Clinical and Experimental, 1997, 46, 1281-1286.	3.4	30
427	Spontaneous remission of childhood epilepsy in two patients with focal extraopercular cortical dysplasia. Brain and Development, 1997, 19, 422-425.	1.1	5
428	Glucose ingestion affects cardiac ANS in healthy subjects with different amounts of body fat.. American Journal of Physiology - Endocrinology and Metabolism, 1997, 273, E471.	3.5	22
429	Negative Myoclonic Status Due to Antiepileptic Drug Tapering: Report of Three Cases. Epilepsia, 1997, 38, 819-823.	5.1	18
430	Association of Fasting Plasma Free Fatty Acid Concentration and Frequency of Ventricular Premature Complexes in Nonischemic Non-Insulin-Dependent Diabetic Patients. American Journal of Cardiology, 1997, 80, 932-937.	1.6	48
431	Epidemiological and socioeconomic aspects of Italian centenarians. Archives of Gerontology and Geriatrics, 1997, 25, 149-157.	3.0	33
432	Long-duration response to levodopa influences the pharmacodynamics of short-duration response in Parkinson's disease. Annals of Neurology, 1997, 42, 245-248.	5.3	63

#	Article	IF	Citations
433	Action palatal tremor in a patient with primary intestinal lymphoma. Movement Disorders, 1997, 12, 794-797.	3.9	12
434	Metabolic changes in elderly cancer patients after glucose ingestion. Cancer, 1997, 79, 177-184.	4.1	8
435	Serum Levels of Insulin-Like Growth Factor-I (IGF-I) and IGF-Binding Protein-3 in Healthy Centenarians: Relationship with Plasma Leptin and Lipid Concentrations, Insulin Action, and Cognitive Function. Journal of Clinical Endocrinology and Metabolism, 1997, 82, 2204-2209.	3.6	63
436	Sequential occurrence of benign partial epilepsy and childhood absence epilepsy in three patients. Brain and Development, 1996, 18, 212-215.	1.1	23
437	Usefulness of focal rhythmic discharges on scalp EEG of patients with focal cortical dysplasia and intractable epilepsy. Electroencephalography and Clinical Neurophysiology, 1996, 98, 243-249.	0.3	170
438	Hyperinsulinemia is associated with ventricular premature complexes. Metabolism: Clinical and Experimental, 1996, 45, 1248-1253.	3.4	5
439	Autosornal recessive hereditary motor and sensory neuropathy with focally folded myelin sheaths. Neurology, 1996, 46, 1318-1318.	1.1	121
440	Localization of a gene responsible for autosomal recessive demyelinating neuropathy with focally folded myelin sheaths to chromosome 11q23 by homozygosity mapping and haplotype sharing. Human Molecular Genetics, 1996, 5, 1051-1054.	2.9	93
441	Glucose tolerance and insulin action in healthy centenarians. American Journal of Physiology - Endocrinology and Metabolism, 1996, 270, E890-E894.	3.5	83
442	Functional Preservation of Benzodiazepine Receptors of the Primary Somatosensory Cortex in Creutzfeldt-Jakob Disease: A Pharmacologic-Evoked Potential Study. Clinical Neuropharmacology, 1996, 19, 87-91.	0.7	6
443	Regression of left ventricular hypertrophy in hypertensive elderly patients with carvedilol. Archives of Gerontology and Geriatrics, 1996, 22, 143-147.	3.0	5
444	The therapy of aged diabetic hypertensives. Archives of Gerontology and Geriatrics, 1996, 22, 275-285.	3.0	0
445	Photic-Induced Epileptic Negative Myoclonus: A Case Report. Epilepsia, 1996, 37, 492-494.	5.1	9
446	Clinical effects of oral theophylline in elderly patients with sick sinus syndrome. Archives of Gerontology and Geriatrics, 1996, 22, 221-224.	3.0	2
447	The oncological therapy in the elderly. Archives of Gerontology and Geriatrics, 1996, 22, 593-598.	3.0	1
448	Significance of mesial temporal atrophy in relation to intracranial ictal and interictal stereo EEG abnormalities. Brain, 1996, 119, 1317-1326.	7.6	48
449	Body composition, body fat distribution, and resting metabolic rate in healthy centenarians. American Journal of Clinical Nutrition, 1995, 62, 746-750.	4.7	71
450	Left ventricular hypertrophy is associated with a stronger impairment of nonâ€oxidative glucose metabolism in hypertensive patients. European Journal of Clinical Investigation, 1995, 25, 529-533.	3.4	28

#	Article	IF	Citations
451	Focal Intermittent Delta Activity in Patients with Mesiotemporal Atrophy: A Reliable Marker of the Epileptogenic Focus. Epilepsia, 1995, 36, 122-129.	5.1	103
452	Intrinsic epileptogenicity of human dysplastic cortex as suggested by corticography and surgical results. Annals of Neurology, 1995, 37, 476-487.	5.3	758
453	Low-dose Iloprost infusion improves insulin action and non-oxidative glucose metabolism in hypertensive patients. European Journal of Clinical Pharmacology, 1995, 48, 333-8.	1.9	6
454	Opposite effects of short- and long-term fatty acid infusion on insulin secretion in healthy subjects. Diabetologia, 1995, 38, 1295-1299.	6.3	189
455	De novo epileptic confusional status in a patient with cobalamin deficiency. Metabolic Brain Disease, 1995, 10, 233-238.	2.9	9
456	Chronic intake of pharmacological doses of vitamin E might be useful in the therapy of elderly patients with coronary heart disease. American Journal of Clinical Nutrition, 1995, 61, 848-852.	4.7	41
457	Metabolic benefits deriving from chronic vitamin C supplementation in aged non-insulin dependent diabetics.. Journal of the American College of Nutrition, 1995, 14, 387-392.	1.8	130
458	The Relation of Spike Foci and of Clinical Seizure Characteristics to Different Patterns of Mesial Temporal Atrophy. Archives of Neurology, 1995, 52, 287-293.	4.5	25
459	Negative myoclonus during valproate-related stupor. Neurophysiological evidence of a cortical non-epileptic origin. Electroencephalography and Clinical Neurophysiology, 1995, 94, 103-108.	0.3	32
460	Relationship between atrophy of the amygdala and ictal fear in temporal lobe epilepsy. Brain, 1994, 117, 739-746.	7.6	170
461	Chlorpromazine versus sleep deprivation in activation of EEG in adult-onset partial epilepsy. Journal of Neurology, 1994, 241, 605-610.	3.6	18
462	Operative Strategies for Patients with Cortical Dysplastic Lesions and Intractable Epilepsy. Epilepsia, 1994, 35, S57-71.	5.1	178
463	Total-body and myocardial substrate oxidation in congestive heart failure. Metabolism: Clinical and Experimental, 1994, 43, 174-179.	3.4	152
464	Antioxidants In adipose tissue and risk of myocardial Infarction. Lancet, The, 1994, 343, 595-596.	13.7	0
465	Insulin Resistance and Hypertension in the Elderly. Drugs and Aging, 1994, 4, 403-409.	2.7	5
466	Lateâ€onset drop attacks in temporal lobe epilepsy. Neurology, 1994, 44, 1074-1074.	1.1	42
467	Different contribution of substrates oxidation to insulin resistance in malnourished elderly patients with cancer. Cancer, 1993, 72, 3106-3113.	4.1	8
468	Functional integrity of benzodiazepine receptors of the geniculo-striate visual pathways in Creutzfeldt-Jakob disease. Journal of Neurology, 1993, 240, 25-27.	3.6	10

#	Article	IF	Citations
469	Metabolic features of patients with and without coronary heart disease but with a superimposable cluster of cardiovascular risk factors. Coronary Artery Disease, 1993, 4, 1085-1092.	0.7	7
470	Effects of Magnesium and Nifedipine Infusions on Insulin Action, Substrate Oxidation, and Blood Pressure in Aged Hypertensive Patients. American Journal of Hypertension, 1993, 6, 920-926.	2.0	20
471	Metabolic and cardiovascular benefits deriving from Î²-adrenergic blockade in chronic congestive heart failure. American Heart Journal, 1992, 123, 103-110.	2.7	59
472	Sensory evoked potentials in herpes simplex encephalitis. Neurophysiologie Clinique, 1991, 21, 301-311.	2.2	0
473	Sleep-Induced Masticatory Myoclonus: A Rare Parasomnia Associated with Insomnia. Sleep, 1991, 14, 80-82.	1.1	29
474	Simvastatin reduces plasma lipid levels and improves insulin action in elderly, non-insulin dependent diabetics. European Journal of Clinical Pharmacology, 1991, 40, 27-31.	1.9	62
475	Triphasic Waves and Cerebral Tumors. European Neurology, 1990, 30, 1-5.	1.4	11
476	Nonmetabolic Causes of Triphasic Waves: A Reappraisal. Clinical EEG (electroencephalography), 1990, 21, 120-125.	0.9	23
477	Relationship between cerebrospinal fluid Î²-endorphin and plasma pituitary-gonadal hormone levels in women. Journal of Endocrinological Investigation, 1990, 13, 149-153.	3.3	24
478	Combined Neurophysiological Studies in Creutzfeldt-Jakob Disease: A Case Report. Clinical EEG (electroencephalography), 1989, 20, 103-110.	0.9	10
479	Purulent meningitis due to spontaneous anterior sacral meningocele perforation. Case report. Italian Journal of Neurological Sciences, 1989, 10, 211-213.	0.1	9
480	Rasmussen encephalitis and related conditions. , 0, , 573-578.		0