

Patrizia Avoni

List of Publications by Year in descending order

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37
papers

2,470
citations

257450

24
h-index

345221

36
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docs citations

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times ranked

2954
citing authors

#	ARTICLE	IF	CITATIONS
1	Presence of Skin α -Synuclein Deposits Discriminates Parkinson's Disease from Progressive Supranuclear Palsy and Corticobasal Syndrome. <i>Journal of Parkinson's Disease</i> , 2022, 12, 585-591.	2.8	9
2	Epilepsy with auditory features: Long-term outcome and predictors of terminal remission. <i>Epilepsia</i> , 2018, 59, 834-843.	5.1	8
3	Subcutaneous immunoglobulin treatment and thromboembolic risk. <i>Annals of Allergy, Asthma and Immunology</i> , 2018, 120, 433-435.	1.0	3
4	Spine Topographical Distribution of Skin α -Synuclein Deposits in Idiopathic Parkinson Disease. <i>Journal of Neuropathology and Experimental Neurology</i> , 2017, 76, 384-389.	1.7	36
5	Syndromic parkinsonism and dementia associated with <i>OPA1</i> missense mutations. <i>Annals of Neurology</i> , 2015, 78, 21-38.	5.3	154
6	Immunotherapy of oneritic stupor in Morvan syndrome: Efficacy documented by actigraphy. <i>Neurology</i> , 2015, 84, 2457-2459.	1.1	12
7	Primary progressive narcolepsy type 1: The other side of the coin. <i>Neurology</i> , 2014, 83, 2189-2190.	1.1	46
8	Small nerve fiber involvement in patients referred for fibromyalgia. <i>Muscle and Nerve</i> , 2014, 49, 757-759.	2.2	90
9	Skin nerve α -synuclein deposits. <i>Neurology</i> , 2014, 82, 1362-1369.	1.1	247
10	Atypical late-onset hereditary spastic paraplegia with thin corpus callosum due to novel compound heterozygous mutations in the SPG11 gene. <i>Journal of Neurology</i> , 2014, 261, 1825-1827.	3.6	12
11	Sleep disorders in patients with spinal cord injury. <i>Sleep Medicine Reviews</i> , 2013, 17, 399-409.	8.5	62
12	Low penetrance of autosomal dominant lateral temporal epilepsy in Italian families without <i>LGI1</i> mutations. <i>Epilepsia</i> , 2013, 54, 1288-1297.	5.1	32
13	Idebenone Treatment In Leber's Hereditary Optic Neuropathy. <i>Brain</i> , 2011, 134, e188-e188.	7.6	192
14	Prognostic factors in patients with mesial temporal lobe epilepsy. <i>Epilepsia</i> , 2009, 50, 41-44.	5.1	51
15	Differential cerebro spinal fluid proteome investigation of Leber hereditary optic neuropathy (LHON) and multiple sclerosis. <i>Journal of Neuroimmunology</i> , 2008, 193, 156-160.	2.3	26
16	The Effect of Entacapone on Levodopa Rate of Absorption and Latency to Motor Response in Patients With Parkinson Disease. <i>Clinical Neuropharmacology</i> , 2008, 31, 267-271.	0.7	4
17	Kinetic-Dynamic Monitoring of Levetiracetam Effects in Patients With Parkinson Disease and Levodopa-Induced Dyskinesias. <i>Clinical Neuropharmacology</i> , 2007, 30, 122-124.	0.7	6
18	Variation in Lamotrigine Plasma Concentrations with Hormonal Contraceptive Monthly Cycles in Patients with Epilepsy. <i>Epilepsia</i> , 2006, 47, 1573-1575.	5.1	85

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19	Pre-symptomatic diagnosis in fatal familial insomnia: serial neurophysiological and 18FDG-PET studies. <i>Brain</i> , 2006, 129, 668-675.	7.6	109
20	A de novo LGI1 mutation in sporadic partial epilepsy with auditory features. <i>Annals of Neurology</i> , 2004, 56, 455-456.	5.3	54
21	Molecular biology of channelopathies: impact on diagnosis and treatment. <i>Expert Review of Neurotherapeutics</i> , 2004, 4, 519-539.	2.8	7
22	Autosomal Dominant Lateral Temporal Epilepsy: Clinical Spectrum, New Epitempin Mutations, and Genetic Heterogeneity in Seven European Families. <i>Epilepsia</i> , 2003, 44, 1289-1297.	5.1	134
23	Topiramate Therapeutic Monitoring in Patients With Epilepsy: Effect of Concomitant Antiepileptic Drugs. <i>Therapeutic Drug Monitoring</i> , 2002, 24, 332-337.	2.0	40
24	Does the prion protein gene 129 codon polymorphism influence sleep? Evidence from a fatal familial insomnia kindred. <i>Clinical Neurophysiology</i> , 2002, 113, 1948-1953.	1.5	13
25	Mitochondrial DNA nucleotide changes C14482G and C14482A in the ND6 gene are pathogenic for Leber's hereditary optic neuropathy. <i>Annals of Neurology</i> , 2002, 51, 774-778.	5.3	50
26	Partial epilepsy with prominent auditory symptoms not linked to chromosome 10q. <i>Epileptic Disorders</i> , 2002, 4, 183-7.	1.3	14
27	Absence of sleep EEG markers in fatal familial insomnia healthy carriers: a spectral analysis study. <i>Clinical Neurophysiology</i> , 2001, 112, 1888-1892.	1.5	26
28	Levodopa Therapy Monitoring in Patients With Parkinson Disease: a Kinetic "Dynamic Approach. <i>Therapeutic Drug Monitoring</i> , 2001, 23, 621-629.	2.0	56
29	Familial continuous motor unit activity and epilepsy. <i>Muscle and Nerve</i> , 2001, 24, 630-633.	2.2	21
30	Congenital encephalomyopathy with epilepsy, chorioretinitis, basal ganglia involvement, and muscle minicores. <i>Annals of Neurology</i> , 2000, 47, 395-399.	5.3	4
31	Clinical Features of Fatal Familial Insomnia: Phenotypic Variability in Relation to a Polymorphism at Codon 129 of the Prion Protein Gene. <i>Brain Pathology</i> , 1998, 8, 515-520.	4.1	110
32	Fatal Familial Insomnia: A Human Model of Prion Disease. , 1998, , 33-35.		0
33	Motor Overactivity and Loss of Motor Circadian Rhythm in Fatal Familial Insomnia: An Actigraphic Study. <i>Sleep</i> , 1997, 20, 739-742.	1.1	38
34	Partial Epilepsy of Long Duration: Changing Semiology with Age. <i>Epilepsia</i> , 1996, 37, 162-164.	5.1	47
35	Dissociated 24-Hour Patterns of Somatotropin and Prolactin in Fatal Familial Insomnia. <i>Neuroendocrinology</i> , 1995, 61, 731-737.	2.5	36
36	Fatal Familial Insomnia, a Prion Disease with a Mutation at Codon 178 of the Prion Protein Gene. <i>New England Journal of Medicine</i> , 1992, 326, 444-449.	27.0	578

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37	Cardiovascular dysautonomia in fatal familial insomnia. <i>Clinical Autonomic Research</i> , 1991, 1, 15-21.	2.5	58