Cesa Lorella Maria Scaglione

List of Publications by Year
in descending order

72 papers	2,713 citations	201385 27 h-index	197535 49 g-index
74 all docs	74 docs citations	74 times ranked	3876 citing authors

#	Article	IF	Citations
1	In Vivo Diagnosis of Synucleinopathies. Neurology, 2021, 96, e2513-e2524.	1.5	63
2	Spread of segmental/multifocal idiopathic adult-onset dystonia to a third body site. Parkinsonism and Related Disorders, 2021, 87, 70-74.	1.1	8
3	Idiopathic <scp>Nonâ€taskâ€Specific</scp> Upper Limb Dystonia, a Neglected Form of Dystonia. Movement Disorders, 2020, 35, 2038-2045.	2.2	21
4	Demographic and clinical determinants of neck pain in idiopathic cervical dystonia. Journal of Neural Transmission, 2020, 127, 1435-1439.	1.4	22
5	A Longitudinal Skin Biopsy Study of Phosphorylated Alpha-Synuclein in a Patient With Parkinson Disease and Orthostatic Hypotension. Journal of Neuropathology and Experimental Neurology, 2020, 79, 813-816.	0.9	7
6	Does acute peripheral trauma contribute to idiopathic adult-onset dystonia?. Parkinsonism and Related Disorders, 2020, 71, 40-43.	1.1	18
7	Clinical pharmacokinetics of pramipexole, ropinirole and rotigotine in patients with Parkinson's disease. Parkinsonism and Related Disorders, 2019, 61, 111-117.	1.1	18
8	Observing movement disorders: best practice proposal in the use of video recording in clinical practice. Neurological Sciences, 2019, 40, 333-338.	0.9	2
9	Clinical and genetic characteristics of late-onset Huntington's disease. Parkinsonism and Related Disorders, 2019, 61, 101-105.	1.1	17
10	Expert recommendations for diagnosing cervical, oromandibular, and limb dystonia. Neurological Sciences, 2019, 40, 89-95.	0.9	44
11	Skin Î±-synuclein deposits differ in clinical variants of synucleinopathy: an in vivo study. Scientific Reports, 2018, 8, 14246.	1.6	75
12	Skin Nerve Phosphorylated Î±-Synuclein Deposits in Parkinson Disease With Orthostatic Hypotension. Journal of Neuropathology and Experimental Neurology, 2018, 77, 942-949.	0.9	40
13	Spread of dystonia in patients with idiopathic adultâ€onset laryngeal dystonia. European Journal of Neurology, 2018, 25, 1341-1344.	1.7	11
14	Reduced Cancer Incidence in Huntington's Disease: Analysis in the Registry Study. Journal of Huntington's Disease, 2018, 7, 209-222.	0.9	14
15	The Italian Dystonia Registry: rationale, design and preliminary findings. Neurological Sciences, 2017, 38, 819-825.	0.9	35
16	Spine Topographical Distribution of Skin Î±-Synuclein Deposits in Idiopathic Parkinson Disease. Journal of Neuropathology and Experimental Neurology, 2017, 76, 384-389.	0.9	36
17	Modulation of the Muscle Activity During Sleep in Cervical Dystonia. Sleep, 2017, 40, .	0.6	22
18	Identification of genetic variants associated with Huntington's disease progression: a genome-wide association study. Lancet Neurology, The, 2017, 16, 701-711.	4.9	248

#	Article	IF	Citations
19	Cognitive decline in Huntington's disease expansion gene carriers. Cortex, 2017, 95, 51-62.	1.1	50
20	BehÃ§et disease presenting with movement disorders and antibasal ganglia antibodies. Autoimmunity Reviews, 2016, 15, 287-288.	2.5	11
21	A patient with PMP22-related hereditary neuropathy and DBH-gene-related dysautonomia. Journal of Neurology, 2015, 262, 2373-2381.	1.8	8
22	Skin biopsy and Iâ€123 MIBG scintigraphy findings in idiopathic Parkinson's disease and parkinsonism: A comparative study. Movement Disorders, 2015, 30, 986-989.	2.2	22
23	Primary familial brain calcification: Genetic analysis and clinical spectrum. Movement Disorders, 2014, 29, 1691-1695.	2.2	95
24	Quality of life in patients with craniocervical dystonia: Italian validation of the â€œCervical Dystonia Impact Profile (CDIP-58)â€•and the â€œCraniocervical Dystonia Questionnaire (CDQ-24)â€• Neurological Sciences, 2014, 35, 1053-1058.	0.9	3
25	Skin nerve Î±-synuclein deposits. Neurology, 2014, 82, 1362-1369.	1.5	247
26	Arylsulphatase A activity in familial parkinsonism: a pathogenetic role?. Journal of Neurology, 2014, 261, 1803-1809.	1.8	8
27	Validation of the Italian version of the Movement Disorder Societyâ€"Unified Parkinsonâ€™s Disease Rating Scale. Neurological Sciences, 2013, 34, 683-687.	0.9	123
28	Loss of temporal retinal nerve fibers in Parkinson disease: a mitochondrial pattern?. European Journal of Neurology, 2013, 20, 198-201.	1.7	92
29	Neurosyphilis orofacial dyskinesia: The candy sign. Movement Disorders, 2013, 28, 246-247.	2.2	20
30	Suicidal ideation in a European Huntington's disease population. Journal of Affective Disorders, 2013, 151, 248-258.	2.0	74
31	The V471A Polymorphism in Autophagy-Related Gene ATG7 Modifies Age at Onset Specifically in Italian Huntington Disease Patients. PLoS ONE, 2013, 8, e68951.	1.1	49
32	Discrepancies in reporting the CAG repeat lengths for Huntington's disease. European Journal of Human Genetics, 2012, 20, 20-26.	1.4	20
33	Group I nonreciprocal inhibition in restless legs syndrome secondary to chronic renal failure. Parkinsonism and Related Disorders, 2012, 18, 362-366.	1.1	15
34	From ritual sword duel to electrophysiology: Hyperactive facial motor nucleus in hemifacial spasm. Movement Disorders, 2012, 27, 927-928.	2.2	1
35	Eating Disorder as a Psychiatric Onset of Juvenile Huntington's Disease. American Journal of Psychiatry, 2011, 168, 1120-1121.	4.0	4
36	Did Goethe Describe Attention Deficit Hyperactivity Disorder. European Neurology, 2011, 65, 70-71.	0.6	3

#	Article	IF	Citations
37	The Effect of a Clinically Practical Exercise on Levodopa Bioavailability and Motor Response in Patients With Parkinson Disease. Clinical Neuropharmacology, 2010, 33, 254-256.	0.2	9
38	Benign tremulous parkinsonism in a patient with dardarin mutation. Movement Disorders, 2009, 24, 1399-1401.	2.2	4
39	Common mutations in the LRRK2 exon 41 are not responsible for essential tremor in Italian patients. Parkinsonism and Related Disorders, 2009, 15, 162-163.	1.1	9
40	MRI and SPECT of midbrain and striatal degeneration in fragile X-associated tremor/ataxia syndrome. Journal of Neurology, 2008, 255, 144-146.	1.8	40
41	Group I nonreciprocal inhibition in primary restless legs syndrome. Movement Disorders, 2008, 23, 96-100.	2.2	14
42	PINK1heterozygous rare variants: prevalence, significance and phenotypic spectrum. Human Mutation, 2008, 29, 565-565.	1.1	74
43	DRD3 Ser9Gly variant is not associated with essential tremor in a series of Italian patients. European Journal of Neurology, 2008, 15, 985-987.	1.7	25
44	Diffusion-weighted brain imaging study of patients with clinical diagnosis of corticobasal degeneration, progressive supranuclear palsy and Parkinson's disease. Brain, 2008, 131, 2690-2700.	3.7	131
45	The Effect of Entacapone on Levodopa Rate of Absorption and Latency to Motor Response in Patients With Parkinson Disease. Clinical Neuropharmacology, 2008, 31, 267-271.	0.2	4
46	Kinetic-Dynamic Monitoring of Levetiracetam Effects in Patients With Parkinson Disease and Levodopa-Induced Dyskinesias. Clinical Neuropharmacology, 2007, 30, 122-124.	0.2	6
47	Diffusion-weighted imaging study of patients with essential tremor. Movement Disorders, 2007, 22, 1182-1185.	2.2	67
48	Transient left ventricular apical ballooning at the onset of multiple system atrophy. Journal of Cardiovascular Medicine, 2006, 7, 631-636.	0.6	4
49	Frequency and phenotypes of LRRK2 G2019S mutation in Italian patients with Parkinson's disease. Movement Disorders, 2006, 21, 1232-1235.	2.2	28
50	A case of fragile X premutation tremor/ataxia syndrome with evidence of mitochondrial dysfunction. Movement Disorders, 2006, 21, 1541-1542.	2.2	9
51	Studies of the Cost-Effectiveness of Social Work Services in Aging: A Review of the Literature. Research on Social Work Practice, 2006, 16, 67-73.	1.1	22
52	PINK1, Parkin, and DJ-1 mutations in Italian patients with early-onset parkinsonism. European Journal of Human Genetics, 2005, 13, 1086-1093.	1.4	132
53	Adult Alexander's disease without leukoencephalopathy. Annals of Neurology, 2005, 58, 813-814.	2.8	28
54	Novel parkin mutations detected in patients with early-onset Parkinson's disease. Movement Disorders, 2005, 20, 424-431.	2.2	60

#	Article	IF	Citations
55	Premutations in theFMR1 gene as a modifying factor inParkin-associated Parkinson's disease?. Movement Disorders, 2005, 20, 1060-1062.	2.2	30
56	REM sleep behaviour disorder in Parkinson?s disease: a questionnaire-based study. Neurological Sciences, 2005, 25, 316-321.	0.9	160
57	Multiparametric MRI in a patient with adult-onset leukoencephalopathy with vanishing white matter. Neurology, 2004, 62, 323-326.	1.5	24
58	Mutational analysis of parkin gene by denaturing high-performance liquid chromatography (DHPLC) in essential tremor. Parkinsonism and Related Disorders, 2004, 10, 357-362.	1.1	20
59	Dopamine Transporter Gene Polymorphism, SPECT Imaging, and Levodopa Response in Patients with Parkinson Disease. Clinical Neuropharmacology, 2004, 27, 111-115.	0.2	31
60	Assessing dopaminergic function in Parkinson?s disease: levodopa kinetic-dynamic modeling and SPECT. Journal of Neurology, 2003, 250, 1475-1481.	1.8	10
61	Levodopa pharmacokinetics and dyskinesias: are there sex-related differences?. Neurological Sciences, 2003, 24, 192-193.	0.9	48
62	Essential tremor is not associated with ?-synuclein gene haplotypes. Movement Disorders, 2003, 18, 823-826.	2.2	19
63	Atypical familial motor neuropathy in patients with mutant TTR Ile68Leu. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2003, 10, 185-189.	1.4	13
64	Late-Onset PANDAS Syndrome with Abdominal Muscle Involvement. European Neurology, 2002, 48, 49-51.	0.6	12
65	Clinical and genetic study of essential tremor in the Italian population. Neurological Sciences, 2001, 22, 39-40.	0.9	24
66	No evidence of association between CAG expansions and essential tremor in a large cohort of Italian patients. Journal of Neural Transmission, 2001, 108, 297-304.	1.4	16
67	Symptomatic unruptured capillary telangiectasia of the brain stem: report of three cases and review of the literature. Journal of Neurology, Neurosurgery and Psychiatry, 2001, 71, 390-393.	0.9	35
68	Deficit of brain and skeletal muscle bioenergetics in progressive supranuclear palsy shown in vivo by phosphorus magnetic resonance spectroscopy. Movement Disorders, 2000, 15, 889-893.	2.2	41
69	Neurophysiological evaluation of areflexia in Holmes-Adie syndrome. Neurophysiologie Clinique, 1999, 29, 255-262.	1.0	8
70	Hemiparkinsonism-hemiatrophy: A new observation. Journal of Neurology, 1998, 245, 180-182.	1.8	5
71	Acute brachial plexus neuropathy as a presenting sign of peripheral nervous system involvement in paraproteinaemia. Acta Neurologica Scandinavica, 1997, 95, 319-320.	1.0	5
72	Nonâ€Hodgkin's lymphoma mimicking amyotrophic lateral sclerosis. European Journal of Neurology, 1996, 3, 479-479.	1.7	0