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	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
2	Skin nerve Î±-synuclein deposits. Neurology, 2014, 82, 1362-1369.	1.5	247
3	REM sleep behaviour disorder in Parkinson?s disease: a questionnaire-based study. Neurological Sciences, 2005, 25, 316-321.	0.9	160
4	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
5	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
6	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
7	Primary familial brain calcification: Genetic analysis and clinical spectrum. Movement Disorders, 2014, 29, 1691-1695.	2.2	95
8	Loss of temporal retinal nerve fibers in Parkinson disease: a mitochondrial pattern?. European Journal of Neurology, 2013, 20, 198-201.	1.7	92
9	Skin Î±-synuclein deposits differ in clinical variants of synucleinopathy: an in vivo study. Scientific Reports, 2018, 8, 14246.	1.6	75
10	PINK1heterozygous rare variants: prevalence, significance and phenotypic spectrum. Human Mutation, 2008, 29, 565-565.	1.1	74
11	Suicidal ideation in a European Huntington's disease population. Journal of Affective Disorders, 2013, 151, 248-258.	2.0	74
12	Diffusion-weighted imaging study of patients with essential tremor. Movement Disorders, 2007, 22, 1182-1185.	2.2	67
13	In Vivo Diagnosis of Synucleinopathies. Neurology, 2021, 96, e2513-e2524.	1.5	63
14	Novel parkin mutations detected in patients with early-onset Parkinson's disease. Movement Disorders, 2005, 20, 424-431.	2.2	60
15	Cognitive decline in Huntington's disease expansion gene carriers. Cortex, 2017, 95, 51-62.	1.1	50
16	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
17	Levodopa pharmacokinetics and dyskinesias: are there sex-related differences?. Neurological Sciences, 2003, 24, 192-193.	0.9	48
18	Expert recommendations for diagnosing cervical, oromandibular, and limb dystonia. Neurological Sciences, 2019, 40, 89-95.	0.9	44

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19	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
20	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
21	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
22	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
23	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
24	The Italian Dystonia Registry: rationale, design and preliminary findings. Neurological Sciences, 2017, 38, 819-825.	0.9	35
25	Dopamine Transporter Gene Polymorphism, SPECT Imaging, and Levodopa Response in Patients with Parkinson Disease. Clinical Neuropharmacology, 2004, 27, 111-115.	0.2	31
26	Premutations in theFMR1 gene as a modifying factor inParkin-associated Parkinson's disease?. Movement Disorders, 2005, 20, 1060-1062.	2.2	30
27	Adult Alexander's disease without leukoencephalopathy. Annals of Neurology, 2005, 58, 813-814.	2.8	28
28	Frequency and phenotypes of LRRK2 G2019S mutation in Italian patients with Parkinson's disease. Movement Disorders, 2006, 21, 1232-1235.	2.2	28
29	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
30	Clinical and genetic study of essential tremor in the Italian population. Neurological Sciences, 2001, 22, 39-40.	0.9	24
31	Multiparametric MRI in a patient with adult-onset leukoencephalopathy with vanishing white matter. Neurology, 2004, 62, 323-326.	1.5	24
32	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
33	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
34	Modulation of the Muscle Activity During Sleep in Cervical Dystonia. Sleep, 2017, 40, .	0.6	22
35	Demographic and clinical determinants of neck pain in idiopathic cervical dystonia. Journal of Neural Transmission, 2020, 127, 1435-1439.	1.4	22
36	Idiopathic <scp>Nonâ€taskâ€Specific</scp> Upper Limb Dystonia, a Neglected Form of Dystonia. Movement Disorders, 2020, 35, 2038-2045.	2.2	21

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37	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
38	Discrepancies in reporting the CAG repeat lengths for Huntington's disease. European Journal of Human Genetics, 2012, 20, 20-26.	1.4	20
39	Neurosyphilis orofacial dyskinesia: The candy sign. Movement Disorders, 2013, 28, 246-247.	2.2	20
40	Essential tremor is not associated with ?-synuclein gene haplotypes. Movement Disorders, 2003, 18, 823-826.	2.2	19
41	Clinical pharmacokinetics of pramipexole, ropinirole and rotigotine in patients with Parkinson's disease. Parkinsonism and Related Disorders, 2019, 61, 111-117.	1.1	18
42	Does acute peripheral trauma contribute to idiopathic adult-onset dystonia?. Parkinsonism and Related Disorders, 2020, 71, 40-43.	1.1	18
43	Clinical and genetic characteristics of late-onset Huntington's disease. Parkinsonism and Related Disorders, 2019, 61, 101-105.	1.1	17
44	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
45	Group I nonreciprocal inhibition in restless legs syndrome secondary to chronic renal failure. Parkinsonism and Related Disorders, 2012, 18, 362-366.	1.1	15
46	Group I nonreciprocal inhibition in primary restless legs syndrome. Movement Disorders, 2008, 23, 96-100.	2.2	14
47	Reduced Cancer Incidence in Huntington's Disease: Analysis in the Registry Study. Journal of Huntington's Disease, 2018, 7, 209-222.	0.9	14
48	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
49	Late-Onset PANDAS Syndrome with Abdominal Muscle Involvement. European Neurology, 2002, 48, 49-51.	0.6	12
50	BehÃ§et disease presenting with movement disorders and antibasal ganglia antibodies. Autoimmunity Reviews, 2016, 15, 287-288.	2.5	11
51	Spread of dystonia in patients with idiopathic adultâ€onset laryngeal dystonia. European Journal of Neurology, 2018, 25, 1341-1344.	1.7	11
52	Assessing dopaminergic function in Parkinson?s disease: levodopa kinetic-dynamic modeling and SPECT. Journal of Neurology, 2003, 250, 1475-1481.	1.8	10
53	A case of fragile X premutation tremor/ataxia syndrome with evidence of mitochondrial dysfunction. Movement Disorders, 2006, 21, 1541-1542.	2.2	9
54	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

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55	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
56	Neurophysiological evaluation of areflexia in Holmes-Adie syndrome. Neurophysiologie Clinique, 1999, 29, 255-262.	1.0	8
57	Arylsulphatase A activity in familial parkinsonism: a pathogenetic role?. Journal of Neurology, 2014, 261, 1803-1809.	1.8	8
58	A patient with PMP22-related hereditary neuropathy and DBH-gene-related dysautonomia. Journal of Neurology, 2015, 262, 2373-2381.	1.8	8
59	Spread of segmental/multifocal idiopathic adult-onset dystonia to a third body site. Parkinsonism and Related Disorders, 2021, 87, 70-74.	1.1	8
60	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
61	Kinetic-Dynamic Monitoring of Levetiracetam Effects in Patients With Parkinson Disease and Levodopa-Induced Dyskinesias. Clinical Neuropharmacology, 2007, 30, 122-124.	0.2	6
62	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
63	Hemiparkinsonism-hemiatrophy: A new observation. Journal of Neurology, 1998, 245, 180-182.	1.8	5
64	Transient left ventricular apical ballooning at the onset of multiple system atrophy. Journal of Cardiovascular Medicine, 2006, 7, 631-636.	0.6	4
65	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
66	Benign tremulous parkinsonism in a patient with dardarin mutation. Movement Disorders, 2009, 24, 1399-1401.	2.2	4
67	Eating Disorder as a Psychiatric Onset of Juvenile Huntington's Disease. American Journal of Psychiatry, 2011, 168, 1120-1121.	4.0	4
68	Did Goethe Describe Attention Deficit Hyperactivity Disorder. European Neurology, 2011, 65, 70-71.	0.6	3
69	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
70	Observing movement disorders: best practice proposal in the use of video recording in clinical practice. Neurological Sciences, 2019, 40, 333-338.	0.9	2
71	From ritual sword duel to electrophysiology: Hyperactive facial motor nucleus in hemifacial spasm. Movement Disorders, 2012, 27, 927-928.	2.2	1
72	Nonâ€Hodgkin's lymphoma mimicking amyotrophic lateral sclerosis. European Journal of Neurology, 1996, 3, 479-479.	1.7	0