Giovanni Cossu

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Mutations in SLC30A10 Cause Parkinsonism and Dystonia with Hypermanganesemia, Polycythemia, and Chronic Liver Disease. American Journal of Human Genetics, 2012, 90, 467-477.	6.2	343
2	The progression of non-motor symptoms in Parkinson's disease and their contribution to motor disability and quality of life. Journal of Neurology, 2012, 259, 2621-2631.	3.6	188
3	Gut Microbiota and Metabolome Alterations Associated with Parkinsonâ \in Ms Disease. MSystems, 2020, 5, .	3.8	161
4	Defective temporal processing of sensory stimuli in DYT1 mutation carriers: a new endophenotype of dystonia?. Brain, 2006, 130, 134-142.	7.6	122
5	A pilot trial of deferiprone for neurodegeneration with brain iron accumulation. Haematologica, 2011, 96, 1708-1711.	3.5	122
6	Neuropathy and levodopa in Parkinson's disease: Evidence from a multicenter study. Movement Disorders, 2013, 28, 1391-1397.	3.9	114
7	LRP10 genetic variants in familial Parkinson's disease and dementia with Lewy bodies: a genome-wide linkage and sequencing study. Lancet Neurology, The, 2018, 17, 597-608.	10.2	101
8	<scp><i>GBA</i>â€Related</scp> Parkinson's Disease: Dissection of Genotype–Phenotype Correlates in a Large Italian Cohort. Movement Disorders, 2020, 35, 2106-2111.	3.9	83
9	Efficacy and safety of deferiprone for the treatment of pantothenate kinase-associated neurodegeneration (PKAN) and neurodegeneration with brain iron accumulation (NBIA): Results from a four years follow-up. Parkinsonism and Related Disorders, 2014, 20, 651-654.	2.2	80
10	Prepulse modulation of the startle reaction and the blink reflex in normal human subjects. Experimental Brain Research, 1999, 129, 49-56.	1.5	79
11	Broadening the phenotype of TARDBP mutations: the TARDBP Ala382Thr mutation and Parkinson's disease in Sardinia. Neurogenetics, 2011, 12, 203-209.	1.4	78
12	Differential induction of dyskinesia and neuroinflammation by pulsatile versus continuous l -DOPA delivery in the 6-OHDA model of Parkinson's disease. Experimental Neurology, 2016, 286, 83-92.	4.1	75
13	Pisa syndrome in Parkinson disease. Neurology, 2015, 85, 1769-1779.	1.1	72
14	Parkinson's disease protein DJ-1 regulates ATP synthase protein components to increase neuronal process outgrowth. Cell Death and Disease, 2019, 10, 469.	6.3	70
15	Restless legs syndrome in multiple sclerosis: A case ontrol study. Movement Disorders, 2009, 24, 697-701.	3.9	64
16	Reversible encephalopathy and axonal neuropathy in Parkinson's disease during duodopa therapy. Movement Disorders, 2009, 24, 2293-2294.	3.9	64
17	Dopaminergic Neuronal Imaging in Genetic Parkinson's Disease: Insights into Pathogenesis. PLoS ONE, 2013, 8, e69190.	2.5	55
18	The `geste antagonistique' induces transient modulation of the blink reflex in human patients with blepharospasm. Neuroscience Letters, 1998, 251, 125-128.	2.1	53

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19	Effects of Physical Rehabilitation Integrated with Rhythmic Auditory Stimulation on Spatio-Temporal and Kinematic Parameters of Gait in Parkinson's Disease. Frontiers in Neurology, 2016, 7, 126.	2.4	52
20	The Use of Footstep Sounds as Rhythmic Auditory Stimulation for Gait Rehabilitation in Parkinson's Disease: A Randomized Controlled Trial. Frontiers in Neurology, 2018, 9, 348.	2.4	51
21	Emergencies in parkinsonism: akinetic crisis, life-threatening dyskinesias, and polyneuropathy during L-Dopa gel treatment. Parkinsonism and Related Disorders, 2009, 15, S233-S236.	2.2	46
22	Which patients discontinue? Issues on Levodopa/carbidopa intestinal gel treatment: Italian multicentre survey of 905 patients with long-term follow-up. Parkinsonism and Related Disorders, 2017, 38, 90-92.	2.2	44
23	Reversible Pisa syndrome (pleurothotonus) due to the cholinesterase inhibitor galantamine: Case report. Movement Disorders, 2004, 19, 1243-1244.	3.9	38
24	The rise and fall of impulse control behavior disorders. Parkinsonism and Related Disorders, 2018, 46, S24-S29.	2.2	34
25	Subthalamic nucleus stimulation and gait in Parkinson's Disease: a not always fruitful relationship. Gait and Posture, 2017, 52, 205-210.	1.4	33
26	Clinical Phenotypes of Parkinson's Disease Associate with Distinct Gut Microbiota and Metabolome Enterotypes. Biomolecules, 2021, 11, 144.	4.0	33
27	Familial psychogenic movement disorders. Movement Disorders, 2013, 28, 1295-1298.	3.9	31
28	Prevalence of primary blepharospasm in Sardinia, Italy: A serviceâ€based survey. Movement Disorders, 2006, 21, 2005-2008.	3.9	30
29	"Smelling and Tasting―Parkinson's Disease: Using Senses to Improve the Knowledge of the Disease. Frontiers in Aging Neuroscience, 2020, 12, 43.	3.4	30
30	Genetic, clinical, and imaging characterization of one patient with late-onset, slowly progressive, pantothenate kinase-associated neurodegeneration. Movement Disorders, 2006, 21, 417-418.	3.9	28
31	6â€ <i>n</i> â€propylthiouracil taste disruption and <i>TAS2R38</i> nontasting form in Parkinson's disease. Movement Disorders, 2018, 33, 1331-1339.	3.9	28
32	Levodopa–carbidopa intrajejunal gel in advanced Parkinson disease with "on―freezing of gait. Neurological Sciences, 2015, 36, 1683-1686.	1.9	25
33	Parkinson's Disease Symptoms Have a Distinct Impact on Caregivers' and Patients' Stress: A Study Assessing the Consequences of the COVIDâ€19 Lockdown. Movement Disorders Clinical Practice, 2020, 7, 865-867.	1.5	25
34	Levodopa and neuropathy risk in patients with Parkinson disease: Effect of COMT inhibition. Parkinsonism and Related Disorders, 2016, 27, 81-84.	2.2	24
35	Odor Identification Performance in Idiopathic Parkinson's Disease Is Associated With Gender and the Genetic Variability of the Olfactory Binding Protein. Chemical Senses, 2019, 44, 311-318.	2.0	23
36	Genetic variants of TAS2R38 bitter taste receptor associate with distinct gut microbiota traits in Parkinson's disease: A pilot study. International Journal of Biological Macromolecules, 2020, 165, 665-674.	7.5	23

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37	GICYF2 mutations are not a frequent cause of familial Parkinson's disease. Parkinsonism and Related Disorders, 2009, 15, 703-705.	2.2	22
38	Idiopathic delayed-onset edema surrounding deep brain stimulation leads: Insights from a case series and systematic literature review. Parkinsonism and Related Disorders, 2016, 32, 108-115.	2.2	22
39	The peripheral nerve involvement in Parkinson Disease: A multifaceted phenomenon. Parkinsonism and Related Disorders, 2016, 25, 17-20.	2.2	22
40	Complex dyskinesias in Parkinson patients on levodopa/carbidopa intestinal gel. Parkinsonism and Related Disorders, 2019, 69, 140-146.	2.2	22
41	Demographic and clinical determinants of neck pain in idiopathic cervical dystonia. Journal of Neural Transmission, 2020, 127, 1435-1439.	2.8	22
42	LRRK2 mutations and Parkinson's disease in Sardinia—A Mediterranean genetic isolate. Parkinsonism and Related Disorders, 2007, 13, 17-21.	2.2	21
43	Quantitative assessment of gait parameters in people with Parkinson's disease in laboratory and clinical setting: Are the measures interchangeable?. Neurology International, 2018, 10, 7729.	2.8	21
44	ldiopathic <scp>Nonâ€ŧaskâ€6pecific</scp> Upper Limb Dystonia, a Neglected Form of Dystonia. Movement Disorders, 2020, 35, 2038-2045.	3.9	21
45	Worldwide barriers to genetic testing for movement disorders. European Journal of Neurology, 2021, 28, 1901-1909.	3.3	21
46	An exome study of Parkinson's disease in Sardinia, a Mediterranean genetic isolate. Neurogenetics, 2015, 16, 55-64.	1.4	20
47	Gut microbiota and metabolome distinctive features in Parkinson disease: Focus on levodopa and levodopaâ€carbidopa intrajejunal gel. European Journal of Neurology, 2021, 28, 1198-1209.	3.3	20
48	Hallervorden Spatz syndrome (pantothenate kinase associated neurodegeneration) in two Sardinian brother with homozygous mutation in PANK 2 gene. Journal of Neurology, 2002, 249, 1599-1600.	3.6	19
49	Hyperkinetic Movement Disorder Emergencies. Current Neurology and Neuroscience Reports, 2017, 17, 6.	4.2	19
50	Does acute peripheral trauma contribute to idiopathic adult-onset dystonia?. Parkinsonism and Related Disorders, 2020, 71, 40-43.	2.2	18
51	Persistent abnormal shoulder elevation after accessory nerve injury and differential diagnosis with post-traumatic focal shoulder-elevation dystonia: Report of a case and literature review. Movement Disorders, 2004, 19, 1109-1111.	3.9	16
52	Mutations inTMEM230are not a common cause of Parkinson's disease. Movement Disorders, 2017, 32, 302-304.	3.9	14
53	Efficacy and safety of deferiprone for the treatment of superficial siderosis: results from a long-term observational study. Neurological Sciences, 2019, 40, 1357-1361.	1.9	13
54	Association between Objectively Measured Physical Activity and Gait Patterns in People with Parkinson's Disease: Results from a 3-Month Monitoring. Parkinson's Disease, 2018, 2018, 1-10.	1.1	12

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55	The arginine growth hormone stimulation test in bradykineticâ€rigid parkinsonisms. Movement Disorders, 2008, 23, 190-194.	3.9	11
56	Parkinsonism and dementia are negative prognostic factors for the outcome of subdural hematoma. Neurological Sciences, 2016, 37, 1299-1303.	1.9	10
57	Quantitative assessment of upper limb functional impairments in people with Parkinson's disease. Clinical Biomechanics, 2018, 57, 137-143.	1.2	10
58	Levodopa–carbidopa intrajejunal infusion in Parkinson's disease: untangling the role of age. Journal of Neurology, 2021, 268, 1728-1737.	3.6	9
59	Correlation between cerebral perfusion and hyperventilation enhanced focal spiking activity. Epilepsy Research, 2000, 40, 79-86.	1.6	8
60	The TANDEM investigation: efficacy and tolerability of levodopa-carbidopa intestinal gel in (LCIG) advanced Parkinson's disease patients. Journal of Neural Transmission, 2020, 127, 881-891.	2.8	8
61	Spread of segmental/multifocal idiopathic adult-onset dystonia to a third body site. Parkinsonism and Related Disorders, 2021, 87, 70-74.	2.2	8
62	Digital work engagement among Italian neurologists. Therapeutic Advances in Chronic Disease, 2021, 12, 204062232110296.	2.5	7
63	Freezing of gait: overview on etiology, treatment, and future directions. Neurological Sciences, 2022, 43, 1627-1639.	1.9	7
64	Deep Brain Stimulation Emergencies: How the New Technologies Could Modify the Current Scenario. Current Neurology and Neuroscience Reports, 2017, 17, 51.	4.2	6
65	Motor and Sensory Features of Cervical Dystonia Subtypes: Data From the Italian Dystonia Registry. Frontiers in Neurology, 2020, 11, 906.	2.4	6
66	Pisa syndrome as presenting symptom of amyotrophic lateral sclerosis. Journal of Neurology, 2011, 258, 2087-2089.	3.6	5
67	Similarities and Differences of Gait Patterns in Women and Men With Parkinson Disease With Mild Disability. Archives of Physical Medicine and Rehabilitation, 2019, 100, 2039-2045.	0.9	5
68	Percutaneous Endoscopic Transgastric Jejunostomy (PEG â€J) Tube Placement for Levodopaâ€Carbidopa Intrajejunal Gel Therapy in the Interventional Radiology Suite: A Longâ€term Followâ€up. Movement Disorders Clinical Practice, 2018, 5, 191-194.	1.5	4
69	BK-virus progressive multifocal leukoencephalitis in a patient with systemic lupus erythematosus. Neurological Sciences, 2018, 39, 1613-1615.	1.9	4
70	Genetic Creutzfeldt-Jakob disease in Sardinia: a case series linked to the PRNP R208H mutation due to a single founder effect. Neurogenetics, 2020, 21, 251-257.	1.4	4
71	Late Blink Reflex Changes in Patients with Pure Sensory Stroke Due to Geniculo-Thalamic Infarct: A Contribution to the Long Loop Theory. Journal of Clinical Neurophysiology, 2004, 21, 105-109.	1.7	3
72	Acute Stereotypic Behavior: Expanding the Spectrum of Movement Disorders Attributed to Vitamin B12 Deficiency. Movement Disorders Clinical Practice, 2020, 7, S63-S64.	1.5	2

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73	Widening the spectrum of secondary headache: intracranial hypotension following a non-aneurysmal subarachnoid hemorrhage. Neurological Sciences, 2019, 40, 2179-2181.	1.9	1
74	Italian survey on intraduodenal levodopa gel treatment in advanced Parkinson disease: State of the art 10 years after marketing. Parkinsonism and Related Disorders, 2016, 22, e97-e98.	2.2	0
75	Long term essential tremor recovery after stroke thalamotomy. Basal Ganglia, 2017, 9, 18-19.	0.3	Ο
76	The Long Way of a "Lost Pigtail†A Unique Complication of Jâ€Tube in Duodopa Therapy. Movement Disorders Clinical Practice, 2018, 5, 101-102.	1.5	0
77	Probable early Lyme neuroborreliosis in a non-endemic area: first reported case in Sardinia. Neurological Sciences, 2019, 40, 1741-1742.	1.9	0
78	Two cases of watershed-pattern reversible encephalopathy syndrome. Journal of the Neurological Sciences, 2021, 429, 118693.	0.6	0