Claudia Ciano

List of Publications by Year
in descending order

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77	2,428	172457	223800
papers	citations	29 h-index	46 g-index

77	77	77	3083
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Hypomyelinating leukodystrophies in adults: Clinical and genetic features. European Journal of Neurology, 2021, 28, 934-944.	3.3	14
2	A Case of Severe Early-Onset Neuropathy Caused by a Compound Heterozygous Deletion of the PMP22 Gene: Clinical and Neurographic Aspects. Neuropediatrics, 2020, 51, 173-177.	0.6	3
3	Expanding the phenotypic spectrum of TRIM2 â€associated Charcotâ€Marieâ€Tooth disease. Journal of the Peripheral Nervous System, 2020, 25, 429-432.	3.1	4
4	Hereditary gelsolin amyloidosis (<scp>HGA</scp>): a neglected cause of bilateral progressive or recurrent facial palsy. Journal of the Peripheral Nervous System, 2017, 22, 59-63.	3.1	8
5	A novel <scp>NDRG1</scp> mutation in a nonâ€Romani patient with <scp>CMT4D</scp>/<scp>HMSN</scp>â€Lom. Journal of the Peripheral Nervous System, 2017, 22, 47-50.	3.1	6
6	Amyotrophic lateral sclerosis causes small fiber pathology. European Journal of Neurology, 2016, 23, 416-420.	3.3	65
7	Screening for SH3TC2 gene mutations in a series of demyelinating recessive Charcotâ€Marieâ€Tooth disease (CMT4). Journal of the Peripheral Nervous System, 2016, 21, 142-149.	3.1	32
8	Nerve conduction velocity in <scp>CMT</scp>1A: what else can we tell?. European Journal of Neurology, 2016, 23, 1566-1571.	3.3	45
9	Monomelic amyotrophy in cervical myelopathy associated with anterior dural sac displacement induced by neck flexion. Journal of Neurology, 2016, 263, 823-825.	3.6	2
10	Hand function assessment in the first years of life in unilateral cerebral palsy: Correlation with neuroimaging and cortico-spinal reorganization. European Journal of Paediatric Neurology, 2016, 20, 114-124.	1.6	15
11	Mutational mechanisms in <scp>MFN2</scp>â€related neuropathy: compound heterozygosity for recessive and semidominant mutations. Journal of the Peripheral Nervous System, 2015, 20, 380-386.	3.1	23
12	Responsiveness of clinical outcome measures in Charcotâ€'Marieâ€'Tooth disease. European Journal of Neurology, 2015, 22, 1556-1563.	3.3	47
13	Childhood onset of acquired neuromyotonia: Association with a ganglioneuroma. Muscle and Nerve, 2015, 51, 620-621.	2.2	1
14	PMP22 messenger RNA levels in skin biopsies: testing the effectiveness of a Charcot-Marie-Tooth 1A biomarker. Brain, 2014, 137, 1614-1620.	7.6	33
15	Overlapping phenotypes in complex spastic paraplegias SPG11, SPG15, SPG35 and SPG48. Brain, 2014, 137, 1907-1920.	7.6	133
16	Xâ€linked Charcotâ€Marieâ€Tooth type 1: strokeâ€like presentation of a novel <scp>GJB1</scp> mutation. Journal of the Peripheral Nervous System, 2014, 19, 183-186.	3.1	20
17	Doubleâ€trouble in pediatric neurology: Myotonia congenita combined with charcotâ€“marieâ€“tooth disease type 1a. Muscle and Nerve, 2014, 50, 145-147.	2.2	6
18	Selected items from the Charcot-Marie-Tooth (CMT) Neuropathy Score and secondary clinical outcome measures serve as sensitive clinical markers of disease severity in CMT1A patients. Neuromuscular Disorders, 2014, 24, 1003-1017.	0.6	25

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19	Subclinical leukodystrophy and infertility in a man with a novel homozygous CLCN2 mutation. Neurology, 2014, 83, 1217-1218.	1.1	42
20	Supratentorial and pontine <scp>MRI</scp> abnormalities characterize recessive spastic ataxia of <scp>C</scp>harlevoixâ€<scp>S</scp>aguenay. A comprehensive study of an <scp>I</scp>talian series. European Journal of Neurology, 2013, 20, 138-146.	3.3	57
21	Giant SEPs and SEP-recovery function in Unverrichtâ€“Lundborg disease. Clinical Neurophysiology, 2013, 124, 1013-1018.	1.5	15
22	Coexistence of Charcotâ€Marieâ€Tooth disease type 1A and antiâ€MAG neuropathy. Journal of the Peripheral Nervous System, 2013, 18, 185-188.	3.1	1
23	Cortical myoclonus in childhood and juvenile onset Huntington's disease. Parkinsonism and Related Disorders, 2012, 18, 794-797.	2.2	22
24	Phenomenology of psychogenic movement disorders in children. Movement Disorders, 2012, 27, 1153-1157.	3.9	39
25	Lower limb areflexia without central and peripheral conduction abnormalities is highly suggestive of Gerstmannâ€“StrÃ¤usslerâ€“Scheinker disease Pro102Leu. Journal of the Neurological Sciences, 2011, 302, 85-88.	0.6	6
26	Co-occurrence of amyotrophic lateral sclerosis and Charcot-Marie-Tooth disease type 2A in a patient with a novel mutation in the mitofusin-2 gene. Neuromuscular Disorders, 2011, 21, 129-131.	0.6	19
27	Ascorbic acid in Charcotâ€“Marieâ€“Tooth disease type 1A (CMT-TRIAAL and CMT-TRAUK): a double-blind randomised trial. Lancet Neurology, The, 2011, 10, 320-328.	10.2	222
28	Characterization of severe action myoclonus in sialidoses. Epilepsy Research, 2011, 94, 86-93.	1.6	24
29	Short and long interval cortical inhibition in patients with Unverricht-Lundborg and Lafora body disease. Epilepsy Research, 2010, 89, 232-237.	1.6	31
30	Painful neuropathy in subclinical hypothyroidism: clinical and neuropathological recovery after hormone replacement therapy. Neurological Sciences, 2009, 30, 149-151.	1.9	44
31	Sleepâ€related tongue biting may not be a sign of epilepsy: A case of sleepâ€related faciomandibular myoclonus. Epilepsia, 2009, 50, 157-159.	5.1	11
32	Novel mutations in the GDAP1 gene in patients affected with early-onset axonal Charcot-Marie-Tooth type 4A. Neuromuscular Disorders, 2009, 19, 476-480.	0.6	23
33	Erythropoietin in amyotrophic lateral sclerosis: A pilot, randomized, double-blind, placebo-controlled study of safety and tolerability. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2009, 10, 410-415.	2.1	41
34	Immunotherapy responsive startle with antibodies to voltage gated potassium channels. BMJ Case Reports, 2009, 2009, bcr0920080988-bcr0920080988.	0.5	3
35	Myoclonusâ€“dystonia syndrome: Clinical presentation, disease course, and genetic features in 11 families. Movement Disorders, 2008, 23, 28-34.	3.9	75
36	The G389R mutation in the MAPT gene presenting as sporadic corticobasal syndrome. Movement Disorders, 2008, 23, 892-895.	3.9	43

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37	A neurophysiological study of myoclonus in patients with DYT11 myoclonusâ€dystonia syndrome. Movement Disorders, 2008, 23, 2041-2048.	3.9	43
38	Polymyography in the diagnosis of childhood onset movement disorders. European Journal of Paediatric Neurology, 2008, 12, 480-483.	1.6	11
39	Reliability of clinical outcome measures in Charcot-Marie-Tooth disease. Neuromuscular Disorders, 2008, 18, 19-26.	0.6	55
40	High-Frequency Rhythmic Cortical Myoclonus in a Long-Surviving Patient With Nonketotic Hypergylcemia. Journal of Child Neurology, 2008, 23, 321-324.	1.4	4
41	Immunotherapy responsive startle with antibodies to voltage gated potassium channels. Journal of Neurology, Neurosurgery and Psychiatry, 2007, 78, 1281-1290.	1.9	9
42	POEMS syndrome: relapse after successful autologous peripheral blood stem cell transplantation. Neuromuscular Disorders, 2007, 17, 980-982.	0.6	25
43	Celiac disease presenting with motor neuropathy: Effect of gluten free-diet. Muscle and Nerve, 2007, 35, 675-677.	2.2	27
44	Movement-related desynchronization-synchronization (ERD/ERS) in patients with Unverrichtâ€“Lundborg disease. NeuroImage, 2006, 33, 161-168.	4.2	26
45	A multicenter, randomized, double-blind, placebo-controlled trial of long-term ascorbic acid treatment in Charcot-Marie-Tooth disease type 1A (CMT-TRIAAL): The study protocol [EudraCT no.: 2006-000032-27]. Pharmacological Research, 2006, 54, 436-441.	7.1	47
46	Rhythmic cortical myoclonus in Niemann-Pick disease type C. Movement Disorders, 2006, 21, 1453-1456.	3.9	22
47	Childhood-onset multifocal motor neuropathy with conduction blocks. Neurology, 2006, 66, 922-924.	1.1	13
48	Propriospinal myoclonus with life threatening tonic spasms as paraneoplastic presentation of breast cancer. Journal of Neurology, Neurosurgery and Psychiatry, 2005, 77, 422-424.	1.9	15
49	Sequential antibodies to potassium channels and glutamic acid decarboxylase in neuromyotonia. Neurology, 2005, 64, 1290-1293.	1.1	30
50	Acute dyspnea due to right phrenic palsy during infusional chemotherapy. Annals of Oncology, 2004, 15, 691-692.	1.2	0
51	Abnormal lysosomal and ubiquitin-proteasome pathways in 19p13.3 distal myopathy. Annals of Neurology, 2004, 56, 133-138.	5.3	22
52	Exteroceptive reflexes in dystonia: A study of the recovery cycle of the R2 component of the blink reflex and of the exteroceptive suppression of the contracting sternocleidomastoid muscle in blepharospasm and torticollis. Movement Disorders, 2004, 9, 183-187.	3.9	35
53	Sensorimotor cortex excitability in Unverrichtâ€“Lundborg disease and Lafora body disease. Neurology, 2004, 63, 2309-2315.	1.1	50
54	Progressive dysarthria: definition and clinical follow-up. Neurological Sciences, 2003, 24, 211-212.	1.9	18

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55	Neurophysiological features in relation to clinical signs in clinically diagnosed corticobasal degeneration. Neurological Sciences, 2003, 24, 16-23.	1.9	26
56	Rhythmic cortical myoclonus in a case of HIV-related encephalopathy. Movement Disorders, 2003, 18, 1533-1538.	3.9	14
57	Movement-activated myoclonus in genetically defined progressive myoclonic epilepsies: EEGâ€“EMG relationship estimated using autoregressive models. Clinical Neurophysiology, 2003, 114, 1041-1052.	1.5	54
58	Does CMT1A homozygosity cause more severe disease with root hypertrophy and higher CSF proteins?. Neurology, 2003, 60, 1721-1722.	1.1	21
59	Myoclonic dystonia as unique presentation of isolated vitamin E deficiency in a young patient. Movement Disorders, 2002, 17, 612-614.	3.9	36
60	Chronic inflammatory demyelinating polyradiculoneuropathy: long-term course and treatment of 60 patients. Neurological Sciences, 2000, 21, 31-37.	1.9	62
61	Unexpected Right Phrenic Nerve Injury During 5-Fluorouracil Continuous Infusion Plus Cisplatin and Vinorelbine in Breast Cancer Patients. Journal of the National Cancer Institute, 2000, 92, 755-755.	6.3	8
62	Cranial nerve involvement in CMT disease type 1 due to early growth response 2 gene mutation. Neurology, 2000, 54, 1696-1698.	1.1	84
63	Heterozygous Null Mutation in the P 0 Gene Associated with Mild Charcot-Marie-Tooth Disease. Annals of the New York Academy of Sciences, 1999, 883, 477-480.	3.8	19
64	Infantile neuroaxonal dystrophy. Neurology, 1999, 52, 1472-1472.	1.1	108
65	Detection of hereditary neuropathy with liability to pressure palsies among patients with acute painless mononeuropathy or plexopathy. , 1998, 21, 1686-1691.		34
66	Myoclonus in corticobasal degeneration. Movement Disorders, 1997, 12, 598-603.	3.9	35
67	Phenotypic heterogeneity in hereditary neuropathy with liability to pressure palsies associated with chromosome 17p11.2-12 deletion. Neurology, 1996, 46, 1133-1137.	1.1	118
68	Unrecognized Charcot-Marie-Tooth Disease. Anesthesia and Analgesia, 1995, 81, 199-201.	2.2	2
69	Unrecognized Charcot-Marie-Tooth Disease. Anesthesia and Analgesia, 1995, 81, 199-201.	2.2	17
70	Intrathecal immune activation in three patients with progressive myoclonic ataxia. Movement Disorders, 1995, 10, 207-210.	3.9	2
71	Serial evoked potentials in multiple sclerosis bouts. Relation to steroid treatment. Italian Journal of Neurological Sciences, 1994, 15, 333-340.	0.1	19
72	Spasmodic dystonic laterocollis in familial cerebellar ataxia. Italian Journal of Neurological Sciences, 1994, 15, 245-248.	0.1	1

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73	Hypo- and Hypersensitivity to Vecuronium in a Patient with Guillain-Barr?? Syndromem. Anesthesia and Analgesia, 1994, 78, 187???189.	2.2	16
74	Adult onset myoclonic Huntington's disease. Movement Disorders, 1993, 8, 201-205.	3.9	42
75	Combined central and peripheral acute demyelination. Italian Journal of Neurological Sciences, 1993, 14, 83-86.	0.1	10
76	HMSN III phenotype due to homozygous expression of a dominant HMSN II gene. Neurology, 1992, 42, 2201-2201.	1.1	42
77	Peripheral nerve conduction velocity in normal infants and children. Italian Journal of Neurological Sciences, 1989, 10, 311-314.	0.1	6