

# Claudia Ciano

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1123589/publications.pdf>

Version: 2024-02-01

77  
papers

2,428  
citations

172457

29  
h-index

223800

46  
g-index

77  
all docs

77  
docs citations

77  
times ranked

3083  
citing authors

#	ARTICLE	IF	CITATIONS
1	Ascorbic acid in Charcot-Marie-Tooth disease type 1A (CMT-TRIAAL and CMT-TRAIUK): a double-blind randomised trial. <i>Lancet Neurology</i> , The, 2011, 10, 320-328.	10.2	222
2	Overlapping phenotypes in complex spastic paraplegias SPG11, SPG15, SPG35 and SPG48. <i>Brain</i> , 2014, 137, 1907-1920.	7.6	133
3	Phenotypic heterogeneity in hereditary neuropathy with liability to pressure palsies associated with chromosome 17p11.2-12 deletion. <i>Neurology</i> , 1996, 46, 1133-1137.	1.1	118
4	Infantile neuroaxonal dystrophy. <i>Neurology</i> , 1999, 52, 1472-1472.	1.1	108
5	Cranial nerve involvement in CMT disease type 1 due to early growth response 2 gene mutation. <i>Neurology</i> , 2000, 54, 1696-1698.	1.1	84
6	Myoclonus-dystonia syndrome: Clinical presentation, disease course, and genetic features in 11 families. <i>Movement Disorders</i> , 2008, 23, 28-34.	3.9	75
7	Amyotrophic lateral sclerosis causes small fiber pathology. <i>European Journal of Neurology</i> , 2016, 23, 416-420.	3.3	65
8	Chronic inflammatory demyelinating polyradiculoneuropathy: long-term course and treatment of 60 patients. <i>Neurological Sciences</i> , 2000, 21, 31-37.	1.9	62
9	Supratentorial and pontine MRI abnormalities characterize recessive spastic ataxia of Charlevoix-Saguenay. A comprehensive study of an Italian series. <i>European Journal of Neurology</i> , 2013, 20, 138-146.	3.3	57
10	Reliability of clinical outcome measures in Charcot-Marie-Tooth disease. <i>Neuromuscular Disorders</i> , 2008, 18, 19-26.	0.6	55
11	Movement-activated myoclonus in genetically defined progressive myoclonic epilepsies: EEG-EMG relationship estimated using autoregressive models. <i>Clinical Neurophysiology</i> , 2003, 114, 1041-1052.	1.5	54
12	Sensorimotor cortex excitability in Unverricht-Lundborg disease and Lafora body disease. <i>Neurology</i> , 2004, 63, 2309-2315.	1.1	50
13	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]. <i>Pharmacological Research</i> , 2006, 54, 436-441.	7.1	47
14	Responsiveness of clinical outcome measures in Charcot-Marie-Tooth disease. <i>European Journal of Neurology</i> , 2015, 22, 1556-1563.	3.3	47
15	Nerve conduction velocity in CMT1A: what else can we tell?. <i>European Journal of Neurology</i> , 2016, 23, 1566-1571.	3.3	45
16	Painful neuropathy in subclinical hypothyroidism: clinical and neuropathological recovery after hormone replacement therapy. <i>Neurological Sciences</i> , 2009, 30, 149-151.	1.9	44
17	The G389R mutation in the MAPT gene presenting as sporadic corticobasal syndrome. <i>Movement Disorders</i> , 2008, 23, 892-895.	3.9	43
18	A neurophysiological study of myoclonus in patients with DYT11 myoclonus-dystonia syndrome. <i>Movement Disorders</i> , 2008, 23, 2041-2048.	3.9	43

#	ARTICLE	IF	CITATIONS
19	Adult onset myoclonic Huntington's disease. <i>Movement Disorders</i> , 1993, 8, 201-205.	3.9	42
20	Subclinical leukodystrophy and infertility in a man with a novel homozygous <i>CLCN2</i> mutation. <i>Neurology</i> , 2014, 83, 1217-1218.	1.1	42
21	HMSN III phenotype due to homozygous expression of a dominant HMSN II gene. <i>Neurology</i> , 1992, 42, 2201-2201.	1.1	42
22	Erythropoietin in amyotrophic lateral sclerosis: A pilot, randomized, double-blind, placebo-controlled study of safety and tolerability. <i>Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders</i> , 2009, 10, 410-415.	2.1	41
23	Phenomenology of psychogenic movement disorders in children. <i>Movement Disorders</i> , 2012, 27, 1153-1157.	3.9	39
24	Myoclonic dystonia as unique presentation of isolated vitamin E deficiency in a young patient. <i>Movement Disorders</i> , 2002, 17, 612-614.	3.9	36
25	Myoclonus in corticobasal degeneration. <i>Movement Disorders</i> , 1997, 12, 598-603.	3.9	35
26	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. <i>Movement Disorders</i> , 2004, 9, 183-187.	3.9	35
27	Detection of hereditary neuropathy with liability to pressure palsies among patients with acute painless mononeuropathy or plexopathy. , 1998, 21, 1686-1691.		34
28	PMP22 messenger RNA levels in skin biopsies: testing the effectiveness of a Charcot-Marie-Tooth 1A biomarker. <i>Brain</i> , 2014, 137, 1614-1620.	7.6	33
29	Screening for <i>SH3TC2</i> gene mutations in a series of demyelinating recessive Charcot-Marie-Tooth disease (CMT4). <i>Journal of the Peripheral Nervous System</i> , 2016, 21, 142-149.	3.1	32
30	Short and long interval cortical inhibition in patients with Unverricht-Lundborg and Lafora body disease. <i>Epilepsy Research</i> , 2010, 89, 232-237.	1.6	31
31	Sequential antibodies to potassium channels and glutamic acid decarboxylase in neuromyotonia. <i>Neurology</i> , 2005, 64, 1290-1293.	1.1	30
32	Celiac disease presenting with motor neuropathy: Effect of gluten free-diet. <i>Muscle and Nerve</i> , 2007, 35, 675-677.	2.2	27
33	Neurophysiological features in relation to clinical signs in clinically diagnosed corticobasal degeneration. <i>Neurological Sciences</i> , 2003, 24, 16-23.	1.9	26
34	Movement-related desynchronization-synchronization (ERD/ERS) in patients with Unverricht-Lundborg disease. <i>NeuroImage</i> , 2006, 33, 161-168.	4.2	26
35	POEMS syndrome: relapse after successful autologous peripheral blood stem cell transplantation. <i>Neuromuscular Disorders</i> , 2007, 17, 980-982.	0.6	25
36	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. <i>Neuromuscular Disorders</i> , 2014, 24, 1003-1017.	0.6	25

#	ARTICLE	IF	CITATIONS
37	Characterization of severe action myoclonus in sialidoses. <i>Epilepsy Research</i> , 2011, 94, 86-93.	1.6	24
38	Novel mutations in the GDAP1 gene in patients affected with early-onset axonal Charcot-Marie-Tooth type 4A. <i>Neuromuscular Disorders</i> , 2009, 19, 476-480.	0.6	23
39	Mutational mechanisms in <i>MFN2</i> -related neuropathy: compound heterozygosity for recessive and semidominant mutations. <i>Journal of the Peripheral Nervous System</i> , 2015, 20, 380-386.	3.1	23
40	Abnormal lysosomal and ubiquitin-proteasome pathways in 19p13.3 distal myopathy. <i>Annals of Neurology</i> , 2004, 56, 133-138.	5.3	22
41	Rhythmic cortical myoclonus in Niemann-Pick disease type C. <i>Movement Disorders</i> , 2006, 21, 1453-1456.	3.9	22
42	Cortical myoclonus in childhood and juvenile onset Huntington's disease. <i>Parkinsonism and Related Disorders</i> , 2012, 18, 794-797.	2.2	22
43	Does CMT1A homozygosity cause more severe disease with root hypertrophy and higher CSF proteins?. <i>Neurology</i> , 2003, 60, 1721-1722.	1.1	21
44	X-linked Charcot-Marie-Tooth type 1: stroke-like presentation of a novel <i>GJB1</i> mutation. <i>Journal of the Peripheral Nervous System</i> , 2014, 19, 183-186.	3.1	20
45	Serial evoked potentials in multiple sclerosis bouts. Relation to steroid treatment. <i>Italian Journal of Neurological Sciences</i> , 1994, 15, 333-340.	0.1	19
46	Heterozygous Null Mutation in the P0 Gene Associated with Mild Charcot-Marie-Tooth Disease. <i>Annals of the New York Academy of Sciences</i> , 1999, 883, 477-480.	3.8	19
47	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. <i>Neuromuscular Disorders</i> , 2011, 21, 129-131.	0.6	19
48	Progressive dysarthria: definition and clinical follow-up. <i>Neurological Sciences</i> , 2003, 24, 211-212.	1.9	18
49	Unrecognized Charcot-Marie-Tooth Disease. <i>Anesthesia and Analgesia</i> , 1995, 81, 199-201.	2.2	17
50	Hypo- and Hypersensitivity to Vecuronium in a Patient with Guillain-Barré Syndrome. <i>Anesthesia and Analgesia</i> , 1994, 78, 187-189.	2.2	16
51	Propriospinal myoclonus with life threatening tonic spasms as paraneoplastic presentation of breast cancer. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2005, 77, 422-424.	1.9	15
52	Giant SEPs and SEP-recovery function in Unverricht-Lundborg disease. <i>Clinical Neurophysiology</i> , 2013, 124, 1013-1018.	1.5	15
53	Hand function assessment in the first years of life in unilateral cerebral palsy: Correlation with neuroimaging and cortico-spinal reorganization. <i>European Journal of Paediatric Neurology</i> , 2016, 20, 114-124.	1.6	15
54	Rhythmic cortical myoclonus in a case of HIV-related encephalopathy. <i>Movement Disorders</i> , 2003, 18, 1533-1538.	3.9	14

#	ARTICLE	IF	CITATIONS
55	Hypomyelinating leukodystrophies in adults: Clinical and genetic features. <i>European Journal of Neurology</i> , 2021, 28, 934-944.	3.3	14
56	Childhood-onset multifocal motor neuropathy with conduction blocks. <i>Neurology</i> , 2006, 66, 922-924.	1.1	13
57	Polymyography in the diagnosis of childhood onset movement disorders. <i>European Journal of Paediatric Neurology</i> , 2008, 12, 480-483.	1.6	11
58	Sleep-related tongue biting may not be a sign of epilepsy: A case of sleep-related faciomandibular myoclonus. <i>Epilepsia</i> , 2009, 50, 157-159.	5.1	11
59	Combined central and peripheral acute demyelination. <i>Italian Journal of Neurological Sciences</i> , 1993, 14, 83-86.	0.1	10
60	Immunotherapy responsive startle with antibodies to voltage gated potassium channels. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2007, 78, 1281-1290.	1.9	9
61	Unexpected Right Phrenic Nerve Injury During 5-Fluorouracil Continuous Infusion Plus Cisplatin and Vinorelbine in Breast Cancer Patients. <i>Journal of the National Cancer Institute</i> , 2000, 92, 755-755.	6.3	8
62	Hereditary gelsolin amyloidosis (<sc>HGA</sc>): a neglected cause of bilateral progressive or recurrent facial palsy. <i>Journal of the Peripheral Nervous System</i> , 2017, 22, 59-63.	3.1	8
63	Peripheral nerve conduction velocity in normal infants and children. <i>Italian Journal of Neurological Sciences</i> , 1989, 10, 311-314.	0.1	6
64	Lower limb areflexia without central and peripheral conduction abnormalities is highly suggestive of Gerstmann-Str�ussler-Scheinker disease Pro102Leu. <i>Journal of the Neurological Sciences</i> , 2011, 302, 85-88.	0.6	6
65	Double-trouble in pediatric neurology: Myotonia congenita combined with charcot-marie-tooth disease type 1a. <i>Muscle and Nerve</i> , 2014, 50, 145-147.	2.2	6
66	A novel <i><sc>NDRG1</sc></i> mutation in a non-Romani patient with <sc>CMT4D</sc>/<sc>HMSN</sc>-Lom. <i>Journal of the Peripheral Nervous System</i> , 2017, 22, 47-50.	3.1	6
67	High-Frequency Rhythmic Cortical Myoclonus in a Long-Surviving Patient With Nonketotic Hyperglycemia. <i>Journal of Child Neurology</i> , 2008, 23, 321-324.	1.4	4
68	Expanding the phenotypic spectrum of TRIM2-associated Charcot-Marie-Tooth disease. <i>Journal of the Peripheral Nervous System</i> , 2020, 25, 429-432.	3.1	4
69	A Case of Severe Early-Onset Neuropathy Caused by a Compound Heterozygous Deletion of the PMP22 Gene: Clinical and Neurographic Aspects. <i>Neuropediatrics</i> , 2020, 51, 173-177.	0.6	3
70	Immunotherapy responsive startle with antibodies to voltage gated potassium channels. <i>BMJ Case Reports</i> , 2009, 2009, bcr0920080988-bcr0920080988.	0.5	3
71	Unrecognized Charcot-Marie-Tooth Disease. <i>Anesthesia and Analgesia</i> , 1995, 81, 199-201.	2.2	2
72	Intrathecal immune activation in three patients with progressive myoclonic ataxia. <i>Movement Disorders</i> , 1995, 10, 207-210.	3.9	2

#	ARTICLE	IF	CITATIONS
73	Monomelic amyotrophy in cervical myelopathy associated with anterior dural sac displacement induced by neck flexion. <i>Journal of Neurology</i> , 2016, 263, 823-825.	3.6	2
74	Spasmodic dystonic laterocollis in familial cerebellar ataxia. <i>Italian Journal of Neurological Sciences</i> , 1994, 15, 245-248.	0.1	1
75	Coexistence of Charcotâ€Marieâ€Tooth disease type 1A and antiâ€MAG neuropathy. <i>Journal of the Peripheral Nervous System</i> , 2013, 18, 185-188.	3.1	1
76	Childhood onset of acquired neuromyotonia: Association with a ganglioneuroma. <i>Muscle and Nerve</i> , 2015, 51, 620-621.	2.2	1
77	Acute dyspnea due to right phrenic palsy during infusional chemotherapy. <i>Annals of Oncology</i> , 2004, 15, 691-692.	1.2	0