

Chiara Pisciotta

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54
papers

1,166
citations

19
h-index

32
g-index

58
ext. papers

1,498
ext. citations

5.6
avg, IF

4.1
L-index

#	Paper	IF	Citations
54	Ascorbic acid in Charcot-Marie-Tooth disease type 1A (CMT-TRIAAL and CMT-TRAUK): a double-blind randomised trial. <i>Lancet Neurology, The</i> , 2011 , 10, 320-8	24.1	184
53	New developments in Charcot-Marie-Tooth neuropathy and related diseases. <i>Current Opinion in Neurology</i> , 2017 , 30, 471-480	7.1	100
52	Functional involvement of central cholinergic circuits and visual hallucinations in Parkinson's disease. <i>Brain</i> , 2009 , 132, 2350-5	11.2	92
51	Novel ATP13A2 (PARK9) homozygous mutation in a family with marked phenotype variability. <i>Neurogenetics</i> , 2011 , 12, 33-9	3	72
50	Charcot-Marie-Tooth disease: frequency of genetic subtypes in a Southern Italy population. <i>Journal of the Peripheral Nervous System</i> , 2014 , 19, 292-8	4.7	50
49	Biallelic mutations in SORD cause a common and potentially treatable hereditary neuropathy with implications for diabetes. <i>Nature Genetics</i> , 2020 , 52, 473-481	36.3	38
48	Nerve conduction velocity in CMT1A: what else can we tell?. <i>European Journal of Neurology</i> , 2016 , 23, 1566-71	6	30
47	Neuropathy. <i>Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn</i> , 2018 , 148, 653-665	3	29
46	Electrophysiological characterisation in hereditary spastic paraplegia type 5. <i>Clinical Neurophysiology</i> , 2011 , 122, 819-22	4.3	28
45	Small-fiber involvement in spinobulbar muscular atrophy (Kennedy's disease). <i>Muscle and Nerve</i> , 2007 , 36, 816-20	3.4	27
44	Cross-sectional analysis of a large cohort with X-linked Charcot-Marie-Tooth disease (CMTX1). <i>Neurology</i> , 2017 , 89, 927-935	6.5	24
43	Novel mutations in provide clues to the pathomechanisms of HSAN-VI. <i>Neurology</i> , 2017 , 88, 2132-2140	6.5	23
42	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, 1003-17	2.9	22
41	Different nerve ultrasound patterns in charcot-marie-tooth types and hereditary neuropathy with liability to pressure palsies. <i>Muscle and Nerve</i> , 2018 , 57, E18-E23	3.4	22
40	Small nerve fiber involvement in CMT1A. <i>Neurology</i> , 2015 , 84, 407-14	6.5	21
39	Charcot-Marie-Tooth disease: New insights from skin biopsy. <i>Neurology</i> , 2015 , 85, 1202-8	6.5	20
38	Short-latency afferent inhibition in patients with Parkinson's disease and freezing of gait. <i>Journal of Neural Transmission</i> , 2015 , 122, 1533-40	4.3	20

37	Expanding the spectrum of genes responsible for hereditary motor neuropathies. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2019 , 90, 1171-1179	5.5	19
36	Altered TDP-43-dependent splicing in HSPB8-related distal hereditary motor neuropathy and myofibrillar myopathy. <i>European Journal of Neurology</i> , 2018 , 25, 154-163	6	19
35	A new Italian FHM2 family: clinical aspects and functional analysis of the disease-associated mutation. <i>Cephalalgia</i> , 2011 , 31, 808-19	6.1	19
34	Somatosensory temporal discrimination threshold is increased in patients with cerebellar atrophy. <i>Cerebellum</i> , 2013 , 12, 456-9	4.3	18
33	Electrophysiological characterization of adult-onset Niemann-Pick type C disease. <i>Journal of the Neurological Sciences</i> , 2015 , 348, 262-5	3.2	18
32	Novel outcome measures for Charcot-Marie-Tooth disease: validation and reliability of the 6-min walk test and StepWatch(®) Activity Monitor and identification of the walking features related to higher quality of life. <i>European Journal of Neurology</i> , 2016 , 23, 1343-50	6	18
31	Absence of Dystrophin Related Protein-2 disrupts Cajal bands in a patient with Charcot-Marie-Tooth disease. <i>Neuromuscular Disorders</i> , 2015 , 25, 786-93	2.9	17
30	Reduced neurofilament expression in cutaneous nerve fibers of patients with CMT2E. <i>Neurology</i> , 2015 , 85, 228-34	6.5	16
29	A longitudinal study of CMT1A using Rasch analysis based CMT neuropathy and examination scores. <i>Neurology</i> , 2020 , 94, e884-e896	6.5	15
28	Postural instability in Charcot-Marie-Tooth 1A disease. <i>Gait and Posture</i> , 2016 , 49, 353-357	2.6	15
27	Influence of comorbidities on the phenotype of patients affected by Charcot-Marie-Tooth neuropathy type 1A. <i>Neuromuscular Disorders</i> , 2013 , 23, 902-6	2.9	14
26	Autoimmune autonomic ganglionopathy: a possible postganglionic neuropathy. <i>Archives of Neurology</i> , 2011 , 68, 504-7		14
25	A multicenter retrospective study of charcot-marie-tooth disease type 4B (CMT4B) associated with mutations in myotubularin-related proteins (MTMRs). <i>Annals of Neurology</i> , 2019 , 86, 55-67	9.4	13
24	GDAP1 mutations in Italian axonal Charcot-Marie-Tooth patients: Phenotypic features and clinical course. <i>Neuromuscular Disorders</i> , 2016 , 26, 26-32	2.9	13
23	Hirayama's disease: an Italian single center experience and review of the literature. <i>Quantitative Imaging in Medicine and Surgery</i> , 2016 , 6, 364-373	3.6	13
22	Autonomic nervous system involvement in a new CMT2B family. <i>Journal of the Peripheral Nervous System</i> , 2012 , 17, 361-4	4.7	12
21	Motor performance deterioration accelerates after 50 years of age in Charcot-Marie-Tooth type 1A patients. <i>European Journal of Neurology</i> , 2018 , 25, 301-306	6	11
20	Early onset Charcot-Marie-Tooth neuropathy type 2A and severe developmental delay: expanding the clinical phenotype of MFN2-related neuropathy. <i>Journal of the Peripheral Nervous System</i> , 2015 , 20, 415-8	4.7	11

19	A novel autosomal dominant GDAP1 mutation in an Italian CMT2 family. <i>Journal of the Peripheral Nervous System</i> , 2012 , 17, 351-5	4.7	10
18	Case of acute motor conduction block neuropathy (AMCBN). <i>Muscle and Nerve</i> , 2009 , 39, 224-6	3.4	10
17	Electrophysiological comparison between males and females in HNPP. <i>Neurological Sciences</i> , 2013 , 34, 1429-32	3.5	8
16	Two families with novel PMP22 point mutations: genotype-phenotype correlation. <i>Journal of the Peripheral Nervous System</i> , 2009 , 14, 208-12	4.7	8
15	Natural history of Charcot-Marie-Tooth disease type 2A: a large international multicentre study. <i>Brain</i> , 2020 , 143, 3589-3602	11.2	8
14	Nine-year case history of monofocal motor neuropathy. <i>Muscle and Nerve</i> , 2008 , 38, 927-9	3.4	6
13	A novel family with axonal Charcot-Marie-Tooth disease caused by a mutation in the EGR2 gene. <i>Journal of the Peripheral Nervous System</i> , 2019 , 24, 219-223	4.7	5
12	Charcot-Marie-Tooth Type 2B: A New Phenotype Associated with a Novel Mutation and Inhibited EGFR Degradation. <i>Cells</i> , 2020 , 9,	7.9	5
11	PFN2 and GAMT as common molecular determinants of axonal Charcot-Marie-Tooth disease. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2018 , 89, 870-878	5.5	5
10	Myelin protein zero gene dose dependent axonal ion-channel dysfunction in a family with Charcot-Marie-Tooth disease. <i>Clinical Neurophysiology</i> , 2020 , 131, 2440-2451	4.3	4
9	Insights into the pathogenesis of ATP1A1-related CMT disease using patient-specific iPSCs. <i>Journal of the Peripheral Nervous System</i> , 2019 , 24, 330-339	4.7	3
8	Pregnancy in Charcot-Marie-Tooth disease: Data from the Italian CMT national registry. <i>Neurology</i> , 2020 , 95, e3180-e3189	6.5	3
7	Thermosensitive hereditary neuropathy with liability to pressure palsy. <i>Muscle and Nerve</i> , 2011 , 43, 448-9	3.4	2
6	Updated review of therapeutic strategies for Charcot-Marie-Tooth disease and related neuropathies. <i>Expert Review of Neurotherapeutics</i> , 2021 , 21, 701-713	4.3	2
5	Are novel outcome measures for Charcot-Marie-Tooth disease sensitive to change? The 6-minute walk test and StepWatch [®] Activity Monitor in a 12-month longitudinal study. <i>Neuromuscular Disorders</i> , 2019 , 29, 310-316	2.9	2
4	Validation of the Italian version of the Charcot-Marie-Tooth disease Pediatric Scale. <i>Journal of the Peripheral Nervous System</i> , 2020 , 25, 138-142	4.7	1
3	Validation of the Italian version of the Charcot-Marie-Tooth Health Index. <i>Journal of the Peripheral Nervous System</i> , 2020 , 25, 292-296	4.7	1
2	A case of congenital cataracts, facial dysmorphisms, neuropathy, and hyperkinetic movement disorder. <i>Movement Disorders</i> , 2013 , 28, 559-60	7	1

- 1 Challenges in Treating Charcot-Marie-Tooth Disease and Related Neuropathies: Current Management and Future Perspectives. *Brain Sciences*, **2021**, 11,

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