Fiore Manganelli

List of Publications by Year
in descending order

169 papers	3,943 citations	136950 32 h-index	189892 50 g-index
176 all docs	176 docs citations	176 times ranked	5518 citing authors

#	Article	IF	Citations
1	Ross syndrome: a rare or a misknown disorder of thermoregulation? A skin innervation study on 12 subjects. Brain, 2006, 129, 2119-2131.	7.6	123
2	Antibodies to neurofascin, contactin-1, and contactin-associated protein 1 in CIDP. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, .	6.0	118
3	Functional involvement of central cholinergic circuits and visual hallucinations in Parkinson's disease. Brain, 2009, 132, 2350-2355.	7.6	115
4	Neuropathy and levodopa in Parkinson's disease: Evidence from a multicenter study. Movement Disorders, 2013, 28, 1391-1397.	3.9	114
5	Atypical CIDP: diagnostic criteria, progression and treatment response. Data from the Italian CIDP Database. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 125-132.	1.9	108
6	Trigeminal Stimulation Elicits a Peripheral Vestibular Imbalance in Migraine Patients. Headache, 2005, 45, 325-331.	3.9	101
7	Biallelic mutations in SORD cause a common and potentially treatable hereditary neuropathy with implications for diabetes. Nature Genetics, 2020, 52, 473-481.	21.4	97
8	Novel ATP13A2 (PARK9) homozygous mutation in a family with marked phenotype variability. Neurogenetics, 2011, 12, 33-39.	1.4	84
9	Monitoring effectiveness and safety of Tafamidis in transthyretin amyloidosis in Italy: a longitudinal multicenter study in a non-endemic area. Journal of Neurology, 2016, 263, 916-924.	3.6	76
10	The effects of prolonged cathodal direct current stimulation on the excitatory and inhibitory circuits of the ipsilateral and contralateral motor cortex. Journal of Neural Transmission, 2012, 119, 1499-1506.	2.8	71
11	Levetiracetam in patients with cortical myoclonus: A clinical and electrophysiological study. Movement Disorders, 2005, 20, 1610-1614.	3.9	66
12	Loss of cutaneous large and small fibers in naive and <scp>l</scp> -dopaâ€“treated PD patients. Neurology, 2017, 89, 776-784.	1.1	66
13	Prevalence and characteristics of peripheral neuropathy in hepatitis C virus population. Journal of Neurology, Neurosurgery and Psychiatry, 2006, 77, 626-629.	1.9	65
14	Charcotâ€Marieâ€Tooth disease: frequency of genetic subtypes in a Southern Italy population. Journal of the Peripheral Nervous System, 2014, 19, 292-298.	3.1	64
15	RFC1 expansions are a common cause of idiopathic sensory neuropathy. Brain, 2021, 144, 1542-1550.	7.6	63
16	Mutations in ATP1A1 Cause Dominant Charcot-Marie-Tooth Type 2. American Journal of Human Genetics, 2018, 102, 505-514.	6.2	59
17	Post-exercise facilitation and depression of motor evoked potentials to transcranial magnetic stimulation: a study in multiple sclerosis. Clinical Neurophysiology, 2004, 115, 2128-2133.	1.5	54
18	ATTRv amyloidosis Italian Registry: clinical and epidemiological data. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2020, 27, 259-265.	3.0	51

#	Article	IF	Citations
19	Brainstem involvement and respiratory failure in COVID-19. Neurological Sciences, 2020, 41, 1663-1665.	1.9	50
20	A cross-sectional study investigating frequency and features of definitely diagnosed diabetic painful polyneuropathy. Pain, 2018, 159, 2658-2666.	4.2	49
21	Responsiveness of clinical outcome measures in Charcotâ^'Marieâ^'Tooth disease. European Journal of Neurology, 2015, 22, 1556-1563.	3.3	47
22	The therapeutic use of non-invasive brain stimulation in multiple sclerosis â€“ a review. Restorative Neurology and Neuroscience, 2017, 35, 497-509.	0.7	46
23	Postganglionic sudomotor denervation in patients with multiple system atrophy. Neurology, 2014, 82, 2223-2229.	1.1	45
24	Nerve conduction velocity in <scp>CMT</scp>1A: what else can we tell?. European Journal of Neurology, 2016, 23, 1566-1571.	3.3	45
25	Nonâ€motor involvement in amyotrophic lateral sclerosis: new insight from nerve and vessel analysis in skin biopsy. Neuropathology and Applied Neurobiology, 2017, 43, 119-132.	3.2	45
26	Novel mutations in dystonin provide clues to the pathomechanisms of HSAN-VI. Neurology, 2017, 88, 2132-2140.	1.1	41
27	Influence of GAA expansion size and disease duration on central nervous system impairment in Friedreich's ataxia: contribution to the understanding of the pathophysiology of the disease. Clinical Neurophysiology, 2000, 111, 1023-1030.	1.5	40
28	Anodal transcranial direct current stimulation of motor cortex does not ameliorate spasticity in multiple sclerosis. Restorative Neurology and Neuroscience, 2015, 33, 487-492.	0.7	39
29	Hereditary transthyretin amyloidosis overview. Neurological Sciences, 2022, 43, 595-604.	1.9	39
30	The Effect of Cerebellar Degeneration on Human Sensori-motor Plasticity. Brain Stimulation, 2015, 8, 1144-1150.	1.6	37
31	Motor cortex cholinergic dysfunction in CADASIL: A transcranial magnetic demonstration. Clinical Neurophysiology, 2008, 119, 351-355.	1.5	35
32	Clinical features and molecular modelling of novel MPZ mutations in demyelinating and axonal neuropathies. European Journal of Human Genetics, 2009, 17, 1129-1134.	2.8	35
33	A multicenter retrospective study of charcotâ€marieâ€tooth disease type 4B (CMT4B) associated with mutations in myotubularinâ€related proteins (MTMRs). Annals of Neurology, 2019, 86, 55-67.	5.3	35
34	Subclinical neurological involvement does not develop if Wilson's disease is treated early. Parkinsonism and Related Disorders, 2016, 24, 15-19.	2.2	34
35	Fast Intracortical Sensory-Motor Integration: A Window Into the Pathophysiology of Parkinsonâ€™s Disease. Frontiers in Human Neuroscience, 2019, 13, 111.	2.0	34
36	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

#	Article	IF	Citations
37	Charcot-Marie-Tooth disease. Neurology, 2015, 85, 1202-1208.	1.1	33
38	Charcotâ€“Marieâ€“Tooth disease type 2C: a distinct genetic entity. Clinical and molecular characterization of the first European family. Neuromuscular Disorders, 2002, 12, 399-404.	0.6	31
39	Smallâ€fiber involvement in spinobulbar muscular atrophy (Kennedy's disease). Muscle and Nerve, 2007, 36, 816-820.	2.2	31
40	Electrophysiological characterisation in hereditary spastic paraplegia type 5. Clinical Neurophysiology, 2011, 122, 819-822.	1.5	31
41	Small nerve fiber involvement in CMT1A. Neurology, 2015, 84, 407-414.	1.1	30
42	Expanding the spectrum of genes responsible for hereditary motor neuropathies. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 1171-1179.	1.9	30
43	<scp>The neuropathy in hereditary transthyretin amyloidosis</scp>: A <scp>narrative review</scp>. Journal of the Peripheral Nervous System, 2021, 26, 155-159.	3.1	30
44	Different nerve ultrasound patterns in charcotâ€marieâ€tooth types and hereditary neuropathy with liability to pressure palsies. Muscle and Nerve, 2018, 57, E18-E23.	2.2	28
45	Increased peptidergic fibers as a potential cutaneous marker of pain in diabetic small fiber neuropathy. Pain, 2021, 162, 778-786.	4.2	28
46	A new Italian FHM2 family: Clinical aspects and functional analysis of the disease-associated mutation. Cephalalgia, 2011, 31, 808-819.	3.9	27
47	Executive functions are impaired in heterozygote patients with oculopharyngeal muscular dystrophy. Journal of Neurology, 2012, 259, 833-837.	3.6	27
48	Alterations of autophagy in the peripheral neuropathy Charcot-Marie-Tooth type 2B. Autophagy, 2018, 14, 1-12.	9.1	27
49	Risk factors for chronic inflammatory demyelinating polyradiculoneuropathy (CIDP): antecedent events, lifestyle and dietary habits. Data from the Italian CIDP Database. European Journal of Neurology, 2020, 27, 136-143.	3.3	27
50	Alteration of the late endocytic pathway in Charcotâ€“Marieâ€“Tooth type 2B disease. Cellular and Molecular Life Sciences, 2021, 78, 351-372.	5.4	27
51	Electrophysiologic characterization in spinocerebellar ataxia 17. Neurology, 2006, 66, 932-934.	1.1	26
52	Novel outcome measures for Charcotâ€'Marieâ€'Tooth disease: validation and reliability of the 6â€min walk test and StepWatch^{â„¢} Activity Monitor and identification of the walking features related to higher quality of life. European Journal of Neurology, 2016, 23, 1343-1350.	3.3	26
53	Multimodal evoked potentials follow up in multiple sclerosis patients under fingolimod therapy. Journal of the Neurological Sciences, 2016, 365, 143-146.	0.6	26
54	Impulse control disorders induced by rasagiline as adjunctive therapy for Parkinson's disease: Report of 2 cases. Parkinsonism and Related Disorders, 2013, 19, 483-484.	2.2	25

#	Article	IF	Citations
55	Phenotypic Variability Among Patients With D4Z4 Reduced Allele Facioscapulohumeral Muscular Dystrophy. JAMA Network Open, 2020, 3, e204040.	5.9	25
56	Central cholinergic dysfunction in the adult form of Niemann Pick disease type C: a further link with Alzheimerâ€™s disease?. Journal of Neurology, 2014, 261, 804-808.	3.6	24
57	A new POLG1 mutation with peo and severe axonal and demyelinating sensoryâ€“motor neuropathy. Journal of Neurology, 2006, 253, 869-874.	3.6	23
58	Anti-GAD antibody ocular flutter: expanding the spectrum of autoimmune ocular motor disorders. Journal of Neurology, 2013, 260, 2675-2677.	3.6	23
59	How to manage with telemedicine people with neuromuscular diseases?. Neurological Sciences, 2021, 42, 3553-3559.	1.9	23
60	Short-latency afferent inhibition in patients with Parkinsonâ€™s disease and freezing of gait. Journal of Neural Transmission, 2015, 122, 1533-1540.	2.8	22
61	Electrophysiological characterization of adult-onset Niemannâ€“Pick type C disease. Journal of the Neurological Sciences, 2015, 348, 262-265.	0.6	22
62	Recommendations for pre-symptomatic genetic testing for hereditary transthyretin amyloidosis in the era of effective therapy: a multicenter Italian consensus. Orphanet Journal of Rare Diseases, 2020, 15, 348.	2.7	22
63	Sural nerve and epidermal vascular abnormalities in a case of POEMS syndrome. European Journal of Neurology, 2006, 13, 99-102.	3.3	20
64	Autonomic nervous system involvement in a new CMT2B family. Journal of the Peripheral Nervous System, 2012, 17, 361-364.	3.1	20
65	Postural instability in Charcot-Marie-Tooth 1A disease. Gait and Posture, 2016, 49, 353-357.	1.4	20
66	Sensitivity and specificity of a commercial ELISA test for anti-MAG antibodies in patients with neuropathy. Journal of Neuroimmunology, 2020, 345, 577288.	2.3	20
67	CSF sphingomyelin: a new biomarker of demyelination in the diagnosis and management of CIDP and GBS. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 303-310.	1.9	20
68	Hirayamaâ€™s disease: an Italian single center experience and review of the literature. Quantitative Imaging in Medicine and Surgery, 2016, 6, 364-373.	2.0	20
69	Multimodal electrophysiologic follow-up study in 3 mutated but presymptomatic members of a spinocerebellar ataxia type 1 (SCA1) family. Neurological Sciences, 2005, 26, 67-71.	1.9	19
70	Autoimmune Autonomic Ganglionopathy. Archives of Neurology, 2011, 68, 504.	4.5	19
71	Somatosensory Temporal Discrimination Threshold Is Increased in Patients with Cerebellar Atrophy. Cerebellum, 2013, 12, 456-459.	2.5	19
72	Atypical clinical and radiological presentation of cryptococcal choroid plexitis in an immunocompetent woman. Journal of the Neurological Sciences, 2013, 334, 180-182.	0.6	19

#	Article	IF	Citations
73	Cognitive correlates of prospective memory in dystonia. Parkinsonism and Related Disorders, 2019, 66, 51-55.	2.2	19
74	Small fiber pathology parallels disease progression in Parkinson disease: a longitudinal study. Acta Neuropathologica, 2018, 136, 501-503.	7.7	19
75	Distal hypoglycemic neuropathy. An insulinoma-associated case, misdiagnosed as temporal lobe epilepsy. Neurophysiologie Clinique, 2003, 33, 223-227.	2.2	18
76	GDAP1 mutations in Italian axonal Charcotâ€“Marieâ€“Tooth patients: Phenotypic features and clinical course. Neuromuscular Disorders, 2016, 26, 26-32.	0.6	18
77	Does acute peripheral trauma contribute to idiopathic adult-onset dystonia?. Parkinsonism and Related Disorders, 2020, 71, 40-43.	2.2	18
78	Neurophysiological evaluation of motor corticospinal pathways by TMS in idiopathic early-onset Parkinsonâ€™s disease. Clinical Neurophysiology, 2011, 122, 546-549.	1.5	17
79	Cognitive profile and 18F-fluorodeoxyglucose PET study in LRRK2 -related Parkinson's disease. Parkinsonism and Related Disorders, 2018, 47, 80-83.	2.2	17
80	Muscle pain syndromes and fibromyalgia: the role of muscle biopsy. Current Opinion in Supportive and Palliative Care, 2018, 12, 382-387.	1.3	17
81	Dealing with immune-mediated neuropathies during COVID-19 outbreak: practical recommendations from the task force of the Italian Society of Neurology (SIN), the Italian Society of Clinical Neurophysiology (SINC) and the Italian Peripheral Nervous System Association (ASNP). Neurological Sciences, 2020, 41, 1345-1348.	1.9	17
82	Inherited neuromyotonia: A clinical and genetic study of a family. Neuromuscular Disorders, 2007, 17, 23-27.	0.6	16
83	A rare mutation in MYH7 gene occurs with overlapping phenotype. Biochemical and Biophysical Research Communications, 2015, 457, 262-266.	2.1	16
84	Upper motor neuron evaluation in multiple sclerosis patients treated with Sativex^{Â®}. Acta Neurologica Scandinavica, 2017, 135, 442-448.	2.1	16
85	Motor performance deterioration accelerates after 50 years of age in Charcotâ€Marieâ€Tooth type 1A patients. European Journal of Neurology, 2018, 25, 301-306.	3.3	16
86	Familial aggregation of white matter lesions in myotonic dystrophy type 1. Neuromuscular Disorders, 2008, 18, 299-305.	0.6	15
87	Case of acute motor conduction block neuropathy (AMCBN). Muscle and Nerve, 2009, 39, 224-226.	2.2	15
88	Relevance of diagnostic investigations in chronic inflammatory demyelinating poliradiculoneuropathy: Data from the Italian CIDP database. Journal of the Peripheral Nervous System, 2020, 25, 152-161.	3.1	15
89	Chronic inflammatory demyelinating polyradiculoneuropathy: can a diagnosis be made in patients not fulfilling electrodiagnostic criteria?. European Journal of Neurology, 2021, 28, 620-629.	3.3	15
90	Acute and chronic inflammatory neuropathies and <scp>COVID</scp>â€19 vaccines: Practical recommendations from the task force of the Italian Peripheral Nervous System Association (<scp>ASNP</scp>). Journal of the Peripheral Nervous System, 2021, 26, 148-154.	3.1	15

#	Article	IF	Citations
91	Abnormal sensorimotor cortex and thalamo-cortical networks in familial adult myoclonic epilepsy type 2: pathophysiology and diagnostic implications. Brain Communications, 2022, 4, fcac037.	3.3	15
92	Early onset Charcotâ€Marieâ€Tooth neuropathy type <scp>2A</scp> and severe developmental delay: expanding the clinical phenotype of <scp>MFN2</scp>â€related neuropathy. Journal of the Peripheral Nervous System, 2015, 20, 415-418.	3.1	14
93	Spinocerebellar ataxia type 2â€”neuronopathy or neuropathy?. Muscle and Nerve, 2019, 60, 271-278.	2.2	14
94	A Novel CAPN1 Mutation Causes a Pure Hereditary Spastic Paraplegia in an Italian Family. Frontiers in Neurology, 2019, 10, 580.	2.4	14
95	Neurophysiological Signatures of Motor Impairment in Patients with Rett Syndrome. Annals of Neurology, 2020, 87, 763-773.	5.3	14
96	A novel autosomal dominant GDAP1 mutation in an Italian CMT2 family. Journal of the Peripheral Nervous System, 2012, 17, 351-355.	3.1	13
97	Cervical dystonia patients display subclinical gait changes. Parkinsonism and Related Disorders, 2017, 43, 97-100.	2.2	13
98	Telemedicine application to headache: a critical review. Neurological Sciences, 2022, 43, 3795-3801.	1.9	13
99	Realâ€life experience with inotersen in hereditary transthyretin amyloidosis with lateâ€onset phenotype: Data from an earlyâ€access program in Italy. European Journal of Neurology, 2022, 29, 2148-2155.	3.3	13
100	Role of <i>MAPT</i> in Pure Motor Neuron Disease: Report of a Recurrent Mutation in Italian Patients. Neurodegenerative Diseases, 2018, 18, 310-314.	1.4	12
101	An altered lipid metabolism characterizes Charcot-Marie-Tooth type 2B peripheral neuropathy. Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids, 2020, 1865, 158805.	2.4	12
102	Normalization of timed neuropsychological tests with the <scp>PATA</scp> rate and nineâ€hole pegboard tests. Journal of Neuropsychology, 2018, 12, 471-483.	1.4	12
103	Cardiovascular Involvement in Transthyretin Cardiac Amyloidosis. Heart Failure Clinics, 2022, 18, 73-87.	2.1	12
104	Early predictive factors of disability in CIDP. Journal of Neurology, 2017, 264, 1939-1944.	3.6	11
105	One-year follow up of three Italian patients with Duchenne muscular dystrophy treated with ataluren: is earlier better?. Therapeutic Advances in Neurological Disorders, 2018, 11, 175628641880958.	3.5	11
106	Six-minute walk test is reliable and sensitive in detecting response to therapy in CIDP. Journal of Neurology, 2019, 266, 860-865.	3.6	11
107	The flavor test is a sensitive tool in identifying the flavor sensorineural dysfunction in Parkinsonâ€™s disease. Neurological Sciences, 2019, 40, 1351-1356.	1.9	11
108	Pregnancy in Charcot-Marie-Tooth disease. Neurology, 2020, 95, e3180-e3189.	1.1	11

#	Article	IF	Citations
109	Electrophysiological comparison between males and females in HNPP. Neurological Sciences, 2013, 34, 1429-1432.	1.9	10
110	Long-term therapy with miglustat and cognitive decline in the adult form of Niemann-Pick disease type C: a case report. Neurological Sciences, 2018, 39, 1015-1019.	1.9	10
111	A novel mutation of myelin protein zero associated with an axonal form of Charcot-Marie-Tooth disease. Journal of Neurology, Neurosurgery and Psychiatry, 2004, 75, 262-5.	1.9	10
112	GDAP1 mutation in autosomal recessive Charcot-Marie-Tooth with pyramidal features. Journal of Neurology, 2006, 253, 1234-1235.	3.6	9
113	Two families with novel PMP22 point mutations: genotypeâ€“phenotype correlation. Journal of the Peripheral Nervous System, 2009, 14, 208-212.	3.1	9
114	In vivo evidence of cortical amyloid deposition in the adult form of Niemann Pick type C. Heliyon, 2019, 5, e02776.	3.2	9
115	Frequency and clinical correlates of anti-nerve antibodies in a large population of CIDP patients included in the Italian database. Neurological Sciences, 2022, 43, 3939-3947.	1.9	9
116	A diagnostic score for antiâ€myelinâ€associatedâ€glycoprotein neuropathy or chronic inflammatory demyelinating polyradiculoneuropathy in patients with antiâ€myelinâ€associatedâ€glycoprotein antibody. European Journal of Neurology, 2023, 30, 501-510.	3.3	9
117	Nineâ€year case history of monofocal motor neuropathy. Muscle and Nerve, 2008, 38, 927-929.	2.2	8
118	Teaching Video Neuro Images : Clonus of the lower jaw. Neurology, 2014, 82, e96.	1.1	8
119	The Role of New Imaging Technologies in the Diagnosis of Cardiac Amyloidosis. Heart Failure Clinics, 2022, 18, 61-72.	2.1	8
120	Small fiber neuropathy in the chronic phase of Chagas disease: a case report. Clinical Autonomic Research, 2013, 23, 149-153.	2.5	7
121	Muscle fiber type disproportion (FTD) in a family with mutations in the LMNA gene. Muscle and Nerve, 2015, 51, 604-608.	2.2	7
122	Does motor cortex plasticity depend on the type of mutation in the leucine-rich repeat kinase 2 gene?. Movement Disorders, 2017, 32, 947-948.	3.9	7
123	Disruption of GABA(A)-mediated intracortical inhibition in patients with chorea-acanthocytosis. Neuroscience Letters, 2017, 654, 107-110.	2.1	7
124	Adult normative values for the PATA Rate Test. Journal of Neurology, 2018, 265, 1102-1105.	3.6	7
125	A novel family with axonal Charcotâ€Marieâ€Tooth disease caused by a mutation in the EGR2 gene. Journal of the Peripheral Nervous System, 2019, 24, 219-223.	3.1	7
126	Brain Plasticity in Charcot-Marie-Tooth Type 1A Patients? A Combined Structural and Diffusion MRI Study. Frontiers in Neurology, 2020, 11, 795.	2.4	7

#	Article	IF	Citations
127	Proximal weakness involvement in the first Italian case of Charcotâ€Marieâ€Tooth 2CC harboring a novel frameshift variant in NEFH. Journal of the Peripheral Nervous System, 2021, 26, 231-234.	3.1	7
128	Electrodiagnosis of Guillain-Barre syndrome in the International GBS Outcome Study: Differences in methods and reference values. Clinical Neurophysiology, 2022, 138, 231-240.	1.5	7
129	Neurophysiological and behavioural correlates of ocrelizumab therapy on manual dexterity in patients with primary progressive multiple sclerosis. Journal of Neurology, 2022, 269, 4791-4801.	3.6	7
130	Mitochondria dysfunction in Charcot Marie Tooth 2B Peripheral Sensory Neuropathy. Communications Biology, 2022, 5, .	4.4	7
131	Are novel outcome measures for Charcotâ€“Marieâ€“Tooth disease sensitive to change? The 6-minute walk test and StepWatchâ„¢ Activity Monitor in a 12-month longitudinal study. Neuromuscular Disorders, 2019, 29, 310-316.	0.6	6
132	The Treatment of Hypersalivation in Rett Syndrome with Botulinum Toxin: Efficacy and Clinical Implications. Neurology and Therapy, 2019, 8, 155-160.	3.2	6
133	Different cortical excitability profiles in hereditary brain iron and copper accumulation. Neurological Sciences, 2020, 41, 679-685.	1.9	6
134	Lichenoid rash: A new side effect of oral Cladribine. Multiple Sclerosis and Related Disorders, 2020, 41, 102023.	2.0	6
135	Distal motor neuropathy associated with novel EMILIN1 mutation. Neurobiology of Disease, 2020, 137, 104757.	4.4	6
136	BDNF polymorphism and interhemispheric balance of motor cortex excitability: a preliminary study. Journal of Neurophysiology, 2022, 127, 204-212.	1.8	6
137	Relationship between high-frequency activity in the cortical sensory and the motor hand areas, and their myelin content. Brain Stimulation, 2022, 15, 717-726.	1.6	6
138	Early changes of myocardial deformation properties in patients with dystrophia myotonica type 1: A three-dimensional Speckle Tracking echocardiographic study. International Journal of Cardiology, 2014, 176, 1094-1096.	1.7	5
139	Electrodiagnostic accuracy in polyneuropathies: supervised learning algorithms as a tool for practitioners. Neurological Sciences, 2020, 41, 3719-3727.	1.9	5
140	The second case of hereditary motor and sensory neuropathy with deafness, mental retardation and absence of large myelinated fibres, detected in the same geographic area as the first family. Journal of Neurology, 1998, 245, 240-244.	3.6	4
141	Clinical Utility of Electrophysiological Evaluation in Crigler-Najjar Syndrome. Neuropediatrics, 2007, 38, 173-178.	0.6	4
142	Isolated intracranial Mycobacterium avium complex granulomas in an immune-competent man. Journal of the Neurological Sciences, 2015, 349, 264-265.	0.6	4
143	Insights into the pathogenesis of ATP1A1 â€related CMT disease using patientâ€specific iPSCs. Journal of the Peripheral Nervous System, 2019, 24, 330-339.	3.1	4
144	A Model to Study Myelinated Fiber Degeneration and Regeneration in Human Skin. Annals of Neurology, 2020, 87, 456-465.	5.3	4

#	Article	IF	Citations
145	Multimodal evaluation of an Italian family with a hereditary spastic paraplegia and POLR3A mutations. Annals of Clinical and Translational Neurology, 2020, 7, 2326-2331.	3.7	4
146	Impact of environmental factors and physical activity on disability and quality of life in CIDP. Journal of Neurology, 2020, 267, 2683-2691.	3.6	4
147	Can we identify hereditary TTR amyloidosis by the screening of carpal tunnel syndrome patients?. Neurological Sciences, 2022, 43, 3435-3438.	1.9	4
148	Mutilating fingertip ulcers in uncontrolled type 1 diabetes mellitus. Neurological Sciences, 2014, 35, 123-124.	1.9	3
149	Seronegative occult HBV reactivation complicated with fulminant acute liver failure after rituximab for chronic inflammatory demyelinating polyneuropathy. Infectious Diseases, 2020, 52, 216-218.	2.8	3
150	Personality traits associated with blepharospasm: A comparison with healthy subjects, patients with facial hemispasm and patients with hyperhidrosis. Journal of Clinical Neuroscience, 2020, 74, 130-134.	1.5	3
151	Diffuse brain connectivity changes in Charcotâ€“Marieâ€“Tooth type 1a patients: a restingâ€state functional magnetic resonance imaging study. European Journal of Neurology, 2021, 28, 305-313.	3.3	3
152	The impact of symptoms on daily life as perceived by patients with Charcot-Marie-Tooth type 1A disease. Neurological Sciences, 2022, 43, 559-563.	1.9	3
153	The neurophysiological lesson from the Italian CIDP database. Neurological Sciences, 2021, , 1.	1.9	3
154	A compound score to screen patients with hereditary transthyretin amyloidosis. Journal of Neurology, 2022, , .	3.6	3
155	Dissective tandem stroke: an endovascular approach. Radiology Case Reports, 2022, 17, 2170-2174.	0.6	3
156	Thermosensitive hereditary neuropathy with liability to pressure palsy. Muscle and Nerve, 2011, 43, 448-449.	2.2	2
157	Sporadic chronic progressive external ophthalmoplegia with single large mitochondrial DNA deletion and neurogenic findings. Journal of Neurology, 2017, 264, 597-599.	3.6	2
158	A case report of limbic encephalitis in a metastatic colon cancer patient during first-line bevacizumab-combined chemotherapy. Medicine (United States), 2018, 97, e0011.	1.0	2
159	Autosomal-dominant transthyretin (TTR)-related amyloidosis is not a frequent CMT2 neuropathy â€œin disguiseâ€. Orphanet Journal of Rare Diseases, 2018, 13, 177.	2.7	2
160	Acute leukocytosis during alemtuzumab treatment in patients with active relapsing-remitting multiple sclerosis. Multiple Sclerosis and Related Disorders, 2019, 28, 98-100.	2.0	2
161	A case of congenital cataracts, facial dysmorphisms, neuropathy, and hyperkinetic movement disorder. Movement Disorders, 2013, 28, 559-560.	3.9	1
162	Chronic inflammatory demyelinating polyneuropathy mimicking an acute painful diabetic neuropathy. Neurological Sciences, 2015, 36, 1509-1510.	1.9	1

#	Article	IF	Citations
163	Pharmacological treatment for familial amyloid neuropathy. The Cochrane Library, 0, , .	2.8	1
164	Generalized anhidrosis as first clinical presentation of systemic lupus erythematosus. Lupus, 2018, 27, 2296-2297.	1.6	1
165	Prolonged distal motor latency of median nerve does not improve diagnostic accuracy for CIDP. Journal of Neurology, 2021, , 1.	3.6	1
166	Primary Progressive Multiple Sclerosis Under Anti-TNFÎ± Treatment: A Case Report. Journal of Central Nervous System Disease, 2020, 12, 117957352097382.	1.9	1
167	Bedside Head Impulse Test: A Useful Tool for Patients With Sensory Ataxia. Neurology: Genetics, 2021, 7, e541.	1.9	1
168	Poems Syndrome With Vasa Nervorum Vasculitis: A Case Report. Journal of the Peripheral Nervous System, 2001, 6, 42-42.	3.1	0
169	Pseudo-orthostatic tremor: description of a not typical case. Neurological Sciences, 2019, 40, 2205-2207.	1.9	0