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	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
2	Hereditary transthyretin amyloidosis overview. Neurological Sciences, 2022, 43, 595-604.	1.9	39
3	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
4	The Role of New Imaging Technologies in the Diagnosis of Cardiac Amyloidosis. Heart Failure Clinics, 2022, 18, 61-72.	2.1	8
5	Cardiovascular Involvement in Transthyretin Cardiac Amyloidosis. Heart Failure Clinics, 2022, 18, 73-87.	2.1	12
6	BDNF polymorphism and interhemispheric balance of motor cortex excitability: a preliminary study. Journal of Neurophysiology, 2022, 127, 204-212.	1.8	6
7	Can we identify hereditary TTR amyloidosis by the screening of carpal tunnel syndrome patients?. Neurological Sciences, 2022, 43, 3435-3438.	1.9	4
8	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
9	Telemedicine application to headache: a critical review. Neurological Sciences, 2022, 43, 3795-3801.	1.9	13
10	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
11	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
12	A compound score to screen patients with hereditary transthyretin amyloidosis. Journal of Neurology, 2022, , .	3.6	3
13	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
14	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
15	Dissective tandem stroke: an endovascular approach. Radiology Case Reports, 2022, 17, 2170-2174.	0.6	3
16	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
17	Mitochondria dysfunction in Charcot Marie Tooth 2B Peripheral Sensory Neuropathy. Communications Biology, 2022, 5, .	4.4	7
18	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

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19	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
20	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
21	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
22	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
23	<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
24	The neurophysiological lesson from the Italian CIDP database. Neurological Sciences, 2021, , 1.	1.9	3
25	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
26	RFC1 expansions are a common cause of idiopathic sensory neuropathy. Brain, 2021, 144, 1542-1550.	7.6	63
27	How to manage with telemedicine people with neuromuscular diseases?. Neurological Sciences, 2021, 42, 3553-3559.	1.9	23
28	Prolonged distal motor latency of median nerve does not improve diagnostic accuracy for CIDP. Journal of Neurology, 2021, , 1.	3.6	1
29	Increased peptidergic fibers as a potential cutaneous marker of pain in diabetic small fiber neuropathy. Pain, 2021, 162, 778-786.	4.2	28
30	Bedside Head Impulse Test: A Useful Tool for Patients With Sensory Ataxia. Neurology: Genetics, 2021, 7, e541.	1.9	1
31	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
32	A Model to Study Myelinated Fiber Degeneration and Regeneration in Human Skin. Annals of Neurology, 2020, 87, 456-465.	5.3	4
33	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
34	Different cortical excitability profiles in hereditary brain iron and copper accumulation. Neurological Sciences, 2020, 41, 679-685.	1.9	6
35	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
36	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

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37	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
38	Pregnancy in Charcot-Marie-Tooth disease. Neurology, 2020, 95, e3180-e3189.	1.1	11
39	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
40	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
41	Phenotypic Variability Among Patients With D4Z4 Reduced Allele Facioscapulohumeral Muscular Dystrophy. JAMA Network Open, 2020, 3, e204040.	5.9	25
42	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
43	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
44	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
45	Brainstem involvement and respiratory failure in COVID-19. Neurological Sciences, 2020, 41, 1663-1665.	1.9	50
46	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
47	Electrodiagnostic accuracy in polyneuropathies: supervised learning algorithms as a tool for practitioners. Neurological Sciences, 2020, 41, 3719-3727.	1.9	5
48	Neurophysiological Signatures of Motor Impairment in Patients with Rett Syndrome. Annals of Neurology, 2020, 87, 763-773.	5.3	14
49	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
50	Antibodies to neurofascin, contactin-1, and contactin-associated protein 1 in CIDP. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, .	6.0	118
51	Lichenoid rash: A new side effect of oral Cladribine. Multiple Sclerosis and Related Disorders, 2020, 41, 102023.	2.0	6
52	Does acute peripheral trauma contribute to idiopathic adult-onset dystonia?. Parkinsonism and Related Disorders, 2020, 71, 40-43.	2.2	18
53	Distal motor neuropathy associated with novel EMILIN1 mutation. Neurobiology of Disease, 2020, 137, 104757.	4.4	6
54	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

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55	Primary Progressive Multiple Sclerosis Under Anti-TNFÎ± Treatment: A Case Report. Journal of Central Nervous System Disease, 2020, 12, 117957352097382.	1.9	1
56	Cognitive correlates of prospective memory in dystonia. Parkinsonism and Related Disorders, 2019, 66, 51-55.	2.2	19
57	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
58	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
59	Spinocerebellar ataxia type 2â€"neuronopathy or neuropathy?. Muscle and Nerve, 2019, 60, 271-278.	2.2	14
60	A Novel CAPN1 Mutation Causes a Pure Hereditary Spastic Paraplegia in an Italian Family. Frontiers in Neurology, 2019, 10, 580.	2.4	14
61	Expanding the spectrum of genes responsible for hereditary motor neuropathies. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 1171-1179.	1.9	30
62	Pseudo-orthostatic tremor: description of a not typical case. Neurological Sciences, 2019, 40, 2205-2207.	1.9	0
63	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
64	Fast Intracortical Sensory-Motor Integration: A Window Into the Pathophysiology of Parkinsonâ€™s Disease. Frontiers in Human Neuroscience, 2019, 13, 111.	2.0	34
65	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
66	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
67	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
68	In vivo evidence of cortical amyloid deposition in the adult form of Niemann Pick type C. Heliyon, 2019, 5, e02776.	3.2	9
69	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
70	The Treatment of Hypersalivation in Rett Syndrome with Botulinum Toxin: Efficacy and Clinical Implications. Neurology and Therapy, 2019, 8, 155-160.	3.2	6
71	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
72	Adult normative values for the PATA Rate Test. Journal of Neurology, 2018, 265, 1102-1105.	3.6	7

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73	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
74	Mutations in ATP1A1 Cause Dominant Charcot-Marie-Tooth Type 2. American Journal of Human Genetics, 2018, 102, 505-514.	6.2	59
75	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
76	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
77	Cognitive profile and 18F-fluorodeoxyglucose PET study in LRRK2 -related Parkinson's disease. Parkinsonism and Related Disorders, 2018, 47, 80-83.	2.2	17
78	Alterations of autophagy in the peripheral neuropathy Charcot-Marie-Tooth type 2B. Autophagy, 2018, 14, 1-12.	9.1	27
79	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
80	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
81	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
82	A cross-sectional study investigating frequency and features of definitely diagnosed diabetic painful polyneuropathy. Pain, 2018, 159, 2658-2666.	4.2	49
83	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
84	Generalized anhidrosis as first clinical presentation of systemic lupus erythematosus. Lupus, 2018, 27, 2296-2297.	1.6	1
85	Muscle pain syndromes and fibromyalgia: the role of muscle biopsy. Current Opinion in Supportive and Palliative Care, 2018, 12, 382-387.	1.3	17
86	Small fiber pathology parallels disease progression in Parkinson disease: a longitudinal study. Acta Neuropathologica, 2018, 136, 501-503.	7.7	19
87	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
88	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
89	Sporadic chronic progressive external ophthalmoplegia with single large mitochondrial DNA deletion and neurogenic findings. Journal of Neurology, 2017, 264, 597-599.	3.6	2
90	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

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91	Novel mutations in dystonin provide clues to the pathomechanisms of HSAN-VI. Neurology, 2017, 88, 2132-2140.	1.1	41
92	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
93	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
94	Early predictive factors of disability in CIDP. Journal of Neurology, 2017, 264, 1939-1944.	3.6	11
95	Disruption of GABA(A)-mediated intracortical inhibition in patients with chorea-acanthocytosis. Neuroscience Letters, 2017, 654, 107-110.	2.1	7
96	Cervical dystonia patients display subclinical gait changes. Parkinsonism and Related Disorders, 2017, 43, 97-100.	2.2	13
97	Upper motor neuron evaluation in multiple sclerosis patients treated with Sativex^{Â®}. Acta Neurologica Scandinavica, 2017, 135, 442-448.	2.1	16
98	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
99	Multimodal evoked potentials follow up in multiple sclerosis patients under fingolimod therapy. Journal of the Neurological Sciences, 2016, 365, 143-146.	0.6	26
100	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
101	Postural instability in Charcot-Marie-Tooth 1A disease. Gait and Posture, 2016, 49, 353-357.	1.4	20
102	Nerve conduction velocity in <scp>CMT</scp>1A: what else can we tell?. European Journal of Neurology, 2016, 23, 1566-1571.	3.3	45
103	Subclinical neurological involvement does not develop if Wilson's disease is treated early. Parkinsonism and Related Disorders, 2016, 24, 15-19.	2.2	34
104	GDAP1 mutations in Italian axonal Charcotâ€“Marieâ€“Tooth patients: Phenotypic features and clinical course. Neuromuscular Disorders, 2016, 26, 26-32.	0.6	18
105	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
106	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
107	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
108	Muscle fiber type disproportion (FTD) in a family with mutations in the LMNA gene. Muscle and Nerve, 2015, 51, 604-608.	2.2	7

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109	Responsiveness of clinical outcome measures in Charcotâ€"Marieâ€"Tooth disease. European Journal of Neurology, 2015, 22, 1556-1563.	3.3	47
110	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
111	Isolated intracranial Mycobacterium avium complex granulomas in an immune-competent man. Journal of the Neurological Sciences, 2015, 349, 264-265.	0.6	4
112	A rare mutation in MYH7 gene occurs with overlapping phenotype. Biochemical and Biophysical Research Communications, 2015, 457, 262-266.	2.1	16
113	Electrophysiological characterization of adult-onset Niemannâ€"Pick type C disease. Journal of the Neurological Sciences, 2015, 348, 262-265.	0.6	22
114	Chronic inflammatory demyelinating polyneuropathy mimicking an acute painful diabetic neuropathy. Neurological Sciences, 2015, 36, 1509-1510.	1.9	1
115	The Effect of Cerebellar Degeneration on Human Sensori-motor Plasticity. Brain Stimulation, 2015, 8, 1144-1150.	1.6	37
116	Small nerve fiber involvement in CMT1A. Neurology, 2015, 84, 407-414.	1.1	30
117	Charcot-Marie-Tooth disease. Neurology, 2015, 85, 1202-1208.	1.1	33
118	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
119	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
120	Teaching Video Neuro Images : Clonus of the lower jaw. Neurology, 2014, 82, e96.	1.1	8
121	Mutilating fingertip ulcers in uncontrolled type 1 diabetes mellitus. Neurological Sciences, 2014, 35, 123-124.	1.9	3
122	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
123	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
124	Postganglionic sudomotor denervation in patients with multiple system atrophy. Neurology, 2014, 82, 2223-2229.	1.1	45
125	Somatosensory Temporal Discrimination Threshold Is Increased in Patients with Cerebellar Atrophy. Cerebellum, 2013, 12, 456-459.	2.5	19
126	Small fiber neuropathy in the chronic phase of Chagas disease: a case report. Clinical Autonomic Research, 2013, 23, 149-153.	2.5	7

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127	Neuropathy and levodopa in Parkinson's disease: Evidence from a multicenter study. Movement Disorders, 2013, 28, 1391-1397.	3.9	114
128	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
129	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
130	Anti-GAD antibody ocular flutter: expanding the spectrum of autoimmune ocular motor disorders. Journal of Neurology, 2013, 260, 2675-2677.	3.6	23
131	Electrophysiological comparison between males and females in HNPP. Neurological Sciences, 2013, 34, 1429-1432.	1.9	10
132	A case of congenital cataracts, facial dysmorphisms, neuropathy, and hyperkinetic movement disorder. Movement Disorders, 2013, 28, 559-560.	3.9	1
133	A novel autosomal dominant GDAP1 mutation in an Italian CMT2 family. Journal of the Peripheral Nervous System, 2012, 17, 351-355.	3.1	13
134	Autonomic nervous system involvement in a new CMT2B family. Journal of the Peripheral Nervous System, 2012, 17, 361-364.	3.1	20
135	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
136	Executive functions are impaired in heterozygote patients with oculopharyngeal muscular dystrophy. Journal of Neurology, 2012, 259, 833-837.	3.6	27
137	Neurophysiological evaluation of motor corticospinal pathways by TMS in idiopathic early-onset Parkinsonâ€™s disease. Clinical Neurophysiology, 2011, 122, 546-549.	1.5	17
138	Electrophysiological characterisation in hereditary spastic paraplegia type 5. Clinical Neurophysiology, 2011, 122, 819-822.	1.5	31
139	Novel ATP13A2 (PARK9) homozygous mutation in a family with marked phenotype variability. Neurogenetics, 2011, 12, 33-39.	1.4	84
140	Thermosensitive hereditary neuropathy with liability to pressure palsy. Muscle and Nerve, 2011, 43, 448-449.	2.2	2
141	A new Italian FHM2 family: Clinical aspects and functional analysis of the disease-associated mutation. Cephalalgia, 2011, 31, 808-819.	3.9	27
142	Autoimmune Autonomic Ganglionopathy. Archives of Neurology, 2011, 68, 504.	4.5	19
143	Functional involvement of central cholinergic circuits and visual hallucinations in Parkinson's disease. Brain, 2009, 132, 2350-2355.	7.6	115
144	Case of acute motor conduction block neuropathy (AMCBN). Muscle and Nerve, 2009, 39, 224-226.	2.2	15

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145	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
146	Two families with novel PMP22 point mutations: genotypeâ€“phenotype correlation. Journal of the Peripheral Nervous System, 2009, 14, 208-212.	3.1	9
147	Nineâ€year case history of monofocal motor neuropathy. Muscle and Nerve, 2008, 38, 927-929.	2.2	8
148	Familial aggregation of white matter lesions in myotonic dystrophy type 1. Neuromuscular Disorders, 2008, 18, 299-305.	0.6	15
149	Motor cortex cholinergic dysfunction in CADASIL: A transcranial magnetic demonstration. Clinical Neurophysiology, 2008, 119, 351-355.	1.5	35
150	Clinical Utility of Electrophysiological Evaluation in Crigler-Najjar Syndrome. Neuropediatrics, 2007, 38, 173-178.	0.6	4
151	Inherited neuromyotonia: A clinical and genetic study of a family. Neuromuscular Disorders, 2007, 17, 23-27.	0.6	16
152	Smallâ€fiber involvement in spinobulbar muscular atrophy (Kennedy's disease). Muscle and Nerve, 2007, 36, 816-820.	2.2	31
153	Prevalence and characteristics of peripheral neuropathy in hepatitis C virus population. Journal of Neurology, Neurosurgery and Psychiatry, 2006, 77, 626-629.	1.9	65
154	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
155	Sural nerve and epidermal vascular abnormalities in a case of POEMS syndrome. European Journal of Neurology, 2006, 13, 99-102.	3.3	20
156	A new POLG1 mutation with peo and severe axonal and demyelinating sensoryâ€“motor neuropathy. Journal of Neurology, 2006, 253, 869-874.	3.6	23
157	GDAP1 mutation in autosomal recessive Charcot-Marie-Tooth with pyramidal features. Journal of Neurology, 2006, 253, 1234-1235.	3.6	9
158	Electrophysiologic characterization in spinocerebellar ataxia 17. Neurology, 2006, 66, 932-934.	1.1	26
159	Trigeminal Stimulation Elicits a Peripheral Vestibular Imbalance in Migraine Patients. Headache, 2005, 45, 325-331.	3.9	101
160	Levetiracetam in patients with cortical myoclonus: A clinical and electrophysiological study. Movement Disorders, 2005, 20, 1610-1614.	3.9	66
161	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
162	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

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163	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
164	Distal hypoglycemic neuropathy. An insulinoma-associated case, misdiagnosed as temporal lobe epilepsy. Neurophysiologie Clinique, 2003, 33, 223-227.	2.2	18
165	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
166	Poems Syndrome With Vasa Nervorum Vasculitis: A Case Report. Journal of the Peripheral Nervous System, 2001, 6, 42-42.	3.1	0
167	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
168	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
169	Pharmacological treatment for familial amyloid neuropathy. The Cochrane Library, 0, , .	2.8	1