Fiore Manganelli

List of Publications by Year in Descending Order

161 papers	2,630 citations	27 h-index	41 g-index
176 ext. papers	3,396 ext. citations	4.8 avg, IF	4.77 L-index

#	Paper	IF	Citations
161	Can we identify hereditary TTR amyloidosis by the screening of carpal tunnel syndrome patients?. *Neurological Sciences*, **2022**, 1	3.5	0
160	Frequency and clinical correlates of anti-nerve antibodies in a large population of CIDP patients included in the Italian database.. *Neurological Sciences*, **2022**, 1	3.5	4
159	Telemedicine application to headache: a critical review.. *Neurological Sciences*, **2022**, 1	3.5	1
158	Abnormal sensorimotor cortex and thalamo-cortical networks in familial adult myoclonic epilepsy type 2: pathophysiology and diagnostic implications.. *Brain Communications*, **2022**, 4, fcac037	4.5	1
157	The Role of New Imaging Technologies in the Diagnosis of Cardiac Amyloidosis. *Heart Failure Clinics* , **2022**, 18, 61-72	3.3	1
156	Cardiovascular Involvement in Transthyretin Cardiac Amyloidosis. *Heart Failure Clinics*, **2022**, 18, 73-87	3.3	3
155	Dissective tandem stroke: an endovascular approach.. *Radiology Case Reports*, **2022**, 17, 2170-2174	1	1
154	Relationship between high-frequency activity in the cortical sensory and the motor hand areas, and their myelin content.. *Brain Stimulation*, **2022**, 15, 717-726	5.1	0
153	Bedside Head Impulse Test: A Useful Tool for Patients With Sensory Ataxia. *Neurology: Genetics*, **2021**, 7, e541	3.8	0
152	Increased peptidergic fibers as a potential cutaneous marker of pain in diabetic small fiber neuropathy. *Pain*, **2021**, 162, 778-786	8	15
151	Acute and chronic inflammatory neuropathies and COVID-19 vaccines: Practical recommendations from the task force of the Italian Peripheral Nervous System Association (ASNP). *Journal of the Peripheral Nervous System*, **2021**, 26, 148-154	4.7	6
150	The impact of symptoms on daily life as perceived by patients with Charcot-Marie-Tooth type 1A disease. *Neurological Sciences*, **2021**, 1	3.5	1
149	The neuropathy in hereditary transthyretin amyloidosis: A narrative review. *Journal of the Peripheral Nervous System*, **2021**, 26, 155-159	4.7	8
148	The neurophysiological lesson from the Italian CIDP database. *Neurological Sciences*, **2021**, 1	3.5	1
147	Proximal weakness involvement in the first Italian case of Charcot-Marie-Tooth 2CC harboring a novel frameshift variant in NEFH. *Journal of the Peripheral Nervous System*, **2021**, 26, 231-234	4.7	3
146	RFC1 expansions are a common cause of idiopathic sensory neuropathy. *Brain*, **2021**, 144, 1542-1550	11.2	11
145	How to manage with telemedicine people with neuromuscular diseases?. *Neurological Sciences*, **2021**, 42, 3553-3559	3.5	7

144	Prolonged distal motor latency of median nerve does not improve diagnostic accuracy for CIDP. *Journal of Neurology*, **2021**, 1	5.5	0
143	Alteration of the late endocytic pathway in Charcot-Marie-Tooth type 2B disease. *Cellular and Molecular Life Sciences*, **2021**, 78, 351-372	10.3	10
142	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	5.5	7
141	Diffuse brain connectivity changes in Charcot-Marie-Tooth type 1a patients: a resting-state functional magnetic resonance imaging study. *European Journal of Neurology*, **2021**, 28, 305-313	6	2
140	Chronic inflammatory demyelinating polyradiculoneuropathy: can a diagnosis be made in patients not fulfilling electrodiagnostic criteria?. *European Journal of Neurology*, **2021**, 28, 620-629	6	9
139	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	4.2	6
138	Phenotypic Variability Among Patients With D4Z4 Reduced Allele Facioscapulohumeral Muscular Dystrophy. *JAMA Network Open*, **2020**, 3, e204040	10.4	14
137	Biallelic mutations in SORD cause a common and potentially treatable hereditary neuropathy with implications for diabetes. *Nature Genetics*, **2020**, 52, 473-481	36.3	38
136	Impact of environmental factors and physical activity on disability and quality of life in CIDP. *Journal of Neurology*, **2020**, 267, 2683-2691	5.5	3
135	Brainstem involvement and respiratory failure in COVID-19. *Neurological Sciences*, **2020**, 41, 1663-1665	3.5	31
134	Sensitivity and specificity of a commercial ELISA test for anti-MAG antibodies in patients with neuropathy. *Journal of Neuroimmunology*, **2020**, 345, 577288	3.5	7
133	Electrodiagnostic accuracy in polyneuropathies: supervised learning algorithms as a tool for practitioners. *Neurological Sciences*, **2020**, 41, 3719-3727	3.5	2
132	Neurophysiological Signatures of Motor Impairment in Patients with Rett Syndrome. *Annals of Neurology*, **2020**, 87, 763-773	9.4	9
131	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	2.2	0
130	Antibodies to neurofascin, contactin-1, and contactin-associated protein 1 in CIDP: Clinical relevance of IgG isotype. *Neurology: Neuroimmunology and NeuroInflammation*, **2020**, 7,	9.1	60
129	Lichenoid rash: A new side effect of oral Cladribine. *Multiple Sclerosis and Related Disorders*, **2020**, 41, 102023	4	3
128	Does acute peripheral trauma contribute to idiopathic adult-onset dystonia?. *Parkinsonism and Related Disorders*, **2020**, 71, 40-43	3.6	10
127	Distal motor neuropathy associated with novel EMILIN1 mutation. *Neurobiology of Disease*, **2020**, 137, 104757	7.5	3

126	RELEVANCE OF DIAGNOSTIC INVESTIGATIONS IN CHRONIC INFLAMMATORY DEMYELINATING POLIRADICULONEUROPATHY: DATA FROM THE ITALIAN CIDP DATABASE. *Journal of the Peripheral Nervous System*, **2020**, 25, 152	4.7	7
125	Primary Progressive Multiple Sclerosis Under Anti-TNF Treatment: A Case Report. *Journal of Central Nervous System Disease*, **2020**, 12, 1179573520973820	4.4	1
124	A Model to Study Myelinated Fiber Degeneration and Regeneration in Human Skin. *Annals of Neurology*, **2020**, 87, 456-465	9.4	2
123	Seronegative occult HBV reactivation complicated with fulminant acute liver failure after rituximab for chronic inflammatory demyelinating polyneuropathy. *Infectious Diseases*, **2020**, 52, 216-218	3.1	1
122	Different cortical excitability profiles in hereditary brain iron and copper accumulation. *Neurological Sciences*, **2020**, 41, 679-685	3.5	3
121	Multimodal evaluation of an Italian family with a hereditary spastic paraplegia and POLR3A mutations. *Annals of Clinical and Translational Neurology*, **2020**, 7, 2326-2331	5.3	4
120	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	5	3
119	Hereditary transthyretin amyloidosis overview. *Neurological Sciences*, **2020**, 1	3.5	11
118	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	2.7	25
117	Pregnancy in Charcot-Marie-Tooth disease: Data from the Italian CMT national registry. *Neurology*, **2020**, 95, e3180-e3189	6.5	3
116	Brain Plasticity in Charcot-Marie-Tooth Type 1A Patients? A Combined Structural and Diffusion MRI Study. *Frontiers in Neurology*, **2020**, 11, 795	4.1	3
115	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	6	15
114	Six-minute walk test is reliable and sensitive in detecting response to therapy in CIDP. *Journal of Neurology*, **2019**, 266, 860-865	5.5	6
113	Spinocerebellar ataxia type 2-neuronopathy or neuropathy?. *Muscle and Nerve*, **2019**, 60, 271-278	3.4	11
112	A Novel Mutation Causes a Pure Hereditary Spastic Paraplegia in an Italian Family. *Frontiers in Neurology*, **2019**, 10, 580	4.1	7
111	Expanding the spectrum of genes responsible for hereditary motor neuropathies. *Journal of Neurology, Neurosurgery and Psychiatry*, **2019**, 90, 1171-1179	5.5	19
110	Pseudo-orthostatic tremor: description of a not typical case. *Neurological Sciences*, **2019**, 40, 2205-2207	3.5	
109	A multicenter retrospective study of charcot-marie-tooth disease type 4B (CMT4B) associated with mutations in myotubularin-related proteins (MTMRs). *Annals of Neurology*, **2019**, 86, 55-67	9.4	13

108	Fast Intracortical Sensory-Motor Integration: A Window Into the Pathophysiology of Parkinson's Disease. *Frontiers in Human Neuroscience*, **2019**, 13, 111	3.3	20
107	A novel family with axonal Charcot-Marie-Tooth disease caused by a mutation in the EGR2 gene. *Journal of the Peripheral Nervous System*, **2019**, 24, 219-223	4.7	5
106	The flavor test is a sensitive tool in identifying the flavor sensorineural dysfunction in Parkinson's disease. *Neurological Sciences*, **2019**, 40, 1351-1356	3.5	7
105	Cognitive correlates of prospective memory in dystonia. *Parkinsonism and Related Disorders*, **2019**, 66, 51-55	3.6	6
104	Insights into the pathogenesis of ATP1A1-related CMT disease using patient-specific iPSCs. *Journal of the Peripheral Nervous System*, **2019**, 24, 330-339	4.7	3
103	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. *Neuromuscular Disorders*, **2019**, 29, 310-316	2.9	2
102	In vivo evidence of cortical amyloid deposition in the adult form of Niemann Pick type C. *Heliyon*, **2019**, 5, e02776	3.6	6
101	Acute leukocytosis during alemtuzumab treatment in patients with active relapsing-remitting multiple sclerosis. *Multiple Sclerosis and Related Disorders*, **2019**, 28, 98-100	4	1
100	The Treatment of Hypersalivation in Rett Syndrome with Botulinum Toxin: Efficacy and Clinical Implications. *Neurology and Therapy*, **2019**, 8, 155-160	4.6	5
99	Atypical CIDP: diagnostic criteria, progression and treatment response. Data from the Italian CIDP Database. *Journal of Neurology, Neurosurgery and Psychiatry*, **2019**, 90, 125-132	5.5	67
98	Adult normative values for the PATA Rate Test. *Journal of Neurology*, **2018**, 265, 1102-1105	5.5	3
97	A case report of limbic encephalitis in a metastatic colon cancer patient during first-line bevacizumab-combined chemotherapy. *Medicine (United States)*, **2018**, 97, e0011	1.8	2
96	Mutations in ATP1A1 Cause Dominant Charcot-Marie-Tooth Type 2. *American Journal of Human Genetics*, **2018**, 102, 505-514	11	36
95	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	3.5	6
94	Motor performance deterioration accelerates after 50 years of age in Charcot-Marie-Tooth type 1A patients. *European Journal of Neurology*, **2018**, 25, 301-306	6	11
93	Muscle pain syndromes and fibromyalgia: the role of muscle biopsy. *Current Opinion in Supportive and Palliative Care*, **2018**, 12, 382-387	2.6	10
92	Normalization of timed neuropsychological tests with the PATA rate and nine-hole pegboard tests. *Journal of Neuropsychology*, **2018**, 12, 471-483	2.6	11
91	Small fiber pathology parallels disease progression in Parkinson disease: a longitudinal study. *Acta Neuropathologica*, **2018**, 136, 501-503	14.3	14

90	Cognitive profile and 18F-fluorodeoxyglucose PET study in LRRK2-related Parkinson's disease. *Parkinsonism and Related Disorders*, **2018**, 47, 80-83	3.6	7
89	Alterations of autophagy in the peripheral neuropathy Charcot-Marie-Tooth type 2B. *Autophagy*, **2018**, 14, 930-941	10.2	22
88	Different nerve ultrasound patterns in charcot-marie-tooth types and hereditary neuropathy with liability to pressure palsies. *Muscle and Nerve*, **2018**, 57, E18-E23	3.4	22
87	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, 1756286418809588	6.6	6
86	A cross-sectional study investigating frequency and features of definitely diagnosed diabetic painful polyneuropathy. *Pain*, **2018**, 159, 2658-2666	8	33
85	Autosomal-dominant transthyretin (TTR)-related amyloidosis is not a frequent CMT2 neuropathy "in disguise". *Orphanet Journal of Rare Diseases*, **2018**, 13, 177	4.2	1
84	Generalized anhidrosis as first clinical presentation of systemic lupus erythematosus. *Lupus*, **2018**, 27, 2296-2297	2.6	0
83	Non-motor involvement in amyotrophic lateral sclerosis: new insight from nerve and vessel analysis in skin biopsy. *Neuropathology and Applied Neurobiology*, **2017**, 43, 119-132	5.2	33
82	Sporadic chronic progressive external ophthalmoplegia with single large mitochondrial DNA deletion and neurogenic findings. *Journal of Neurology*, **2017**, 264, 597-599	5.5	2
81	Does motor cortex plasticity depend on the type of mutation in the leucine-rich repeat kinase 2 gene?. *Movement Disorders*, **2017**, 32, 947-948	7	6
80	Novel mutations in provide clues to the pathomechanisms of HSAN-VI. *Neurology*, **2017**, 88, 2132-2140	6.5	23
79	The therapeutic use of non-invasive brain stimulation in multiple sclerosis - a review. *Restorative Neurology and Neuroscience*, **2017**, 35, 497-509	2.8	32
78	Loss of cutaneous large and small fibers in naive and l-dopa-treated PD patients. *Neurology*, **2017**, 89, 776-784	6.5	49
77	Early predictive factors of disability in CIDP. *Journal of Neurology*, **2017**, 264, 1939-1944	5.5	7
76	Disruption of GABA(A)-mediated intracortical inhibition in patients with chorea-acanthocytosis. *Neuroscience Letters*, **2017**, 654, 107-110	3.3	6
75	Cervical dystonia patients display subclinical gait changes. *Parkinsonism and Related Disorders*, **2017**, 43, 97-100	3.6	7
74	Upper motor neuron evaluation in multiple sclerosis patients treated with Sativex. *Acta Neurologica Scandinavica*, **2017**, 135, 442-448	3.8	12
73	Postural instability in Charcot-Marie-Tooth 1A disease. *Gait and Posture*, **2016**, 49, 353-357	2.6	15

72	Nerve conduction velocity in CMT1A: what else can we tell?. *European Journal of Neurology*, **2016**, 23, 1566-71	6	30
71	Pharmacological treatment for familial amyloid neuropathy. *The Cochrane Library*, **2016**,	5.2	1
70	Subclinical neurological involvement does not develop if Wilson's disease is treated early. *Parkinsonism and Related Disorders*, **2016**, 24, 15-9	3.6	25
69	GDAP1 mutations in Italian axonal Charcot-Marie-Tooth patients: Phenotypic features and clinical course. *Neuromuscular Disorders*, **2016**, 26, 26-32	2.9	13
68	Hirayama's disease: an Italian single center experience and review of the literature. *Quantitative Imaging in Medicine and Surgery*, **2016**, 6, 364-373	3.6	13
67	Electromyography **2016**, 21-37		
66	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. *European Journal of Neurology*, **2016**, 23, 1343-50	6	18
65	Multimodal evoked potentials follow up in multiple sclerosis patients under fingolimod therapy. *Journal of the Neurological Sciences*, **2016**, 365, 143-6	3.2	21
64	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	5.5	62
63	Chronic inflammatory demyelinating polyneuropathy mimicking an acute painful diabetic neuropathy. *Neurological Sciences*, **2015**, 36, 1509-10	3.5	1
62	The Effect of Cerebellar Degeneration on Human Sensori-motor Plasticity. *Brain Stimulation*, **2015**, 8, 1144-50	5.1	27
61	Small nerve fiber involvement in CMT1A. *Neurology*, **2015**, 84, 407-14	6.5	21
60	Charcot-Marie-Tooth disease: New insights from skin biopsy. *Neurology*, **2015**, 85, 1202-8	6.5	20
59	Anodal transcranial direct current stimulation of motor cortex does not ameliorate spasticity in multiple sclerosis. *Restorative Neurology and Neuroscience*, **2015**, 33, 487-92	2.8	27
58	Early onset Charcot-Marie-Tooth neuropathy type 2A and severe developmental delay: expanding the clinical phenotype of MFN2-related neuropathy. *Journal of the Peripheral Nervous System*, **2015**, 20, 415-8	4.7	11
57	Muscle fiber type disproportion (FTD) in a family with mutations in the LMNA gene. *Muscle and Nerve*, **2015**, 51, 604-8	3.4	6
56	Responsiveness of clinical outcome measures in Charcot-Marie-Tooth disease. *European Journal of Neurology*, **2015**, 22, 1556-63	6	33
55	Short-latency afferent inhibition in patients with Parkinson's disease and freezing of gait. *Journal of Neural Transmission*, **2015**, 122, 1533-40	4.3	20

54	Isolated intracranial Mycobacterium avium complex granulomas in an immune-competent man. *Journal of the Neurological Sciences*, **2015**, 349, 264-5	3.2	4
53	A rare mutation in MYH7 gene occurs with overlapping phenotype. *Biochemical and Biophysical Research Communications*, **2015**, 457, 262-6	3.4	11
52	Electrophysiological characterization of adult-onset Niemann-Pick type C disease. *Journal of the Neurological Sciences*, **2015**, 348, 262-5	3.2	18
51	Central cholinergic dysfunction in the adult form of Niemann Pick disease type C: a further link with Alzheimer's disease?. *Journal of Neurology*, **2014**, 261, 804-8	5.5	21
50	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, 1094-6	3.2	5
49	Postganglionic sudomotor denervation in patients with multiple system atrophy. *Neurology*, **2014**, 82, 2223-9	6.5	39
48	PMP22 messenger RNA levels in skin biopsies: testing the effectiveness of a Charcot-Marie-Tooth 1A biomarker. *Brain*, **2014**, 137, 1614-20	11.2	23
47	Charcot-Marie-Tooth disease: frequency of genetic subtypes in a Southern Italy population. *Journal of the Peripheral Nervous System*, **2014**, 19, 292-8	4.7	50
46	Teaching video neuroimages: clonus of the lower jaw: an old sign that comes back. *Neurology*, **2014**, 82, e96	6.5	7
45	Mutilating fingertip ulcers in uncontrolled type 1 diabetes mellitus. *Neurological Sciences*, **2014**, 35, 123-4	3.5	1
44	Somatosensory temporal discrimination threshold is increased in patients with cerebellar atrophy. *Cerebellum*, **2013**, 12, 456-9	4.3	18
43	Small fiber neuropathy in the chronic phase of Chagas disease: a case report. *Clinical Autonomic Research*, **2013**, 23, 149-53	4.3	5
42	Neuropathy and levodopa in Parkinson's disease: evidence from a multicenter study. *Movement Disorders*, **2013**, 28, 1391-7	7	86
41	Impulse control disorders induced by rasagiline as adjunctive therapy for Parkinson's disease: report of 2 cases. *Parkinsonism and Related Disorders*, **2013**, 19, 483-4	3.6	22
40	Atypical clinical and radiological presentation of cryptococcal choroid plexitis in an immunocompetent woman. *Journal of the Neurological Sciences*, **2013**, 334, 180-2	3.2	14
39	Anti-GAD antibody ocular flutter: expanding the spectrum of autoimmune ocular motor disorders. *Journal of Neurology*, **2013**, 260, 2675-7	5.5	22
38	Electrophysiological comparison between males and females in HNPP. *Neurological Sciences*, **2013**, 34, 1429-32	3.5	8
37	A case of congenital cataracts, facial dysmorphisms, neuropathy, and hyperkinetic movement disorder. *Movement Disorders*, **2013**, 28, 559-60	7	1

36	A novel autosomal dominant GDAP1 mutation in an Italian CMT2 family. *Journal of the Peripheral Nervous System*, **2012**, 17, 351-5	4.7	10
35	Autonomic nervous system involvement in a new CMT2B family. *Journal of the Peripheral Nervous System*, **2012**, 17, 361-4	4.7	12
34	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, 1499-506	4.3	60
33	Executive functions are impaired in heterozygote patients with oculopharyngeal muscular dystrophy. *Journal of Neurology*, **2012**, 259, 833-7	5.5	22
32	Neurophysiological evaluation of motor corticospinal pathways by TMS in idiopathic early-onset Parkinson's disease. *Clinical Neurophysiology*, **2011**, 122, 546-549	4.3	12
31	Electrophysiological characterisation in hereditary spastic paraplegia type 5. *Clinical Neurophysiology*, **2011**, 122, 819-22	4.3	28
30	Novel ATP13A2 (PARK9) homozygous mutation in a family with marked phenotype variability. *Neurogenetics*, **2011**, 12, 33-9	3	72
29	Thermosensitive hereditary neuropathy with liability to pressure palsy. *Muscle and Nerve*, **2011**, 43, 448-9	3.4	2
28	A new Italian FHM2 family: clinical aspects and functional analysis of the disease-associated mutation. *Cephalalgia*, **2011**, 31, 808-19	6.1	19
27	Autoimmune autonomic ganglionopathy: a possible postganglionic neuropathy. *Archives of Neurology*, **2011**, 68, 504-7		14
26	Functional involvement of central cholinergic circuits and visual hallucinations in Parkinson's disease. *Brain*, **2009**, 132, 2350-5	11.2	92
25	Case of acute motor conduction block neuropathy (AMCBN). *Muscle and Nerve*, **2009**, 39, 224-6	3.4	10
24	Clinical features and molecular modelling of novel MPZ mutations in demyelinating and axonal neuropathies. *European Journal of Human Genetics*, **2009**, 17, 1129-34	5.3	25
23	Two families with novel PMP22 point mutations: genotype-phenotype correlation. *Journal of the Peripheral Nervous System*, **2009**, 14, 208-12	4.7	8
22	Familial aggregation of white matter lesions in myotonic dystrophy type 1. *Neuromuscular Disorders* , **2008**, 18, 299-305	2.9	15
21	Motor cortex cholinergic dysfunction in CADASIL: a transcranial magnetic demonstration. *Clinical Neurophysiology*, **2008**, 119, 351-5	4.3	33
20	Nine-year case history of monofocal motor neuropathy. *Muscle and Nerve*, **2008**, 38, 927-9	3.4	6
19	Small-fiber involvement in spinobulbar muscular atrophy (Kennedy's disease). *Muscle and Nerve*, **2007**, 36, 816-20	3.4	27

18	Clinical utility of electrophysiological evaluation in Crigler-Najjar syndrome. *Neuropediatrics*, **2007**, 38, 173-8	1.6	3
17	Inherited neuromyotonia: a clinical and genetic study of a family. *Neuromuscular Disorders*, **2007**, 17, 23-7	2.9	14
16	Electrophysiologic characterization in spinocerebellar ataxia 17. *Neurology*, **2006**, 66, 932-4	6.5	23
15	Prevalence and characteristics of peripheral neuropathy in hepatitis C virus population. *Journal of Neurology, Neurosurgery and Psychiatry*, **2006**, 77, 626-9	5.5	52
14	Ross syndrome: a rare or a misknown disorder of thermoregulation? A skin innervation study on 12 subjects. *Brain*, **2006**, 129, 2119-31	11.2	100
13	Sural nerve and epidermal vascular abnormalities in a case of POEMS syndrome. *European Journal of Neurology*, **2006**, 13, 99-102	6	15
12	A new POLG1 mutation with peo and severe axonal and demyelinating sensory-motor neuropathy. *Journal of Neurology*, **2006**, 253, 869-74	5.5	20
11	GDAP1 mutation in autosomal recessive Charcot-Marie-Tooth with pyramidal features. *Journal of Neurology*, **2006**, 253, 1234-5	5.5	9
10	Trigeminal stimulation elicits a peripheral vestibular imbalance in migraine patients. *Headache*, **2005**, 45, 325-31	4.2	84
9	Levetiracetam in patients with cortical myoclonus: a clinical and electrophysiological study. *Movement Disorders*, **2005**, 20, 1610-4	7	51
8	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	3.5	19
7	Post-exercise facilitation and depression of motor evoked potentials to transcranial magnetic stimulation: a study in multiple sclerosis. *Clinical Neurophysiology*, **2004**, 115, 2128-33	4.3	48
6	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	5.5	9
5	Distal hypoglycemic neuropathy. An insulinoma-associated case, misdiagnosed as temporal lobe epilepsy. *Neurophysiologie Clinique*, **2003**, 33, 223-7	2.7	15
4	Charcot-Marie-Tooth disease type 2C: a distinct genetic entity. Clinical and molecular characterization of the first European family. *Neuromuscular Disorders*, **2002**, 12, 399-404	2.9	24
3	Poems Syndrome With Vasa Nervorum Vasculitis: A Case Report. *Journal of the Peripheral Nervous System*, **2001**, 6, 42-42	4.7	
2	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, 1023-30	4.3	37
1	A second family with 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, 240-4	5.5	4