

# Shahram Attarian

## List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

134  
papers

4,788  
citations

31  
h-index

64  
g-index

152  
ext. papers

6,256  
ext. citations

5.2  
avg, IF

5.11  
L-index

#	Paper	IF	Citations
134	Multidisciplinary rehabilitation is relevant in severe myasthenia gravis: An observation. <i>Annals of Physical and Rehabilitation Medicine</i> , <b>2021</b> , 65, 101593	3.8	
133	A multicenter cross-sectional French study of the impact of COVID-19 on neuromuscular diseases. <i>Orphanet Journal of Rare Diseases</i> , <b>2021</b> , 16, 450	4.2	2
132	A double-blind, placebo-controlled, randomized trial of PXT3003 for the treatment of Charcot-Marie-Tooth type 1A. <i>Orphanet Journal of Rare Diseases</i> , <b>2021</b> , 16, 433	4.2	1
131	Visualization of Gray Matter Atrophy and Anterior Corticospinal Tract Signal Hyperintensity in Amyotrophic Lateral Sclerosis Using 7T MRI. <i>Neurology</i> , <b>2021</b> , 96, 1094-1095	6.5	
130	Genotype-phenotype correlation in French patients with myelin protein zero gene-related inherited neuropathy. <i>European Journal of Neurology</i> , <b>2021</b> , 28, 2913-2921	6	1
129	Charcot-Marie-Tooth disease misdiagnosed as chronic inflammatory demyelinating polyradiculoneuropathy: An international multicentric retrospective study. <i>European Journal of Neurology</i> , <b>2021</b> , 28, 2846-2854	6	5
128	Prevalence of spasticity and spasticity-related pain among patients with Amyotrophic Lateral Sclerosis. <i>Revue Neurologique</i> , <b>2021</b> , 177, 694-698	3	2
127	European Academy of Neurology/Peripheral Nerve Society guideline on diagnosis and treatment of chronic inflammatory demyelinating polyradiculoneuropathy: Report of a joint Task Force-Second revision. <i>Journal of the Peripheral Nervous System</i> , <b>2021</b> , 26, 242-268	4.7	23
126	Safety, efficacy, and tolerability of efgartigimod in patients with generalised myasthenia gravis (ADAPT): a multicentre, randomised, placebo-controlled, phase 3 trial. <i>Lancet Neurology</i> , <b>2021</b> , 20, 526-536	24.1	29
125	Refining NGS diagnosis of muscular disorders. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , <b>2021</b> , 92, 223-225	5.5	1
124	Long-term safety and efficacy of patisiran for hereditary transthyretin-mediated amyloidosis with polyneuropathy: 12-month results of an open-label extension study. <i>Lancet Neurology</i> , <b>2021</b> , 20, 49-59	24.1	23
123	Patient-reported impact of Charcot-Marie-Tooth disease: protocol for a real-world digital lifestyle study. <i>Neurodegenerative Disease Management</i> , <b>2021</b> , 11, 21-33	2.8	0
122	Combination of serum and CSF neurofilament-light and neuroinflammatory biomarkers to evaluate ALS. <i>Scientific Reports</i> , <b>2021</b> , 11, 703	4.9	11
121	Impact of Coronavirus Disease 2019 in a French Cohort of Myasthenia Gravis. <i>Neurology</i> , <b>2021</b> , 96, e21096-21104	6.2	14
120	European Academy of Neurology/Peripheral Nerve Society guideline on diagnosis and treatment of chronic inflammatory demyelinating polyradiculoneuropathy: Report of a joint Task Force-Second revision. <i>European Journal of Neurology</i> , <b>2021</b> , 28, 3556-3583	6	22
119	Motor unit number index: A potential electrophysiological biomarker for pediatric spinal muscular atrophy. <i>Muscle and Nerve</i> , <b>2021</b> , 64, 445-453	3.4	1
118	Assessing the upper motor neuron in amyotrophic lateral sclerosis using the triple stimulation technique: A multicenter prospective study. <i>Clinical Neurophysiology</i> , <b>2021</b> , 132, 2551-2557	4.3	0

117	Expanding the spectrum of VEXAS syndrome: association with acute-onset CIDP. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , <b>2021</b> ,	5.5	0
116	Single-fiber studies for assigning pathogenicity of eight mitochondrial DNA variants associated with mitochondrial diseases. <i>Human Mutation</i> , <b>2020</b> , 41, 1394-1406	4.7	1
115	Quantitative assessment of sciatic nerve changes in Charcot-Marie-Tooth type 1A patients using magnetic resonance neurography. <i>European Journal of Neurology</i> , <b>2020</b> , 27, 1382-1389	6	3
114	A New Point Mutation in the PMP22 Gene in a Family Suffering From Atypical HNPP. <i>Journal of Neuromuscular Diseases</i> , <b>2020</b> , 7, 505-510	5	
113	Measuring Outcomes in Adults with Spinal Muscular Atrophy - Challenges and Future Directions - Meeting Report. <i>Journal of Neuromuscular Diseases</i> , <b>2020</b> , 7, 523-534	5	16
112	Type 1 FSHD with 6-10 Repeated Units: Factors Underlying Severity in Index Cases and Disease Penetrance in Their Relatives Attention. <i>International Journal of Molecular Sciences</i> , <b>2020</b> , 21,	6.3	2
111	Quantitative sudomotor test helps differentiate transthyretin familial amyloid polyneuropathy from chronic inflammatory demyelinating polyneuropathy. <i>Clinical Neurophysiology</i> , <b>2020</b> , 131, 1129-1133	4.3	4
110	Quantitative muscle MRI study of patients with sporadic inclusion body myositis. <i>Muscle and Nerve</i> , <b>2020</b> , 61, 496-503	3.4	6
109	Fat fraction distribution in lower limb muscles of patients with CMT1A: A quantitative MRI study. <i>Neurology</i> , <b>2020</b> , 94, e1480-e1487	6.5	8
108	Electrophysiological features of chronic inflammatory demyelinating polyradiculoneuropathy associated with IgG4 antibodies targeting neurofascin 155 or contactin 1 glycoproteins. <i>Clinical Neurophysiology</i> , <b>2020</b> , 131, 921-927	4.3	18
107	Novel CAPN3 variant associated with an autosomal dominant calpainopathy. <i>Neuropathology and Applied Neurobiology</i> , <b>2020</b> , 46, 564-578	5.2	6
106	Selective inhibition of anti-MAG IgM autoantibody binding to myelin by an antigen-specific glycopolymer. <i>Journal of Neurochemistry</i> , <b>2020</b> , 154, 486-501	6	7
105	Guidance for the care of neuromuscular patients during the COVID-19 pandemic outbreak from the French Rare Health Care for Neuromuscular Diseases Network. <i>Revue Neurologique</i> , <b>2020</b> , 176, 507-515	3	47
104	Ulnar neuropathy at the elbow: Reappraisal of the wrist-upper arm latency difference between ulnar and median nerves. <i>Clinical Neurophysiology</i> , <b>2020</b> , 131, 372-376	4.3	1
103	Comparison of MRI and motor evoked potential with triple stimulation technique for the detection of brachial plexus abnormalities in multifocal motor neuropathy. <i>Muscle and Nerve</i> , <b>2020</b> , 61, 325-329	3.4	1
102	Motor unit number index as an individual biomarker: Reference limits of intra-individual variability over time in healthy subjects. <i>Clinical Neurophysiology</i> , <b>2020</b> , 131, 2209-2215	4.3	2
101	Antibodies against the node of Ranvier: a real-life evaluation of incidence, clinical features and response to treatment based on a prospective analysis of 1500 sera. <i>Journal of Neurology</i> , <b>2020</b> , 267, 3664-3672	5.5	31
100	Application of muscle ultrasound for the evaluation of patients with amyotrophic lateral sclerosis: An observational cross-sectional study. <i>Muscle and Nerve</i> , <b>2020</b> , 62, 516-521	3.4	0

99	Evolving Immunologic Perspectives in Chronic Inflammatory Demyelinating Polyneuropathy. <i>Journal of Inflammation Research</i> , <b>2020</b> , 13, 543-549	4.8	4
98	Hereditary neuropathy with liability to pressure palsies. <i>Journal of Neurology</i> , <b>2020</b> , 267, 2198-2206	5.5	15
97	Management challenges for chronic dysimmune neuropathies during the COVID-19 pandemic. <i>Muscle and Nerve</i> , <b>2020</b> , 62, 34-40	3.4	14
96	SMCHD1 is involved in de novo methylation of the DUX4-encoding D4Z4 macrosatellite. <i>Nucleic Acids Research</i> , <b>2019</b> , 47, 2822-2839	20.1	23
95	Expanding the importance of HMERF titinopathy: new mutations and clinical aspects. <i>Journal of Neurology</i> , <b>2019</b> , 266, 680-690	5.5	16
94	Quantitative Brain Sodium MRI Depicts Corticospinal Impairment in Amyotrophic Lateral Sclerosis. <i>Radiology</i> , <b>2019</b> , 292, 422-428	20.5	10
93	Relevance of anti-HNK1 antibodies in the management of anti-MAG neuropathies. <i>Journal of Neurology</i> , <b>2019</b> , 266, 1973-1979	5.5	9
92	Deciphering the complexity of the 4q and 10q subtelomeres by molecular combing in healthy individuals and patients with facioscapulohumeral dystrophy. <i>Journal of Medical Genetics</i> , <b>2019</b> , 56, 590-601	5.8	14
91	Characteristics of patients with vitamin B12-responsive neuropathy: a case series with systematic repeated electrophysiological assessment. <i>Neurological Research</i> , <b>2019</b> , 41, 569-576	2.7	14
90	Early diagnosis of ATTR amyloidosis through targeted follow-up of identified carriers of TTR gene mutations. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , <b>2019</b> , 26, 3-9	2.7	55
89	An overview of motor unit number index reproducibility in amyotrophic lateral sclerosis. <i>Iranian Journal of Neurology</i> , <b>2019</b> , 18, 119-126	0.6	0
88	Guillain-Barré syndrome subtypes: A clinical electrophysiological study of 100 patients. <i>Revue Neurologique</i> , <b>2019</b> , 175, 73-80	3	4
87	Wish to die and reasons for living among patients with amyotrophic lateral sclerosis. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , <b>2019</b> , 20, 68-73	3.6	12
86	Upper limb onset of hereditary transthyretin amyloidosis is common in non-endemic areas. <i>European Journal of Neurology</i> , <b>2019</b> , 26, 497-e36	6	17
85	Motor unit number index correlates with disability in Charcot-Marie-Tooth disease. <i>Clinical Neurophysiology</i> , <b>2018</b> , 129, 1390-1396	4.3	17
84	Guillain-Barré syndrome subtype diagnosis: A prospective multicentric European study. <i>Muscle and Nerve</i> , <b>2018</b> , 58, 23	3.4	15
83	Chronic inflammatory demyelinating polyneuropathy and malignancy: A systematic review. <i>Muscle and Nerve</i> , <b>2018</b> , 57, 875-883	3.4	17
82	Subcutaneous immunoglobulin for maintenance treatment in chronic inflammatory demyelinating polyneuropathy (PATH): a randomised, double-blind, placebo-controlled, phase 3 trial. <i>Lancet Neurology</i> , <b>2018</b> , 17, 35-46	24.1	146

81	Detection of proximal conduction blocks using a triple stimulation technique improves the early diagnosis of Guillain-Barré syndrome. <i>Clinical Neurophysiology</i> , <b>2018</b> , 129, 127-132	4.3	11
80	The utility of motor unit number index: A systematic review. <i>Neurophysiologie Clinique</i> , <b>2018</b> , 48, 251-259.	7	19
79	Regional variation of Guillain-Barré syndrome. <i>Brain</i> , <b>2018</b> , 141, 2866-2877	11.2	96
78	Molecular diagnosis of inherited peripheral neuropathies by targeted next-generation sequencing: molecular spectrum delineation. <i>BMJ Open</i> , <b>2018</b> , 8, e021632	3	20
77	Patisiran, an RNAi Therapeutic, for Hereditary Transthyretin Amyloidosis. <i>New England Journal of Medicine</i> , <b>2018</b> , 379, 11-21	59.2	1163
76	Global motor unit number index sum score for assessing the loss of lower motor neurons in amyotrophic lateral sclerosis. <i>Muscle and Nerve</i> , <b>2017</b> , 56, 202-206	3.4	14
75	Predictive factors of efficacy of rituximab in patients with anti-MAG neuropathy. <i>Journal of the Neurological Sciences</i> , <b>2017</b> , 377, 144-148	3.2	20
74	Disorders of motor neurons manifested by hyperactivity. <i>Revue Neurologique</i> , <b>2017</b> , 173, 345-351	3	5
73	Motor unit number index (MUNIX) in patients with anti-MAG neuropathy. <i>Clinical Neurophysiology</i> , <b>2017</b> , 128, 1264-1269	4.3	10
72	Autoantibodies to nodal isoforms of neurofascin in chronic inflammatory demyelinating polyneuropathy. <i>Brain</i> , <b>2017</b> , 140, 1851-1858	11.2	120
71	Motor-evoked potential gain is a helpful test for the detection of corticospinal tract dysfunction in amyotrophic lateral sclerosis. <i>Clinical Neurophysiology</i> , <b>2017</b> , 128, 357-364	4.3	5
70	Monitoring the short-term effect of intravenous immunoglobulins in multifocal motor neuropathy using motor unit number index. <i>Clinical Neurophysiology</i> , <b>2017</b> , 128, 235-240	4.3	12
69	Long-term follow-up of MRI changes in thigh muscles of patients with Facioscapulohumeral dystrophy: A quantitative study. <i>PLoS ONE</i> , <b>2017</b> , 12, e0183825	3.7	24
68	Region-specific impairment of the cervical spinal cord (SC) in amyotrophic lateral sclerosis: A preliminary study using SC templates and quantitative MRI (diffusion tensor imaging/inhomogeneous magnetization transfer). <i>NMR in Biomedicine</i> , <b>2017</b> , 30, e3801	4.4	25
67	Neuropatie motorie multifocali con blocco della conduzione. <i>EMC - Neurologia</i> , <b>2017</b> , 17, 1-7	0	
66	New strategy for improving the diagnostic sensitivity of repetitive nerve stimulation in myasthenia gravis. <i>Muscle and Nerve</i> , <b>2017</b> , 55, 532-538	3.4	20
65	Determinants of health-related quality of life in anti-MAG neuropathy: a cross-sectional multicentre European study. <i>Journal of the Peripheral Nervous System</i> , <b>2017</b> , 22, 27-33	4.7	9
64	Anti-NF155 chronic inflammatory demyelinating polyradiculoneuropathy strongly associates to HLA-DRB15. <i>Journal of Neuroinflammation</i> , <b>2017</b> , 14, 224	10.1	43

63	Molecular combing reveals complex 4q35 rearrangements in Facioscapulohumeral dystrophy. <i>Human Mutation</i> , <b>2017</b> , 38, 1432-1441	4.7	22
62	Motor unit number index (MUNIX): Is it relevant in chronic inflammatory demyelinating polyradiculoneuropathy (CIDP)?. <i>Clinical Neurophysiology</i> , <b>2016</b> , 127, 1891-4	4.3	21
61	Hereditary and inflammatory neuropathies: a review of reported associations, mimics and misdiagnoses. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , <b>2016</b> , 87, 1051-60	5.5	45
60	Internal modeling of upcoming speech: A causal role of the right posterior cerebellum in non-motor aspects of language production. <i>Cortex</i> , <b>2016</b> , 81, 203-14	3.8	26
59	Multiple acyl-CoA dehydrogenase deficiency (MADD) as a cause of late-onset treatable metabolic disease. <i>Revue Neurologique</i> , <b>2016</b> , 172, 231-41	3	31
58	Muscle MRI of facioscapulohumeral dystrophy (FSHD): A growing demand and a promising approach. <i>Revue Neurologique</i> , <b>2016</b> , 172, 566-571	3	11
57	Low penetrance in facioscapulohumeral muscular dystrophy type 1 with large pathological D4Z4 alleles: a cross-sectional multicenter study. <i>Orphanet Journal of Rare Diseases</i> , <b>2015</b> , 10, 2	4.2	22
56	Myofibrillar myopathies: State of the art, present and future challenges. <i>Revue Neurologique</i> , <b>2015</b> , 171, 715-29	3	26
55	Paraneoplastic subacute lower motor neuron syndrome associated with solid cancer. <i>Journal of the Neurological Sciences</i> , <b>2015</b> , 358, 413-6	3.2	13
54	Dysferlinopathy in Iran: Clinical and genetic report. <i>Journal of the Neurological Sciences</i> , <b>2015</b> , 359, 256-9	3.2	7
53	A meta-analysis of randomized double-blind clinical trials in CMT1A to assess the change from baseline in CMTNS and ONLS scales after one year of treatment. <i>Orphanet Journal of Rare Diseases</i> , <b>2015</b> , 10, 74	4.2	10
52	Fibromyalgia-like symptoms associated with irritable bowel syndrome: A challenging diagnosis of late-onset Pompe disease. <i>Muscle and Nerve</i> , <b>2015</b> , 52, 300-4	3.4	13
51	Proximal conduction block in the pharyngeal-cervical-brachial variant of Guillain-Barré Syndrome. <i>Muscle and Nerve</i> , <b>2015</b> , 52, 1102-6	3.4	5
50	Muscle Quantitative MR Imaging and Clustering Analysis in Patients with Facioscapulohumeral Muscular Dystrophy Type 1. <i>PLoS ONE</i> , <b>2015</b> , 10, e0132717	3.7	26
49	Identification of variants in the 4q35 gene FAT1 in patients with a facioscapulohumeral dystrophy-like phenotype. <i>Human Mutation</i> , <b>2015</b> , 36, 443-53	4.7	29
48	Triple-stimulation technique improves the diagnosis of chronic inflammatory demyelinating polyradiculoneuropathy. <i>Muscle and Nerve</i> , <b>2015</b> , 51, 541-8	3.4	14
47	Quantitative multiplex PCR of short fluorescent fragments for the detection of large intragenic POLG rearrangements in a large French cohort. <i>European Journal of Human Genetics</i> , <b>2014</b> , 22, 542-50	5.3	25
46	Primary neurolymphomatosis diagnosis and treatment: a retrospective study. <i>Journal of the Neurological Sciences</i> , <b>2014</b> , 342, 178-81	3.2	22

45	Association between structural and functional corticospinal involvement in amyotrophic lateral sclerosis assessed by diffusion tensor MRI and triple stimulation technique. <i>Muscle and Nerve</i> , <b>2014</b> , 49, 551-7	3.4	19
44	An exploratory randomised double-blind and placebo-controlled phase 2 study of a combination of baclofen, naltrexone and sorbitol (PXT3003) in patients with Charcot-Marie-Tooth disease type 1A. <i>Orphanet Journal of Rare Diseases</i> , <b>2014</b> , 9, 199	4.2	72
43	Clinical spectrum and gender differences in a large cohort of Charcot-Marie-Tooth type 1A patients. <i>Journal of the Neurological Sciences</i> , <b>2014</b> , 336, 155-60	3.2	16
42	Constitutive activation of the calcium sensor STIM1 causes tubular-aggregate myopathy. <i>American Journal of Human Genetics</i> , <b>2013</b> , 92, 271-8	11	133
41	Dysregulation of 4q35- and muscle-specific genes in fetuses with a short D4Z4 array linked to facio-scapulo-humeral dystrophy. <i>Human Molecular Genetics</i> , <b>2013</b> , 22, 4206-14	5.6	38
40	Deregulation of the protocadherin gene FAT1 alters muscle shapes: implications for the pathogenesis of facioscapulohumeral dystrophy. <i>PLoS Genetics</i> , <b>2013</b> , 9, e1003550	6	58
39	Novel electromyographic monitoring technique for prevention of right phrenic nerve palsy during cryoballoon ablation. <i>Circulation: Arrhythmia and Electrophysiology</i> , <b>2013</b> , 6, 1109-14	6.4	41
38	Recommendations for the management of facioscapulohumeral muscular dystrophy in 2011. <i>Revue Neurologique</i> , <b>2012</b> , 168, 910-8	3	13
37	Motor cortical reorganization is present after a single attack of multiple sclerosis devoid of cortico-spinal dysfunction. <i>Magnetic Resonance Materials in Physics, Biology, and Medicine</i> , <b>2011</b> , 24, 77-84	2.8	9
36	Clinical features and follow-up of four new cases of facial-onset sensory and motor neuronopathy. <i>Muscle and Nerve</i> , <b>2011</b> , 43, 136-40	3.4	16
35	Response to treatment in patients with Lewis-Sumner syndrome. <i>Muscle and Nerve</i> , <b>2011</b> , 44, 179-84	3.4	31
34	Molecular combing reveals allelic combinations in facioscapulohumeral dystrophy. <i>Annals of Neurology</i> , <b>2011</b> , 70, 627-33	9.4	33
33	Neurologic disorders and hepatitis E, France, 2010. <i>Emerging Infectious Diseases</i> , <b>2011</b> , 17, 1510-2	10.2	45
32	Chronic ataxic neuropathies associated with anti-GD1b IgM antibodies: response to IVIg therapy. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , <b>2010</b> , 81, 61-4	5.5	40
31	Acute motor and sensory axonal neuropathy and concomitant encephalopathy during tumor necrosis factor-alpha antagonist therapy. <i>Journal of the Neurological Sciences</i> , <b>2010</b> , 291, 103-6	3.2	21
30	Excitability of the lower-limb area of the motor cortex in Parkinson's disease. <i>Neurophysiologie Clinique</i> , <b>2010</b> , 40, 201-8	2.7	16
29	ALS with respiratory onset: clinical features and effects of non-invasive ventilation on the prognosis. <i>Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders</i> , <b>2010</b> , 11, 379-82		49
28	A motor cortex excitability and gait analysis on Parkinsonian patients. <i>Movement Disorders</i> , <b>2010</b> , 25, 2747-55	7	23

27	Effect of ascorbic acid in patients with Charcot-Marie-Tooth disease type 1A: a multicentre, randomised, double-blind, placebo-controlled trial. <i>Lancet Neurology, The</i> , <b>2009</b> , 8, 1103-10	24.1	103
26	Analysis of the DYSF mutational spectrum in a large cohort of patients. <i>Human Mutation</i> , <b>2009</b> , 30, E345-7	7.5	78
25	Changes in cortically induced inhibition in amyotrophic lateral sclerosis with time. <i>Muscle and Nerve</i> , <b>2009</b> , 39, 310-7	3.4	13
24	Periodic salbutamol in facioscapulohumeral muscular dystrophy: a randomized controlled trial. <i>Archives of Physical Medicine and Rehabilitation</i> , <b>2009</b> , 90, 1094-101	2.8	24
23	Enteral and parenteral nutrition in the later stages of ALS: an observational study. <i>Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders</i> , <b>2009</b> , 10, 42-6		24
22	Motor evoked potentials in clinically isolated syndrome suggestive of multiple sclerosis. <i>Multiple Sclerosis Journal</i> , <b>2009</b> , 15, 355-62	5	17
21	Rapid identification of mitochondrial DNA (mtDNA) mutations in neuromuscular disorders by using surveyor strategy. <i>Mitochondrion</i> , <b>2008</b> , 8, 136-45	4.9	19
20	Enhancement of single motor unit inhibitory responses to transcranial magnetic stimulation in amyotrophic lateral sclerosis. <i>Experimental Brain Research</i> , <b>2008</b> , 189, 229-42	2.3	7
19	Progression of cortical and spinal dysfunctions over time in amyotrophic lateral sclerosis. <i>Muscle and Nerve</i> , <b>2008</b> , 37, 364-75	3.4	22
18	Triple-stimulation technique in multifocal neuropathy with conduction block. <i>Muscle and Nerve</i> , <b>2007</b> , 35, 632-6	3.4	31
17	Magnetic stimulation including the triple-stimulation technique in amyotrophic lateral sclerosis. <i>Muscle and Nerve</i> , <b>2007</b> , 36, 55-61	3.4	27
16	Phenotypic study in 40 patients with dysferlin gene mutations: high frequency of atypical phenotypes. <i>Archives of Neurology</i> , <b>2007</b> , 64, 1176-82		196
15	Prospective clinical and electrophysiological follow-up on a multiple sclerosis population treated with interferon beta-1 a: a pilot study. <i>Multiple Sclerosis Journal</i> , <b>2007</b> , 13, 348-56	5	23
14	Development of a French isometric strength normative database for adults using quantitative muscle testing. <i>Archives of Physical Medicine and Rehabilitation</i> , <b>2007</b> , 88, 1289-97	2.8	75
13	Assessment of cortico-spinal tract impairment in multiple system atrophy using transcranial magnetic stimulation. <i>Clinical Neurophysiology</i> , <b>2007</b> , 118, 815-23	4.3	32
12	Cortical versus spinal dysfunction in amyotrophic lateral sclerosis. <i>Muscle and Nerve</i> , <b>2006</b> , 33, 677-90	3.4	20
11	Covariation of corticospinal efficiency and silent period in motoneuron diseases. <i>Muscle and Nerve</i> , <b>2006</b> , 34, 178-88	3.4	10
10	Multifocal motor neuropathy with and without conduction block: a single entity?. <i>Neurology</i> , <b>2006</b> , 67, 592-6	6.5	95



9	Transcranial magnetic stimulation in lower motor neuron diseases. <i>Clinical Neurophysiology</i> , <b>2005</b> , 116, 35-42	4.3	45
8	Lewis-Sumner syndrome and multifocal motor neuropathy. <i>Muscle and Nerve</i> , <b>2005</b> , 31, 88-94	3.4	69
7	Magnetic stimulation using a triple-stimulation technique in patients with multifocal neuropathy without conduction block. <i>Muscle and Nerve</i> , <b>2005</b> , 32, 710-4	3.4	30
6	Treatment of myasthenia gravis exacerbation with intravenous immunoglobulin: a randomized double-blind clinical trial. <i>Archives of Neurology</i> , <b>2005</b> , 62, 1689-93		138
5	MUSK, a new target for mutations causing congenital myasthenic syndrome. <i>Human Molecular Genetics</i> , <b>2004</b> , 13, 3229-40	5.6	157
4	Neonatal lower motor neuron syndrome associated with maternal neuropathy with anti-GM1 IgG. <i>Neurology</i> , <b>2004</b> , 63, 379-81	6.5	15
3	Terminal latency index and modified F ratio in distinction of chronic demyelinating neuropathies. <i>Clinical Neurophysiology</i> , <b>2001</b> , 112, 457-63	4.3	60
2	Transcranial magnetic stimulation (TMS): compared sensitivity of different motor response parameters in ALS. <i>Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders: Official Publication of the World Federation of Neurology, Research Group on Motor Neuron Diseases</i> , <b>2000</b> , 1		32
1	Microinfusion of the metabotropic glutamate receptor agonist 1S,3R-1-aminocyclopentane-1,3-dicarboxylic acid into the nucleus accumbens induces dopamine-dependent locomotor activation in the rat. <i>European Journal of Neuroscience</i> , <b>1997</b> , 9, 809-16	3.5	38