Shahram Attarian

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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	Patisiran, an RNAi Therapeutic, for Hereditary Transthyretin Amyloidosis. *New England Journal of Medicine*, **2018**, 379, 11-21	59.2	1163
133	Phenotypic study in 40 patients with dysferlin gene mutations: high frequency of atypical phenotypes. *Archives of Neurology*, **2007**, 64, 1176-82		196
132	MUSK, a new target for mutations causing congenital myasthenic syndrome. *Human Molecular Genetics*, **2004**, 13, 3229-40	5.6	157
131	Subcutaneous immunoglobulin for maintenance treatment in chronic inflammatory demyelinating polyneuropathy (PATH): a randomised, double-blind, placebo-controlled, phase 3 trial. *Lancet Neurology, The*, **2018**, 17, 35-46	24.1	146
130	Treatment of myasthenia gravis exacerbation with intravenous immunoglobulin: a randomized double-blind clinical trial. *Archives of Neurology*, **2005**, 62, 1689-93		138
129	Constitutive activation of the calcium sensor STIM1 causes tubular-aggregate myopathy. *American Journal of Human Genetics*, **2013**, 92, 271-8	11	133
128	Autoantibodies to nodal isoforms of neurofascin in chronic inflammatory demyelinating polyneuropathy. *Brain*, **2017**, 140, 1851-1858	11.2	120
127	Effect of ascorbic acid in patients with Charcot-Marie-Tooth disease type 1A: a multicentre, randomised, double-blind, placebo-controlled trial. *Lancet Neurology, The*, **2009**, 8, 1103-10	24.1	103
126	Regional variation of Guillain-Barr□syndrome. *Brain*, **2018**, 141, 2866-2877	11.2	96
125	Multifocal motor neuropathy with and without conduction block: a single entity?. *Neurology*, **2006**, 67, 592-6	6.5	95
124	Analysis of the DYSF mutational spectrum in a large cohort of patients. *Human Mutation*, **2009**, 30, E345-75	4.7	78
123	Development of a French isometric strength normative database for adults using quantitative muscle testing. *Archives of Physical Medicine and Rehabilitation*, **2007**, 88, 1289-97	2.8	75
122	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. *Orphanet Journal of Rare Diseases*, **2014**, 9, 199	4.2	72
121	Lewis-Sumner syndrome and multifocal motor neuropathy. *Muscle and Nerve*, **2005**, 31, 88-94	3.4	69
120	Terminal latency index and modified F ratio in distinction of chronic demyelinating neuropathies. *Clinical Neurophysiology*, **2001**, 112, 457-63	4.3	60
119	Deregulation of the protocadherin gene FAT1 alters muscle shapes: implications for the pathogenesis of facioscapulohumeral dystrophy. *PLoS Genetics*, **2013**, 9, e1003550	6	58
118	Early diagnosis of ATTR amyloidosis through targeted follow-up of identified carriers of TTR gene mutations. *Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis*, **2019**, 26, 3-9	2.7	55

117	ALS with respiratory onset: clinical features and effects of non-invasive ventilation on the prognosis. *Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders*, **2010**, 11, 379-82		49
116	Guidance for the care of neuromuscular patients during the COVID-19 pandemic outbreak from the French Rare Health Care for Neuromuscular Diseases Network. *Revue Neurologique*, **2020**, 176, 507-515	3	47
115	Neurologic disorders and hepatitis E, France, 2010. *Emerging Infectious Diseases*, **2011**, 17, 1510-2	10.2	45
114	Transcranial magnetic stimulation in lower motor neuron diseases. *Clinical Neurophysiology*, **2005**, 116, 35-42	4.3	45
113	Hereditary and inflammatory neuropathies: a review of reported associations, mimics and misdiagnoses. *Journal of Neurology, Neurosurgery and Psychiatry*, **2016**, 87, 1051-60	5.5	45
112	Anti-NF155 chronic inflammatory demyelinating polyradiculoneuropathy strongly associates to HLA-DRB15. *Journal of Neuroinflammation*, **2017**, 14, 224	10.1	43
111	Novel electromyographic monitoring technique for prevention of right phrenic nerve palsy during cryoballoon ablation. *Circulation: Arrhythmia and Electrophysiology*, **2013**, 6, 1109-14	6.4	41
110	Chronic ataxic neuropathies associated with anti-GD1b IgM antibodies: response to IVIg therapy. *Journal of Neurology, Neurosurgery and Psychiatry*, **2010**, 81, 61-4	5.5	40
109	Dysregulation of 4q35- and muscle-specific genes in fetuses with a short D4Z4 array linked to facio-scapulo-humeral dystrophy. *Human Molecular Genetics*, **2013**, 22, 4206-14	5.6	38
108	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. *European Journal of Neuroscience*, **1997**, 9, 809-16	3.5	38
107	Molecular combing reveals allelic combinations in facioscapulohumeral dystrophy. *Annals of Neurology*, **2011**, 70, 627-33	9.4	33
106	Assessment of cortico-spinal tract impairment in multiple system atrophy using transcranial magnetic stimulation. *Clinical Neurophysiology*, **2007**, 118, 815-23	4.3	32
105	Transcranial magnetic stimulation (TMS): compared sensitivity of different motor response parameters in ALS. *Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders: Official Publication of the World Federation of Neurology, Research Group on Motor Neuron Diseases*, **2000**, 1 Suppl 2, S45-9		32
104	Response to treatment in patients with Lewis-Sumner syndrome. *Muscle and Nerve*, **2011**, 44, 179-84	3.4	31
103	Triple-stimulation technique in multifocal neuropathy with conduction block. *Muscle and Nerve*, **2007**, 35, 632-6	3.4	31
102	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. *Journal of Neurology*, **2020**, 267, 3664-3672	5.5	31
101	Multiple acyl-CoA dehydrogenase deficiency (MADD) as a cause of late-onset treatable metabolic disease. *Revue Neurologique*, **2016**, 172, 231-41	3	31
100	Magnetic stimulation using a triple-stimulation technique in patients with multifocal neuropathy without conduction block. *Muscle and Nerve*, **2005**, 32, 710-4	3.4	30

99	Identification of variants in the 4q35 gene FAT1 in patients with a facioscapulohumeral dystrophy-like phenotype. *Human Mutation*, **2015**, 36, 443-53	4.7	29
98	Safety, efficacy, and tolerability of efgartigimod in patients with generalised myasthenia gravis (ADAPT): a multicentre, randomised, placebo-controlled, phase 3 trial. *Lancet Neurology, The*, **2021**, 20, 526-536	24.1	29
97	Magnetic stimulation including the triple-stimulation technique in amyotrophic lateral sclerosis. *Muscle and Nerve*, **2007**, 36, 55-61	3.4	27
96	Myofibrillar myopathies: State of the art, present and future challenges. *Revue Neurologique*, **2015**, 171, 715-29	3	26
95	Muscle Quantitative MR Imaging and Clustering Analysis in Patients with Facioscapulohumeral Muscular Dystrophy Type 1. *PLoS ONE*, **2015**, 10, e0132717	3.7	26
94	Internal modeling of upcoming speech: A causal role of the right posterior cerebellum in non-motor aspects of language production. *Cortex*, **2016**, 81, 203-14	3.8	26
93	Quantitative multiplex PCR of short fluorescent fragments for the detection of large intragenic POLG rearrangements in a large French cohort. *European Journal of Human Genetics*, **2014**, 22, 542-50	5.3	25
92	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). *NMR in Biomedicine*, **2017**, 30, e3801	4.4	25
91	Long-term follow-up of MRI changes in thigh muscles of patients with Facioscapulohumeral dystrophy: A quantitative study. *PLoS ONE*, **2017**, 12, e0183825	3.7	24
90	Periodic salbutamol in facioscapulohumeral muscular dystrophy: a randomized controlled trial. *Archives of Physical Medicine and Rehabilitation*, **2009**, 90, 1094-101	2.8	24
89	Enteral and parenteral nutrition in the later stages of ALS: an observational study. *Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders*, **2009**, 10, 42-6		24
88	SMCHD1 is involved in de novo methylation of the DUX4-encoding D4Z4 macrosatellite. *Nucleic Acids Research*, **2019**, 47, 2822-2839	20.1	23
87	A motor cortex excitability and gait analysis on Parkinsonian patients. *Movement Disorders*, **2010**, 25, 2747-55	7	23
86	Prospective clinical and electrophysiological follow-up on a multiple sclerosis population treated with interferon beta-1 a: a pilot study. *Multiple Sclerosis Journal*, **2007**, 13, 348-56	5	23
85	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. *Journal of the Peripheral Nervous System*, **2021**, 26, 242-268	4.7	23
84	Long-term safety and efficacy of patisiran for hereditary transthyretin-mediated amyloidosis with polyneuropathy: 12-month results of an open-label extension study. *Lancet Neurology, The*, **2021**, 20, 49-59	24.1	23
83	Low penetrance in facioscapulohumeral muscular dystrophy type 1 with large pathological D4Z4 alleles: a cross-sectional multicenter study. *Orphanet Journal of Rare Diseases*, **2015**, 10, 2	4.2	22
82	Primary neurolymphomatosis diagnosis and treatment: a retrospective study. *Journal of the Neurological Sciences*, **2014**, 342, 178-81	3.2	22

81	Progression of cortical and spinal dysfunctions over time in amyotrophic lateral sclerosis. *Muscle and Nerve*, **2008**, 37, 364-75	3.4	22
80	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. *European Journal of Neurology*, **2021**, 28, 3556-3583	6	22
79	Molecular combing reveals complex 4q35 rearrangements in Facioscapulohumeral dystrophy. *Human Mutation*, **2017**, 38, 1432-1441	4.7	22
78	Motor unit number index (MUNIX): Is it relevant in chronic inflammatory demyelinating polyradiculoneuropathy (CIDP)?. *Clinical Neurophysiology*, **2016**, 127, 1891-4	4.3	21
77	Acute motor and sensory axonal neuropathy and concomitant encephalopathy during tumor necrosis factor-alpha antagonist therapy. *Journal of the Neurological Sciences*, **2010**, 291, 103-6	3.2	21
76	Predictive factors of efficacy of rituximab in patients with anti-MAG neuropathy. *Journal of the Neurological Sciences*, **2017**, 377, 144-148	3.2	20
75	New strategy for improving the diagnostic sensitivity of repetitive nerve stimulation in myasthenia gravis. *Muscle and Nerve*, **2017**, 55, 532-538	3.4	20
74	Cortical versus spinal dysfunction in amyotrophic lateral sclerosis. *Muscle and Nerve*, **2006**, 33, 677-90	3.4	20
73	Molecular diagnosis of inherited peripheral neuropathies by targeted next-generation sequencing: molecular spectrum delineation. *BMJ Open*, **2018**, 8, e021632	3	20
72	Association between structural and functional corticospinal involvement in amyotrophic lateral sclerosis assessed by diffusion tensor MRI and triple stimulation technique. *Muscle and Nerve*, **2014**, 49, 551-7	3.4	19
71	Rapid identification of mitochondrial DNA (mtDNA) mutations in neuromuscular disorders by using surveyor strategy. *Mitochondrion*, **2008**, 8, 136-45	4.9	19
70	The utility of motor unit number index: A systematic review. *Neurophysiologie Clinique*, **2018**, 48, 251-259	2.7	19
69	Electrophysiological features of chronic inflammatory demyelinating polyradiculoneuropathy associated with IgG4 antibodies targeting neurofascin 155 or contactin 1 glycoproteins. *Clinical Neurophysiology*, **2020**, 131, 921-927	4.3	18
68	Motor unit number index correlates with disability in Charcot-Marie-Tooth disease. *Clinical Neurophysiology*, **2018**, 129, 1390-1396	4.3	17
67	Motor evoked potentials in clinically isolated syndrome suggestive of multiple sclerosis. *Multiple Sclerosis Journal*, **2009**, 15, 355-62	5	17
66	Upper limb onset of hereditary transthyretin amyloidosis is common in non-endemic areas. *European Journal of Neurology*, **2019**, 26, 497-e36	6	17
65	Chronic inflammatory demyelinating polyneuropathy and malignancy: A systematic review. *Muscle and Nerve*, **2018**, 57, 875-883	3.4	17
64	Expanding the importance of HMERF titinopathy: new mutations and clinical aspects. *Journal of Neurology*, **2019**, 266, 680-690	5.5	16

63	Measuring Outcomes in Adults with Spinal Muscular Atrophy - Challenges and Future Directions - Meeting Report. *Journal of Neuromuscular Diseases*, **2020**, 7, 523-534	5	16
62	Clinical spectrum and gender differences in a large cohort of Charcot-Marie-Tooth type 1A patients. *Journal of the Neurological Sciences*, **2014**, 336, 155-60	3.2	16
61	Clinical features and follow-up of four new cases of facial-onset sensory and motor neuronopathy. *Muscle and Nerve*, **2011**, 43, 136-40	3.4	16
60	Excitability of the lower-limb area of the motor cortex in Parkinson's disease. *Neurophysiologie Clinique*, **2010**, 40, 201-8	2.7	16
59	Guillain-Barré syndrome subtype diagnosis: A prospective multicentric European study. *Muscle and Nerve*, **2018**, 58, 23	3.4	15
58	Neonatal lower motor neuron syndrome associated with maternal neuropathy with anti-GM1 IgG. *Neurology*, **2004**, 63, 379-81	6.5	15
57	Hereditary neuropathy with liability to pressure palsies. *Journal of Neurology*, **2020**, 267, 2198-2206	5.5	15
56	Global motor unit number index sum score for assessing the loss of lower motor neurons in amyotrophic lateral sclerosis. *Muscle and Nerve*, **2017**, 56, 202-206	3.4	14
55	Deciphering the complexity of the 4q and 10q subtelomeres by molecular combing in healthy individuals and patients with facioscapulohumeral dystrophy. *Journal of Medical Genetics*, **2019**, 56, 590-601	5.8	14
54	Characteristics of patients with vitamin B12-responsive neuropathy: a case series with systematic repeated electrophysiological assessment. *Neurological Research*, **2019**, 41, 569-576	2.7	14
53	Triple-stimulation technique improves the diagnosis of chronic inflammatory demyelinating polyradiculoneuropathy. *Muscle and Nerve*, **2015**, 51, 541-8	3.4	14
52	Management challenges for chronic dysimmune neuropathies during the COVID-19 pandemic. *Muscle and Nerve*, **2020**, 62, 34-40	3.4	14
51	Impact of Coronavirus Disease 2019 in a French Cohort of Myasthenia Gravis. *Neurology*, **2021**, 96, e2109-e2120	6.5	14
50	Paraneoplastic subacute lower motor neuron syndrome associated with solid cancer. *Journal of the Neurological Sciences*, **2015**, 358, 413-6	3.2	13
49	Fibromyalgia-like symptoms associated with irritable bowel syndrome: A challenging diagnosis of late-onset Pompe disease. *Muscle and Nerve*, **2015**, 52, 300-4	3.4	13
48	Recommendations for the management of facioscapulohumeral muscular dystrophy in 2011. *Revue Neurologique*, **2012**, 168, 910-8	3	13
47	Changes in cortically induced inhibition in amyotrophic lateral sclerosis with time. *Muscle and Nerve*, **2009**, 39, 310-7	3.4	13
46	Monitoring the short-term effect of intravenous immunoglobulins in multifocal motor neuropathy using motor unit number index. *Clinical Neurophysiology*, **2017**, 128, 235-240	4.3	12

45	Wish to die and reasons for living among patients with amyotrophic lateral sclerosis. *Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration*, **2019**, 20, 68-73	3.6	12
44	Muscle MRI of facioscapulohumeral dystrophy (FSHD): A growing demand and a promising approach. *Revue Neurologique*, **2016**, 172, 566-571	3	11
43	Combination of serum and CSF neurofilament-light and neuroinflammatory biomarkers to evaluate ALS. *Scientific Reports*, **2021**, 11, 703	4.9	11
42	Detection of proximal conduction blocks using a triple stimulation technique improves the early diagnosis of Guillain-Barr□syndrome. *Clinical Neurophysiology*, **2018**, 129, 127-132	4.3	11
41	Motor unit number index (MUNIX) in patients with anti-MAG neuropathy. *Clinical Neurophysiology*, **2017**, 128, 1264-1269	4.3	10
40	Quantitative Brain Sodium MRI Depicts Corticospinal Impairment in Amyotrophic Lateral Sclerosis. *Radiology*, **2019**, 292, 422-428	20.5	10
39	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. *Orphanet Journal of Rare Diseases*, **2015**, 10, 74	4.2	10
38	Covariation of corticospinal efficiency and silent period in motoneuron diseases. *Muscle and Nerve*, **2006**, 34, 178-88	3.4	10
37	Relevance of anti-HNK1 antibodies in the management of anti-MAG neuropathies. *Journal of Neurology*, **2019**, 266, 1973-1979	5.5	9
36	Determinants of health-related quality of life in anti-MAG neuropathy: a cross-sectional multicentre European study. *Journal of the Peripheral Nervous System*, **2017**, 22, 27-33	4.7	9
35	Motor cortical reorganization is present after a single attack of multiple sclerosis devoid of cortico-spinal dysfunction. *Magnetic Resonance Materials in Physics, Biology, and Medicine*, **2011**, 24, 77-84	2.8	9
34	Fat fraction distribution in lower limb muscles of patients with CMT1A: A quantitative MRI study. *Neurology*, **2020**, 94, e1480-e1487	6.5	8
33	Dysferlinopathy in Iran: Clinical and genetic report. *Journal of the Neurological Sciences*, **2015**, 359, 256-9	3.2	7
32	Selective inhibition of anti-MAG IgM autoantibody binding to myelin by an antigen-specific glycopolymer. *Journal of Neurochemistry*, **2020**, 154, 486-501	6	7
31	Enhancement of single motor unit inhibitory responses to transcranial magnetic stimulation in amyotrophic lateral sclerosis. *Experimental Brain Research*, **2008**, 189, 229-42	2.3	7
30	Quantitative muscle MRI study of patients with sporadic inclusion body myositis. *Muscle and Nerve*, **2020**, 61, 496-503	3.4	6
29	Novel CAPN3 variant associated with an autosomal dominant calpainopathy. *Neuropathology and Applied Neurobiology*, **2020**, 46, 564-578	5.2	6
28	Disorders of motor neurons manifested by hyperactivity. *Revue Neurologique*, **2017**, 173, 345-351	3	5

27	Motor-evoked potential gain is a helpful test for the detection of corticospinal tract dysfunction in amyotrophic lateral sclerosis. *Clinical Neurophysiology*, **2017**, 128, 357-364	4.3	5
26	Proximal conduction block in the pharyngeal-cervical-brachial variant of Guillain-Barr syndrome. *Muscle and Nerve*, **2015**, 52, 1102-6	3.4	5
25	Charcot-Marie-Tooth disease misdiagnosed as chronic inflammatory demyelinating polyradiculoneuropathy: An international multicentric retrospective study. *European Journal of Neurology*, **2021**, 28, 2846-2854	6	5
24	Quantitative sudomotor test helps differentiate transthyretin familial amyloid polyneuropathy from chronic inflammatory demyelinating polyneuropathy. *Clinical Neurophysiology*, **2020**, 131, 1129-1135	4.3	4
23	Evolving Immunologic Perspectives in Chronic Inflammatory Demyelinating Polyneuropathy. *Journal of Inflammation Research*, **2020**, 13, 543-549	4.8	4
22	Guillain-Barr syndrome subtypes: A clinical electrophysiological study of 100 patients. *Revue Neurologique*, **2019**, 175, 73-80	3	4
21	Quantitative assessment of sciatic nerve changes in Charcot-Marie-Tooth type 1A patients using magnetic resonance neurography. *European Journal of Neurology*, **2020**, 27, 1382-1389	6	3
20	Type 1 FSHD with 6-10 Repeated Units: Factors Underlying Severity in Index Cases and Disease Penetrance in Their Relatives Attention. *International Journal of Molecular Sciences*, **2020**, 21,	6.3	2
19	A multicenter cross-sectional French study of the impact of COVID-19 on neuromuscular diseases. *Orphanet Journal of Rare Diseases*, **2021**, 16, 450	4.2	2
18	Motor unit number index as an individual biomarker: Reference limits of intra-individual variability over time in healthy subjects. *Clinical Neurophysiology*, **2020**, 131, 2209-2215	4.3	2
17	Prevalence of spasticity and spasticity-related pain among patients with Amyotrophic Lateral Sclerosis. *Revue Neurologique*, **2021**, 177, 694-698	3	2
16	Single-fiber studies for assigning pathogenicity of eight mitochondrial DNA variants associated with mitochondrial diseases. *Human Mutation*, **2020**, 41, 1394-1406	4.7	1
15	A double-blind, placebo-controlled, randomized trial of PXT3003 for the treatment of Charcot-Marie-Tooth type 1A. *Orphanet Journal of Rare Diseases*, **2021**, 16, 433	4.2	1
14	Ulnar neuropathy at the elbow: Reappraisal of the wrist-upper arm latency difference between ulnar and median nerves. *Clinical Neurophysiology*, **2020**, 131, 372-376	4.3	1
13	Comparison of MRI and motor evoked potential with triple stimulation technique for the detection of brachial plexus abnormalities in multifocal motor neuropathy. *Muscle and Nerve*, **2020**, 61, 325-329	3.4	1
12	Genotype-phenotype correlation in French patients with myelin protein zero gene-related inherited neuropathy. *European Journal of Neurology*, **2021**, 28, 2913-2921	6	1
11	Refining NGS diagnosis of muscular disorders. *Journal of Neurology, Neurosurgery and Psychiatry*, **2021**, 92, 223-225	5.5	1
10	Motor unit number index: A potential electrophysiological biomarker for pediatric spinal muscular atrophy. *Muscle and Nerve*, **2021**, 64, 445-453	3.4	1

9	An overview of motor unit number index reproducibility in amyotrophic lateral sclerosis. *Iranian Journal of Neurology*, **2019**, 18, 119-126	0.6	0
8	Application of muscle ultrasound for the evaluation of patients with amyotrophic lateral sclerosis: An observational cross-sectional study. *Muscle and Nerve*, **2020**, 62, 516-521	3.4	0
7	Patient-reported impact of Charcot-Marie-Tooth disease: protocol for a real-world digital lifestyle study. *Neurodegenerative Disease Management*, **2021**, 11, 21-33	2.8	0
6	Assessing the upper motor neuron in amyotrophic lateral sclerosis using the triple stimulation technique: A multicenter prospective study. *Clinical Neurophysiology*, **2021**, 132, 2551-2557	4.3	0
5	Expanding the spectrum of VEXAS syndrome: association with acute-onset CIDP. *Journal of Neurology, Neurosurgery and Psychiatry*, **2021**,	5.5	0
4	A New Point Mutation in the PMP22 Gene in a Family Suffering From Atypical HNPP. *Journal of Neuromuscular Diseases*, **2020**, 7, 505-510	5	
3	Neuropatie motorie multifocali con blocco della conduzione. *EMC - Neurologia*, **2017**, 17, 1-7	0	
2	Multidisciplinary rehabilitation is relevant in severe myasthenia gravis: An observation. *Annals of Physical and Rehabilitation Medicine*, **2021**, 65, 101593	3.8	
1	Visualization of Gray Matter Atrophy and Anterior Corticospinal Tract Signal Hyperintensity in Amyotrophic Lateral Sclerosis Using 7T MRI. *Neurology*, **2021**, 96, 1094-1095	6.5	