Shin J Oh

List of Publications by Year in descending order

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		172457	138484
98	3,735	29	58
papers	citations	h-index	g-index
99	99	99	4790
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Treatment and Management of Disorders of the Neuromuscular Junction., 2022,, 446-491.		1
2	Nodal conduction block: A unifying concept. Muscle and Nerve, 2021, 63, 178-180.	2.2	8
3	Improving the efficacy of exome sequencing at a quaternary care referral centre: novel mutations, clinical presentations and diagnostic challenges in rare neurogenetic diseases. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 1186-1196.	1.9	9
4	Nodal conduction block and reversible conduction failure are not electrophysiological markers for axonal loss. Clinical Neurophysiology, 2021, 132, 2932-2933.	1.5	2
5	Neuromuscular junction disorders beyond myasthenia gravis. Current Opinion in Neurology, 2021, Publish Ahead of Print, 648-657.	3.6	4
6	Assessment of the compound muscle action potential amplitude return time between exercises or tests in the repetitive nerve stimulation test for Lambertâ€Eaton myasthenic syndrome. Muscle and Nerve, 2020, 62, 742-745.	2.2	1
7	Chronic inflammatory axonal polyneuropathy. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 1175-1180.	1.9	12
8	Amifampridine for the treatment of Lambert-Eaton myasthenic syndrome. Expert Review of Clinical Immunology, 2019, 15, 991-1007.	3.0	7
9	Amifampridine Phosphate (Firdapse) Is Effective in a Confirmatory Phase 3 Clinical Trial in LEMS. Journal of Clinical Neuromuscular Disease, 2019, 20, 111-119.	0.7	17
10	Monosialosyl Antibody in a Case Mimicking CANOMAD Syndrome. Journal of Clinical Neuromuscular Disease, 2019, 21, 53-54.	0.7	2
11	Repetitive nerve stimulation test in myasthenic crisis. Muscle and Nerve, 2019, 59, 544-548.	2.2	17
12	High abnormal rate in the repetitive nerve stimulation test in acute onset myasthenia gravis. Clinical Neurophysiology, 2018, 129, 1337-1338.	1.5	2
13	Congenital Myasthenic Syndromes. Neurologic Clinics, 2018, 36, 367-378.	1.8	27
14	Lambert-Eaton Myasthenic Syndrome. Neurologic Clinics, 2018, 36, 379-394.	1.8	80
15	3,4â€diaminopyridine in Lambertâ€Eaton myasthenic syndrome: Concerns regarding presentation of previous studies. Muscle and Nerve, 2018, 57, E130.	2.2	1
16	Diagnostic value of the nearâ€nerve needle sensory nerve conduction in sensory inflammatory demyelinating polyneuropathy. Muscle and Nerve, 2018, 57, 414-418.	2.2	2
17	Multifocal sensory demyelinating neuropathy: Report of a case. Muscle and Nerve, 2017, 56, 825-828.	2.2	3
18	Distinguishing Features of the Repetitive Nerve Stimulation Test Between Lambert–Eaton Myasthenic Syndrome and Myasthenia Gravis, 50-Year Reappraisal. Journal of Clinical Neuromuscular Disease, 2017, 19, 66-75.	0.7	9

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19	Perry Syndrome: A Distinctive Type of TDP-43 Proteinopathy. Journal of Neuropathology and Experimental Neurology, 2017, 76, 676-682.	1.7	50
20	Ocular LEMS or MLOS. Muscle and Nerve, 2016, 54, 981-982.	2.2	5
21	Diagnostic markers of axonal degeneration and demyelination in sensory nerve conduction. Muscle and Nerve, 2016, 53, 866-871.	2.2	8
22	Amifampridine phosphate (Firdapse ^{\hat{A}^{\otimes}}) is effective and safe in a phase 3 clinical trial in LEMS. Muscle and Nerve, 2016, 53, 717-725.	2.2	51
23	Myotonic dystrophy type 1 presenting with asymmetric winged scapulae. Muscle and Nerve, 2016, 54, 339-340.	2.2	1
24	Myasthenia gravis Lambertâ€Eaton overlap syndrome. Muscle and Nerve, 2016, 53, 20-26.	2.2	25
25	Paraneoplastic myeloneuropathy in a man with breast cancer. Muscle and Nerve, 2015, 52, 685-686.	2.2	4
26	Transforming Growth Factor Beta (TGF- \hat{l}^2) is a Muscle Biomarker of Disease Progression in ALS and Correlates with Smad Expression. PLoS ONE, 2015, 10, e0138425.	2.5	44
27	On-nerve needle nerve conduction study in the sural nerve: A new technique for evaluation of peripheral neuropathy. Clinical Neurophysiology, 2015, 126, 1811-1816.	1.5	4
28	Asymptomatic vasculitic neuropathy. Muscle and Nerve, 2015, 52, 34-38.	2.2	8
29	Post-exercise exhaustion in Lambert–Eaton myasthenic syndrome. Clinical Neurophysiology, 2014, 125, 411-414.	1.5	8
30	Oneâ€minute exercise is best for evaluation of postexercise exhaustion in myasthenia gravis. Muscle and Nerve, 2014, 50, 413-416.	2.2	8
31	Smads as muscle biomarkers in amyotrophic lateral sclerosis. Annals of Clinical and Translational Neurology, 2014, 1, 778-787.	3.7	23
32	Vasculitic Neuropathy., 2014,, 156-177.		1
33	Update on Amifampridine as a Drug of Choice in Lambert-Eaton Myasthenic Syndrome. US Neurology, 2014, 10, i.	0.2	2
34	Singleâ€fiber EMG and clinical correlation in Lambertâ€Eaton myasthenic syndrome. Muscle and Nerve, 2013, 47, 664-667.	2.2	18
35	Different Characteristic Phenotypes According to Antibody in Myasthenia Gravis. Journal of Clinical Neuromuscular Disease, 2012, 14, 57-65.	0.7	18
36	Clinical utility of sensory nerve conduction of medial femoral cutaneous nerve. Muscle and Nerve, 2012, 45, 195-199.	2.2	12

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37	Treatment and Management of Disorders of the Neuromuscular Junction., 2011,, 307-342.		2
38	Intraoperative onâ€nerve nerve conduction study and conversion factor in the sural nerve. Muscle and Nerve, 2010, 42, 373-378.	2.2	4
39	Racial differences in myasthenia gravis in Alabama. Muscle and Nerve, 2009, 39, 328-332.	2.2	41
40	3,4â€Diaminopyridine is more effective than placebo in a randomized, doubleâ€blind, crossâ€over drug study in LEMS. Muscle and Nerve, 2009, 40, 795-800.	2.2	84
41	Tenâ€second exercise is superior to 30â€second exercise for postâ€exercise facilitation in diagnosing Lambert–Eaton myasthenic syndrome. Muscle and Nerve, 2008, 37, 572-575.	2.2	51
42	Statins may aggravate myasthenia gravis. Muscle and Nerve, 2008, 38, 1101-1107.	2.2	63
43	Neuropathies of the foot. Clinical Neurophysiology, 2007, 118, 954-980.	1.5	25
44	Electrophysiological differences in seropositive and seronegative Lambert–Eaton myasthenic syndrome. Muscle and Nerve, 2007, 35, 178-183.	2.2	47
45	Chapter 41 Diseases of the nerves in the pelvic girdle and lower limbs. Handbook of Clinical Neurophysiology, 2006, 7, 859-892.	0.0	1
46	Repetitive nerve stimulation of facial muscles in musk antibody–positive myasthenia gravis. Muscle and Nerve, 2006, 33, 500-504.	2.2	76
47	Exercise-induced cramp, myoglobinuria, and tubular aggregates in phosphoglycerate mutase deficiency. Muscle and Nerve, 2006, 34, 572-576.	2.2	34
48	Electrophysiological diagnostic criteria of Lambert-Eaton myasthenic syndrome. Muscle and Nerve, 2005, 32, 515-520.	2.2	138
49	Anti-Hu antibody neuropathy: a clinical, electrophysiological, and pathological study. Clinical Neurophysiology, 2005, 116, 28-34.	1.5	72
50	MG and LEMS overlap syndrome: case report with electrophysiological and immunological evidence. Clinical Neurophysiology, 2005, 116, 1167-1171.	1.5	40
51	Myokymia, neuromyotonia, dermatomyositis, and voltage-gated K+ channel antibodies. Muscle and Nerve, 2003, 27, 757-760.	2.2	24
52	Primary axonal degeneration in tarsal tunnel syndrome: Fact or fiction?. Muscle and Nerve, 2002, 25, 301-302.	2.2	5
53	Near-nerve needle sensory conduction study of the medial calcaneal nerve: New method and report of four cases of medial calcaneal neuropathy. Muscle and Nerve, 2002, 26, 654-658.	2.2	12
54	Neurolymphomatosis associated with muscle and cerebral involvement caused by natural killer cell lymphoma: a case report and review of literature. Journal of the Peripheral Nervous System, 2001, 6, 197-203.	3.1	36

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55	Distal sensory nerve conduction of the superficial peroneal nerve: New method and its clinical application. Muscle and Nerve, 2001, 24, 689-694.	2.2	42
56	Primary LAMP-2 deficiency causes X-linked vacuolar cardiomyopathy and myopathy (Danon disease). Nature, 2000, 406, 906-910.	27.8	865
57	New near-nerve needle nerve conduction technique: Differentiating epicondylar from cubital tunnel ulnar neuropathy. , 1999, 22, 718-723.		19
58	Lateral plantar neuropathy. Muscle and Nerve, 1999, 22, 1234-1238.	2.2	28
59	ENTRAPMENT NEUROPATHIES OF THE TIBIAL (POSTERIOR TIBIAL) NERVE. Neurologic Clinics, 1999, 17, 593-615.	1.8	93
60	Wide Spectrum of Symptomatic Treatment in Lambert-Eaton Myasthenic Syndrome. Annals of the New York Academy of Sciences, 1998, 841, 827-831.	3.8	14
61	Single-fiber electromyography improvement with 3,4-diaminopyridine in Lambert-Eaton myasthenic syndrome. , 1998, 21, 1107-1108.		10
62	Low-dose guanidine and pyridostigmine: relatively safe and effective long-term symptomatic therapy in Lambert-Eaton myasthenic syndrome., 1997, 20, 1146-1152.		62
63	Anti-Hu-associated paraneoplastic sensory neuronopathy responding to early aggressive immunotherapy: Report of two cases and review of literature., 1997, 20, 1576-1582.		60
64	Antiâ∈Huâ€associated paraneoplastic sensory neuronopathy responding to early aggressive immunotherapy: Report of two cases and review of literature. Muscle and Nerve, 1997, 20, 1576-1582.	2.2	2
65	Case of the month: Isaacs' syndrome associated with chronic inflammatory demyelinating polyneuropathy., 1996, 19, 210-215.		29
66	Combined motor and sensory median-ulnar anastomosis: Report of an electrophysiologically proven case., 1996, 19, 231-233.		11
67	Electrophysiological and clinical correlations in the Lambert-Eaton myasthenic syndrome. , 1996, 19, 903-906.		19
68	Letters to the editor. Muscle and Nerve, 1996, 19, 1361-1366.	2.2	14
69	Modified trichrome staining technique of the nerve to determine proximal nerve viability. Microsurgery, 1995, 16, 129-132.	1.3	12
70	A macro-EMG study in charonic demyelinating neuropathy. Muscle and Nerve, 1995, 18, 348-350.	2.2	2
71	Electrophysiological studies in Joplin's neuroma. Muscle and Nerve, 1995, 18, 671-672.	2.2	20
72	Peroneal nerve repetitive nerve stimulation test: Its value in diagnosis of myasthenia gravis and Lambert-Eaton myasthenic syndrome. Muscle and Nerve, 1995, 18, 867-873.	2.2	22

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73	Double anastomosis of median-ulnar and ulnar-median nerves: Report of an electrophysiologically proven case. Muscle and Nerve, 1995, 18, 1332-1334.	2.2	6
74	Early appearance of aging phenomenon in the interdigital nerves of the foot. Muscle and Nerve, 1994, 17, 58-63.	2.2	21
75	Tendon-reflex testing in chronic demyelinating polyneuropathy. Muscle and Nerve, 1994, 17, 145-150.	2.2	51
76	Letters to the editor. Muscle and Nerve, 1994, 17, 245-253.	2.2	3
77	What is the best diagnostic index of conduction block and temporal dispersion?. Muscle and Nerve, 1994, 17, 489-493.	2.2	91
78	Myokymia-cramp syndrome: Evidence of hyperexcitable peripheral nerve. Muscle and Nerve, 1994, 17, 1065-1067.	2.2	12
79	Primary respiratory failure as the presenting symptom in Lambert-Eaton myasthenic syndrome. Muscle and Nerve, 1993, 16, 712-715.	2.2	25
80	Electrophysiological Characteristics in Seronegative Myasthenia Gravis. Annals of the New York Academy of Sciences, 1993, 681, 584-587.	3.8	19
81	Letters to the editor. Muscle and Nerve, 1992, 15, 255-258.	2.2	19
82	Letters to the editor. Muscle and Nerve, 1992, 15, 513-523.	2.2	13
83	Letters to the editor. Muscle and Nerve, 1992, 15, 513-523. Diagnostic sensitivity of the laboratory tests in myasthenia gravis. Muscle and Nerve, 1992, 15, 720-724.	2,2	185
83	Diagnostic sensitivity of the laboratory tests in myasthenia gravis. Muscle and Nerve, 1992, 15, 720-724.	2.2	185
83	Diagnostic sensitivity of the laboratory tests in myasthenia gravis. Muscle and Nerve, 1992, 15, 720-724. Paraneoplastic vasculitic neuropathy: A treatable neuropathy. Muscle and Nerve, 1991, 14, 152-156. Electrophysiological improvement following decompression surgery in tarsal tunnel syndrome.	2.2	185 70
83 84 85	Diagnostic sensitivity of the laboratory tests in myasthenia gravis. Muscle and Nerve, 1992, 15, 720-724. Paraneoplastic vasculitic neuropathy: A treatable neuropathy. Muscle and Nerve, 1991, 14, 152-156. Electrophysiological improvement following decompression surgery in tarsal tunnel syndrome. Muscle and Nerve, 1991, 14, 407-410. Edrophonium responsiveness not necessarily diagnostic of myasthenia gravis. Muscle and Nerve, 1990,	2.2 2.2	185 70 21
83 84 85 86	Diagnostic sensitivity of the laboratory tests in myasthenia gravis. Muscle and Nerve, 1992, 15, 720-724. Paraneoplastic vasculitic neuropathy: A treatable neuropathy. Muscle and Nerve, 1991, 14, 152-156. Electrophysiological improvement following decompression surgery in tarsal tunnel syndrome. Muscle and Nerve, 1991, 14, 407-410. Edrophonium responsiveness not necessarily diagnostic of myasthenia gravis. Muscle and Nerve, 1990, 13, 187-191.	2.2 2.2 2.2 2.2	185 70 21 61
83 84 85 86	Diagnostic sensitivity of the laboratory tests in myasthenia gravis. Muscle and Nerve, 1992, 15, 720-724. Paraneoplastic vasculitic neuropathy: A treatable neuropathy. Muscle and Nerve, 1991, 14, 152-156. Electrophysiological improvement following decompression surgery in tarsal tunnel syndrome. Muscle and Nerve, 1991, 14, 407-410. Edrophonium responsiveness not necessarily diagnostic of myasthenia gravis. Muscle and Nerve, 1990, 13, 187-191. Electrophysiological spectrum of inclusion body myositis. Muscle and Nerve, 1990, 13, 949-951.	2.2 2.2 2.2 2.2	185 70 21 61 74

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91	Diverse electrophysiological spectrum of the Lambert-Eaton myasthenic syndrome. Muscle and Nerve, 1989, 12, 464-469.	2.2	48
92	Tomaculous neuropathy presenting as acute recurrent polyneuropathy. Annals of Neurology, 1989, 26, 98-100.	5.3	26
93	SFEMG improvement with remission in the cancer-associated Lambert-Eaton myasthenic syndrome. Muscle and Nerve, 1989, 12, 844-848.	2.2	13
94	Electrophysiological diagnosis of interdigital neuropathy of the foot. Muscle and Nerve, 1984, 7, 218-225.	2.2	50
95	A benign form of reducing body myopathy. Muscle and Nerve, 1983, 6, 278-282.	2.2	25
96	Electrophysiological and clinical correlation in myasthenia gravis. Annals of Neurology, 1982, 12, 348-354.	5.3	89
97	Tarsal tunnel syndrome: Electrophysiological study. Annals of Neurology, 1979, 5, 327-330.	5.3	132
98	Color Atlas of Nerve Biopsy Pathology. , 0, , .		21