

David Lacomis

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.

80 papers	2,517 citations	24 h-index	49 g-index
89 ext. papers	2,972 ext. citations	3.7 avg, IF	5.23 L-index

#	Paper	IF	Citations
80	Granulomatous Myositis Associated With Acetylcholine Receptor Antibodies Without Clinical Myasthenia. <i>Journal of Clinical Neuromuscular Disease</i> , 2021 , 23, 49-52	1.1	
79	What Is in the Myopathy Literature?. <i>Journal of Clinical Neuromuscular Disease</i> , 2021 , 23, 66-74	1.1	
78	What Is in the Neuromuscular Junction Literature?. <i>Journal of Clinical Neuromuscular Disease</i> , 2021 , 22, 147-154	1.1	1
77	Lambert-Eaton Myasthenic Syndrome and Dermatomyositis With Anti-TIF1-gamma Autoantibody: A Unique Association of Autoimmune Neuromuscular Conditions Without Malignancy. <i>Journal of Clinical Neuromuscular Disease</i> , 2021 , 22, 164-168	1.1	0
76	SDH Subunit C Regulates Muscle Oxygen Consumption and Fatigability in an Animal Model of Pulmonary Emphysema. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2021 , 65, 259-271	5.7	3
75	Neuronal mitochondrial dysfunction in sporadic amyotrophic lateral sclerosis is developmentally regulated. <i>Scientific Reports</i> , 2021 , 11, 18916	4.9	3
74	Effects of mexiletine on hyperexcitability in sporadic amyotrophic lateral sclerosis: Preliminary findings from a small phase II randomized controlled trial. <i>Muscle and Nerve</i> , 2021 , 63, 371-383	3.4	3
73	Longitudinal biomarkers in amyotrophic lateral sclerosis. <i>Annals of Clinical and Translational Neurology</i> , 2020 , 7, 1103-1116	5.3	24
72	What Is in the Neuromuscular Junction Literature?. <i>Journal of Clinical Neuromuscular Disease</i> , 2020 , 21, 195-204	1.1	0
71	Making the Diagnosis of Myositis: Electrodiagnostic Testing 2020 , 99-108		
70	Making the Diagnosis of Myositis: Muscle Biopsy and Interpretation 2020 , 109-124		
69	What is in the Myopathy Literature?. <i>Journal of Clinical Neuromuscular Disease</i> , 2020 , 22, 77-83	1.1	1
68	IL-13-driven pulmonary emphysema leads to skeletal muscle dysfunction attenuated by endurance exercise. <i>Journal of Applied Physiology</i> , 2020 , 128, 134-148	3.7	10
67	Late-onset peripheral neuropathy in patients with wild type transthyretin amyloidosis (wtATTR). <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2020 , 27, 142-143	2.7	5
66	Differential tractography as a track-based biomarker for neuronal injury. <i>NeuroImage</i> , 2019 , 202, 116131	7.9	24
65	-associated congenital fiber-type disproportion and cardiomyopathy with variants in additional neuromuscular disease genes; the dilemma of panel testing. <i>Journal of Physical Education and Sports Management</i> , 2019 , 5,	2.8	1
64	What is in the Myopathy Literature?. <i>Journal of Clinical Neuromuscular Disease</i> , 2019 , 21, 7-13	1.1	

63	Clinical spectrum of neuromuscular complications after immune checkpoint inhibition. <i>Neuromuscular Disorders</i> , 2019 , 29, 127-133	2.9	32
62	What is in the Myopathy Literature?. <i>Journal of Clinical Neuromuscular Disease</i> , 2018 , 19, 131-134	1.1	
61	Duchenne muscular dystrophy caused by a novel deep intronic DMD mutation. <i>Muscle and Nerve</i> , 2018 , 57, E136-E138	3.4	4
60	Electrodiagnostic studies in the intensive care unit: A comparison study 2 decades later. <i>Muscle and Nerve</i> , 2018 , 57, 772-776	3.4	3
59	What is in the Myopathy Literature?. <i>Journal of Clinical Neuromuscular Disease</i> , 2018 , 19, 217-223	1.1	
58	Lambert-Eaton myasthenic syndrome: mouse passive-transfer model illuminates disease pathology and facilitates testing therapeutic leads. <i>Annals of the New York Academy of Sciences</i> , 2018 , 1412, 73-81	6.5	8
57	What is in the Neuromuscular Junction Literature?. <i>Journal of Clinical Neuromuscular Disease</i> , 2018 , 20, 76-84	1.1	
56	Reported direct aminopyridine effects on voltage-gated calcium channels is a high-dose pharmacological off-target effect of no clinical relevance. <i>Journal of Biological Chemistry</i> , 2018 , 293, 16100	5.4	4
55	What is in the Literature?. <i>Journal of Clinical Neuromuscular Disease</i> , 2017 , 18, 165-175	1.1	
54	Poly(GP) proteins are a useful pharmacodynamic marker for -associated amyotrophic lateral sclerosis. <i>Science Translational Medicine</i> , 2017 , 9,	17.5	128
53	Clinical and neuropathological features of ALS/FTD with TIA1 mutations. <i>Acta Neuropathologica Communications</i> , 2017 , 5, 96	7.3	27
52	TIA1 Mutations in Amyotrophic Lateral Sclerosis and Frontotemporal Dementia Promote Phase Separation and Alter Stress Granule Dynamics. <i>Neuron</i> , 2017 , 95, 808-816.e9	13.9	341
51	Proximal Limb Weakness in a Patient with Celiac Disease: Copper Deficiency, Gluten Sensitivity, or Both as the Underlying Cause?. <i>Case Reports in Neurological Medicine</i> , 2016 , 2016, 5415949	0.7	3
50	Exposure to hazardous air pollutants and the risk of amyotrophic lateral sclerosis. <i>Environmental Pollution</i> , 2015 , 197, 181-186	9.3	41
49	Letter to the editor on "Exposure to hazardous air pollutants and the risk of amyotrophic lateral sclerosis". <i>Environmental Pollution</i> , 2015 , 207, 432-3	9.3	
48	Receptor for Advanced Glycation End Products and its Inflammatory Ligands are Upregulated in Amyotrophic Lateral Sclerosis. <i>Frontiers in Cellular Neuroscience</i> , 2015 , 9, 485	6.1	38
47	Advanced diffusion MRI fiber tracking in neurosurgical and neurodegenerative disorders and neuroanatomical studies: A review. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2014 , 1842, 2286-2297	6.9	73
46	Localized scleroderma and regional inflammatory myopathy. <i>Neuromuscular Disorders</i> , 2014 , 24, 425-30	2.9	8

45	Use of diffusion spectrum imaging in preliminary longitudinal evaluation of amyotrophic lateral sclerosis: development of an imaging biomarker. <i>Frontiers in Human Neuroscience</i> , 2014 , 8, 270	3.3	19
44	Life factors affecting depression and burden in amyotrophic lateral sclerosis caregivers. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2014 , 15, 292-7	3.6	30
43	Critical illness polyneuropathy 2014 , 273-275		
42	Complete reversal of Lambert-Eaton myasthenic syndrome synaptic impairment by the combined use of a K ⁺ channel blocker and a Ca ²⁺ channel agonist. <i>Journal of Physiology</i> , 2014 , 592, 3687-96	3.9	14
41	Environmental and occupational risk factors for amyotrophic lateral sclerosis: a case-control study. <i>Neurodegenerative Diseases</i> , 2014 , 14, 31-8	2.3	47
40	Acute Neuromuscular Weakness in the Intensive Care Unit 2014 , 1515-1532		
39	Electrophysiology of neuromuscular disorders in critical illness. <i>Muscle and Nerve</i> , 2013 , 47, 452-63	3.4	56
38	What's in the literature?. <i>Journal of Clinical Neuromuscular Disease</i> , 2013 , 15, 34-42	1.1	
37	Electrodiagnostic approach to the patient with suspected myopathy. <i>Neurologic Clinics</i> , 2012 , 30, 641-604.5	4.5	14
36	Neurosarcoidosis. <i>Current Neuropharmacology</i> , 2011 , 9, 429-36	7.6	86
35	Neuromuscular disorders in critically ill patients: review and update. <i>Journal of Clinical Neuromuscular Disease</i> , 2011 , 12, 197-218	1.1	35
34	Post-translational protein modifications of transthyretin in amyotrophic lateral sclerosis. <i>FASEB Journal</i> , 2010 , 24, 568.10	0.9	
33	Cystatin C: expression and activity in amyotrophic lateral sclerosis. <i>FASEB Journal</i> , 2010 , 24, 568.11	0.9	
32	Neuromuscular pathology case. <i>Journal of Clinical Neuromuscular Disease</i> , 2008 , 10, 79-82	1.1	2
31	Severe hydroxychloroquine myopathy. <i>Muscle and Nerve</i> , 2008 , 38, 1206-10	3.4	56
30	Biomarkers for ALS disease progression. <i>FASEB Journal</i> , 2008 , 22, 173.6	0.9	
29	Cystatin C: A potential surrogate biomarker in amyotrophic lateral sclerosis. <i>FASEB Journal</i> , 2008 , 22, 173.7	0.9	
28	Approach to vasculitic neuropathies. <i>Journal of Clinical Neuromuscular Disease</i> , 2007 , 9, 265-76	1.1	26

27	Neuromuscular pathology unknown. <i>Journal of Clinical Neuromuscular Disease</i> , 2005 , 6, 191-2	1.1	
26	Myasthenic crisis. <i>Neurocritical Care</i> , 2005 , 3, 189-94	3.3	45
25	Neuropathy and Fabry's disease. <i>Muscle and Nerve</i> , 2005 , 31, 102-7	3.4	25
24	Clinical utility of peripheral nerve biopsy. <i>Current Neurology and Neuroscience Reports</i> , 2005 , 5, 41-7	6.6	8
23	The utility of muscle biopsy. <i>Current Neurology and Neuroscience Reports</i> , 2004 , 4, 81-6	6.6	8
22	Evaluation of the patient with foot pain: when is the cause small-fiber neuropathy?. <i>Journal of Clinical Neuromuscular Disease</i> , 2004 , 6, 24-39	1.1	3
21	Small-fiber neuropathy. <i>Muscle and Nerve</i> , 2002 , 26, 173-88	3.4	297
20	Hypothyroid myopathy. <i>Journal of Clinical Neuromuscular Disease</i> , 2002 , 4, 100-1	1.1	
19	Electrodiagnostic approach to the patient with suspected myopathy. <i>Neurologic Clinics</i> , 2002 , 20, 587-604	3.5	12
18	Critical illness neuromyopathies. <i>Advances in Neurology</i> , 2002 , 88, 325-35		4
17	The use of percutaneous needle muscle biopsy in the diagnosis of myopathy. <i>Current Rheumatology Reports</i> , 2000 , 2, 225-9	4.9	16
16	Giant cell arteritis presenting with proximal weakness and skeletal muscle vasculitis. <i>Muscle and Nerve</i> , 1999 , 22, 142-4	3.4	7
15	Comparison of different modalities for detection of small fiber neuropathy. <i>Clinical Neurophysiology</i> , 1999 , 110, 1909-12	4.3	75
14	Causes of neuromuscular weakness in the intensive care unit: a study of ninety-two patients. <i>Muscle and Nerve</i> , 1998 , 21, 610-7	3.4	141
13	Oculopharyngeal muscular dystrophy: non-French-Canadian pedigrees. <i>Muscle and Nerve</i> , 1998 , 21, 816-8	3.4	5
12	Mononeuropathies associated with liver transplantation. <i>Muscle and Nerve</i> , 1998 , 21, 896-901	3.4	16
11	Neuropathy associated with hyperoxaluria: improvement after combined renal and liver transplantations. <i>Brain Pathology</i> , 1998 , 8, 247-51	6	9
10	October 1996--rapidly progressive weakness. <i>Brain Pathology</i> , 1997 , 7, 837-8	6	

9	Small fiber neuropathy and vasculitis. <i>Arthritis and Rheumatism</i> , 1997 , 40, 1173-7		40
8	Percutaneous needle muscle biopsy in the evaluation of patients with suspected inflammatory myopathy. <i>Arthritis and Rheumatism</i> , 1997 , 40, 1886-91		23
7	Vacuolar myopathies in adults with myalgias: value of paraspinal muscle investigation. <i>Muscle and Nerve</i> , 1997 , 20, 1321-3	3-4	4
6	Case of the month. June 1996--anorexia nervosa. <i>Brain Pathology</i> , 1996 , 6, 535-6	6	8
5	Acute myopathy of intensive care: clinical, electromyographic, and pathological aspects. <i>Annals of Neurology</i> , 1996 , 40, 645-54	9-4	231
4	Mushroom myopathy. <i>Muscle and Nerve</i> , 1996 , 19, 790-2	3-4	7
3	Disseminated histoplasmosis presenting as myositis and fasciitis in a patient with dermatomyositis. <i>Muscle and Nerve</i> , 1995 , 18, 531-5	3-4	24
2	A reversible paralytic syndrome with anti-GD1b antibodies following influenza immunization. <i>Muscle and Nerve</i> , 1995 , 18, 1199-201	3-4	3
1	Acute myopathy and neuropathy in status asthmaticus: case report and literature review. <i>Muscle and Nerve</i> , 1993 , 16, 84-90	3-4	111