Christopher Jon Klein

List of Publications by Year
in descending order

195	8,819	36271 51	53190 85
papers	citations	h-index	g-index

196	196	196	8930
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Autoimmune encephalitis epidemiology and a comparison to infectious encephalitis. Annals of Neurology, 2018, 83, 166-177.	2.8	479
2	Mutations in DNMT1 cause hereditary sensory neuropathy with dementia and hearing loss. Nature Genetics, 2011, 43, 595-600.	9.4	342
3	Autoimmune Epilepsy. Archives of Neurology, 2012, 69, 582.	4.9	324
4	Neurological Complications Associated With Antiâ€“Programmed Death 1 (PD-1) Antibodies. JAMA Neurology, 2017, 74, 1216.	4.5	244
5	Expanded phenotypes and outcomes among 256 <scp>LGI</scp>1/<scp>CASPR</scp>2â€<scp>I</scp>g<scp>G</scp>â€“positive patients. Annals of Neurology, 2017, 82, 79-92.	2.8	242
6	Scapuloperoneal spinal muscular atrophy and CMT2C are allelic disorders caused by alterations in TRPV4. Nature Genetics, 2010, 42, 165-169.	9.4	232
7	Post-surgical inflammatory neuropathy. Brain, 2010, 133, 2866-2880.	3.7	203
8	Signs and symptoms versus nerve conduction studies to diagnose diabetic sensorimotor polyneuropathy: Cl vs. NPhys trial. Muscle and Nerve, 2010, 42, 157-164.	1.0	191
9	Insights From LGI1 and CASPR2 Potassium Channel Complex Autoantibody Subtyping. JAMA Neurology, 2013, 70, 229.	4.5	170
10	Voltage-Gated Potassium Channel Autoimmunity Mimicking Creutzfeldt-Jakob Disease. Archives of Neurology, 2008, 65, 1341-6.	4.9	166
11	Basal ganglia T1 hyperintensity in LGI1-autoantibody faciobrachial dystonic seizures. Neurology: Neuroimmunology and NeuroInflammation, 2015, 2, e161.	3.1	163
12	Chronic pain as a manifestation of potassium channel-complex autoimmunity. Neurology, 2012, 79, 1136-1144.	1.5	154
13	Guillain-BarrÃ© Syndrome. Mayo Clinic Proceedings, 2017, 92, 467-479.	1.4	152
14	LGI1, CASPR2 and related antibodies: a molecular evolution of the phenotypes. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 526-534.	0.9	146
15	Imaging features of copper deficiency myelopathy: a study of 25 cases. Neuroradiology, 2006, 48, 78-83.	1.1	126
16	Incidence and disease burden of chemotherapy-induced peripheral neuropathy in a population-based cohort. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 636-641.	0.9	109
17	Batch effect correction for genome-wide methylation data with Illumina Infinium platform. BMC Medical Genomics, 2011, 4, 84.	0.7	108
18	Diabetic cervical radiculoplexus neuropathy: a distinct syndrome expanding the spectrum of diabetic radiculoplexus neuropathies. Brain, 2012, 135, 3074-3088.	3.7	103

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19	Mass Spectrometricâ€“Based Proteomic Analysis of Amyloid Neuropathy Type in Nerve Tissue. Archives of Neurology, 2011, 68, 195-9.	4.9	101
20	MRI Findings in Autoimmune Voltage-Gated Potassium Channel Complex Encephalitis with Seizures: One Potential Etiology for Mesial Temporal Sclerosis. American Journal of Neuroradiology, 2014, 35, 84-89.	1.2	100
21	CSF herpes virus and autoantibody profiles in the evaluation of encephalitis. Neurology: Neuroimmunology and NeuroInflammation, 2016, 3, e245.	3.1	96
22	Positive neuropathic sensory symptoms as endpoints in diabetic neuropathy trials. Journal of the Neurological Sciences, 2001, 189, 3-5.	0.3	95
23	Lenalidomide therapy in a patient with POEMS syndrome. Blood, 2007, 110, 1075-1076.	0.6	95
24	Meralgia paresthetica. Neurology, 2011, 77, 1538-1542.	1.5	95
25	Defects of mutant DNMT1 are linked to a spectrum of neurological disorders. Brain, 2015, 138, 845-861.	3.7	94
26	Neurologic autoimmunity and immune checkpoint inhibitors. Neurology, 2020, 95, e2442-e2452.	1.5	94
27	Purkinje Cell Cytoplasmic Autoantibody Type 1 Accompaniments. Archives of Neurology, 2011, 68, 1282.	4.9	92
28	Characterization of translational frame exception patients in Duchenne/Becker muscular dystrophy. Human Molecular Genetics, 1993, 2, 737-744.	1.4	89
29	Autoimmune CRMP5 neuropathy phenotype and outcome defined from 105 cases. Neurology, 2018, 90, e103-e110.	1.5	86
30	Modeling Chronic Glycemic Exposure Variables as Correlates and Predictors of Microvascular Complications of Diabetes. Diabetes Care, 2006, 29, 2282-2288.	4.3	85
31	Inflammation and neuropathic attacks in hereditary brachial plexus neuropathy. Journal of Neurology, Neurosurgery and Psychiatry, 2002, 73, 45-50.	0.9	83
32	P/Q- and N-type calcium-channel antibodies: Oncological, neurological, and serological accompaniments. Muscle and Nerve, 2016, 54, 220-227.	1.0	83
33	The gene for HMSN2C maps to 12q23-24. Neurology, 2003, 60, 1151-1156.	1.5	80
34	Association of Long-term Opioid Therapy With Functional Status, Adverse Outcomes, and Mortality Among Patients With Polyneuropathy. JAMA Neurology, 2017, 74, 773.	4.5	80
35	The Classification of Autosomal Recessive Cerebellar Ataxias: a Consensus Statement from the Society for Research on the Cerebellum and Ataxias Task Force. Cerebellum, 2019, 18, 1098-1125.	1.4	80
36	Impaired Glycemia and Diabetic Polyneuropathy. Diabetes Care, 2012, 35, 584-591.	4.3	78

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37	Impairments and comorbidities of polyneuropathy revealed by population-based analyses. Neurology, 2015, 84, 1644-1651.	1.5	77
38	Hereditary Motor and Sensory Neuropathies: An Overview of Clinical, Genetic, Electrophysiologic, and Pathologic Features. , 2005, , 1623-1658.		71
39	Monotonicity of Nerve Tests in Diabetes: Subclinical nerve dysfunction precedes diagnosis of polyneuropathy. Diabetes Care, 2005, 28, 2192-2200.	4.3	71
40	TRPV4 mutations and cytotoxic hypercalcemia in axonal Charcot-Marie-Tooth neuropathies. Neurology, 2011, 76, 887-894.	1.5	71
41	A case of sensory ataxia as the presenting manifestation of neurosarcoidosis. Muscle and Nerve, 2011, 43, 900-905.	1.0	70
42	Predictors of neural-specific autoantibodies and immunotherapy response in patients with cognitive dysfunction. Journal of Neuroimmunology, 2018, 323, 62-72.	1.1	68
43	Sequencing of <scp>C</scp>harcotâ€“<scp>M</scp>arieâ€“<scp>T</scp>ooth disease genes in a toxic polyneuropathy. Annals of Neurology, 2014, 76, 727-737.	2.8	63
44	Natural history of spinal-bulbar muscular atrophy. Neurology, 2008, 70, 1967-1971.	1.5	62
45	Inherited neuropathies: Clinical overview and update. Muscle and Nerve, 2013, 48, 604-622.	1.0	62
46	Neural Autoantibody Clusters Aid Diagnosis of Cancer. Clinical Cancer Research, 2014, 20, 3862-3869.	3.2	62
47	Identification of Transthyretin Variants by Sequential Proteomic and Genomic Analysis. Clinical Chemistry, 2004, 50, 1544-1552.	1.5	61
48	Does impaired glucose metabolism cause polyneuropathy? Review of previous studies and design of a prospective controlled populationâ€based study. Muscle and Nerve, 2007, 36, 536-541.	1.0	61
49	Polyneuropathy improvement following autologous stem cell transplantation for POEMS syndrome. Neurology, 2015, 84, 1981-1987.	1.5	61
50	DNMT1 mutation hot spot causes varied phenotypes of HSAN1 with dementia and hearing loss. Neurology, 2013, 80, 824-828.	1.5	60
51	Retrospective study of a TTR FAP cohort to modify NIS + 7 for therapeutic trials. Journal of the Neurological Sciences, 2014, 344, 121-128.	0.3	60
52	Aberrant signature methylome by DNMT1 hot spot mutation in hereditary sensory and autonomic neuropathy 1E. Epigenetics, 2014, 9, 1184-1193.	1.3	55
53	Neurology Individualized Medicine: When to Use Next-Generation Sequencing Panels. Mayo Clinic Proceedings, 2017, 92, 292-305.	1.4	55
54	The Neuropathies of WaldenstrÃ¶m's Macroglobulinemia (WM) and IgM-MGUS. Canadian Journal of Neurological Sciences, 2011, 38, 289-295.	0.3	53

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55	Genomic analysis reveals frequent TRAF7 mutations in intraneural perineuriomas. Annals of Neurology, 2017, 81, 316-321.	2.8	53
56	LGI1 and CASPR2 neurological autoimmunity in children. Annals of Neurology, 2018, 84, 473-480.	2.8	53
57	Later-Onset Fabry Disease. Archives of Neurology, 2006, 63, 453.	4.9	52
58	Target-enrichment sequencing and copy number evaluation in inherited polyneuropathy. Neurology, 2016, 86, 1762-1771.	1.5	52
59	Infrequent SCN9A mutations in congenital insensitivity to pain and erythromelalgia. Journal of Neurology, Neurosurgery and Psychiatry, 2013, 84, 386-391.	0.9	49
60	Systematic review of autosomal recessive ataxias and proposal for a classification. Cerebellum and Ataxias, 2017, 4, 3.	1.9	49
61	LGI1 antibody encephalitis: acute treatment comparisons and outcome. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, 309-315.	0.9	48
62	Adult polyglucosan body disease: Case description of an expanding genetic and clinical syndrome. Muscle and Nerve, 2004, 29, 323-328.	1.0	47
63	A trial of proficiency of nerve conduction: Greater standardization still needed. Muscle and Nerve, 2013, 48, 369-374.	1.0	44
64	Psychiatric Autoimmunity: N-Methyl-d-Aspartate Receptor IgG and Beyond. Psychosomatics, 2015, 56, 227-241.	2.5	44
65	Pain and the immune system: emerging concepts of IgG-mediated autoimmune pain and immunotherapies. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 177-188.	0.9	44
66	Application of whole exome sequencing in undiagnosed inherited polyneuropathies. Journal of Neurology, Neurosurgery and Psychiatry, 2014, 85, 1265-1272.	0.9	43
67	Chronic meralgia paresthetica and neurectomy. Neurology, 2014, 82, 1551-1555.	1.5	43
68	Autoimmune gait disturbance accompanying adaptor protein-3B2-IgG. Neurology, 2019, 93, e954-e963.	1.5	43
69	Assessing mNIS+7_{Ionis} and international neurologists' proficiency in a familial amyloidotic polyneuropathy trial. Muscle and Nerve, 2017, 56, 901-911.	1.0	42
70	Amphiphysin-IgG autoimmune neuropathy. Neurology, 2019, 93, e1873-e1880.	1.5	41
71	Crampâ€fasciculation syndrome in patients with and without neural autoantibodies. Muscle and Nerve, 2014, 49, 351-356.	1.0	40
72	Central Nervous System Histoplasmosis Mimicking a Brain Tumor: Difficulties in Diagnosis and Treatment. Mayo Clinic Proceedings, 1999, 74, 803-807.	1.4	39

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73	Full recovery after acute hemorrhagic leukoencephalitis (Hurst's disease). Journal of Neurology, 2000, 247, 977-979.	1.8	39
74	Peripheral neuropathy incidence in inflammatory bowel disease. Neurology, 2013, 80, 1693-1697.	1.5	39
75	Multicenter trial of the proficiency of smart quantitative sensation tests. Muscle and Nerve, 2014, 49, 645-653.	1.0	39
76	Psychiatric Manifestations of Voltage-Gated Potassium-Channel Complex Autoimmunity. Journal of Neuropsychiatry and Clinical Neurosciences, 2011, 23, 425-433.	0.9	37
77	Antibodies to myelinâ€associated glycoprotein (antiâ€Mag) in IgM amyloidosis may influence expression of neuropathy in rare patients. Muscle and Nerve, 2008, 37, 490-495.	1.0	35
78	Elevated <scp>LGI</scp>1â€IgG <scp>CSF</scp> index predicts worse neurological outcome. Annals of Clinical and Translational Neurology, 2018, 5, 646-650.	1.7	35
79	â€œUnequivocally Abnormalâ€•vs â€œUsualâ€•Signs and Symptoms for Proficient Diagnosis of Diabetic Polyneuropathy. Archives of Neurology, 2012, 69, 1609.	4.9	33
80	Striational antibodies in a paraneoplastic context. Muscle and Nerve, 2013, 47, 585-587.	1.0	33
81	Proficiency of nerve conduction using standard methods and reference values (cl. NPhys Trial 4). Muscle and Nerve, 2014, 50, 900-908.	1.0	32
82	Isolated amyloidosis presenting with lumbosacral radiculoplexopathy: description of two cases and pathogenic review. Journal of the Peripheral Nervous System, 2006, 11, 346-352.	1.4	31
83	Elevated creatine kinase and transaminases in asymptomatic SBMA. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2007, 8, 62-64.	2.3	29
84	Autoimmune/Paraneoplastic Encephalitis Antibody Biomarkers: Frequency, Age, and Sex Associations. Mayo Clinic Proceedings, 2022, 97, 547-559.	1.4	29
85	Ubiquitin ligase defect by DCAF8 mutation causes HMSN2 with giant axons. Neurology, 2014, 82, 873-878.	1.5	28
86	Two-stage Study of Familial Prostate Cancer by Whole-exome Sequencing and Custom Capture Identifies 10 Novel Genes Associated with the Risk of Prostate Cancer. European Urology, 2021, 79, 353-361.	0.9	28
87	Antiepileptic drug therapy in autoimmune epilepsy associated with antibodies targeting the leucineâ€rich gliomaâ€inactivated protein 1. Epilepsia Open, 2018, 3, 348-356.	1.3	26
88	Paraneoplastic Myeloneuropathies. Neurology, 2021, 96, e632-e639.	1.5	26
89	SPTLC1 and RAB7 mutation analysis in dominantly inherited and idiopathic sensory neuropathies. Journal of Neurology, Neurosurgery and Psychiatry, 2005, 76, 1022-1024.	0.9	25
90	Progressive Multifocal Leukoencephalopathy: a rare infectious complication following allogeneic hematopoietic cell transplantation (<scp>HCT</scp>). European Journal of Haematology, 2014, 92, 83-87.	1.1	25

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91	Recurrent Genomic Alterations in Soft Tissue Perineuriomas. American Journal of Surgical Pathology, 2018, 42, 1708-1714.	2.1	25
92	Collapsin Response-Mediator Protein 5â€“Associated Retinitis, Vitritis, and Optic Disc Edema. Ophthalmology, 2020, 127, 221-229.	2.5	25
93	Phenotypic presentations of paraneoplastic neuropathies associated with MAP1B-IgG. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 328-330.	0.9	25
94	Large Kindred Evaluation of Mitofusin 2 Novel Mutation, Extremes of Neurologic Presentations, and Preserved Nerve Mitochondria. Archives of Neurology, 2011, 68, 1295.	4.9	24
95	Epidermal nerve fibers. Neurology, 2012, 79, 2187-2193.	1.5	24
96	The Inherited Neuropathies. Neurologic Clinics, 2007, 25, 173-207.	0.8	23
97	Potassium channel complex autoimmunity induced by inhaled brain tissue aerosol. Annals of Neurology, 2012, 71, 417-426.	2.8	23
98	Amyloidâ€like IgM deposition neuropathy: a distinct clinicoâ€pathologic and proteomic profiled disorder. Journal of the Peripheral Nervous System, 2012, 17, 182-190.	1.4	23
99	IgM-monoclonal gammopathy neuropathy and tremor: A first epidemiologic case control study. Parkinsonism and Related Disorders, 2012, 18, 748-752.	1.1	22
100	Brachial Plexus Neuritis Associated With Antiâ€“Programmed Cell Death-1 Antibodies: Report of 2 Cases. Mayo Clinic Proceedings Innovations, Quality & Outcomes, 2017, 1, 192-197.	1.2	22
101	Neurologic Course, Endocrine Dysfunction and Triplet Repeat Size in Spinal Bulbar Muscular Atrophy. Canadian Journal of Neurological Sciences, 2004, 31, 378-382.	0.3	21
102	Quantitative Sensation Testing. , 2005, , 1063-1093.		21
103	Clinicalâ€pathologic correlations in voltageâ€gated Kv1 potassium channel complexâ€subtyped autoimmune painful polyneuropathy. Muscle and Nerve, 2017, 55, 520-525.	1.0	20
104	T-cell neurolymphomatosis involving cauda equina and sciatic nerves. Neurology, 2009, 72, 98-98.	1.5	19
105	Case of mononeuritis multiplex onset with rituximab therapy for WaldenstrÃ¶m's macroglobulinemia. Journal of the Neurological Sciences, 2007, 260, 240-243.	0.3	18
106	Small Fiber Neuropathy Incidence, Prevalence, Longitudinal Impairments, and Disability. Neurology, 2021, 97, e2236-e2247.	1.5	18
107	Novel myelin protein zero mutation (Arg36Trp) in a patient with acute onset painful neuropathy. Neuromuscular Disorders, 2006, 16, 308-310.	0.3	17
108	Autoimmune-mediated peripheral neuropathies and autoimmune pain. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2016, 133, 417-446.	1.0	17

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109	Transthyretin amyloid neuropathy has earlier neural involvement but better prognosis than primary amyloid counterpart: an answer to the paradox?. Annals of Neurology, 2016, 80, 401-411.	2.8	17
110	Guillain-BarrÃ© Syndrome in a Patient With Evidence of Recent SARS-CoV-2 Infection. Mayo Clinic Proceedings, 2020, 95, 1799-1801.	1.4	17
111	Neurofascin-155 Immunoglobulin Subtypes. Neurology, 2021, 97, .	1.5	17
112	Epigenetic regulation: Basic concepts and relevance to neurologic disease. Neurology, 2014, 82, 1833-1840.	1.5	16
113	Office immunotherapy in chronic inflammatory demyelinating polyneuropathy and multifocal motor neuropathy. Muscle and Nerve, 2015, 52, 488-497.	1.0	16
114	The utility of genetic testing in neuromuscular disease: A consensus statement from the AANEM on the clinical utility of genetic testing in diagnosis of neuromuscular disease. Muscle and Nerve, 2016, 54, 1007-1009.	1.0	16
115	Contactin-1 autoimmunity. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, e771.	3.1	15
116	Charcot-Marie-Tooth Disease and Other Hereditary Neuropathies. CONTINUUM Lifelong Learning in Neurology, 2020, 26, 1224-1256.	0.4	15
117	Surgical and postpartum hereditary brachial plexus attacks and prophylactic immunotherapy. Muscle and Nerve, 2013, 47, 23-27.	1.0	14
118	Clinical, physiological and pathological characterisation of the sensory predominant peripheral neuropathy in copper deficiency. Journal of Neurology, Neurosurgery and Psychiatry, 2017, 88, 839-845.	0.9	14
119	Onionâ€bulb patterns predict acquired or inherited demyelinating polyneuropathy. Muscle and Nerve, 2019, 59, 665-670.	1.0	14
120	Improving accuracy of myasthenia gravis autoantibody testing by reflex algorithm. Neurology, 2020, 95, e3002-e3011.	1.5	14
121	GFAP IgG associated inflammatory polyneuropathy. Journal of Neuroimmunology, 2020, 343, 577233.	1.1	14
122	Genetic testing in inherited peripheral neuropathies. Journal of the Peripheral Nervous System, 2005, 10, 77-84.	1.4	13
123	Expanded teased nerve fibre pathological conditions in disease association. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 138-140.	0.9	13
124	Comparison of immune checkpoint inhibitor-related neuropathies among patients with neuroendocrine and non-neuroendocrine tumours. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, 112-114.	0.9	13
125	Hypertrophic nerves producing myelopathy in fulminant CIDP. Neurology, 2010, 75, 750-750.	1.5	12
126	Neuronal intracellular transport and neurodegenerative disease. Neurology, 2011, 76, 1015-1020.	1.5	12

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127	Composite ganglioside autoantibodies and immune treatment response in MMN and MADSAM. Muscle and Nerve, 2018, 57, 1000-1005.	1.0	12
128	Genetic variation in the B7-1 gene in patients with multiple sclerosis. Journal of Neuroimmunology, 2000, 105, 184-188.	1.1	10
129	Bulbospinal Muscular Atrophy. Archives of Neurology, 2004, 61, 1324-6.	4.9	10
130	Pathology and molecular genetics of inherited neuropathy. Journal of the Neurological Sciences, 2004, 220, 141-143.	0.3	10
131	HSANs: Clinical Features, Pathologic Classification, and Molecular Genetics. , 2005, , 1809-1844.		10
132	SEPT9 Mutations and a Conserved 17q25 Sequence in Sporadic and Hereditary Brachial Plexus Neuropathy. Archives of Neurology, 2009, 66, 238-43.	4.9	10
133	Copy number analysis reveals a novel multiexon deletion of the COLQ gene in congenital myasthenia. Neurology: Genetics, 2016, 2, e117.	0.9	10
134	gsSKAT: Rapid gene set analysis and multiple testing correction for rareâ€variant association studies using weighted linear kernels. Genetic Epidemiology, 2017, 41, 297-308.	0.6	9
135	Blink R1 latency utility in diagnosis and treatment assessment of polyradiculoneuropathyâ€organomegalyâ€endocrinopathyâ€monoclonal proteinâ€skin changes and chronic inflammatory demyelinating polyradiculoneuropathy. Muscle and Nerve, 2018, 57, E8-E13.	1.0	9
136	Recurrent Brachial Neuritis Attacks in Presentation of B-Cell Lymphoma. Mayo Clinic Proceedings Innovations, Quality & Outcomes, 2018, 2, 382-386.	1.2	9
137	DNA methylation patterns in human iPSC-derived sensory neuronal differentiation. Epigenetics, 2019, 14, 927-937.	1.3	9
138	Filamentous tangles with nemaline rods in MYH2 myopathy: a novel phenotype. Acta Neuropathologica Communications, 2021, 9, 79.	2.4	9
139	Diagnostic modelling and therapeutic monitoring of immune-mediated necrotizing myopathy: role of electrical myotonia. Brain Communications, 2020, 2, fcaa191.	1.5	9
140	Mutant C. elegans mitofusin leads to selective removal of mtDNA heteroplasmic deletions across generations to maintain fitness. BMC Biology, 2022, 20, 40.	1.7	9
141	Indifference rather than insensitivity to pain. Annals of Neurology, 2003, 53, 417-418.	2.8	8
142	Nerve Tests Expressed as Percentiles, Normal Deviates, and Composite Scores. , 2005, , 971-983.		8
143	Immunotherapy-responsive pain in an abattoir worker with fluctuating potassium channel-complex IgG. Neurology, 2012, 79, 1824-1825.	1.5	8
144	Mass spectrometry analysis reveals nonâ€mutated apolipoprotein a1 lumbosacral radiculoplexus amyloidoma. Muscle and Nerve, 2012, 46, 817-822.	1.0	8

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145	Blink reflex role in algorithmic genetic testing of inherited polyneuropathies. Muscle and Nerve, 2017, 55, 316-322.	1.0	8
146	Polyneuropathies and chronic inflammatory demyelinating polyradiculoneuropathy in multiple sclerosis. Multiple Sclerosis and Related Disorders, 2019, 30, 284-290.	0.9	8
147	Association of intraneural perineurioma with neurofibromatosis type 2. Acta Neurochirurgica, 2020, 162, 1891-1897.	0.9	8
148	Expanding the Spectrum of Chronic Immune Sensory Polyradiculopathy. Neurology, 2021, 96, e2078-e2089.	1.5	8
149	Genetic variants related to successful migraine prophylaxis with verapamil. Molecular Genetics & Genomic Medicine, 2021, 9, e1680.	0.6	8
150	Incidence and prevalence of immuneâ€mediated necrotizing myopathy in adults in Olmsted County, Minnesota. Muscle and Nerve, 2022, 65, 541-546.	1.0	8
151	LRP4-IgG service line testing in seronegative myasthenia gravis and controls. Journal of Neuroimmunology, 2022, 368, 577895.	1.1	8
152	Disseminated sporotrichosis presenting with granulomatous inflammatory multiple mononeuropathies. Muscle and Nerve, 2007, 36, 866-872.	1.0	7
153	Pediatric Nerve Biopsy Diagnostic and Treatment Utility inÂTertiary Care Referral. Pediatric Neurology, 2016, 58, 3-11.	1.0	7
154	Expanded genetic insight and clinical experience of DNMT1-complex disorder. Neurology: Genetics, 2020, 6, e456.	0.9	7
155	Clinical Utility of Striational Antibodies in Paraneoplastic and Myasthenia Gravis Paraneoplastic Panels. Neurology, 2021, , 10.1212/WNL.0000000000012050.	1.5	7
156	Hereditary Brachial Plexus Neuropathy. , 2005, , 1753-1767.		7
157	Inclusion body myositis: correlation of clinical outcomes with histopathology, electromyography and laboratory findings. Rheumatology, 2022, 61, 2504-2511.	0.9	7
158	Wild-type TTR neuropathy with cardiomyopathy presenting with burning feet. Neurology, 2017, 88, 1101-1102.	1.5	6
159	Subacute demyelinating polyradiculoneuropathy complicating Epsteinâ€“Barr virus infection in GATA2 haploinsufficiency. Muscle and Nerve, 2018, 57, 150-156.	1.0	6
160	Expanded neuromuscular morbidity in Hodgkin lymphoma after radiotherapy. Brain Communications, 2020, 2, fcaa050.	1.5	6
161	Novel SOD1 mutation discovered in atypical ALS by whole exome sequencing. Journal of Neurology, Neurosurgery and Psychiatry, 2013, 84, 943-944.	0.9	5
162	Post hoc Analysis for Detecting Individual Rare Variant Risk Associations Using Probit Regression Bayesian Variable Selection Methods in Caseâ€Control Sequencing Studies. Genetic Epidemiology, 2016, 40, 461-469.	0.6	5

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163	Mitochondrial cerebellar ataxia, renal failure, neuropathy, and encephalopathy (MCARNE). Neurology: Genetics, 2019, 5, e314.	0.9	5
164	Plantar Ulcers and Neuropathic Arthropathies: Associated Diseases, Polyneuropathy Correlates, and Risk Covariates. Advances in Skin and Wound Care, 2019, 32, 168-175.	0.5	5
165	Pure Motor Onset and IgM-Gammopathy Occurrence in Multifocal Acquired Demyelinating Sensory and Motor Neuropathy. Neurology, 2021, 97, .	1.5	5
166	Cancer and immune-mediated necrotizing myopathy: a longitudinal referral case-controlled outcomes evaluation. Rheumatology, 2022, 62, 281-289.	0.9	5
167	Cryoglobulinemic vasculitis in a patient with CREST syndrome. Journal of Clinical Neuroscience, 2014, 21, 1821-1823.	0.8	4
168	Bilateral Facial and Trigeminal Nerve Hypertrophy in a Patient With Polyneuropathy. Otology and Neurotology, 2016, 37, e404-e406.	0.7	4
169	Somatotopic heat pain thresholds and intraepidermal nerve fibers in health. Muscle and Nerve, 2018, 58, 509-516.	1.0	4
170	Fatal TTR amyloidosis with neuropathy from domino liver p.Val71Ala transplant. Neurology: Genetics, 2019, 5, e351.	0.9	4
171	IgM-gammopathy strongly favours immune treatable MMN and MADSAM over ALS. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 324-326.	0.9	4
172	IVIG and Small Fiber Neuropathy. Neurology, 2021, 96, 929-930.	1.5	4
173	HMSN II (CMT2) and Miscellaneous Inherited System Atrophies of Nerve Axon: Clinicalâ€“Molecular Genetic Correlates. , 2005, , 1717-1751.		4
174	Solving the Riddle of the Sphinx May Provide New Insights Into Diabetes and Polyneuropathy. Diabetes, 2015, 64, 706-708.	0.3	3
175	Novel hemizygous nonsense mutation in DRP2 is associated with inherited neuropathy. Neurology: Genetics, 2018, 4, e220.	0.9	3
176	Teaching Video NeuroImages: Acquired focal neuromyotonia in LGI-1 autoimmunity. Neurology, 2018, 90, e1636-e1637.	1.5	3
177	Multifocal neuropathy as the presenting symptom of Purkinje cell cytoplasmic autoantibody-1. Muscle and Nerve, 2013, 48, 827-831.	1.0	2
178	Radiation-induced spinal nerve root cavernous malformations as a rare cause of radiculopathy. Neurology, 2017, 89, 2299-2300.	1.5	2
179	Inherited neuropathy precision classification. Neurology, 2018, 90, 445-446.	1.5	2
180	CASPR2-IgGâ€“Associated Autoimmune Encephalitis: Unusual Cause of Delirium. Psychosomatics, 2020, 61, 774-778.	2.5	2

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