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	Treatment and Management of Autosomal Recessive Cerebellar Ataxias: Current Advances and Future Perspectives. CNS and Neurological Disorders - Drug Targets, 2023, 22, 678-697.	0.8	1
2	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
3	Inclusion body myositis: correlation of clinical outcomes with histopathology, electromyography and laboratory findings. Rheumatology, 2022, 61, 2504-2511.	0.9	7
4	LGI1 antibody encephalitis: acute treatment comparisons and outcome. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, 309-315.	0.9	48
5	Incidence and prevalence of immuneâ€mediated necrotizing myopathy in adults in Olmsted County, Minnesota. Muscle and Nerve, 2022, 65, 541-546.	1.0	8
6	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
7	Autoimmune/Paraneoplastic Encephalitis Antibody Biomarkers: Frequency, Age, and Sex Associations. Mayo Clinic Proceedings, 2022, 97, 547-559.	1.4	29
8	Cancer and immune-mediated necrotizing myopathy: a longitudinal referral case-controlled outcomes evaluation. Rheumatology, 2022, 62, 281-289.	0.9	5
9	Letter to editor: Orofacial overgrowth with peripheral nerve enlargement and perineuriomatous pseudo-onion bulb proliferations is part of the PIK3CA-related overgrowth spectrum. Human Genetics and Genomics Advances, 2022, 3, 100110.	1.0	1
10	LRP4-IgG service line testing in seronegative myasthenia gravis and controls. Journal of Neuroimmunology, 2022, 368, 577895.	1.1	8
11	64-Year-Old Woman With Aphasia and Troponin Elevation. Mayo Clinic Proceedings, 2022, , .	1.4	0
12	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
13	Paraneoplastic Myeloneuropathies. Neurology, 2021, 96, e632-e639.	1.5	26
14	IVIG and Small Fiber Neuropathy. Neurology, 2021, 96, 929-930.	1.5	4
15	Expanding the Spectrum of Chronic Immune Sensory Polyradiculopathy. Neurology, 2021, 96, e2078-e2089.	1.5	8
16	Filamentous tangles with nemaline rods in MYH2 myopathy: a novel phenotype. Acta Neuropathologica Communications, 2021, 9, 79.	2.4	9
17	Genetic variants related to successful migraine prophylaxis with verapamil. Molecular Genetics & Genomic Medicine, 2021, 9, e1680.	0.6	8
18	Clinical Utility of Striational Antibodies in Paraneoplastic and Myasthenia Gravis Paraneoplastic Panels. Neurology, 2021, , 10.1212/WNL.0000000000012050.	1.5	7

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19	003â€…Autoimmune encephalitis antibody biomarkers: frequency, age and sex associations. , 2021, , .		0
20	Pure Motor Onset and IgM-Gammopathy Occurrence in Multifocal Acquired Demyelinating Sensory and Motor Neuropathy. Neurology, 2021, 97, .	1.5	5
21	Neurofascin-155 Immunoglobulin Subtypes. Neurology, 2021, 97, .	1.5	17
22	Small Fiber Neuropathy Incidence, Prevalence, Longitudinal Impairments, and Disability. Neurology, 2021, 97, e2236-e2247.	1.5	18
23	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
24	Collapsin Response-Mediator Protein 5â€“Associated Retinitis, Vitritis, and Optic Disc Edema. Ophthalmology, 2020, 127, 221-229.	2.5	25
25	IgM-gammopathy strongly favours immune treatable MMN and MADSAM over ALS. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 324-326.	0.9	4
26	Phenotypic presentations of paraneoplastic neuropathies associated with MAP1B-IgG. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 328-330.	0.9	25
27	Expanded neuromuscular morbidity in Hodgkin lymphoma after radiotherapy. Brain Communications, 2020, 2, fcaa050.	1.5	6
28	Guillain-BarrÃ© Syndrome in a Patient With Evidence of Recent SARS-CoV-2 Infection. Mayo Clinic Proceedings, 2020, 95, 1799-1801.	1.4	17
29	Expanded genetic insight and clinical experience of DNMT1-complex disorder. Neurology: Genetics, 2020, 6, e456.	0.9	7
30	Improving accuracy of myasthenia gravis autoantibody testing by reflex algorithm. Neurology, 2020, 95, e3002-e3011.	1.5	14
31	Neurologic autoimmunity and immune checkpoint inhibitors. Neurology, 2020, 95, e2442-e2452.	1.5	94
32	Contactin-1 autoimmunity. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, e771.	3.1	15
33	CASPR2-IgGâ€“Associated Autoimmune Encephalitis: Unusual Cause of Delirium. Psychosomatics, 2020, 61, 774-778.	2.5	2
34	Association of intraneural perineurioma with neurofibromatosis type 2. Acta Neurochirurgica, 2020, 162, 1891-1897.	0.9	8
35	Phrenic neuropathy water immersion dyspnea. Neurology, 2020, 94, e1314-e1319.	1.5	1
36	GFAP IgG associated inflammatory polyneuropathy. Journal of Neuroimmunology, 2020, 343, 577233.	1.1	14

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37	Charcot-Marie-Tooth Disease and Other Hereditary Neuropathies. CONTINUUM Lifelong Learning in Neurology, 2020, 26, 1224-1256.	0.4	15
38	Diagnostic modelling and therapeutic monitoring of immune-mediated necrotizing myopathy: role of electrical myotonia. Brain Communications, 2020, 2, fcaa191.	1.5	9
39	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
40	Amphiphysin-IgG autoimmune neuropathy. Neurology, 2019, 93, e1873-e1880.	1.5	41
41	Embolized Spinal Dural AV Fistula Repairs Syringomyelia and Polyradiculopathy. Mayo Clinic Proceedings, 2019, 94, 1904-1906.	1.4	1
42	Autoimmune gait disturbance accompanying adaptor protein-3B2-IgG. Neurology, 2019, 93, e954-e963.	1.5	43
43	DNA methylation patterns in human iPSC-derived sensory neuronal differentiation. Epigenetics, 2019, 14, 927-937.	1.3	9
44	Polyneuropathies and chronic inflammatory demyelinating polyradiculoneuropathy in multiple sclerosis. Multiple Sclerosis and Related Disorders, 2019, 30, 284-290.	0.9	8
45	Mitochondrial cerebellar ataxia, renal failure, neuropathy, and encephalopathy (MCARNE). Neurology: Genetics, 2019, 5, e314.	0.9	5
46	Onionâ€bulb patterns predict acquired or inherited demyelinating polyneuropathy. Muscle and Nerve, 2019, 59, 665-670.	1.0	14
47	Fatal TTR amyloidosis with neuropathy from domino liver p.Val71Ala transplant. Neurology: Genetics, 2019, 5, e351.	0.9	4
48	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
49	Targeted gene approach with biochemical assay confirms ABCD1 mutation of X-linked adrenoleukodystrophy in a 62-year-old man with gait imbalance. Neuromuscular Disorders, 2019, 29, 146-149.	0.3	1
50	Expanded teased nerve fibre pathological conditions in disease association. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 138-140.	0.9	13
51	Novel hemizygous nonsense mutation in DRP2 is associated with inherited neuropathy. Neurology: Genetics, 2018, 4, e220.	0.9	3
52	Inherited neuropathy precision classification. Neurology, 2018, 90, 445-446.	1.5	2
53	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
54	Composite ganglioside autoantibodies and immune treatment response in MMN and MADSAM. Muscle and Nerve, 2018, 57, 1000-1005.	1.0	12

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55	Autoimmune encephalitis epidemiology and a comparison to infectious encephalitis. Annals of Neurology, 2018, 83, 166-177.	2.8	479
56	Teaching Video NeuroImages: Acquired focal neuromyotonia in LGI-1 autoimmunity. Neurology, 2018, 90, e1636-e1637.	1.5	3
57	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
58	Somatotopic heat pain thresholds and intraepidermal nerve fibers in health. Muscle and Nerve, 2018, 58, 509-516.	1.0	4
59	Subacute demyelinating polyradiculoneuropathy complicating Epsteinâ€“Barr virus infection in GATA2 haploinsufficiency. Muscle and Nerve, 2018, 57, 150-156.	1.0	6
60	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
61	LGI1, CASPR2 and related antibodies: a molecular evolution of the phenotypes. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 526-534.	0.9	146
62	Autoimmune CRMP5 neuropathy phenotype and outcome defined from 105 cases. Neurology, 2018, 90, e103-e110.	1.5	86
63	Recurrent Genomic Alterations in Soft Tissue Perineuriomas. American Journal of Surgical Pathology, 2018, 42, 1708-1714.	2.1	25
64	Recurrent Brachial Neuritis Attacks in Presentation of B-Cell Lymphoma. Mayo Clinic Proceedings Innovations, Quality & Outcomes, 2018, 2, 382-386.	1.2	9
65	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
66	LGI1 and CASPR2 neurological autoimmunity in children. Annals of Neurology, 2018, 84, 473-480.	2.8	53
67	Predictors of neural-specific autoantibodies and immunotherapy response in patients with cognitive dysfunction. Journal of Neuroimmunology, 2018, 323, 62-72.	1.1	68
68	Assessing mNIS+7_{Ionis} and international neurologists' proficiency in a familial amyloidotic polyneuropathy trial. Muscle and Nerve, 2017, 56, 901-911.	1.0	42
69	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
70	Guillain-BarrÃ© Syndrome. Mayo Clinic Proceedings, 2017, 92, 467-479.	1.4	152
71	Neurology Individualized Medicine: When to Use Next-Generation Sequencing Panels. Mayo Clinic Proceedings, 2017, 92, 292-305.	1.4	55
72	Systematic review of autosomal recessive ataxias and proposal for a classification. Cerebellum and Ataxias, 2017, 4, 3.	1.9	49

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73	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
74	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
75	Wild-type TTR neuropathy with cardiomyopathy presenting with burning feet. Neurology, 2017, 88, 1101-1102.	1.5	6
76	Genomic analysis reveals frequent TRAF7 mutations in intraneural perineuriomas. Annals of Neurology, 2017, 81, 316-321.	2.8	53
77	Neurological Complications Associated With Antiâ€“Programmed Death 1 (PD-1) Antibodies. JAMA Neurology, 2017, 74, 1216.	4.5	244
78	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
79	Radiation-induced spinal nerve root cavernous malformations as a rare cause of radiculopathy. Neurology, 2017, 89, 2299-2300.	1.5	2
80	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
81	Clinicalâ€pathologic correlations in voltageâ€gated Kv1 potassium channel complexâ€subtyped autoimmune painful polyneuropathy. Muscle and Nerve, 2017, 55, 520-525.	1.0	20
82	Blink reflex role in algorithmic genetic testing of inherited polyneuropathies. Muscle and Nerve, 2017, 55, 316-322.	1.0	8
83	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
84	Bilateral Facial and Trigeminal Nerve Hypertrophy in a Patient With Polyneuropathy. Otology and Neurotology, 2016, 37, e404-e406.	0.7	4
85	Target-enrichment sequencing and copy number evaluation in inherited polyneuropathy. Neurology, 2016, 86, 1762-1771.	1.5	52
86	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
87	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
88	Targeting sequence domain: a captain at the helm precisely steering DNMT1 through maintenance methylation?. Epigenomics, 2016, 8, 737-740.	1.0	1
89	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
90	Copy number analysis reveals a novel multiexon deletion of the COLQ gene in congenital myasthenia. Neurology: Genetics, 2016, 2, e117.	0.9	10

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91	P/Q- and N-type calcium-channel antibodies: Oncological, neurological, and serological accompaniments. Muscle and Nerve, 2016, 54, 220-227.	1.0	83
92	Thirty Years Later, Case Closed: A Case of PMP22 Triplication From Anticipation. Mayo Clinic Proceedings, 2016, 91, 687-688.	1.4	1
93	CSF herpes virus and autoantibody profiles in the evaluation of encephalitis. Neurology: Neuroimmunology and NeuroInflammation, 2016, 3, e245.	3.1	96
94	Pediatric Nerve Biopsy Diagnostic and Treatment Utility inÂTertiary Care Referral. Pediatric Neurology, 2016, 58, 3-11.	1.0	7
95	Office immunotherapy in chronic inflammatory demyelinating polyneuropathy and multifocal motor neuropathy. Muscle and Nerve, 2015, 52, 488-497.	1.0	16
96	Basal ganglia T1 hyperintensity in LGI1-autoantibody faciobrachial dystonic seizures. Neurology: Neuroimmunology and NeuroInflammation, 2015, 2, e161.	3.1	163
97	Psychiatric Autoimmunity: N-Methyl-d-Aspartate Receptor IgG and Beyond. Psychosomatics, 2015, 56, 227-241.	2.5	44
98	Impairments and comorbidities of polyneuropathy revealed by population-based analyses. Neurology, 2015, 84, 1644-1651.	1.5	77
99	Polyneuropathy improvement following autologous stem cell transplantation for POEMS syndrome. Neurology, 2015, 84, 1981-1987.	1.5	61
100	Defects of mutant DNMT1 are linked to a spectrum of neurological disorders. Brain, 2015, 138, 845-861.	3.7	94
101	Solving the Riddle of the Sphinx May Provide New Insights Into Diabetes and Polyneuropathy. Diabetes, 2015, 64, 706-708.	0.3	3
102	Peripheral Nerve Amyloidosis. Current Clinical Pathology, 2015, , 437-450.	0.0	1
103	Application of whole exome sequencing in undiagnosed inherited polyneuropathies. Journal of Neurology, Neurosurgery and Psychiatry, 2014, 85, 1265-1272.	0.9	43
104	Neural Autoantibody Clusters Aid Diagnosis of Cancer. Clinical Cancer Research, 2014, 20, 3862-3869.	3.2	62
105	Ubiquitin ligase defect by DCAF8 mutation causes HMSN2 with giant axons. Neurology, 2014, 82, 873-878.	1.5	28
106	Langerhans cell activation in diabetic small fiber polyneuropathy. Neurology: Neuroimmunology and NeuroInflammation, 2014, 1, e42.	3.1	1
107	Chronic meralgia paresthetica and neurectomy. Neurology, 2014, 82, 1551-1555.	1.5	43
108	Epigenetic regulation: Basic concepts and relevance to neurologic disease. Neurology, 2014, 82, 1833-1840.	1.5	16

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109	Multicenter trial of the proficiency of smart quantitative sensation tests. Muscle and Nerve, 2014, 49, 645-653.	1.0	39
110	Aberrant signature methylome by DNMT1 hot spot mutation in hereditary sensory and autonomic neuropathy 1E. Epigenetics, 2014, 9, 1184-1193.	1.3	55
111	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
112	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
113	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
114	Crampâ€fasciculation syndrome in patients with and without neural autoantibodies. Muscle and Nerve, 2014, 49, 351-356.	1.0	40
115	Cryoglobulinemic vasculitis in a patient with CREST syndrome. Journal of Clinical Neuroscience, 2014, 21, 1821-1823.	0.8	4
116	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
117	Proficiency of nerve conduction using standard methods and reference values (cl. NPhys Trial 4). Muscle and Nerve, 2014, 50, 900-908.	1.0	32
118	Inherited neuropathies: Clinical overview and update. Muscle and Nerve, 2013, 48, 604-622.	1.0	62
119	Reply. Muscle and Nerve, 2013, 48, 625-625.	1.0	1
120	Surgical and postpartum hereditary brachial plexus attacks and prophylactic immunotherapy. Muscle and Nerve, 2013, 47, 23-27.	1.0	14
121	Novel SOD1 mutation discovered in atypical ALS by whole exome sequencing. Journal of Neurology, Neurosurgery and Psychiatry, 2013, 84, 943-944.	0.9	5
122	Insights From LGI1 and CASPR2 Potassium Channel Complex Autoantibody Subtyping. JAMA Neurology, 2013, 70, 229.	4.5	170
123	DNMT1 mutation hot spot causes varied phenotypes of HSAN1 with dementia and hearing loss. Neurology, 2013, 80, 824-828.	1.5	60
124	Infrequent SCN9A mutations in congenital insensitivity to pain and erythromelalgia. Journal of Neurology, Neurosurgery and Psychiatry, 2013, 84, 386-391.	0.9	49
125	Peripheral neuropathy incidence in inflammatory bowel disease. Neurology, 2013, 80, 1693-1697.	1.5	39
126	A trial of proficiency of nerve conduction: Greater standardization still needed. Muscle and Nerve, 2013, 48, 369-374.	1.0	44

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127	Multifocal neuropathy as the presenting symptom of Purkinje cell cytoplasmic autoantibody-1. Muscle and Nerve, 2013, 48, 827-831.	1.0	2
128	Striational antibodies in a paraneoplastic context. Muscle and Nerve, 2013, 47, 585-587.	1.0	33
129	Immunotherapy-responsive pain in an abattoir worker with fluctuating potassium channel-complex IgG. Neurology, 2012, 79, 1824-1825.	1.5	8
130	Autoimmune Epilepsy. Archives of Neurology, 2012, 69, 582.	4.9	324
131	Impaired Glycemia and Diabetic Polyneuropathy. Diabetes Care, 2012, 35, 584-591.	4.3	78
132	Diabetic cervical radiculoplexus neuropathy: a distinct syndrome expanding the spectrum of diabetic radiculoplexus neuropathies. Brain, 2012, 135, 3074-3088.	3.7	103
133	Epidermal nerve fibers. Neurology, 2012, 79, 2187-2193.	1.5	24
134	Chronic pain as a manifestation of potassium channel-complex autoimmunity. Neurology, 2012, 79, 1136-1144.	1.5	154
135	â€œUnequivocally Abnormalâ€•vs â€œUsualâ€•Signs and Symptoms for Proficient Diagnosis of Diabetic Polyneuropathy. Archives of Neurology, 2012, 69, 1609.	4.9	33
136	Mass spectrometry analysis reveals nonâ€mutated apolipoprotein a1 lumbosacral radiculoplexus amyloidoma. Muscle and Nerve, 2012, 46, 817-822.	1.0	8
137	IgM-monoclonal gammopathy neuropathy and tremor: A first epidemiologic case control study. Parkinsonism and Related Disorders, 2012, 18, 748-752.	1.1	22
138	Potassium channel complex autoimmunity induced by inhaled brain tissue aerosol. Annals of Neurology, 2012, 71, 417-426.	2.8	23
139	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
140	Peripheral Nerve Amyloidosis. , 2012, , 361-374.		1
141	Psychiatric Manifestations of Voltage-Gated Potassium-Channel Complex Autoimmunity. Journal of Neuropsychiatry and Clinical Neurosciences, 2011, 23, 425-433.	0.9	37
142	Purkinje Cell Cytoplasmic Autoantibody Type 1 Accompaniments. Archives of Neurology, 2011, 68, 1282.	4.9	92
143	The Neuropathies of WaldenstrÃ¶m's Macroglobulinemia (WM) and IgM-MGUS. Canadian Journal of Neurological Sciences, 2011, 38, 289-295.	0.3	53
144	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

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145	Mutations in DNMT1 cause hereditary sensory neuropathy with dementia and hearing loss. Nature Genetics, 2011, 43, 595-600.	9.4	342
146	Batch effect correction for genome-wide methylation data with Illumina Infinium platform. BMC Medical Genomics, 2011, 4, 84.	0.7	108
147	A case of sensory ataxia as the presenting manifestation of neurosarcoidosis. Muscle and Nerve, 2011, 43, 900-905.	1.0	70
148	Mass Spectrometricâ€“Based Proteomic Analysis of Amyloid Neuropathy Type in Nerve Tissue. Archives of Neurology, 2011, 68, 195-9.	4.9	101
149	Meralgia paresthetica. Neurology, 2011, 77, 1538-1542.	1.5	95
150	Neuronal intracellular transport and neurodegenerative disease. Neurology, 2011, 76, 1015-1020.	1.5	12
151	TRPV4 mutations and cytotoxic hypercalcemia in axonal Charcot-Marie-Tooth neuropathies. Neurology, 2011, 76, 887-894.	1.5	71
152	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
153	Scapuloperoneal spinal muscular atrophy and CMT2C are allelic disorders caused by alterations in TRPV4. Nature Genetics, 2010, 42, 165-169.	9.4	232
154	Hypertrophic nerves producing myelopathy in fulminant CIDP. Neurology, 2010, 75, 750-750.	1.5	12
155	Post-surgical inflammatory neuropathy. Brain, 2010, 133, 2866-2880.	3.7	203
156	T-cell neurolymphomatosis involving cauda equina and sciatic nerves. Neurology, 2009, 72, 98-98.	1.5	19
157	SEPT9 Mutations and a Conserved 17q25 Sequence in Sporadic and Hereditary Brachial Plexus Neuropathy. Archives of Neurology, 2009, 66, 238-43.	4.9	10
158	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
159	Voltage-Gated Potassium Channel Autoimmunity Mimicking Creutzfeldt-Jakob Disease. Archives of Neurology, 2008, 65, 1341-6.	4.9	166
160	Natural history of spinal-bulbar muscular atrophy. Neurology, 2008, 70, 1967-1971.	1.5	62
161	Elevated creatine kinase and transaminases in asymptomatic SBMA. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2007, 8, 62-64.	2.3	29
162	Lenalidomide therapy in a patient with POEMS syndrome. Blood, 2007, 110, 1075-1076.	0.6	95

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163	The Inherited Neuropathies. Neurologic Clinics, 2007, 25, 173-207.	0.8	23
164	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
165	Disseminated sporotrichosis presenting with granulomatous inflammatory multiple mononeuropathies. Muscle and Nerve, 2007, 36, 866-872.	1.0	7
166	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
167	Novel myelin protein zero mutation (Arg36Trp) in a patient with acute onset painful neuropathy. Neuromuscular Disorders, 2006, 16, 308-310.	0.3	17
168	Genetic Basis of Neurologic and Neuromuscular Diseases. , 2006, , 267-280.		0
169	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
170	Imaging features of copper deficiency myelopathy: a study of 25 cases. Neuroradiology, 2006, 48, 78-83.	1.1	126
171	Later-Onset Fabry Disease. Archives of Neurology, 2006, 63, 453.	4.9	52
172	Modeling Chronic Glycemic Exposure Variables as Correlates and Predictors of Microvascular Complications of Diabetes. Diabetes Care, 2006, 29, 2282-2288.	4.3	85
173	Genetic testing in inherited peripheral neuropathies. Journal of the Peripheral Nervous System, 2005, 10, 77-84.	1.4	13
174	Nerve Tests Expressed as Percentiles, Normal Deviates, and Composite Scores. , 2005, , 971-983.		8
175	Quantitative Sensation Testing. , 2005, , 1063-1093.		21
176	Hereditary Motor and Sensory Neuropathies: An Overview of Clinical, Genetic, Electrophysiologic, and Pathologic Features. , 2005, , 1623-1658.		71
177	HSANs: Clinical Features, Pathologic Classification, and Molecular Genetics. , 2005, , 1809-1844.		10
178	Monotonicity of Nerve Tests in Diabetes: Subclinical nerve dysfunction precedes diagnosis of polyneuropathy. Diabetes Care, 2005, 28, 2192-2200.	4.3	71
179	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
180	HMSN II (CMT2) and Miscellaneous Inherited System Atrophies of Nerve Axon: Clinicalâ€“Molecular Genetic Correlates. , 2005, , 1717-1751.		4

#	Article	IF	Citations
181	Hereditary Brachial Plexus Neuropathy. , 2005, , 1753-1767.		7
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