Giuseppe Lauria

List of Publications by Year
in descending order

300	22,015	9786	11308
papers	citations	73 h-index	136 g-index

311	311	311	18716
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Phosphorylated TDP-43 aggregates in peripheral motor nerves of patients with amyotrophic lateral sclerosis. Brain, 2022, 145, 276-284.	7.6	22
2	Corneal nerve loss is related to the severity of painful diabetic neuropathy. European Journal of Neurology, 2022, 29, 286-294.	3.3	13
3	Mutations associated with hypokalemic periodic paralysis: from hotspot regions to complete analysis of CACNA1S and SCN4A genes. Neurogenetics, 2022, 23, 19-25.	1.4	8
4	Laser evoked potentials in fibromyalgia with peripheral small fiber involvement. Clinical Neurophysiology, 2022, 135, 96-106.	1.5	6
5	Review of techniques useful for the assessment of sensory small fiber neuropathies: Report from an IFCN expert group. Clinical Neurophysiology, 2022, 136, 13-38.	1.5	21
6	Frequency and clinical correlates of anti-nerve antibodies in a large population of CIDP patients included in the Italian database. Neurological Sciences, 2022, 43, 3939-3947.	1.9	9
7	<scp>COVID</scp>â€19 and the peripheral nervous system. A 2â€year review from the pandemic to the vaccine era. Journal of the Peripheral Nervous System, 2022, 27, 4-30.	3.1	45
8	Skin biopsy and small fibre neuropathies: facts and thoughts 30 years later. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, 915-918.	1.9	12
9	DNAJB2â€related Charcotâ€Marieâ€Tooth disease type 2: Pathomechanism insights and phenotypic spectrum widening. European Journal of Neurology, 2022, 29, 2056-2065.	3.3	7
10	Peripheral Ion Channel Gene Screening in Painful- and Painless-Diabetic Neuropathy. International Journal of Molecular Sciences, 2022, 23, 7190.	4.1	9
11	Facial Onset Sensory and Motor Neuronopathy. Neurology: Clinical Practice, 2021, 11, 147-157.	1.6	16
12	Amyotrophic lateral sclerosis patientsâ€™ and caregivers' distress and loneliness during COVID-19 lockdown. Journal of Neurology, 2021, 268, 420-423.	3.6	33
13	Cognitive reserve is associated with altered clinical expression in amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2021, 22, 237-247.	1.7	12
14	Diagnosis of Neuropathy and Risk Factors for Corneal Nerve Loss in Type 1 and Type 2 Diabetes: A Corneal Confocal Microscopy Study. Diabetes Care, 2021, 44, 150-156.	8.6	60
15	Life with chronic pain during COVID-19 lockdown: the case of patients with small fibre neuropathy and chronic migraine. Neurological Sciences, 2021, 42, 389-397.	1.9	26
16	Pathogenic Huntingtin Repeat Expansions in Patients with Frontotemporal Dementia and Amyotrophic Lateral Sclerosis. Neuron, 2021, 109, 448-460.e4.	8.1	56
17	Chronic inflammatory demyelinating polyradiculoneuropathy: can a diagnosis be made in patients not fulfilling electrodiagnostic criteria?. European Journal of Neurology, 2021, 28, 620-629.	3.3	15
18	Cluster headache not responsive to sumatriptan: A retrospective study. Cephalalgia, 2021, 41, 117-121.	3.9	3

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19	Non-extensitivity and criticality of atomic hydropathicity around a voltage-gated sodium channelâ€™s pore: a modeling study. Journal of Biological Physics, 2021, 47, 61-77.	1.5	3
20	SARS-COV-2 comorbidity network and outcome in hospitalized patients in Crema, Italy. PLoS ONE, 2021, 16, e0248498.	2.5	30
21	Hydropathicity-based prediction of pain-causing NaV1.7 variants. BMC Bioinformatics, 2021, 22, 212.	2.6	5
22	The unfolded protein response in amyotrophic later sclerosis: results of a phase 2 trial. Brain, 2021, 144, 2635-2647.	7.6	33
23	The neurophysiological lesson from the Italian CIDP database. Neurological Sciences, 2021, , 1.	1.9	3
24	Cognitive and behavioural impairment in amyotrophic lateral sclerosis: A landmark of the disease? A mini review of longitudinal studies. Neuroscience Letters, 2021, 754, 135898.	2.1	15
25	Dysregulation of Muscle-Specific MicroRNAs as Common Pathogenic Feature Associated with Muscle Atrophy in ALS, SMA and SBMA: Evidence from Animal Models and Human Patients. International Journal of Molecular Sciences, 2021, 22, 5673.	4.1	14
26	Reader Response: In Vivo Distribution of Î±-Synuclein in Multiple Tissues and Biofluids in Parkinson Disease. Neurology, 2021, 96, 964-965.	1.1	4
27	Prolonged distal motor latency of median nerve does not improve diagnostic accuracy for CIDP. Journal of Neurology, 2021, , 1.	3.6	1
28	A novel gain-of-function sodium channel Î²2 subunit mutation in idiopathic small fiber neuropathy. Journal of Neurophysiology, 2021, 126, 827-839.	1.8	5
29	Syncope and COVID-19 disease â€“ A systematic review. Autonomic Neuroscience: Basic and Clinical, 2021, 235, 102872.	2.8	11
30	Two independent mouse lines carrying the Nav1.7 I228M gain-of-function variant display dorsal root ganglion neuron hyperexcitability but a minimal pain phenotype. Pain, 2021, 162, 1758-1770.	4.2	9
31	The unfolded protein response in amyotrophic later sclerosis: Results of a phase 2 trial. Journal of the Neurological Sciences, 2021, 429, 117702.	0.6	0
32	Congenital insensitivity to pain. Pain, 2021, Publish Ahead of Print, .	4.2	6
33	Common and rare variant association analyses in amyotrophic lateral sclerosis identify 15 risk loci with distinct genetic architectures and neuron-specific biology. Nature Genetics, 2021, 53, 1636-1648.	21.4	223
34	Lacosamide Inhibition of NaV1.7 Channels Depends on its Interaction With the Voltage Sensor Domain and the Channel Pore. Frontiers in Pharmacology, 2021, 12, 791740.	3.5	5
35	Trigeminal Neuralgia: Channels, Pathophysiology, and Therapeutic Challenges. Headache, 2020, , 209-219.	0.4	0
36	Risk factors for chronic inflammatory demyelinating polyradiculoneuropathy (CIDP): antecedent events, lifestyle and dietary habits. Data from the Italian CIDP Database. European Journal of Neurology, 2020, 27, 136-143.	3.3	27

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37	Reversible cerebellar MRI hyperintensities and ataxia associated with hypomagnesemia: a case report with review of the literature. Neurological Sciences, 2020, 41, 961-963.	1.9	4
38	The small fiber neuropathy NaV1.7 I228M mutation: impaired neurite integrity via bioenergetic and mitotoxic mechanisms, and protection by dexpramipexole. Journal of Neurophysiology, 2020, 123, 645-657.	1.8	9
39	Idiopathic distal sensory polyneuropathy. Neurology, 2020, 95, 1005-1014.	1.1	49
40	Computational pipeline to probe NaV1.7 gain-of-function variants in neuropathic painful syndromes. Scientific Reports, 2020, 10, 17930.	3.3	3
41	Clinical diagnosis and management of small fiber neuropathy: an update on best practice. Expert Review of Neurotherapeutics, 2020, 20, 967-980.	2.8	31
42	Placebo effect in chronic inflammatory demyelinating polyneuropathy: The <scp>PATH</scp> study and a systematic review. Journal of the Peripheral Nervous System, 2020, 25, 230-237.	3.1	15
43	Syncope at SARS-CoV-2 onset. Autonomic Neuroscience: Basic and Clinical, 2020, 229, 102734.	2.8	17
44	Cortical thinning trajectories across disease stages and cognitive impairment in amyotrophic lateral sclerosis. Cortex, 2020, 131, 284-294.	2.4	18
45	Frequency of diabetes and other comorbidities in chronic inflammatory demyelinating polyradiculoneuropathy and their impact on clinical presentation and response to therapy. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 1092-1099.	1.9	22
46	Evaluation of molecular inversion probe versus TruSeqÂ® custom methods for targeted next-generation sequencing. PLoS ONE, 2020, 15, e0238467.	2.5	17
47	Impact of environmental factors and physical activity on disability and quality of life in CIDP. Journal of Neurology, 2020, 267, 2683-2691.	3.6	4
48	Peripheral and central nervous system correlates in fibromyalgia. European Journal of Pain, 2020, 24, 1537-1547.	2.8	19
49	GI symptoms as early signs of COVID-19 in hospitalised Italian patients. Gut, 2020, 69, 1547-1548.	12.1	50
50	No efficacy of transcranial direct current stimulation on chronic migraine with medication overuse: A double blind, randomised clinical trial. Cephalalgia, 2020, 40, 1202-1211.	3.9	29
51	Sensitivity and specificity of a commercial ELISA test for anti-MAG antibodies in patients with neuropathy. Journal of Neuroimmunology, 2020, 345, 577288.	2.3	20
52	Focus on the heterogeneity of amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 485-495.	1.7	32
53	Sorting Rare ALS Genetic Variants by Targeted Re-Sequencing Panel in Italian Patients: OPTN, VCP, and SQSTM1 Variants Account for 3% of Rare Genetic Forms. Journal of Clinical Medicine, 2020, 9, 412.	2.4	24
54	Antibodies to neurofascin, contactin-1, and contactin-associated protein 1 in CIDP. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, .	6.0	118

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55	Corneal confocal microscopy detects small nerve fibre damage in patients with painful diabetic neuropathy. Scientific Reports, 2020, 10, 3371.	3.3	41
56	Differential effect of lacosamide on Nav1.7 variants from responsive and non-responsive patients with small fibre neuropathy. Brain, 2020, 143, 771-782.	7.6	31
57	Relevance of diagnostic investigations in chronic inflammatory demyelinating poliradiculoneuropathy: Data from the Italian CIDP database. Journal of the Peripheral Nervous System, 2020, 25, 152-161.	3.1	15
58	Charcot-Marie-Tooth Type 2B: A New Phenotype Associated with a Novel RAB7A Mutation and Inhibited EGFR Degradation. Cells, 2020, 9, 1028.	4.1	20
59	Diabetic Neuropathy Is Characterized by Progressive Corneal Nerve Fiber Loss in the Central and Inferior Whorl Regions. , 2020, 61, 48.		26
60	Cumulative hydropathic topology of a voltageâ€gated sodium channel at atomic resolution. Proteins: Structure, Function and Bioinformatics, 2020, 88, 1319-1328.	2.6	3
61	Corneal confocal microscopy compared with quantitative sensory testing and nerve conduction for diagnosing and stratifying the severity of diabetic peripheral neuropathy. BMJ Open Diabetes Research and Care, 2020, 8, e001801.	2.8	15
62	Role of Human-Induced Pluripotent Stem Cell-Derived Spinal Cord Astrocytes in the Functional Maturation of Motor Neurons in a Multielectrode Array System. Stem Cells Translational Medicine, 2019, 8, 1272-1285.	3.3	34
63	Diagnostic criteria for small fibre neuropathy in clinical practice and research. Brain, 2019, 142, 3728-3736.	7.6	111
64	A message from the new Editorâ€inâ€Chief. Journal of the Peripheral Nervous System, 2019, 24, 234-234.	3.1	0
65	Late-onset and fast progressive neuropathy and cardiomyopathy in Val32Ala transthyretin gene mutation. Neurological Sciences, 2019, 40, 1267-1269.	1.9	4
66	Efficacy and safety of IVIG in CIDP: Combined data of the PRIMA and PATH studies. Journal of the Peripheral Nervous System, 2019, 24, 48-55.	3.1	17
67	Restabilization treatment after intravenous immunoglobulin withdrawal in chronic inflammatory demyelinating polyneuropathy: Results from the preâ€randomization phase of the Polyneuropathy And Treatment with Hizentra study. Journal of the Peripheral Nervous System, 2019, 24, 72-79.	3.1	13
68	Headache, chest pain, and multiplex cranial neuropathy. Neurological Sciences, 2019, 40, 1477-1480.	1.9	0
69	A case of Ciguatera poisoning with paradoxical dysaesthesia and degenerative features at skin biopsy. Journal of the Neurological Sciences, 2019, 403, 112-113.	0.6	2
70	Cognitive Syndromes and C9orf72 Mutation Are Not Related to Cerebellar Degeneration in Amyotrophic Lateral Sclerosis. Frontiers in Neuroscience, 2019, 13, 440.	2.8	10
71	Development of MRC Centre MRI calf muscle fat fraction protocol as a sensitive outcome measure in Hereditary Sensory Neuropathy Type 1. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 895-906.	1.9	17
72	A gain-of-function sodium channelÎ²2-subunit mutation in painful diabetic neuropathy. Molecular Pain, 2019, 15, 174480691984980.	2.1	38

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73	International Clinical Collaboration in Headache Medicine: The International Visiting Scholars Program. Headache, 2019, 59, 446-449.	3.9	1
74	A Delphi consensus statement of the Neuropathic Pain Special Interest Group of the Italian Neurological Society on pharmacoresistant neuropathic pain. Neurological Sciences, 2019, 40, 1425-1431.	1.9	5
75	Smallâ€fiber neuropathy: Expanding the clinical pain universe. Journal of the Peripheral Nervous System, 2019, 24, 19-33.	3.1	71
76	Expression of pathogenic SCN9A mutations in the zebrafish: A model to study small-fiber neuropathy. Experimental Neurology, 2019, 311, 257-264.	4.1	16
77	Cortical correlates of behavioural change in amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 380-386.	1.9	24
78	Yield of peripheral sodium channels gene screening in pure small fibre neuropathy. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 342-352.	1.9	47
79	Atypical CIDP: diagnostic criteria, progression and treatment response. Data from the Italian CIDP Database. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 125-132.	1.9	108
80	Skin nerve Î±-synuclein deposits in a parkinsonian patient with heterozygous parkin mutation. Parkinsonism and Related Disorders, 2019, 60, 182-183.	2.2	7
81	Greater corneal nerve loss at the inferior whorl is related to the presence of diabetic neuropathy and painful diabetic neuropathy. Scientific Reports, 2018, 8, 3283.	3.3	74
82	Intravenous versus subcutaneous immunoglobulin â€“ Authors' reply. Lancet Neurology, The, 2018, 17, 393-394.	10.2	0
83	Genome-wide Analyses Identify KIF5A as a Novel ALS Gene. Neuron, 2018, 97, 1268-1283.e6.	8.1	517
84	Adult leukoencephalopathies with prominent infratentorial involvement can be caused by Erdheimâ€“Chester disease. Journal of Neurology, 2018, 265, 273-284.	3.6	17
85	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.	10.2	193
86	Rapamycin treatment for amyotrophic lateral sclerosis. Medicine (United States), 2018, 97, e11119.	1.0	96
87	A cross-sectional study investigating frequency and features of definitely diagnosed diabetic painful polyneuropathy. Pain, 2018, 159, 2658-2666.	4.2	49
88	Functioning and quality of life in patients with neuropathy associated with anti-MAG antibodies. Journal of Neurology, 2018, 265, 2927-2933.	3.6	12
89	Bilateral Radiation-Induced Hypoglossal Nerve Palsy Responsive to Steroid Treatment. Journal of		

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91	A novel SCN9A splicing mutation in a compound heterozygous girl with congenital insensitivity to pain, hyposmia and hypogeusia. Journal of the Peripheral Nervous System, 2018, 23, 202-206.	3.1	25
92	Cortical markers of cognitive syndromes in amyotrophic lateral sclerosis. NeuroImage: Clinical, 2018, 19, 675-682.	2.7	24
93	COL6A5variants in familial neuropathic chronic itch. Brain, 2017, 140, aww343.	7.6	25
94	Network topology of NaV1.7 mutations in sodium channel-related painful disorders. BMC Systems Biology, 2017, 11, 28.	3.0	29
95	Genetic Counseling Dilemmas for a Patient with Sporadic Amyotrophic Lateral Sclerosis, Frontotemporal Degeneration & Parkinson's Disease. Journal of Genetic Counseling, 2017, 26, 442-446.	1.6	3
96	Botulinum Toxin for Burning Mouth Syndrome. Annals of Internal Medicine, 2017, 166, 762.	3.9	10
97	Is firstly diagnosed ALS really ALS? Results of a population-based study with long-term follow-up. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2017, 18, 221-226.	1.7	4
98	New technologies for the assessment of neuropathies. Nature Reviews Neurology, 2017, 13, 203-216.	10.1	90
99	Pain in amyotrophic lateral sclerosis. Lancet Neurology, The, 2017, 16, 144-157.	10.2	97
100	The diagnostic challenge of small fibre neuropathy: clinical presentations, evaluations, and causes. Lancet Neurology, The, 2017, 16, 934-944.	10.2	215
101	Protein misfolding, amyotrophic lateral sclerosis and guanabenz: protocol for a phase II RCT with futility design (ProMISe trial). BMJ Open, 2017, 7, e015434.	1.9	14
102	Small fibre neuropathy. Current Opinion in Neurology, 2017, 30, 490-499.	3.6	116
103	Therapeutic potential of Mesenchymal Stem Cells for the treatment of diabetic peripheral neuropathy. Experimental Neurology, 2017, 288, 75-84.	4.1	21
104	Anti-NF155 chronic inflammatory demyelinating polyradiculoneuropathy strongly associates to HLA-DRB15. Journal of Neuroinflammation, 2017, 14, 224.	7.2	50
105	Amyotrophic lateral sclerosis causes small fiber pathology. European Journal of Neurology, 2016, 23, 416-420.	3.3	65
106	Unraveling gene expression profiles in peripheral motor nerve from amyotrophic lateral sclerosis patients: insights into pathogenesis. Scientific Reports, 2016, 6, 39297.	3.3	24
107	TBK1 is associated with ALS and ALS-FTD in Sardinian patients. Neurobiology of Aging, 2016, 43, 180.e1-180.e5.	3.1	40
108	Beyond the consensus criteria: multiple cognitive profiles in amyotrophic lateral sclerosis?. Cortex, 2016, 81, 162-167.	2.4	45

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109	NEK1 variants confer susceptibility to amyotrophic lateral sclerosis. Nature Genetics, 2016, 48, 1037-1042.	21.4	218
110	A painful neuropathy-associated Nav1.7 mutant leads to time-dependent degeneration of small-diameter axons associated with intracellular Ca²⁺dysregulation and decrease in ATP levels. Molecular Pain, 2016, 12, 174480691667447.	2.1	23
111	Correlation of the patient's reported outcome Inflammatoryâ€<scp>RODS</scp> with an objective metric in immuneâ€mediated neuropathies. European Journal of Neurology, 2016, 23, 1248-1253.	3.3	33
112	Mutant SOD1 accumulation in sensory neurons does not associate with endoplasmic reticulum stress features: Implications for differential vulnerability of sensory and motor neurons to SOD1 toxicity. Neuroscience Letters, 2016, 627, 107-114.	2.1	6
113	Small fiber neuropathy is a common feature of Ehlers-Danlos syndromes. Neurology, 2016, 87, 155-159.	1.1	90
114	Does ability to walk reflect general functionality in inflammatory neuropathies?. Journal of the Peripheral Nervous System, 2016, 21, 74-81.	3.1	13
115	ATNX2 is not a regulatory gene in Italian amyotrophic lateral sclerosis patients with C9ORF72 GGGGCC expansion. Neurobiology of Aging, 2016, 39, 218.e5-218.e8.	3.1	6
116	Degenerative neuromuscular diseases: crucial gene and cell machinery discoveries. Lancet Neurology, The, 2016, 15, 12-13.	10.2	1
117	ALS mouse model SOD1^{G93A}displays early pathology of sensory small fibers associated to accumulation of a neurotoxic splice variant of peripherin. Human Molecular Genetics, 2016, 25, 1588-1599.	2.9	39
118	Cerebellar ataxia, neuropathy, and vestibular areflexia syndrome: a slowly progressive disorder with stereotypical presentation. Journal of Neurology, 2016, 263, 245-249.	3.6	13
119	SMART (stroke-like migraine attack after radiation therapy) syndrome: a case report with review of the literature. Neurological Sciences, 2016, 37, 157-161.	1.9	16
120	ClCâ€1 mutations in myotonia congenita patients: insights into molecular gating mechanisms and genotypeâ€“phenotype correlation. Journal of Physiology, 2015, 593, 4181-4199.	2.9	24
121	Ca²⁺ toxicity due to reverse Na⁺/Ca²⁺ exchange contributes to degeneration of neurites of DRG neurons induced by a neuropathy-associated Nav1.7 mutation. Journal of Neurophysiology, 2015, 114, 1554-1564.	1.8	41
122	Comparing the <scp>NIS</scp> vs. <scp>MRC</scp> and <scp>INCAT</scp> sensory scale through Rasch analyses. Journal of the Peripheral Nervous System, 2015, 20, 277-288.	3.1	27
123	Epidermal innervation morphometry by immunofluorescence and bright-field microscopy. Journal of the Peripheral Nervous System, 2015, 20, 387-391.	3.1	30
124	Grip strength comparison in immuneâ€mediated neuropathies: Vigorimeter vs. Jamar. Journal of the Peripheral Nervous System, 2015, 20, 269-276.	3.1	28
125	Impairment measures versus inflammatory <scp>RODS</scp> in <scp>GBS</scp> and <scp>CIDP</scp>: a responsiveness comparison. Journal of the Peripheral Nervous System, 2015, 20, 289-295.	3.1	30
126	Side and time variability of intraepidermal nerve fiber density. Neurology, 2015, 84, 2368-2371.	1.1	29

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127	A case of papilloedema associated with Chiari I malformation. Journal of the Neurological Sciences, 2015, 353, 183-184.	0.6	5
128	Late reoperations after acute aortic dissection repair: Single-center experience. Asian Cardiovascular and Thoracic Annals, 2015, 23, 787-794.	0.5	26
129	Exome sequencing in amyotrophic lateral sclerosis identifies risk genes and pathways. Science, 2015, 347, 1436-1441.	12.6	823
130	Lowering Plasma 1-Deoxysphingolipids Improves Neuropathy in Diabetic Rats. Diabetes, 2015, 64, 1035-1045.	0.6	69
131	Advances in diagnostics and outcome measures in peripheral neuropathies. Neuroscience Letters, 2015, 596, 3-13.	2.1	25
132	HFE p.H63D polymorphism does not influence ALS phenotype and survival. Neurobiology of Aging, 2015, 36, 2906.e7-2906.e11.	3.1	8
133	Erythropoietin in amyotrophic lateral sclerosis: a multicentre, randomised, double blind, placebo controlled, phase III study. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 879-886.	1.9	32
134	The MITOS system predicts long-term survival in amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 1180-1185.	1.9	42
135	TUBA4A gene analysis in sporadic amyotrophic lateral sclerosis: identification of novel mutations. Journal of Neurology, 2015, 262, 1376-1378.	3.6	44
136	The Domain II S4-S5 Linker in Nav1.9: A Missense Mutation Enhances Activation, Impairs Fast Inactivation, and Produces Human Painful Neuropathy. NeuroMolecular Medicine, 2015, 17, 158-169.	3.4	70
137	CHCH10 mutations in an Italian cohort of familial and sporadic amyotrophic lateral sclerosis patients. Neurobiology of Aging, 2015, 36, 1767.e3-1767.e6.	3.1	44
138	ATXN2 is a modifier of phenotype in ALS patients of Sardinian ancestry. Neurobiology of Aging, 2015, 36, 2906.e1-2906.e5.	3.1	19
139	Neurological Pain. , 2015, , 489-505.		0
140	Neutralization of Schwann Cell-Secreted VEGF Is Protective to In Vitro and In Vivo Experimental Diabetic Neuropathy. PLoS ONE, 2014, 9, e108403.	2.5	26
141	PMP22 messenger RNA levels in skin biopsies: testing the effectiveness of a Charcot-Marie-Tooth 1A biomarker. Brain, 2014, 137, 1614-1620.	7.6	33
142	Pentraxin-3 and VEGF in POEMS syndrome: A 2-year longitudinal study. Journal of Neuroimmunology, 2014, 277, 189-192.	2.3	7
143	Small fibers, large impact: Quality of life in smallâ€fiber neuropathy. Muscle and Nerve, 2014, 49, 329-336.	2.2	102
144	<scp>LRP</scp> 4 antibodies in serum and <scp>CSF</scp> from amyotrophic lateral sclerosis patients. Annals of Clinical and Translational Neurology, 2014, 1, 80-87.	3.7	94

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145	Changing outcome in inflammatory neuropathies. Neurology, 2014, 83, 2124-2132.	1.1	89
146	Heterozygous D90A-SOD1 mutation in a patient with facial onset sensory motor neuronopathy (FOSMN) syndrome: a bridge to amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2014, 85, 1009-1011.	1.9	32
147	Letter to the Editor. Pain, 2014, 155, 1177-1178.	4.2	0
148	Non-paraneoplastic voltage-gated calcium channels antibody-mediated cerebellar ataxia responsive to IVIG treatment. Journal of the Neurological Sciences, 2014, 336, 169-170.	0.6	13
149	Valproate induced hyperammonemic encephalopathy successfully treated with levocarnitine. Journal of Clinical Neuroscience, 2014, 21, 690-691.	1.5	16
150	Mutations in the Matrin 3 gene cause familial amyotrophic lateral sclerosis. Nature Neuroscience, 2014, 17, 664-666.	14.8	398
151	A case of Bingâ€“Neel syndrome presenting as spinal cord compression. Journal of the Neurological Sciences, 2014, 346, 345-347.	0.6	5
152	Selected items from the Charcot-Marie-Tooth (CMT) Neuropathy Score and secondary clinical outcome measures serve as sensitive clinical markers of disease severity in CMT1A patients. Neuromuscular Disorders, 2014, 24, 1003-1017.	0.6	25
153	Exome-wide Rare Variant Analysis Identifies TUBA4A Mutations Associated with Familial ALS. Neuron, 2014, 84, 324-331.	8.1	308
154	The Role of Sodium Channels in Painful Diabetic and Idiopathic Neuropathy. Current Diabetes Reports, 2014, 14, 538.	4.2	33
155	Sodium channel genes in pain-related disorders: phenotypeâ€“genotype associations and recommendations for clinical use. Lancet Neurology, The, 2014, 13, 1152-1160.	10.2	148
156	Guillainâ^'BarrÃ© syndrome following measles infection: case report and review of the literature. Neurological Sciences, 2014, 35, 2017-2018.	1.9	5
157	Gain-of-function mutations in sodium channel NaV1.9 in painful neuropathy. Brain, 2014, 137, 1627-1642.	7.6	242
158	Longâ€term survival in amyotrophic lateral sclerosis: A populationâ€based study. Annals of Neurology, 2014, 75, 287-297.	5.3	141
159	Paroxysmal itch caused by gain-of-function Nav1.7 mutation. Pain, 2014, 155, 1702-1707.	4.2	78
160	Approach to Small Fiber Neuropathy. , 2014, , 507-517.		2
161	Epidermal Nerve Fibers. , 2014, , 76-79.		2
162	Epithelial expression of vanilloid and cannabinoid receptors: a potential role in burning mouth syndrome pathogenesis. Histology and Histopathology, 2014, 29, 523-33.	0.7	34

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163	Islet Transplantation and Insulin Administration Relieve Long-Term Complications and Rescue the Residual Endogenous Pancreatic Î² Cells. American Journal of Pathology, 2013, 183, 1527-1538.	3.8	8
164	Ischemic stroke as clinical onset of POEMS syndrome. Journal of Neurology, 2013, 260, 3178-3181.	3.6	17
165	A case of late herpes simplex encephalitis relapse. Journal of Clinical Virology, 2013, 58, 269-270.	3.1	12
166	Pure autonomic failure with cold induced sweating. Autonomic Neuroscience: Basic and Clinical, 2013, 176, 98-100.	2.8	3
167	Screening of the PFN1 gene in sporadic amyotrophic lateral sclerosis and in frontotemporal dementia. Neurobiology of Aging, 2013, 34, 1517.e9-1517.e10.	3.1	35
168	Dermal innervation in healthy subjects and small fiber neuropathy patients: a stereological reappraisal. Journal of the Peripheral Nervous System, 2013, 18, 48-53.	3.1	22
169	Millsâ€™ syndrome: an italian case and revision of the literature. Neurological Sciences, 2013, 34, 255-256.	1.9	6
170	Small Nerve Fiber Pathology in Critical Illness. PLoS ONE, 2013, 8, e75696.	2.5	46
171	Outcome measures in peripheral neuropathies. Current Opinion in Neurology, 2012, 25, 556-563.	3.6	18
172	Small fibre neuropathy. Current Opinion in Neurology, 2012, 25, 542-549.	3.6	94
173	Recurrent Horner Syndrome and Persistent Trigeminal Artery. Neurologist, 2012, 18, 406-408.	0.7	0
174	Functional profiles of SCN9A variants in dorsal root ganglion neurons and superior cervical ganglion neurons correlate with autonomic symptoms in small fibre neuropathy. Brain, 2012, 135, 2613-2628.	7.6	90
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