## Giuseppe Lauria

List of Publications by Year in descending order

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300 papers 22,015 citations

9786 73 h-index 136 g-index

311 all docs

311 docs citations

times ranked

311

18716 citing authors

#	Article	IF	CITATIONS
1	Diabetic Neuropathies: Update on Definitions, Diagnostic Criteria, Estimation of Severity, and Treatments. Diabetes Care, 2010, 33, 2285-2293.	8.6	1,963
2	Frequency of the C9orf72 hexanucleotide repeat expansion in patients with amyotrophic lateral sclerosis and frontotemporal dementia: a cross-sectional study. Lancet Neurology, The, 2012, 11, 323-330.	10.2	1,039
3	Exome sequencing in amyotrophic lateral sclerosis identifies risk genes and pathways. Science, 2015, 347, 1436-1441.	12.6	823
4	The diagnostic criteria for small fibre neuropathy: from symptoms to neuropathology. Brain, 2008, 131, 1912-1925.	7.6	707
5	European Federation of Neurological Societies/Peripheral Nerve Society Guideline on the use of skin biopsy in the diagnosis of small fiber neuropathy. Report of a joint task force of the European Feâ€deration of Neurological Societies and the Peripheral Nerve Society. European Journal of Neurology. 2010. 17. 903.	3.3	651
6	Gain of function Na <sub>V</sub> 1.7 mutations in idiopathic small fiber neuropathy. Annals of Neurology, 2012, 71, 26-39.	5.3	518
7	Genome-wide Analyses Identify KIF5A as a Novel ALS Gene. Neuron, 2018, 97, 1268-1283.e6.	8.1	517
8	EFNS guidelines on the use of skin biopsy in the diagnosis of peripheral neuropathy. European Journal of Neurology, 2005, 12, 747-758.	3.3	489
9	Intraepidermal nerve fiber density at the distal leg: a worldwide normative reference study. Journal of the Peripheral Nervous System, 2010, 15, 202-207.	3.1	462
10	Mutations in the Matrin 3 gene cause familial amyotrophic lateral sclerosis. Nature Neuroscience, 2014, 17, 664-666.	14.8	398
11	Gain-of-function Na <sub>v</sub> 1.8 mutations in painful neuropathy. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 19444-19449.	7.1	369
12	Trigeminal small-fiber sensory neuropathy causes burning mouth syndrome. Pain, 2005, 115, 332-337.	4.2	356
13	Practice Parameter: Evaluation of distal symmetric polyneuropathy: Role of autonomic testing, nerve biopsy, and skin biopsy (an evidence-based review) [RETIRED]. Neurology, 2009, 72, 177-184.	1.1	331
14	Exome-wide Rare Variant Analysis Identifies TUBA4A Mutations Associated with Familial ALS. Neuron, 2014, 84, 324-331.	8.1	308
15	Practice Parameter: Evaluation of distal symmetric polyneuropathy: Role of laboratory and genetic testing (an evidence-based review). Neurology, 2009, 72, 185-192.	1.1	265
16	Gain-of-function mutations in sodium channel NaV1.9 in painful neuropathy. Brain, 2014, 137, 1627-1642.	7.6	242
17	Erythropoietin both protects from and reverses experimental diabetic neuropathy. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 823-828.	7.1	238
18	Modifying the Medical Research Council grading system through Rasch analyses. Brain, 2012, 135, 1639-1649.	7.6	224

#	Article	IF	CITATIONS
19	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
20	Ascorbic acid in Charcot–Marie–Tooth disease type 1A (CMT-TRIAAL and CMT-TRAUK): a double-blind randomised trial. Lancet Neurology, The, 2011, 10, 320-328.	10.2	222
21	Axonal swellings predict the degeneration of epidermal nerve fibers in painful neuropathies. Neurology, 2003, 61, 631-636.	1.1	220
22	NEK1 variants confer susceptibility to amyotrophic lateral sclerosis. Nature Genetics, 2016, 48, 1037-1042.	21.4	218
23	The diagnostic challenge of small fibre neuropathy: clinical presentations, evaluations, and causes. Lancet Neurology, The, 2017, 16, 934-944.	10.2	215
24	Intraepidermal nerve fiber density and its application in sarcoidosis. Neurology, 2009, 73, 1142-1148.	1.1	206
25	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
26	Small-fibre neuropathiesâ€"advances in diagnosis, pathophysiology and management. Nature Reviews Neurology, 2012, 8, 369-379.	10.1	187
27	Clinical characteristics of patients with familial amyotrophic lateral sclerosis carrying the pathogenic GGGGCC hexanucleotide repeat expansion of C9ORF72. Brain, 2012, 135, 784-793.	7.6	182
28	Progesterone and its derivatives are neuroprotective agents in experimental diabetic neuropathy: A multimodal analysis. Neuroscience, 2007, 144, 1293-1304.	2.3	175
29	Epidermal innervation: changes with aging, topographic location, and in sensory neuropathy. Journal of the Neurological Sciences, 1999, 164, 172-178.	0.6	167
30	Alpha-lipoic acid prevents mitochondrial damage and neurotoxicity in experimental chemotherapy neuropathy. Experimental Neurology, 2008, 214, 276-284.	4.1	158
31	Randomised controlled trial of methotrexate for chronic inflammatory demyelinating polyradiculoneuropathy (RMC trial): a pilot, multicentre study. Lancet Neurology, The, 2009, 8, 158-164.	10.2	155
32	Sensory neuron diseases. Lancet Neurology, The, 2005, 4, 349-361.	10.2	151
33	European Federation of Neurological Societies/Peripheral Nerve Society Guideline on the use of skin biopsy in the diagnosis of small fiber neuropathy. Report of a joint task force of the European Federation of Neurological Societies and the Peripheral Nerve Society. Journal of the Peripheral Nervous System. 2010. 15, 79-92.	3.1	151
34	Sodium channel genes in pain-related disorders: phenotype–genotype associations and recommendations for clinical use. Lancet Neurology, The, 2014, 13, 1152-1160.	10.2	148
35	Longâ€term survival in amyotrophic lateral sclerosis: A populationâ€based study. Annals of Neurology, 2014, 75, 287-297.	5.3	141
36	Intraepidermal nerve fiber density in rat foot pad: neuropathologic–neurophysiologic correlation. Journal of the Peripheral Nervous System, 2005, 10, 202-208.	3.1	132

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37	Skin biopsy as a diagnostic tool in peripheral neuropathy. Nature Clinical Practice Neurology, 2007, 3, 546-557.	2.5	125
38	Small fibre neuropathy: role in the diagnosis of diabetic sensorimotor polyneuropathy. Diabetes/Metabolism Research and Reviews, 2011, 27, 678-684.	4.0	123
39	Revised normative values for grip strength with the Jamar dynamometer. Journal of the Peripheral Nervous System, 2011, 16, 47-50.	3.1	118
40	Antibodies to neurofascin, contactin-1, and contactin-associated protein 1 in CIDP. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, .	6.0	118
41	Skin biopsy in the management of peripheral neuropathy. Lancet Neurology, The, 2007, 6, 632-642.	10.2	117
42	Small fibre neuropathy. Current Opinion in Neurology, 2017, 30, 490-499.	3 <b>.</b> 6	116
43	Expression of capsaicin receptor immunoreactivity in human peripheral nervous system and in painful neuropathies. Journal of the Peripheral Nervous System, 2006, 11, 262-271.	3.1	114
44	Plasma deoxysphingolipids: a novel class of biomarkers for the metabolic syndrome?. Diabetologia, 2012, 55, 421-431.	6.3	113
45	Skin biopsy for the diagnosis of peripheral neuropathy. Histopathology, 2009, 54, 273-285.	2.9	112
46	Diagnostic criteria for small fibre neuropathy in clinical practice and research. Brain, 2019, 142, 3728-3736.	7.6	111
47	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
48	Small fibre neuropathies. Current Opinion in Neurology, 2005, 18, 591-597.	3.6	105
49	Small fibers, large impact: Quality of life in smallâ€fiber neuropathy. Muscle and Nerve, 2014, 49, 329-336.	2.2	102
50	lgM deposits on skin nerves in anti-myelin-associated glycoprotein neuropathy. Annals of Neurology, 2005, 57, 180-187.	5 <b>.</b> 3	100
51	Neuropathological alterations in diabetic truncal neuropathy: evaluation by skin biopsy. Journal of Neurology, Neurosurgery and Psychiatry, 1998, 65, 762-766.	1.9	97
52	Pain in amyotrophic lateral sclerosis. Lancet Neurology, The, 2017, 16, 144-157.	10.2	97
53	Rapamycin treatment for amyotrophic lateral sclerosis. Medicine (United States), 2018, 97, e11119.	1.0	96
54	Skin biopsy: a new tool for diagnosing peripheral neuropathy. BMJ: British Medical Journal, 2007, 334, 1159-1162.	2.3	94

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55	Intra- and Interfamily Phenotypic Diversity in Pain Syndromes Associated with a Gain-of-Function Variant of Na <sub>V</sub> 1.7. Molecular Pain, 2011, 7, 1744-8069-7-92.	2.1	94
56	Small fibre neuropathy. Current Opinion in Neurology, 2012, 25, 542-549.	3.6	94
57	<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
58	Incidence and Prognosis of Stroke in the Belluno Province, Italy. Stroke, 1995, 26, 1787-1793.	2.0	93
59	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
60	Small fiber neuropathy is a common feature of Ehlers-Danlos syndromes. Neurology, 2016, 87, 155-159.	1.1	90
61	New technologies for the assessment of neuropathies. Nature Reviews Neurology, 2017, 13, 203-216.	10.1	90
62	Changing outcome in inflammatory neuropathies. Neurology, 2014, 83, 2124-2132.	1.1	89
63	Effects of Manidipine and Delapril in Hypertensive Patients With Type 2 Diabetes Mellitus. Hypertension, 2011, 58, 776-783.	2.7	86
64	Na <sub>v</sub> 1.7-related small fiber neuropathy. Neurology, 2012, 78, 1635-1643.	1.1	86
65	Protective Effect of Erythropoietin and Its Carbamylated Derivative in Experimental Cisplatin Peripheral Neurotoxicity. Clinical Cancer Research, 2006, 12, 2607-2612.	7.0	85
66	Cranial nerve involvement in CMT disease type 1 due to early growth response 2 gene mutation. Neurology, 2000, 54, 1696-1698.	1.1	84
67	Neuroprotective effects of a ligand of translocator protein-18kDa (Ro5-4864) in experimental diabetic neuropathy. Neuroscience, 2009, 164, 520-529.	2.3	82
68	Clinical and magnetic resonance imaging findings in chronic sensory ganglionopathies. Annals of Neurology, 2000, 47, 104-109.	5.3	79
69	Neuroactive steroids and peripheral neuropathy. Brain Research Reviews, 2008, 57, 460-469.	9.0	79
70	Paroxysmal itch caused by gain-of-function Nav1.7 mutation. Pain, 2014, 155, 1702-1707.	4.2	78
71	Evaluation of distal symmetric polyneuropathy: The role of autonomic testing, nerve biopsy, and skin biopsy (an evidenceâ€based review). Muscle and Nerve, 2009, 39, 106-115.	2.2	76
72	C9ORF72 hexanucleotide repeat expansions in the Italian sporadic ALS population. Neurobiology of Aging, 2012, 33, 1848.e15-1848.e20.	3.1	76

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73	Epidermal nerve fiber density in sensory ganglionopathies: Clinical and neurophysiologic correlations. Muscle and Nerve, 2001, 24, 1034-1039.	2.2	75
74	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
75	Worsening after rituximab treatment in anti-mag neuropathy. Muscle and Nerve, 2005, 32, 378-379.	2.2	72
76	Smallâ€fiber neuropathy: Expanding the clinical pain universe. Journal of the Peripheral Nervous System, 2019, 24, 19-33.	3.1	71
77	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
78	Small nerve fibres, small hands and small feet: a new syndrome of pain, dysautonomia and acromesomelia in a kindred with a novel NaV1.7 mutation. Brain, 2012, 135, 345-358.	7.6	69
79	Lowering Plasma 1-Deoxysphingolipids Improves Neuropathy in Diabetic Rats. Diabetes, 2015, 64, 1035-1045.	0.6	69
80	Tubule and neurofilament immunoreactivity in human hairy skin: Markers for intraepidermal nerve fibers. Muscle and Nerve, 2004, 30, 310-316.	2.2	66
81	Amyotrophic lateral sclerosis causes small fiber pathology. European Journal of Neurology, 2016, 23, 416-420.	3.3	65
82	Evaluation of distal symmetric polyneuropathy: The role of laboratory and genetic testing (an) Tj ETQq0 0 0 rgB	Γ/Overlock 2.2	10 Tf 50 382
83	Practice Parameter: The Evaluation of Distal Symmetric Polyneuropathy: The Role of Laboratory and Genetic Testing (An Evidenceâ€Based Review). PM and R, 2009, 1, 5-13.	1.6	60
84	Practice Parameter: The Evaluation of Distal Symmetric Polyneuropathy: The Role of Autonomic Testing, Nerve Biopsy, and Skin Biopsy (An Evidenceâ€Based Review). PM and R, 2009, 1, 14-22.	1.6	60
85	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
86	Cisplatin-induced peripheral neuropathy: Neuroprotection by erythropoietin without affecting tumour growth. European Journal of Cancer, 2007, 43, 710-717.	2.8	58
87	Testosterone derivatives are neuroprotective agents in experimental diabetic neuropathy. Cellular and Molecular Life Sciences, 2007, 64, 1158-1168.	5.4	58
88	Docetaxelâ€induced peripheral neuropathy: protective effects of dihydroprogesterone and progesterone in an experimental model. Journal of the Peripheral Nervous System, 2009, 14, 36-44.	3.1	58
89	Excellent response to steroid treatment in anti-GAD cerebellar ataxia. Lancet Neurology, The, 2003, 2,	10.0	
	634-635.	10.2	57

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91	Pathogenic Huntingtin Repeat Expansions in Patients with Frontotemporal Dementia and Amyotrophic Lateral Sclerosis. Neuron, 2021, 109, 448-460.e4.	8.1	56
92	Improving fatigue assessment in immuneâ€mediated neuropathies: the modified Raschâ€built fatigue severity scale. Journal of the Peripheral Nervous System, 2009, 14, 268-278.	3.1	54
93	The Multiple Sclerosis Knowledge Questionnaire: a self-administered instrument for recently diagnosed patients. Multiple Sclerosis Journal, 2010, 16, 100-111.	3.0	50
94	Unmyelinated and myelinated skin nerve damage in Guillain–Barré syndrome: Correlation with pain and recovery. Pain, 2012, 153, 399-409.	4.2	50
95	Anti-NF155 chronic inflammatory demyelinating polyradiculoneuropathy strongly associates to HLA-DRB15. Journal of Neuroinflammation, 2017, 14, 224.	7.2	50
96	GI symptoms as early signs of COVID-19 in hospitalised Italian patients. Gut, 2020, 69, 1547-1548.	12.1	50
97	A cross-sectional study investigating frequency and features of definitely diagnosed diabetic painful polyneuropathy. Pain, 2018, 159, 2658-2666.	4.2	49
98	Idiopathic distal sensory polyneuropathy. Neurology, 2020, 95, 1005-1014.	1.1	49
99	Erythropoietin: not just about erythropoiesis. Lancet, The, 2010, 375, 2142.	13.7	48
100	Morphometry of dermal nerve fibers in human skin. Neurology, 2011, 77, 242-249.	1.1	48
101	A multicenter, randomized, double-blind, placebo-controlled trial of long-term ascorbic acid treatment in Charcot-Marie-Tooth disease type 1A (CMT-TRIAAL): The study protocol [EudraCT no.: 2006-000032-27]. Pharmacological Research, 2006, 54, 436-441.	7.1	47
102	Four novel cases of periaxin-related neuropathy and review of the literature. Neurology, 2010, 75, 1830-1838.	1.1	47
103	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
104	Neurophysiological Diagnosis of Acquired Sensory Ganglionopathies. European Neurology, 2003, 50, 146-152.	1.4	46
105	Small Nerve Fiber Pathology in Critical Illness. PLoS ONE, 2013, 8, e75696.	2.5	46
106	Small Fiber Neuropathy: Is Skin Biopsy the Holy Grail?. Current Diabetes Reports, 2012, 12, 384-392.	4.2	45
107	Beyond the consensus criteria: multiple cognitive profiles in amyotrophic lateral sclerosis?. Cortex, 2016, 81, 162-167.	2.4	45
108	<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

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109	Painful neuropathy in subclinical hypothyroidism: clinical and neuropathological recovery after hormone replacement therapy. Neurological Sciences, 2009, 30, 149-151.	1.9	44
110	TUBA4A gene analysis in sporadic amyotrophic lateral sclerosis: identification of novel mutations. Journal of Neurology, 2015, 262, 1376-1378.	3.6	44
111	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
112	The MITOS system predicts long-term survival in amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 1180-1185.	1.9	42
113	Erythropoietin in amyotrophic lateral sclerosis: A pilot, randomized, double-blind, placebo-controlled study of safety and tolerability. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2009, 10, 410-415.	2.1	41
114	Beneficial Effects of PKF275-055, a Novel, Selective, Orally Bioavailable, Long-Acting Dipeptidyl Peptidase IV Inhibitor in Streptozotocin-Induced Diabetic Peripheral Neuropathy. Journal of Pharmacology and Experimental Therapeutics, 2012, 340, 64-72.	2.5	41
115	Ca <sup>2+</sup> toxicity due to reverse Na <sup>+</sup> /Ca <sup>2+</sup> 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
116	Corneal confocal microscopy detects small nerve fibre damage in patients with painful diabetic neuropathy. Scientific Reports, 2020, 10, 3371.	3.3	41
117	TBK1 is associated with ALS and ALS-FTD in Sardinian patients. Neurobiology of Aging, 2016, 43, 180.e1-180.e5.	3.1	40
118	Hydroxytyrosol Attenuates Peripheral Neuropathy in Streptozotocin-Induced Diabetes in Rats. Journal of Agricultural and Food Chemistry, 2012, 60, 5859-5865.	5.2	39
119	ALS mouse model SOD1 <sup>G93A</sup> 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
120	Altered peripheral myelination in mice lacking GABAB receptors. Molecular and Cellular Neurosciences, 2008, 37, 599-609.	2.2	38
121	A gain-of-function sodium channel $\langle b \rangle \hat{l}^2 \langle b \rangle 2$ -subunit mutation in painful diabetic neuropathy. Molecular Pain, 2019, 15, 174480691984980.	2.1	38
122	Innervation of the human epidermis. A historical review. Italian Journal of Neurological Sciences, 1999, 20, 63-70.	0.1	37
123	Incidence of transient global amnesia in the Belluno province, Italy: 1985 through 1995. Acta Neurologica Scandinavica, 1997, 95, 303-310.	2.1	35
124	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
125	Regression of diabetic complications by islet transplantation in the rat. Diabetologia, 2009, 52, 2653-2661.	6.3	34
126	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

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127	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
128	Experimental epothilone B neurotoxicity: Results of in vitro and in vivo studies. Neurobiology of Disease, 2009, 35, 270-277.	4.4	33
129	"Burning Tongue―and "Burning Tip― The Diagnostic Challenge of the Burning Mouth Syndrome. Clinical Journal of Pain, 2010, 26, 528-532.	1.9	33
130	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
131	The Role of Sodium Channels in Painful Diabetic and Idiopathic Neuropathy. Current Diabetes Reports, 2014, 14, 538.	4.2	33
132	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
133	Amyotrophic lateral sclerosis patients' and caregivers' distress and loneliness during COVID-19 lockdown. Journal of Neurology, 2021, 268, 420-423.	3.6	33
134	The unfolded protein response in amyotrophic later sclerosis: results of a phase 2 trial. Brain, 2021, 144, 2635-2647.	7.6	33
135	Posterior reversible encephalopathy syndrome as the initial manifestation of Guillainâ€Barré syndrome: case report and review of the literature. Journal of the Peripheral Nervous System, 2012, 17, 356-360.	3.1	32
136	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
137	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
138	Focus on the heterogeneity of amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 485-495.	1.7	32
139	Botulinum Toxin Treatment for Oropharyngeal Dysphagia Associated With Diabetic Neuropathy. Diabetes Care, 2006, 29, 2650-2653.	8.6	31
140	Clinical diagnosis and management of small fiber neuropathy: an update on best practice. Expert Review of Neurotherapeutics, 2020, 20, 967-980.	2.8	31
141	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
142	Temporal Trend and Factors Associated with Delayed Hospital Admission of Stroke Patients. Neuroepidemiology, 1999, 18, 255-264.	2.3	30
143	Epidermal innervation morphometry by immunofluorescence and bright-field microscopy. Journal of the Peripheral Nervous System, 2015, 20, 387-391.	3.1	30
144	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

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145	SARS-COV-2 comorbidity network and outcome in hospitalized patients in Crema, Italy. PLoS ONE, 2021, 16, e0248498.	2.5	30
146	An epidemiological study of multiple sclerosis in central Sardinia, Italy. Acta Neurologica Scandinavica, 1998, 98, 391-394.	2.1	29
147	Side and time variability of intraepidermal nerve fiber density. Neurology, 2015, 84, 2368-2371.	1.1	29
148	Network topology of NaV1.7 mutations in sodium channel-related painful disorders. BMC Systems Biology, 2017, 11, 28.	3.0	29
149	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
150	Pain and autonomic dysfunction in patients with sarcoidosis and small fibre neuropathy. Journal of Neurology, 2010, 257, 2086-2090.	3.6	28
151	Grip strength comparison in immuneâ€mediated neuropathies: Vigorimeter vs. Jamar. Journal of the Peripheral Nervous System, 2015, 20, 269-276.	3.1	28
152	Celiac disease presenting with motor neuropathy: Effect of gluten free-diet. Muscle and Nerve, 2007, 35, 675-677.	2.2	27
153	ATAXIN2 CAG-repeat length in Italian patients with amyotrophic lateral sclerosis: risk factor or variant phenotype? Implication for genetic testing and counseling. Neurobiology of Aging, 2012, 33, 1847.e15-1847.e21.	3.1	27
154	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
155	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
156	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
157	Late reoperations after acute aortic dissection repair: Single-center experience. Asian Cardiovascular and Thoracic Annals, 2015, 23, 787-794.	0.5	26
158	Diabetic Neuropathy Is Characterized by Progressive Corneal Nerve Fiber Loss in the Central and Inferior Whorl Regions., 2020, 61, 48.		26
159	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
160	The somatosensory blink reflex in upper and lower brainstem lesions. Muscle and Nerve, 2011, 43, 196-202.	2.2	25
161	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
162	Advances in diagnostics and outcome measures in peripheral neuropathies. Neuroscience Letters, 2015, 596, 3-13.	2.1	25

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163	<i>COL6A5</i> variants in familial neuropathic chronic itch. Brain, 2017, 140, aww343.	7.6	25
164	A novel <i>SCN9A</i> 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
165	ClCâ $\in$ 1 mutations in myotonia congenita patients: insights into molecular gating mechanisms and genotypeâ $\in$ "phenotype correlation. Journal of Physiology, 2015, 593, 4181-4199.	2.9	24
166	Unraveling gene expression profiles in peripheral motor nerve from amyotrophic lateral sclerosis patients: insights into pathogenesis. Scientific Reports, 2016, 6, 39297.	3.3	24
167	Cortical markers of cognitive syndromes in amyotrophic lateral sclerosis. NeuroImage: Clinical, 2018, 19, 675-682.	2.7	24
168	Cortical correlates of behavioural change in amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 380-386.	1.9	24
169	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
170	Skin biopsy in painful and immuneâ€mediated neuropathies. Journal of the Peripheral Nervous System, 2012, 17, 38-45.	3.1	23
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