Marco Luigetti

List of Publications by Year in descending order

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210 papers	7,824 citations	117453 34 h-index	60497 81 g-index
213 all docs	213 docs citations	213 times ranked	9199 citing authors

#	Article	IF	Citations
1	Nusinersen versus Sham Control in Infantile-Onset Spinal Muscular Atrophy. New England Journal of Medicine, 2017, 377, 1723-1732.	13.9	1,533
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.	4.9	1,039
3	Genome-wide Analyses Identify KIF5A as a Novel ALS Gene. Neuron, 2018, 97, 1268-1283.e6.	3.8	517
4	Mutations in the Matrin 3 gene cause familial amyotrophic lateral sclerosis. Nature Neuroscience, 2014, 17, 664-666.	7.1	398
5	Clinical characteristics of patients with familial amyotrophic lateral sclerosis carrying the pathogenic GGGGCC hexanucleotide repeat expansion of C9ORF72. Brain, 2012, 135, 784-793.	3.7	182
6	Randomised controlled trial of methotrexate for chronic inflammatory demyelinating polyradiculoneuropathy (RMC trial): a pilot, multicentre study. Lancet Neurology, The, 2009, 8, 158-164.	4.9	155
7	Heterogeneity of root and nerve ultrasound pattern in CIDP patients. Clinical Neurophysiology, 2014, 125, 160-165.	0.7	142
8	P525L FUS mutation is consistently associated with a severe form of juvenile Amyotrophic Lateral Sclerosis. Neuromuscular Disorders, 2012, 22, 73-75.	0.3	124
9	Antibodies to neurofascin, contactin-1, and contactin-associated protein 1 in CIDP. Neurology: Neuroimmunology and NeuroInflammation, 2020, 7, .	3.1	118
10	Atypical CIDP: diagnostic criteria, progression and treatment response. Data from the Italian CIDP Database. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 125-132.	0.9	108
11	Rituximab in patients with chronic inflammatory demyelinating polyradiculoneuropathy: a report of 13 cases and review of the literature. Journal of Neurology, Neurosurgery and Psychiatry, 2011, 82, 306-308.	0.9	106
12	Large Proportion of Amyotrophic Lateral Sclerosis Cases in Sardinia Due to a Single Founder Mutation of the TARDBP Gene. Archives of Neurology, 2011, 68, 594.	4.9	104
13	Early diagnosis of ATTR amyloidosis through targeted follow-up of identified carriers ofTTRgene mutations*. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 3-9.	1.4	102
14	Contribution of major amyotrophic lateral sclerosis genes to the etiology of sporadic disease. Neurology, 2012, 79, 66-72.	1.5	99
15	Genetic counselling in ALS: facts, uncertainties and clinical suggestions. Journal of Neurology, Neurosurgery and Psychiatry, 2014, 85, 478-485.	0.9	99
16	Mutations in the 3â€² untranslated region of FUS causing FUS overexpression are associated with amyotrophic lateral sclerosis. Human Molecular Genetics, 2013, 22, 4748-4755.	1.4	94
17	Long-term safety and efficacy of patisiran for hereditary transthyretin-mediated amyloidosis with polyneuropathy: 12-month results of an open-label extension study. Lancet Neurology, The, 2021, 20, 49-59.	4.9	93
18	Low Sensitivity of Bone Scintigraphy in Detecting Phe64Leu Mutation-Related Transthyretin Cardiac Amyloidosis. JACC: Cardiovascular Imaging, 2020, 13, 1314-1321.	2.3	82

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19	Natural history of young-adult amyotrophic lateral sclerosis. Neurology, 2008, 71, 876-881.	1.5	81
20	Diagnosis and Treatment of Hereditary Transthyretin Amyloidosis (hATTR) Polyneuropathy: Current Perspectives on Improving Patient Care . Therapeutics and Clinical Risk Management, 2020, Volume 16, 109-123.	0.9	78
21	C9ORF72 hexanucleotide repeat expansions in the Italian sporadic ALS population. Neurobiology of Aging, 2012, 33, 1848.e15-1848.e20.	1.5	76
22	Monitoring effectiveness and safety of Tafamidis in transthyretin amyloidosis in Italy: a longitudinal multicenter study in a non-endemic area. Journal of Neurology, 2016, 263, 916-924.	1.8	76
23	Frequency and time to relapse after discontinuing 6-month therapy with IVIg or pulsed methylprednisolone in CIDP. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 729-734.	0.9	70
24	Repeated courses of granulocyte colonyâ€stimulating factor in amyotrophic lateral sclerosis: Clinical and biological results from a prospective multicenter study. Muscle and Nerve, 2011, 43, 189-195.	1.0	64
25	Ultrasound visualization of nerve morphological alteration at the site of conduction block. Muscle and Nerve, 2009, 40, 1068-1070.	1.0	54
26	ATTRv amyloidosis Italian Registry: clinical and epidemiological data. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2020, 27, 259-265.	1.4	51
27	CHCH10 mutations in an Italian cohort of familial and sporadic amyotrophic lateral sclerosis patients. Neurobiology of Aging, 2015, 36, 1767.e3-1767.e6.	1.5	44
28	TTR-related amyloid neuropathy: clinical, electrophysiological and pathological findings in 15 unrelated patients. Neurological Sciences, 2013, 34, 1057-1063.	0.9	43
29	Rare missense variants of neuronal nicotinic acetylcholine receptor altering receptor function are associated with sporadic amyotrophic lateral sclerosis. Human Molecular Genetics, 2009, 18, 3997-4006.	1.4	42
30	A novel HSPB1 mutation in an Italian patient with CMT2/dHMN phenotype. Journal of the Neurological Sciences, 2010, 298, 114-117.	0.3	42
31	Primary fibroblasts cultures reveal TDP-43 abnormalities in amyotrophic lateral sclerosis patients with and without SOD1 mutations. Neurobiology of Aging, 2015, 36, 2005.e5-2005.e13.	1.5	42
32	TBK1 is associated with ALS and ALS-FTD in Sardinian patients. Neurobiology of Aging, 2016, 43, 180.e1-180.e5.	1.5	40
33	Hereditary transthyretin amyloidosis overview. Neurological Sciences, 2022, 43, 595-604.	0.9	39
34	Nerve ultrasound in hereditary transthyretin amyloidosis: red flags and possible progression biomarkers. Journal of Neurology, 2021, 268, 189-198.	1.8	38
35	Peripheral neuropathy is a common manifestation of mitochondrial diseases: a singleâ€centre experience. European Journal of Neurology, 2016, 23, 1020-1027.	1.7	36
36	6MWT can identify type 3 SMA patients with neuromuscular junction dysfunction. Neuromuscular Disorders, 2017, 27, 879-882.	0.3	36

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37	Sporadic late-onset nemaline myopathy: clinical, pathology and imaging findings in a single center cohort. Journal of Neurology, 2018, 265, 542-551.	1.8	36
38	Assessment of neurological manifestations in hospitalized patients with COVIDâ€19. European Journal of Neurology, 2020, 27, 2322-2328.	1.7	36
39	A multicenter retrospective study of charcotâ€marieâ€tooth disease type 4B (CMT4B) associated with mutations in myotubularinâ€related proteins (MTMRs). Annals of Neurology, 2019, 86, 55-67.	2.8	35
40	Nerve ultrasound findings in neuropathy associated with antiâ€myelinâ€associated glycoprotein antibodies. European Journal of Neurology, 2015, 22, 193-202.	1.7	34
41	Epstein-Barr virus antibodies in serum and cerebrospinal fluid from Multiple sclerosis, Chronic Inflammatory Demyelinating Polyradiculoneuropathy and Amyotrophic Lateral Sclerosis. Journal of Neuroimmunology, 2010, 225, 149-152.	1.1	33
42	Bilateral thalamic stroke transiently reduces arousals and NREM sleep instability. Journal of the Neurological Sciences, 2011, 300, 151-154.	0.3	33
43	Ocular Involvement in Hereditary Amyloidosis. Genes, 2021, 12, 955.	1.0	33
44	SOD1 G93D sporadic amyotrophic lateral sclerosis (SALS) patient with rapid progression and concomitant novel ANG variant. Neurobiology of Aging, 2011, 32, 1924.e15-1924.e18.	1.5	32
45	Uncovering amyotrophic lateral sclerosis phenotypes: Clinical features and long-term follow-up of upper motor neuron-dominant ALS. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2011, 12, 278-282.	2.3	32
46	Ultrasound evaluation in transthyretinâ€related amyloid neuropathy. Muscle and Nerve, 2014, 50, 372-376.	1.0	32
47	Clinical, neurophysiological and pathological findings of HNPP patients with 17p12 deletion: A single-centre experience. Journal of the Neurological Sciences, 2014, 341, 46-50.	0.3	32
48	pSTAT1, pSTAT3, and Tâ€bet as markers of disease activity in chronic inflammatory demyelinating polyradiculoneuropathy. Journal of the Peripheral Nervous System, 2009, 14, 107-117.	1.4	31
49	Primary multifocal lymphoma of peripheral nervous system: Case report and review of the literature. Muscle and Nerve, 2014, 50, 1016-1022.	1.0	30
50	Different nerve ultrasound patterns in charcotâ€marieâ€tooth types and hereditary neuropathy with liability to pressure palsies. Muscle and Nerve, 2018, 57, E18-E23.	1.0	28
51	Sudoscan in the evaluation and follow-up of patients and carriers with TTR mutations: experience from an Italian Centre. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2018, 25, 242-246.	1.4	28
52	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.	1.7	27
53	Improvement of obstructive sleep apneas caused by hydrocephalus associated with Chiari malformation Type II following surgery. Journal of Neurosurgery: Pediatrics, 2010, 6, 336-339.	0.8	25
54	Efficacy of lenalidomide plus dexamethasone for POEMS syndrome relapsed after autologous peripheral stemâ€cell transplantation. American Journal of Hematology, 2012, 87, 641-642.	2.0	24

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55	hATTR Pathology: Nerve Biopsy Results from Italian Referral Centers. Brain Sciences, 2020, 10, 780.	1.1	24
56	Light chain deposition in peripheral nerve as a cause of mononeuritis multiplex in WaldenstrÃ¶m's macroglobulinaemia. Journal of the Neurological Sciences, 2010, 291, 89-91.	0.3	23
57	Non-coding variants contribute to the clinical heterogeneity of TTR amyloidosis. European Journal of Human Genetics, 2017, 25, 1055-1060.	1.4	23
58	Classification of familial amyotrophic lateral sclerosis by family history: effects on frequency of genes mutation. Journal of Neurology, Neurosurgery and Psychiatry, 2012, 83, 1201-1203.	0.9	22
59	Brain MRI in global hypoxiaâ€“ischemia: a map of selective vulnerability. Acta Neurologica Belgica, 2012, 112, 105-107.	0.5	22
60	Persistence of Abnormal Electrophysiological Findings after Carpal Tunnel Release. Journal of Reconstructive Microsurgery, 2013, 29, 511-516.	1.0	22
61	Effect of mexiletine on transitory depression of compound motor action potential in recessive myotonia congenita. Clinical Neurophysiology, 2015, 126, 399-403.	0.7	22
62	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.	0.9	22
63	Recommendations for pre-symptomatic genetic testing for hereditary transthyretin amyloidosis in the era of effective therapy: a multicenter Italian consensus. Orphanet Journal of Rare Diseases, 2020, 15, 348.	1.2	22
64	Heterozygous SOD1 D90A mutation presenting as slowly progressive predominant upper motor neuron amyotrophic lateral sclerosis. Neurological Sciences, 2009, 30, 517-520.	0.9	21
65	AL amyloid neuropathy mimicking a chronic inflammatory demyelinating polyneuropathy. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2012, 19, 53-55.	1.4	20
66	ATXN2 is a modifier of phenotype in ALS patients of Sardinian ancestry. Neurobiology of Aging, 2015, 36, 2906.e1-2906.e5.	1.5	19
67	Nerve conduction studies of the sural nerve: Normative data from a single-center experience. Clinical Neurophysiology, 2012, 123, 1891-1892.	0.7	18
68	Clinical and pathological heterogeneity in a series of 31 patients with IgM-related neuropathy. Journal of the Neurological Sciences, 2012, 319, 75-80.	0.3	18
69	Most recent common ancestor ofTTRVal30Met mutation in Italian population and its potential role in genotype-phenotype correlation. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2015, 22, 73-78.	1.4	18
70	Charcot-Marie-Tooth type 2 and distal hereditary motor neuropathy: Clinical, neurophysiological and genetic findings from a single-centre experience. Clinical Neurology and Neurosurgery, 2016, 144, 67-71.	0.6	18
71	Renal Involvement in Hereditary Transthyretin Amyloidosis: An Italian Single-Centre Experience. Brain Sciences, 2021, 11, 980.	1.1	18
72	Autonomic Dysfunction during Acute SARS-CoV-2 Infection: A Systematic Review. Journal of Clinical Medicine, 2022, 11, 3883.	1.0	18

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73	Sural nerve pathology in ALS patients: a single-centre experience. Neurological Sciences, 2012, 33, 1095-1099.	0.9	17
74	Clinicalâ€“neurophysiological correlations in a series of patients with IgM-related neuropathy. Clinical Neurophysiology, 2013, 124, 1899-1903.	0.7	17
75	Population diversity of the genetically determined TTR expression in human tissues and its implications in TTR amyloidosis. BMC Genomics, 2017, 18, 254.	1.2	17
76	SEIPIN S90L Mutation in an Italian family with CMT2/dHMN and pyramidal signs. Muscle and Nerve, 2010, 42, 448-451.	1.0	16
77	A novel homozygous mutation in the <scp>MTMR2</scp> gene in two siblings with â€˜hypermyelinating neuropathyâ€™. Journal of the Peripheral Nervous System, 2013, 18, 192-194.	1.4	16
78	Triple A syndrome: A novel compound heterozygous mutation in the AAAS gene in an Italian patient without adrenal insufficiency. Journal of the Neurological Sciences, 2010, 290, 150-152.	0.3	15
79	Influence of comorbidities on the phenotype of patients affected by Charcotâ€“Marieâ€“Tooth neuropathy type 1A. Neuromuscular Disorders, 2013, 23, 902-906.	0.3	15
80	A novel LITAF/SIMPLE mutation within a family with a demyelinating form of Charcotâ€“Marieâ€“Tooth disease. Journal of the Neurological Sciences, 2014, 343, 183-186.	0.3	15
81	Admission neurophysiological abnormalities in Guillainâ€“BarrÃ© syndrome: A single-center experience. Clinical Neurology and Neurosurgery, 2015, 135, 6-10.	0.6	15
82	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.	1.4	15
83	Chronic inflammatory demyelinating polyradiculoneuropathy: can a diagnosis be made in patients not fulfilling electrodiagnostic criteria?. European Journal of Neurology, 2021, 28, 620-629.	1.7	15
84	Neurofilament light chain as aÂdisease severity biomarker in ATTRv: data from a single-centre experience. Neurological Sciences, 2022, 43, 2845-2848.	0.9	15
85	A new singleâ€nucleotide deletion of PMP22 in an HNPP family without recurrent palsies. Muscle and Nerve, 2008, 38, 1060-1064.	1.0	14
86	Usefulness of F-18 FDG PET/CT in the Follow-up of POEMS Syndrome After Autologous Peripheral Blood Stem Cell Transplantation. Clinical Nuclear Medicine, 2012, 37, 181-183.	0.7	14
87	Small fibre neuropathy in mitochondrial diseases explored with sudoscan. Clinical Neurophysiology, 2018, 129, 1618-1623.	0.7	14
88	Phenome-wide association study of TTR and RBP4 genes in 361,194 individuals reveals novel insights in the genetics of hereditary and wildtype transthyretin amyloidoses. Human Genetics, 2019, 138, 1331-1340.	1.8	14
89	Causes of chronic neuropathies: a single-center experience. Neurological Sciences, 2019, 40, 1611-1617.	0.9	14
90	Instrumental Evaluation of COVID-19 Related Dysautonomia in Non-Critically-Ill Patients: An Observational, Cross-Sectional Study. Journal of Clinical Medicine, 2021, 10, 5861.	1.0	14

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91	A case of CMT 1B due to Val 102/fs null mutation of the MPZ gene presenting as hyperCKemia. Clinical Neurology and Neurosurgery, 2010, 112, 794-797.	0.6	13
92	Teaching NeuroImages: Peroneal intraneural ganglion cyst: A rare cause of drop foot in a child. Neurology, 2012, 78, e46-e47.	1.5	13
93	Frontotemporal dementia, Parkinsonism and lower motor neuron involvement in a patient with C9ORF72 expansion. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2013, 14, 66-69.	1.1	13
94	Epigenetic profiling of Italian patients identified methylation sites associated with hereditary transthyretin amyloidosis. Clinical Epigenetics, 2020, 12, 176.	1.8	13
95	Progressive brachial plexus enlargement in hereditary transthyretin amyloidosis. Journal of Neurology, 2022, 269, 1905-1912.	1.8	13
96	Realâ€life experience with inotersen in hereditary transthyretin amyloidosis with lateâ€onset phenotype: Data from an earlyâ€access program in Italy. European Journal of Neurology, 2022, 29, 2148-2155.	1.7	13
97	SOD1 G93D mutation presenting as paucisymptomatic amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2009, 10, 479-482.	2.3	12
98	D11Y SOD1 mutation and benign ALS: A consistent genotype-phenotype correlation. Journal of the Neurological Sciences, 2011, 309, 31-33.	0.3	12
99	A novel compound heterozygousALS2mutation in two Italian siblings with juvenile amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2013, 14, 470-472.	1.1	12
100	Oculopharyngeal muscular dystrophy: Clinical and neurophysiological features. Clinical Neurophysiology, 2015, 126, 2406-2408.	0.7	12
101	Sural nerve biopsy in peripheral neuropathies: 30-year experience from a single center. Neurological Sciences, 2020, 41, 341-346.	0.9	12
102	Sporadic hereditary neuropathies misdiagnosed as chronic inflammatory demyelinating polyradiculoneuropathy: Pitfalls and red flags. Journal of the Peripheral Nervous System, 2020, 25, 19-26.	1.4	12
103	Restless Leg Syndrome in Different Types of Demyelinating Neuropathies: A Single-Center Pilot Study. Journal of Clinical Sleep Medicine, 2013, 09, 945-949.	1.4	12
104	Autonomic dysfunction in non-critically ill COVID-19 patients during the acute phase of disease: an observational, cross-sectional study. Neurological Sciences, 2022, 43, 4635-4643.	0.9	12
105	A novel L67P SOD1 mutation in an Italian ALS patient. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2011, 12, 150-152.	2.3	11
106	Clinical features and outcome of patients with autoimmune cerebellar ataxia evaluated with the Scale for the Assessment and Rating of Ataxia. European Journal of Neurology, 2022, 29, 564-572.	1.7	11
107	Neuropathy with Predominant Small Fiber Involvement Associated with Abnormal Anti-MAG Titer. Internal Medicine, 2010, 49, 2627-2629.	0.3	10
108	MRI findings of crossed cerebellar diaschisis in a case of Rasmussenâ€™s encephalitis. Journal of Neurology, 2010, 257, 1748-1750.	1.8	10

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109	Low rate repetitive nerve stimulation in Lambert-Eaton myasthenic syndrome: Peculiar characteristics of decremental pattern from a single-centre experience. Clinical Neurophysiology, 2013, 124, 825-826.	0.7	10
110	Longitudinal assessments in discordant twins with SMA. Neuromuscular Disorders, 2017, 27, 890-893.	0.3	10
111	Does albuterol have an effect on neuromuscular junction dysfunction in spinal muscular atrophy?. Neuromuscular Disorders, 2018, 28, 863-864.	0.3	10
112	Pathological Findings in Chronic Inflammatory Demyelinating Polyradiculoneuropathy: A Single-Center Experience. Brain Sciences, 2020, 10, 383.	1.1	10
113	Skin biopsy and quantitative sensory assessment in an Italian cohort of ATTRv patients with polyneuropathy and asymptomatic carriers: possible evidence of early non-length dependent denervation. Neurological Sciences, 2022, 43, 1359-1364.	0.9	10
114	Gastrointestinal Manifestations in Hereditary Transthyretin Amyloidosis: a Single-Centre Experience. Journal of Gastrointestinal and Liver Diseases, 2020, 29, 339-343.	0.5	10
115	Patisiran in hereditary transthyretin-mediated amyloidosis. Lancet Neurology, The, 2021, 20, 21-23.	4.9	9
116	ATTRv in Lazio-Italy: A High-Prevalence Region in a Non-Endemic Country. Genes, 2021, 12, 829.	1.0	9
117	Teaching Neuro Images : The full-blown neuroimaging of Wernicke encephalopathy. Neurology, 2009, 72, e115.	1.5	8
118	Gadolinium enhancement of the lumbar leptomeninges and roots in a case of ALS. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2010, 11, 412-413.	2.3	8
119	Clinical, electrophysiological and pathological findings of a patient with CMT2 due to the p.Ala738Val mitofusin 2 mutation. Journal of the Neurological Sciences, 2011, 307, 168-170.	0.3	8
120	Creutzfeldtâ€“Jakob disease manifesting as stroke mimic in a 78-year-old patient: Pitfalls and tips in the diagnosis. Journal of the Neurological Sciences, 2014, 346, 343-344.	0.3	8
121	HFE p.H63D polymorphism does not influence ALS phenotype and survival. Neurobiology of Aging, 2015, 36, 2906.e7-2906.e11.	1.5	8
122	Patisiran in hATTR Amyloidosis: Six-Month Latency Period before Efficacy. Brain Sciences, 2021, 11, 515.	1.1	8
123	Psychosocial burden and professional and social support in patients with hereditary transthyretin amyloidosis (ATTRv) and their relatives in Italy. Orphanet Journal of Rare Diseases, 2021, 16, 163.	1.2	8
124	Phenotypic Differences of Glu89Gln Genotype in ATTR Amyloidosis From Endemic Loci: Update From THAOS. Cardiology and Therapy, 2021, 10, 481-490.	1.1	8
125	Ocular Involvement in Hereditary Transthyretin Amyloidosis: A Case Series Describing Novel Potential Biomarkers. Genes, 2021, 12, 927.	1.0	8
126	Replication of association of CHRNA4 rare variants with sporadic amyotrophic lateral sclerosis: The Italian multicentre study. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2012, 13, 580-584.	2.3	7

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127	MRI Neurography Findings in Patients with Idiopathic Brachial Plexopathy: Correlations with Clinical-neurophysiological Data in Eight Consecutive Cases. Internal Medicine, 2013, 52, 2031-2039.	0.3	7
128	Nerve ultrasound in patients with CMT1C: Description of 3 cases. Muscle and Nerve, 2015, 51, 781-782.	1.0	7
129	Movement disorders phenomenology in focal motor seizures. Parkinsonism and Related Disorders, 2019, 61, 161-165.	1.1	7
130	Muscle MRI as a Useful Biomarker in Hereditary Transthyretin Amyloidosis: A Pilot Study. Genes, 2021, 12, 1786.	1.0	7
131	Cerebellar degeneration and ocular myasthenia gravis in a patient with recurring ovarian carcinoma. Neurological Sciences, 2010, 31, 79-81.	0.9	6
132	Teaching Neuro Images : Neurogenic thoracic outlet syndrome. Neurology, 2012, 79, e11.	1.5	6
133	Multimodal CT imaging of a posterior fossa stroke. Neurological Sciences, 2012, 33, 215-216.	0.9	6
134	Inherited neuropathies and deafness caused by a PMP22 point mutation: a case report and a review of the literature. Neurological Sciences, 2013, 34, 1705-1707.	0.9	6
135	Muscle cramps and weakness after teriparatide therapy: A new drugâ€induced myopathy?. Muscle and Nerve, 2013, 47, 615-615.	1.0	6
136	Clinical, electrophysiological and pathological findings in a patient with Charcotâ€“Marieâ€“Tooth disease 4D caused by the NDRG1 Lom mutation. Journal of the Neurological Sciences, 2014, 345, 271-273.	0.3	6
137	Myasthenia gravis with presynaptic neurophysiological signs: Two case reports and literature review. Neuromuscular Disorders, 2015, 25, 646-650.	0.3	6
138	Nerve ultrasound in CMT2E/CMT1F due to NEFL mutation: Confirmation of an axonal pathology. Clinical Neurophysiology, 2016, 127, 2990-2991.	0.7	6
139	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.	1.5	6
140	Autologous peripheral blood stem cell transplantation and the role of lenalidomide in patients affected by poems syndrome. Hematological Oncology, 2018, 36, 392-398.	0.8	6
141	Reassessing IVIg therapy in chronic inflammatory demyelinating polyradiculoneuropathy during COVID-19: a chance to verify the need for chronic maintenance therapy. Neurological Sciences, 2021, 42, 787-789.	0.9	6
142	Mycosis Fungoides as a Cause of Severe Obstructive Sleep Apnea. Internal Medicine, 2011, 50, 1753-1755.	0.3	5
143	A novel LITAF/SIMPLE variant within a family with minimal demyelinating Charcotâ€“Marieâ€“Tooth disease. Neurological Sciences, 2014, 35, 2005-2007.	0.9	5
144	Post-Encephalitic Parkinsonism and Sleep Disorder Responsive to Immunological Treatment. Clinical EEG and Neuroscience, 2016, 47, 324-329.	0.9	5

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145	Recurrent miller fisher: a new case report and a literature review. Clinica Terapeutica, 2017, 168, e208-e213.	0.2	5
146	NEUROLOGIC IMPROVEMENT AFTER PERIPHERAL BLOOD STEM CELL TRANSPLANTATION IN POEMS. Neurology, 2009, 73, 1165-1166.	1.5	4
147	Vertebral Artery Dissection Presenting With Isolated Occipital Headache. Headache, 2010, 50, 1378-1380.	1.8	4
148	Posterior ischaemic myelopathy associated with cocaine abuse. Internal Medicine Journal, 2010, 40, 732-733.	0.5	4
149	Founder effect hypothesis of D11Y SOD1 mutation in Italian amyotrophic lateral sclerosis patients. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2012, 13, 241-242.	2.3	4
150	Rasmussen encephalitis: an unusual cause for intractable seizures in elderly. Neurological Sciences, 2014, 35, 143-145.	0.9	4
151	Teaching NeuroImages. Neurology, 2019, 92, e1000-e1001.	1.5	4
152	Teaching Neuro Image : MRI of diabetic lumbar plexopathy treated with local steroid injection. Neurology, 2009, 72, e32-3.	1.5	3
153	TEACHING NEURO IMAGES : THE FULL-BLOWN NEUROIMAGING OF WERNICKE ENCEPHALOPATHY. Neurology, 2010, 74, 527-528.	1.5	3
154	Progressive ascending myelopathy: atypical forms of multiple sclerosis or what else?. Journal of Neurology, 2011, 258, 1965-1970.	1.8	3
155	Mitochondrial neuropathy: considerations on pathogenesis. European Journal of Neurology, 2016, 23, e55.	1.7	3
156	Restless legs syndrome: A new comorbidity in idiopathic pulmonary fibrosis. Respiratory Medicine, 2020, 170, 105982.	1.3	3
157	The neurophysiological lesson from the Italian CIDP database. Neurological Sciences, 2021, , 1.	0.9	3
158	ANCA-related vasculitic neuropathy mimicking motor neuron disease. Acta Neurologica Belgica, 2008, 108, 109-11.	0.5	3
159	Wernicke's encephalopathy following chronic diarrhoea. Acta Neurologica Belgica, 2011, 111, 257.	0.5	3
160	A compound score to screen patients with hereditary transthyretin amyloidosis. Journal of Neurology, 2022, , .	1.8	3
161	A novel GJB1 mutation in an Italian patient with Charcotâ€“Marieâ€“Tooth disease and pyramidal signs. Muscle and Nerve, 2011, 44, 613-615.	1.0	2
162	Peripheral neuropathy and 46XY gonadal dysgenesis: Confirmation of a heterogeneous entity. Clinical Neurology and Neurosurgery, 2012, 114, 748-750.	0.6	2

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163	Cranial botulism. Neuromuscular Disorders, 2012, 22, 995-996.	0.3	2
164	An Uncommon Cause of Headache Resolution: Spontaneous Ventriculostomy in Obstructive Hydrocephalus. Headache, 2013, 53, 1356-1357.	1.8	2
165	Distinct lymphocytes subsets in IgM-related neuropathy: clinical-immunological correlations. Neurological Sciences, 2015, 36, 303-308.	0.9	2
166	‘White Nails'. European Neurology, 2015, 73, 89-89.	0.6	2
167	A Case of Hemiabdominal Myoclonus. Clinical EEG and Neuroscience, 2015, 46, 331-334.	0.9	2
168	Central conduction abnormalities in patients receiving levodopa-carbidopa intestinal gel infusion. Neurological Sciences, 2017, 38, 1869-1872.	0.9	2
169	Small Fibre Involvement in Multifocal Motor Neuropathy Explored with Sudoscan: A Single-Centre Experience. Diagnostics, 2020, 10, 755.	1.3	2
170	Uncommon pathological findings in sural nerve biopsy from a patient with Churg-Strauss related multiple mononeuropathy. Acta ReumatolÃ³gica Portuguesa, 2013, 38, 286-9.	0.2	2
171	Abnormal vascular smooth muscle cell proliferation in sural nerve biopsy from a patient with sensorimotor axonal neuropathy. Neuropathology, 2011, 31, 197-198.	0.7	1
172	Teaching Neuro Images : Cochleitis. Neurology, 2011, 77, e109.	1.5	1
173	Hirayama flexion myelopathy. Acta Neurologica Belgica, 2013, 113, 543-544.	0.5	1
174	Bilateral thoracic long nerve involvement in motor multifocal neuropathy. Journal of Neurology, Neurosurgery and Psychiatry, 2013, 84, 584-584.	0.9	1
175	Posterior interosseous nerve syndrome due to radioulnar joint cyst. Muscle and Nerve, 2013, 48, 842-843.	1.0	1
176	Lower motor neuron involvement in longitudinally extensive transverse myelitis with and without aquaporin-4 antibodies. Clinical Neurophysiology, 2014, 125, 1925-1926.	0.7	1
177	Subacute combined degeneration. Acta Neurologica Belgica, 2014, 114, 221-222.	0.5	1
178	Dysautonomia as Onset Symptom of Myotonic Dystrophy Type 2. European Neurology, 2018, 79, 166-170.	0.6	1
179	PERIPHERAL NERVOUS SYSTEM INVOLVEMENT IN LYMPHOPROLIFERATIVE DISORDERS. Mediterranean Journal of Hematology and Infectious Diseases, 2018, 10, e2018057.	0.5	1
180	Progressive axonal polyneuropathy in a mitochondrial disorder: an uncommon association with familial amyloid neuropathy. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2018, 25, 261-262.	1.4	1

#	Article	IF	Citations
181	Nine syndrome. Acta Neurologica Belgica, 2019, 119, 475-476.	0.5	1
182	Isolated light chain deposition disease neuropathy in a patient with multiple myeloma. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2020, 27, 67-68.	1.4	1
183	Prolonged distal motor latency of median nerve does not improve diagnostic accuracy for CIDP. Journal of Neurology, 2021, , 1.	1.8	1
184	Fahr's Syndrome in a Sporadic Case. Neurology India, 2019, 67, 1573.	0.2	1
185	Somatosensory Evoked Potentials of Inferior Alveolar Nerve: confirmation of a possible non-invasive neurophysiological approach. Clinica Terapeutica, 2016, 167, 25-8.	0.2	1
186	Generation of an induced pluripotent stem cell line (UCSCi002-A) from a patient with a variant in TARDBP gene associated with familial amyotrophic lateral sclerosis and frontotemporal dementia. Stem Cell Research, 2022, 62, 102825.	0.3	1
187	NATURAL HISTORY OF YOUNG-ADULT AMYOTROPHIC LATERAL SCLEROSIS. Neurology, 2009, 73, 648-650.	1.5	0
188	Retinal Detachment with an Unusual Shape. Internal Medicine, 2009, 48, 1777-1778.	0.3	0
189	Spontaneous regression of a midbrain lesion in a patient with chronic transtentorial herniation: is it a pre-syrinx?. Journal of Neurology, 2010, 257, 848-850.	1.8	0
190	Pseudoperipheral tongue weakness. Journal of Neurology, Neurosurgery and Psychiatry, 2010, 81, 1024-1025.	0.9	0
191	Spontaneous Bilateral Internal Carotid Artery Dissection Presenting With Right Hemifacial Pain. Headache, 2012, 52, 1574-1575.	1.8	0
192	Nocturnal Cramps. JAMA Neurology, 2013, 70, 792.	4.5	0
193	Benedictine hand of â€˜centralâ€™ origin. Emergency Medicine Journal, 2013, 30, 173-173.	0.4	0
194	Vanishing aneurysm. Journal of NeuroInterventional Surgery, 2013, 5, e26-e26.	2.0	0
195	Lumbosacral multiradiculopathy responsive to antibiotic therapy: description of four patients with lumbar spondylosis and a superimposed Lyme disease. Acta Neurologica Belgica, 2014, 114, 297-301.	0.5	0
196	Reply to â€œThe significance of significance: Overstating in the setting of many comparisonsâ€• Clinical Neurophysiology, 2014, 125, 1073-1074.	0.7	0
197	An Uncommon Diagnostic Algorithm for a Common Condition. Internal Medicine, 2015, 54, 1679-1679.	0.3	0
198	Skin Changes in POEMS Syndrome. European Neurology, 2015, 73, 112-112.	0.6	0

#	Article	IF	Citations
199	Restless Legs Syndrome and lateralized periodic movements due to a spinal schwannoma. Sleep and Biological Rhythms, 2015, 13, 106-108.	0.5	0
200	SjÃ¶gren's syndrome presenting with isolated sensory axonal polyneuropathy. International Journal of Rheumatic Diseases, 2017, 20, 2165-2167.	0.9	0
201	Chlorambucil Plus Rituximab as Front-Line Therapy for Elderly Patients with Comorbidities Affected by Waldenstrom Macroglobulinemia: A Single Center Experience. Journal of Blood & Lymph, 2018, 08, .	0.0	0
202	Imaging Features of Varicella Zoster Virus Cranial Multiple Mononeuropathies. European Neurology, 2018, 79, 315-316.	0.6	0
203	Reply to â€œWork-up for mitochondrial small fiber neuropathy requires application of skin biopsiesâ€. Clinical Neurophysiology, 2018, 129, 2053.	0.7	0
204	Clinical Health Care during the COVID-19 Pandemic. Journal of Clinical Medicine, 2020, 9, 3559.	1.0	0
205	Pediatric Motor Inflammatory Neuropathy: The Role of Antiphospholipid Antibodies. Brain Sciences, 2020, 10, 156.	1.1	0
206	Letter: Allograft Nerve Repair Reduces Postoperative Neuropathic Pain Following Nerve Biopsy. Neurosurgery, 2021, 88, E568-E568.	0.6	0
207	Nerve Biopsy in Peripheral Neuropathies: Not All Water Is under the Bridge. Brain Sciences, 2021, 11, 550.	1.1	0
208	Thr124Met myelin protein zero mutation mimicking motor neuron disease. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2021, , 1-6.	1.1	0
209	Proximal basilar artery fenestration with bridging artery appearance. Acta Neurologica Belgica, 2009, 109, 243.	0.5	0
210	Guillainâ€“BarrÃ© syndrome from an emergency department view: how to better predict the outcome?. Neurological Research, 2022, , 1-5.	0.6	0