

Giulia Bisogni

List of Publications by Year in descending order

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Version: 2024-02-01

46
papers

1,293
citations

361413

20
h-index

377865

34
g-index

48
all docs

48
docs citations

48
times ranked

2015
citing authors

#	ARTICLE	IF	CITATIONS
1	Progressive brachial plexus enlargement in hereditary transthyretin amyloidosis. <i>Journal of Neurology</i> , 2022, 269, 1905-1912.	3.6	13
2	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. <i>Stem Cell Research</i> , 2022, 62, 102825.	0.7	1
3	Nerve ultrasound in hereditary transthyretin amyloidosis: red flags and possible progression biomarkers. <i>Journal of Neurology</i> , 2021, 268, 189-198.	3.6	38
4	Long-term safety and efficacy of patisiran for hereditary transthyretin-mediated amyloidosis with polyneuropathy: 12-month results of an open-label extension study. <i>Lancet Neurology</i> , The, 2021, 20, 49-59.	10.2	93
5	Novel variants and cellular studies on patientsâ€™ primary fibroblasts support a role for NEK1 missense variants in ALS pathogenesis. <i>Human Molecular Genetics</i> , 2021, 30, 65-71.	2.9	7
6	Patisiran in hATTR Amyloidosis: Six-Month Latency Period before Efficacy. <i>Brain Sciences</i> , 2021, 11, 515.	2.3	8
7	Psychosocial burden and professional and social support in patients with hereditary transthyretin amyloidosis (ATTRv) and their relatives in Italy. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 163.	2.7	8
8	Thr124Met myelin protein zero mutation mimicking motor neuron disease. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2021, , 1-6.	1.7	0
9	Generation of an induced pluripotent stem cell line (UCSCi001-A) from a patient with early-onset amyotrophic lateral sclerosis carrying a FUS variant. <i>Stem Cell Research</i> , 2021, 55, 102461.	0.7	0
10	Association of Variants in the <i>SPTLC1</i> Gene With Juvenile Amyotrophic Lateral Sclerosis. <i>JAMA Neurology</i> , 2021, 78, 1236.	9.0	46
11	Reassessing IVIg therapy in chronic inflammatory demyelinating polyradiculoneuropathy during COVID-19: a chance to verify the need for chronic maintenance therapy. <i>Neurological Sciences</i> , 2021, 42, 787-789.	1.9	6
12	A longitudinal study defined circulating microRNAs as reliable biomarkers for disease prognosis and progression in ALS human patients. <i>Cell Death Discovery</i> , 2021, 7, 4.	4.7	36
13	Isolated light chain deposition disease neuropathy in a patient with multiple myeloma. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2020, 27, 67-68.	3.0	1
14	Sural nerve biopsy in peripheral neuropathies: 30-year experience from a single center. <i>Neurological Sciences</i> , 2020, 41, 341-346.	1.9	12
15	High-Throughput Genetic Testing in ALS: The Challenging Path of Variant Classification Considering the ACMG Guidelines. <i>Genes</i> , 2020, 11, 1123.	2.4	15
16	ATTRv amyloidosis Italian Registry: clinical and epidemiological data. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2020, 27, 259-265.	3.0	51
17	hATTR Pathology: Nerve Biopsy Results from Italian Referral Centers. <i>Brain Sciences</i> , 2020, 10, 780.	2.3	24
18	Recommendations for pre-symptomatic genetic testing for hereditary transthyretin amyloidosis in the era of effective therapy: a multicenter Italian consensus. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 348.	2.7	22

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19	Pathological Findings in Chronic Inflammatory Demyelinating Polyradiculoneuropathy: A Single-Center Experience. <i>Brain Sciences</i> , 2020, 10, 383.	2.3	10
20	<p>Diagnosis and Treatment of Hereditary Transthyretin Amyloidosis (hATTR) Polyneuropathy: Current Perspectives on Improving Patient Care<p>. <i>Therapeutics and Clinical Risk Management</i> , 2020, Volume 16, 109-123.	2.0	78
21	Antibodies to neurofascin, contactin-1, and contactin-associated protein 1 in CIDP. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2020, 7, .	6.0	118
22	SOD1 p.D12Y variant is associated with amyotrophic lateral sclerosis/distal myopathy spectrum. <i>European Journal of Neurology</i> , 2020, 27, 1304-1309.	3.3	4
23	Gastrointestinal Manifestations in Hereditary Transthyretin Amyloidosis: a Single-Centre Experience. <i>Journal of Gastrointestinal and Liver Diseases</i> , 2020, 29, 339-343.	0.9	10
24	Small Fibre Involvement in Multifocal Motor Neuropathy Explored with Sudoscan: A Single-Centre Experience. <i>Diagnostics</i> , 2020, 10, 755.	2.6	2
25	Coexistence of variants in TBK1 and in other ALS-related genes elucidates an oligogenic model of pathogenesis in sporadic ALS. <i>Neurobiology of Aging</i> , 2019, 84, 239.e9-239.e14.	3.1	21
26	ATXN1 intermediate-length polyglutamine expansions are associated with amyotrophic lateral sclerosis. <i>Neurobiology of Aging</i> , 2018, 64, 157.e1-157.e5.	3.1	34
27	Sudoscan in the evaluation and follow-up of patients and carriers with TTR mutations: experience from an Italian Centre. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2018, 25, 242-246.	3.0	28
28	Progressive axonal polyneuropathy in a mitochondrial disorder: an uncommon association with familial amyloid neuropathy. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2018, 25, 261-262.	3.0	1
29	Matrin 3 variants are frequent in Italian ALS patients. <i>Neurobiology of Aging</i> , 2017, 49, 218.e1-218.e7.	3.1	35
30	Monitoring effectiveness and safety of Tafamidis in transthyretin amyloidosis in Italy: a longitudinal multicenter study in a non-endemic area. <i>Journal of Neurology</i> , 2016, 263, 916-924.	3.6	76
31	Charcot-Marie-Tooth type 2 and distal hereditary motor neuropathy: Clinical, neurophysiological and genetic findings from a single-centre experience. <i>Clinical Neurology and Neurosurgery</i> , 2016, 144, 67-71.	1.4	18
32	Skin Changes in POEMS Syndrome. <i>European Neurology</i> , 2015, 73, 112-112.	1.4	0
33	Primary fibroblasts cultures reveal TDP-43 abnormalities in amyotrophic lateral sclerosis patients with and without SOD1 mutations. <i>Neurobiology of Aging</i> , 2015, 36, 2005.e5-2005.e13.	3.1	42
34	Distinct lymphocytes subsets in IgM-related neuropathy: clinical-immunological correlations. <i>Neurological Sciences</i> , 2015, 36, 303-308.	1.9	2
35	‘White Nails'. <i>European Neurology</i> , 2015, 73, 89-89.	1.4	2
36	<i>ATXN2</i> polyQ intermediate repeats are a modifier of ALS survival. <i>Neurology</i> , 2015, 84, 251-258.	1.1	52

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37	Ultrasound evaluation in transthyretin-related amyloid neuropathy. <i>Muscle and Nerve</i> , 2014, 50, 372-376.	2.2	32
38	Clinical, neurophysiological and pathological findings of HNPP patients with 17p12 deletion: A single-centre experience. <i>Journal of the Neurological Sciences</i> , 2014, 341, 46-50.	0.6	32
39	Clinical, electrophysiological and pathological findings in a patient with Charcot-Marie-Tooth disease 4D caused by the NDRG1 Lom mutation. <i>Journal of the Neurological Sciences</i> , 2014, 345, 271-273.	0.6	6
40	TTR-related amyloid neuropathy: clinical, electrophysiological and pathological findings in 15 unrelated patients. <i>Neurological Sciences</i> , 2013, 34, 1057-1063.	1.9	43
41	Mutations in the 3' untranslated region of FUS causing FUS overexpression are associated with amyotrophic lateral sclerosis. <i>Human Molecular Genetics</i> , 2013, 22, 4748-4755.	2.9	94
42	Restless Leg Syndrome in Different Types of Demyelinating Neuropathies: A Single-Center Pilot Study. <i>Journal of Clinical Sleep Medicine</i> , 2013, 09, 945-949.	2.6	12
43	Sural nerve pathology in ALS patients: a single-centre experience. <i>Neurological Sciences</i> , 2012, 33, 1095-1099.	1.9	17
44	Contribution of major amyotrophic lateral sclerosis genes to the etiology of sporadic disease. <i>Neurology</i> , 2012, 79, 66-72.	1.1	99
45	D11Y SOD1 mutation and benign ALS: A consistent genotype-phenotype correlation. <i>Journal of the Neurological Sciences</i> , 2011, 309, 31-33.	0.6	12
46	Uncovering amyotrophic lateral sclerosis phenotypes: Clinical features and long-term follow-up of upper motor neuron-dominant ALS. <i>Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders</i> , 2011, 12, 278-282.	2.1	32