Eleonora Dalla Bella

List of Publications by Year
in descending order

28	799	516710 16	501196 28
papers	citations	h-index	g-index

29	29	29	1394
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Rapamycin treatment for amyotrophic lateral sclerosis. Medicine (United States), 2018, 97, e11119.	1.0	96
2	Small fiber neuropathy is a common feature of Ehlers-Danlos syndromes. Neurology, 2016, 87, 155-159.	1.1	90
3	Amyotrophic lateral sclerosis causes small fiber pathology. European Journal of Neurology, 2016, 23, 416-420.	3.3	65
4	The Multiple Sclerosis Knowledge Questionnaire: a self-administered instrument for recently diagnosed patients. Multiple Sclerosis Journal, 2010, 16, 100-111.	3.0	50
5	Beyond the consensus criteria: multiple cognitive profiles in amyotrophic lateral sclerosis?. Cortex, 2016, 81, 162-167.	2.4	45
6	The MITOS system predicts long-term survival in amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 1180-1185.	1.9	42
7	Amyotrophic lateral sclerosis patientsâ€™ and caregivers' distress and loneliness during COVID-19 lockdown. Journal of Neurology, 2021, 268, 420-423.	3.6	33
8	The unfolded protein response in amyotrophic later sclerosis: results of a phase 2 trial. Brain, 2021, 144, 2635-2647.	7.6	33
9	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
10	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
11	Side and time variability of intraepidermal nerve fiber density. Neurology, 2015, 84, 2368-2371.	1.1	29
12	Development and validation of a patient self-assessed questionnaire on satisfaction with communication of the multiple sclerosis diagnosis. Multiple Sclerosis Journal, 2010, 16, 1237-1247.	3.0	27
13	Cortical markers of cognitive syndromes in amyotrophic lateral sclerosis. NeuroImage: Clinical, 2018, 19, 675-682.	2.7	24
14	Cortical correlates of behavioural change in amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 380-386.	1.9	24
15	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
16	Cortical thinning trajectories across disease stages and cognitive impairment in amyotrophic lateral sclerosis. Cortex, 2020, 131, 284-294.	2.4	18
17	Ischemic stroke as clinical onset of POEMS syndrome. Journal of Neurology, 2013, 260, 3178-3181.	3.6	17
18	Facial Onset Sensory and Motor Neuronopathy. Neurology: Clinical Practice, 2021, 11, 147-157.	1.6	16

#	Article	IF	Citations
19	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
20	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
21	Cerebellar ataxia, neuropathy, and vestibular areflexia syndrome: a slowly progressive disorder with stereotypical presentation. Journal of Neurology, 2016, 263, 245-249.	3.6	13
22	Experience of an information aid for newly diagnosed multiple sclerosis patients: a qualitative study on the SIMSâ€Trial. Health Expectations, 2014, 17, 36-48.	2.6	12
23	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
24	Cognitive Syndromes and C9orf72 Mutation Are Not Related to Cerebellar Degeneration in Amyotrophic Lateral Sclerosis. Frontiers in Neuroscience, 2019, 13, 440.	2.8	10
25	The first case of the TARDBP p.G294V mutation in a homozygous state: is a single pathogenic allele sufficient to cause ALS?. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 273-279.	1.7	10
26	Myelin protein zero <scp>Arg36Gly</scp> mutation with very late onset and rapidly progressive painful neuropathy. Journal of the Peripheral Nervous System, 2012, 17, 422-425.	3.1	9
27	Late-onset and fast progressive neuropathy and cardiomyopathy in Val32Ala transthyretin gene mutation. Neurological Sciences, 2019, 40, 1267-1269.	1.9	4
28	Spinal and bulbar muscular atrophy and Charcotâ€“Marieâ€“Tooth type 1A: Co-existence of two rare neuromuscular genetic diseases in the same patient. Neuromuscular Disorders, 2015, 25, 800-801.	0.6	3