

# Eleonora Dalla Bella

## List of Publications by Year in descending order

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28  
papers

799  
citations

516710

16  
h-index

501196

28  
g-index

29  
all docs

29  
docs citations

29  
times ranked

1394  
citing authors

#	ARTICLE	IF	CITATIONS
1	Facial Onset Sensory and Motor Neuronopathy. <i>Neurology: Clinical Practice</i> , 2021, 11, 147-157.	1.6	16
2	Amyotrophic lateral sclerosis patients' and caregivers' distress and loneliness during COVID-19 lockdown. <i>Journal of Neurology</i> , 2021, 268, 420-423.	3.6	33
3	Cognitive reserve is associated with altered clinical expression in amyotrophic lateral sclerosis. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2021, 22, 237-247.	1.7	12
4	The unfolded protein response in amyotrophic later sclerosis: results of a phase 2 trial. <i>Brain</i> , 2021, 144, 2635-2647.	7.6	33
5	Cognitive and behavioural impairment in amyotrophic lateral sclerosis: A landmark of the disease? A mini review of longitudinal studies. <i>Neuroscience Letters</i> , 2021, 754, 135898.	2.1	15
6	The first case of the <i>TARDBP</i> p.G294V mutation in a homozygous state: is a single pathogenic allele sufficient to cause ALS?. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2020, 21, 273-279.	1.7	10
7	Cortical thinning trajectories across disease stages and cognitive impairment in amyotrophic lateral sclerosis. <i>Cortex</i> , 2020, 131, 284-294.	2.4	18
8	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. <i>Journal of Clinical Medicine</i> , 2020, 9, 412.	2.4	24
9	Late-onset and fast progressive neuropathy and cardiomyopathy in Val32Ala transthyretin gene mutation. <i>Neurological Sciences</i> , 2019, 40, 1267-1269.	1.9	4
10	Cognitive Syndromes and C9orf72 Mutation Are Not Related to Cerebellar Degeneration in Amyotrophic Lateral Sclerosis. <i>Frontiers in Neuroscience</i> , 2019, 13, 440.	2.8	10
11	Cortical correlates of behavioural change in amyotrophic lateral sclerosis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2019, 90, 380-386.	1.9	24
12	Rapamycin treatment for amyotrophic lateral sclerosis. <i>Medicine (United States)</i> , 2018, 97, e11119.	1.0	96
13	Cortical markers of cognitive syndromes in amyotrophic lateral sclerosis. <i>NeuroImage: Clinical</i> , 2018, 19, 675-682.	2.7	24
14	Protein misfolding, amyotrophic lateral sclerosis and guanabenz: protocol for a phase II RCT with futility design (ProMiSe trial). <i>BMJ Open</i> , 2017, 7, e015434.	1.9	14
15	Amyotrophic lateral sclerosis causes small fiber pathology. <i>European Journal of Neurology</i> , 2016, 23, 416-420.	3.3	65
16	Beyond the consensus criteria: multiple cognitive profiles in amyotrophic lateral sclerosis?. <i>Cortex</i> , 2016, 81, 162-167.	2.4	45
17	Small fiber neuropathy is a common feature of Ehlers-Danlos syndromes. <i>Neurology</i> , 2016, 87, 155-159.	1.1	90
18	Cerebellar ataxia, neuropathy, and vestibular areflexia syndrome: a slowly progressive disorder with stereotypical presentation. <i>Journal of Neurology</i> , 2016, 263, 245-249.	3.6	13

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19	Side and time variability of intraepidermal nerve fiber density. <i>Neurology</i> , 2015, 84, 2368-2371.	1.1	29
20	Erythropoietin in amyotrophic lateral sclerosis: a multicentre, randomised, double blind, placebo controlled, phase III study. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2015, 86, 879-886.	1.9	32
21	The MITOS system predicts long-term survival in amyotrophic lateral sclerosis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2015, 86, 1180-1185.	1.9	42
22	Spinal and bulbar muscular atrophy and Charcot-Marie-Tooth type 1A: Co-existence of two rare neuromuscular genetic diseases in the same patient. <i>Neuromuscular Disorders</i> , 2015, 25, 800-801.	0.6	3
23	Heterozygous D90A-SOD1 mutation in a patient with facial onset sensory motor neuronopathy (FOSMN) syndrome: a bridge to amyotrophic lateral sclerosis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2014, 85, 1009-1011.	1.9	32
24	Experience of an information aid for newly diagnosed multiple sclerosis patients: a qualitative study on the SIMS trial. <i>Health Expectations</i> , 2014, 17, 36-48.	2.6	12
25	Ischemic stroke as clinical onset of POEMS syndrome. <i>Journal of Neurology</i> , 2013, 260, 3178-3181.	3.6	17
26	Myelin protein zero <sc>Arg36Gly</sc> mutation with very late onset and rapidly progressive painful neuropathy. <i>Journal of the Peripheral Nervous System</i> , 2012, 17, 422-425.	3.1	9
27	The Multiple Sclerosis Knowledge Questionnaire: a self-administered instrument for recently diagnosed patients. <i>Multiple Sclerosis Journal</i> , 2010, 16, 100-111.	3.0	50
28	Development and validation of a patient self-assessed questionnaire on satisfaction with communication of the multiple sclerosis diagnosis. <i>Multiple Sclerosis Journal</i> , 2010, 16, 1237-1247.	3.0	27