Fabiola De Marchi

List of Publications by Year
in descending order

33	816	623734	552781
papers	citations	14 h-index	26 g-index

33	33	33	1130
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Study protocol on the safety and feasibility of a normocaloric ketogenic diet in people with amyotrophic lateral sclerosis. Nutrition, 2022, 94, 111525.	2.4	7
2	Neurological emergency at the COVID-19 pandemic: report from a referral hospital in Eastern Piedmont, Italy. Neurological Sciences, 2022, 43, 2195.	1.9	2
3	The Potential Role of Peripheral Oxidative Stress on the Neurovascular Unit in Amyotrophic Lateral Sclerosis Pathogenesis: A Preliminary Report from Human and In Vitro Evaluations. Biomedicines, 2022, 10, 691.	3.2	8
4	The Role of Tau beyond Alzheimerâ€™s Disease: A Narrative Review. Biomedicines, 2022, 10, 760.	3.2	12
5	Characterization of the p.L145F and p.S135N Mutations in SOD1: Impact on the Metabolism of Fibroblasts Derived from Amyotrophic Lateral Sclerosis Patients. Antioxidants, 2022, 11, 815.	5.1	3
6	Telehealth approach for amyotrophic lateral sclerosis patients: the experience during COVIDâ€19 pandemic. Acta Neurologica Scandinavica, 2021, 143, 489-496.	2.1	19
7	Exploring the use of educational materials for increasing participation in a stretching program: a quality improvement project in people with motor neuron disease. European Journal of Physical and Rehabilitation Medicine, 2021, 57, 78-84.	2.2	1
8	Telehealth in Neurodegenerative Diseases: Opportunities and Challenges for Patients and Physicians. Brain Sciences, 2021, 11, 237.	2.3	34
9	Patient reported outcomes in ALS: characteristics of the self-entry ALS Functional Rating Scale-revised and the Activities-specific Balance Confidence Scale. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2021, 22, 467-477.	1.7	10
10	Cognitive dysfunction in amyotrophic lateral sclerosis: can we predict it?. Neurological Sciences, 2021, 42, 2211-2222.	1.9	16
11	Accelerated Early Progression of Amyotrophic Lateral Sclerosis over the COVID-19 Pandemic. Brain Sciences, 2021, 11, 1291.	2.3	15
12	C9ORF72 Repeat Expansion Affects the Proteome of Primary Skin Fibroblasts in ALS. International Journal of Molecular Sciences, 2021, 22, 10385.	4.1	6
13	Do ecological factors influence the clinical presentation of amyotrophic lateral sclerosis?. Journal of the Neurological Sciences, 2021, 429, 117704.	0.6	0
14	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
15	Immunity in amyotrophic lateral sclerosis: blurred lines between excessive inflammation and inefficient immune responses. Brain Communications, 2020, 2, fcaa124.	3.3	53
16	Generation of an induced pluripotent stem cell line, CSSi011-A (6534), from an Amyotrophic lateral sclerosis patient with heterozygous L145F mutation in SOD1 gene. Stem Cell Research, 2020, 47, 101924.	0.7	2
17	The NEALS primary lateral sclerosis registry. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 74-81.	1.7	5
18	Detection of White Matter Ultrastructural Changes for Amyotrophic Lateral Sclerosis Characterization: A Diagnostic Study from Dti-Derived Data. Brain Sciences, 2020, 10, 996.	2.3	5

#	Article	IF	Citations
19	Decline of cognitive and behavioral functions in amyotrophic lateral sclerosis: a longitudinal study. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 373-379.	1.7	40
20	A prospective longitudinal study on the microbiota composition in amyotrophic lateral sclerosis. BMC Medicine, 2020, 18, 153.	5.5	78
21	Reply to: Amyotrophic lateral sclerosis with depression, cognitive impairment, and mortality. Acta Neurologica Scandinavica, 2020, 142, 86-87.	2.1	0
22	Patient reported outcome measures (PROMs) in amyotrophic lateral sclerosis. Journal of Neurology, 2020, 267, 1754-1759.	3.6	15
23	ALS phenotype is influenced by age, sex, and genetics. Neurology, 2020, 94, e802-e810.	1.1	99
24	Telemedicine and technological devices for amyotrophic lateral sclerosis in the era of COVID-19. Neurological Sciences, 2020, 41, 1365-1367.	1.9	23
25	Depression and risk of cognitive dysfunctions in amyotrophic lateral sclerosis. Acta Neurologica Scandinavica, 2019, 139, 438-445.	2.1	20
26	Integration of a palliative care specialist in an amyotrophic lateral sclerosis clinic: Observations from one center. Muscle and Nerve, 2019, 60, 137-140.	2.2	30
27	Adjusted cost analysis of video televisits for the care of people with amyotrophic lateral sclerosis. Muscle and Nerve, 2019, 60, 147-154.	2.2	23
28	A pilot trial of RNS60 in amyotrophic lateral sclerosis. Muscle and Nerve, 2019, 59, 303-308.	2.2	29
29	Ptosis and bulbar onset: an unusual phenotype of familial ALS?. Neurological Sciences, 2018, 39, 377-378.	1.9	5
30	Chronic obstructive pulmonary disease may complicate Alzheimerâ€™s disease: a comorbidity problem. Neurological Sciences, 2018, 39, 1585-1589.	1.9	18
31	Headache in immigrant patients: similarities and differences with Italian population. Neurological Sciences, 2018, 39, 749-752.	1.9	2
32	Marchiafava-Bignami Disease with frontal cortex involvement and late onset, long-lasting psychiatric symptoms: a case report. Rivista Di Psichiatria, 2016, 51, 79-82.	0.6	2
33	Predicting Cognitive Decline in ParkinsonÃ¢â‚¬â„¢s Disease: Can We Ask the Genes?. Frontiers in Neurology, 2014, 5, 224.	2.4	11