

Fabiola De Marchi

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7274978/publications.pdf>

Version: 2024-02-01

33
papers

816
citations

623734

14
h-index

552781

26
g-index

33
all docs

33
docs citations

33
times ranked

1130
citing authors

#	ARTICLE	IF	CITATIONS
1	Common and rare variant association analyses in amyotrophic lateral sclerosis identify 15 risk loci with distinct genetic architectures and neuron-specific biology. <i>Nature Genetics</i> , 2021, 53, 1636-1648.	21.4	223
2	ALS phenotype is influenced by age, sex, and genetics. <i>Neurology</i> , 2020, 94, e802-e810.	1.1	99
3	A prospective longitudinal study on the microbiota composition in amyotrophic lateral sclerosis. <i>BMC Medicine</i> , 2020, 18, 153.	5.5	78
4	Immunity in amyotrophic lateral sclerosis: blurred lines between excessive inflammation and inefficient immune responses. <i>Brain Communications</i> , 2020, 2, fcaa124.	3.3	53
5	Decline of cognitive and behavioral functions in amyotrophic lateral sclerosis: a longitudinal study. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2020, 21, 373-379.	1.7	40
6	Telehealth in Neurodegenerative Diseases: Opportunities and Challenges for Patients and Physicians. <i>Brain Sciences</i> , 2021, 11, 237.	2.3	34
7	Integration of a palliative care specialist in an amyotrophic lateral sclerosis clinic: Observations from one center. <i>Muscle and Nerve</i> , 2019, 60, 137-140.	2.2	30
8	A pilot trial of RNS60 in amyotrophic lateral sclerosis. <i>Muscle and Nerve</i> , 2019, 59, 303-308.	2.2	29
9	Adjusted cost analysis of video televisits for the care of people with amyotrophic lateral sclerosis. <i>Muscle and Nerve</i> , 2019, 60, 147-154.	2.2	23
10	Telemedicine and technological devices for amyotrophic lateral sclerosis in the era of COVID-19. <i>Neurological Sciences</i> , 2020, 41, 1365-1367.	1.9	23
11	Depression and risk of cognitive dysfunctions in amyotrophic lateral sclerosis. <i>Acta Neurologica Scandinavica</i> , 2019, 139, 438-445.	2.1	20
12	Telehealth approach for amyotrophic lateral sclerosis patients: the experience during COVID-19 pandemic. <i>Acta Neurologica Scandinavica</i> , 2021, 143, 489-496.	2.1	19
13	Chronic obstructive pulmonary disease may complicate Alzheimer's disease: a comorbidity problem. <i>Neurological Sciences</i> , 2018, 39, 1585-1589.	1.9	18
14	Cognitive dysfunction in amyotrophic lateral sclerosis: can we predict it?. <i>Neurological Sciences</i> , 2021, 42, 2211-2222.	1.9	16
15	Patient reported outcome measures (PROMs) in amyotrophic lateral sclerosis. <i>Journal of Neurology</i> , 2020, 267, 1754-1759.	3.6	15
16	Accelerated Early Progression of Amyotrophic Lateral Sclerosis over the COVID-19 Pandemic. <i>Brain Sciences</i> , 2021, 11, 1291.	2.3	15
17	The Role of Tau beyond Alzheimer's Disease: A Narrative Review. <i>Biomedicines</i> , 2022, 10, 760.	3.2	12
18	Predicting Cognitive Decline in Parkinson's Disease: Can We Ask the Genes?. <i>Frontiers in Neurology</i> , 2014, 5, 224.	2.4	11

#	ARTICLE	IF	CITATIONS
19	Patient reported outcomes in ALS: characteristics of the self-entry ALS Functional Rating Scale-revised and the Activities-specific Balance Confidence Scale. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2021, 22, 467-477.	1.7	10
20	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. <i>Biomedicines</i> , 2022, 10, 691.	3.2	8
21	Study protocol on the safety and feasibility of a normocaloric ketogenic diet in people with amyotrophic lateral sclerosis. <i>Nutrition</i> , 2022, 94, 111525.	2.4	7
22	C9ORF72 Repeat Expansion Affects the Proteome of Primary Skin Fibroblasts in ALS. <i>International Journal of Molecular Sciences</i> , 2021, 22, 10385.	4.1	6
23	Ptosis and bulbar onset: an unusual phenotype of familial ALS?. <i>Neurological Sciences</i> , 2018, 39, 377-378.	1.9	5
24	The NEALS primary lateral sclerosis registry. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2020, 21, 74-81.	1.7	5
25	Detection of White Matter Ultrastructural Changes for Amyotrophic Lateral Sclerosis Characterization: A Diagnostic Study from Dti-Derived Data. <i>Brain Sciences</i> , 2020, 10, 996.	2.3	5
26	Characterization of the p.L145F and p.S135N Mutations in SOD1: Impact on the Metabolism of Fibroblasts Derived from Amyotrophic Lateral Sclerosis Patients. <i>Antioxidants</i> , 2022, 11, 815.	5.1	3
27	Generation of an induced pluripotent stem cell line, CSSi011-A (6534), from an Amyotrophic lateral sclerosis patient with heterozygous L145F mutation in SOD1 gene. <i>Stem Cell Research</i> , 2020, 47, 101924.	0.7	2
28	Headache in immigrant patients: similarities and differences with Italian population. <i>Neurological Sciences</i> , 2018, 39, 749-752.	1.9	2
29	Neurological emergency at the COVID-19 pandemic: report from a referral hospital in Eastern Piedmont, Italy. <i>Neurological Sciences</i> , 2022, 43, 2195.	1.9	2
30	Marchiafava-Bignami Disease with frontal cortex involvement and late onset, long-lasting psychiatric symptoms: a case report. <i>Rivista Di Psichiatria</i> , 2016, 51, 79-82.	0.6	2
31	Exploring the use of educational materials for increasing participation in a stretching program: a quality improvement project in people with motor neuron disease. <i>European Journal of Physical and Rehabilitation Medicine</i> , 2021, 57, 78-84.	2.2	1
32	Reply to: Amyotrophic lateral sclerosis with depression, cognitive impairment, and mortality. <i>Acta Neurologica Scandinavica</i> , 2020, 142, 86-87.	2.1	0
33	Do ecological factors influence the clinical presentation of amyotrophic lateral sclerosis?. <i>Journal of the Neurological Sciences</i> , 2021, 429, 117704.	0.6	0