

Martina de Majo

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9347785/publications.pdf>

Version: 2024-02-01

7
papers

351
citations

1478505

6
h-index

1720034

7
g-index

7
all docs

7
docs citations

7
times ranked

746
citing authors

#	ARTICLE	IF	CITATIONS
1	Tropism of SARS-CoV-2 for human cortical astrocytes. Proceedings of the National Academy of Sciences of the United States of America, 2022, 119, .	7.1	77
2	A recessive S174X mutation in Optineurin causes amyotrophic lateral sclerosis through a loss of function via allele-specific nonsense-mediated decay. Neurobiology of Aging, 2021, 106, 1-6.	3.1	3
3	An update on human astrocytes and their role in development and disease. Glia, 2020, 68, 685-704.	4.9	46
4	ALS-associated missense and nonsense TBK1 mutations can both cause loss of kinase function. Neurobiology of Aging, 2018, 71, 266.e1-266.e10.	3.1	59
5	Mutations in the vesicular trafficking protein annexin A11 are associated with amyotrophic lateral sclerosis. Science Translational Medicine, 2017, 9, .	12.4	129
6	C9ORF72 and UBQLN2 mutations are causes of amyotrophic lateral sclerosis in New Zealand: a genetic and pathologic study using banked human brain tissue. Neurobiology of Aging, 2017, 49, 214.e1-214.e5.	3.1	18
7	The CHCHD10 P34S variant is not associated with ALS in a UK cohort of familial and sporadic patients. Neurobiology of Aging, 2015, 36, 2908.e17-2908.e18.	3.1	19