

Maria Elena Erro

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7040623/publications.pdf>

Version: 2024-02-01

14
papers

986
citations

759233

12
h-index

794594

19
g-index

23
all docs

23
docs citations

23
times ranked

1454
citing authors

#	ARTICLE	IF	CITATIONS
1	Frequency, symptoms, risk factors, and outcomes of autoimmune encephalitis after herpes simplex encephalitis: a prospective observational study and retrospective analysis. <i>Lancet Neurology</i> , The, 2018, 17, 760-772.	10.2	422
2	Autoimmune post-herpes simplex encephalitis of adults and teenagers. <i>Neurology</i> , 2015, 85, 1736-1743.	1.1	226
3	Isolated pontine infarcts: etiopathogenic mechanisms. <i>European Journal of Neurology</i> , 2005, 12, 984-988.	3.3	50
4	Myasthenia gravis: Sleep quality, quality of life, and disease severity. <i>Muscle and Nerve</i> , 2012, 46, 174-180.	2.2	50
5	Frequency and Characterization of Movement Disorders in Anti-IgLON5 Disease. <i>Neurology</i> , 2021, 97, .	1.1	50
6	Anti-IgLON5 disease. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2020, 7, .	6.0	43
7	CRTC1 gene is differentially methylated in the human hippocampus in Alzheimer's disease. <i>Alzheimer's Research and Therapy</i> , 2016, 8, 15.	6.2	28
8	Striatal input from the ventrobasal complex of the rat thalamus. <i>Histochemistry and Cell Biology</i> , 2001, 115, 447-454.	1.7	23
9	Globular glial tauopathy caused by MAPT P301T mutation: clinical and neuropathological findings. <i>Journal of Neurology</i> , 2019, 266, 2396-2405.	3.6	22
10	PLD3 epigenetic changes in the hippocampus of Alzheimer's disease. <i>Clinical Epigenetics</i> , 2018, 10, 116.	4.1	21
11	Parkinson's Disease Case Ascertainment in the EPIC Cohort: The NeuroEPIC4PD Study. <i>Neurodegenerative Diseases</i> , 2015, 15, 331-338.	1.4	16
12	Acute carotid obliteration: a new vascular manifestation in POEMS syndrome. <i>European Journal of Neurology</i> , 2003, 10, 383-384.	3.3	12
13	Reversible Parkinsonism after accidental oral intake of mancozeb. <i>Movement Disorders</i> , 2011, 26, 557-558.	3.9	4
14	Progressive cognitive impairment and familial spastic paraparesis due to PRESENILIN 1 mutation: anatomoclinical characterization. <i>Journal of Neurology</i> , 2022, , 1.	3.6	1