

Emanuela Dazzo

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1856835/publications.pdf>

Version: 2024-02-01

7
papers

211
citations

1478505

6
h-index

1720034

7
g-index

7
all docs

7
docs citations

7
times ranked

437
citing authors

#	ARTICLE	IF	CITATIONS
1	Heterozygous Reelin Mutations Cause Autosomal-Dominant Lateral Temporal Epilepsy. American Journal of Human Genetics, 2015, 96, 992-1000.	6.2	94
2	The novel S59P mutation in the TNFRSF1A gene identified in an adult onset TNF receptor associated periodic syndrome (TRAPS) constitutively activates NF- κ B pathway. Arthritis Research and Therapy, 2015, 17, 93.	3.5	43
3	The clinical phenotype of autosomal dominant lateral temporal lobe epilepsy related to reelin mutations. Epilepsy and Behavior, 2017, 68, 103-107.	1.7	31
4	Mutations in <i>MICAL4</i> cause autosomal dominant lateral temporal epilepsy. Annals of Neurology, 2018, 83, 483-493.	5.3	25
5	Autosomal dominant lateral temporal lobe epilepsy associated with a novel reelin mutation. Epileptic Disorders, 2020, 22, 443-448.	1.3	8
6	Epilepsy-causing Reelin mutations result in impaired secretion and intracellular degradation of mutant proteins. Human Molecular Genetics, 2022, 31, 665-673.	2.9	7
7	LGI1 tumor tissue expression and serum autoantibodies in patients with primary malignant glioma. Clinical Neurology and Neurosurgery, 2018, 170, 27-33.	1.4	3