

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 | Epilepsy-causing Reelin mutations result in impaired secretion and intracellular degradation of mutant proteins. <i>Human Molecular Genetics</i> , 2022, 31, 665-673. | 2.9 | 7 |
| 2 | Autosomal dominant lateral temporal lobe epilepsy associated with a novel reelin mutation. <i>Epileptic Disorders</i> , 2020, 22, 443-448. | 1.3 | 8 |
| 3 | LGI1 tumor tissue expression and serum autoantibodies in patients with primary malignant glioma. <i>Clinical Neurology and Neurosurgery</i> , 2018, 170, 27-33. | 1.4 | 3 |
| 4 | Mutations in <i>MICAL1</i> cause autosomal dominant lateral temporal epilepsy. <i>Annals of Neurology</i> , 2018, 83, 483-493. | 5.3 | 25 |
| 5 | The clinical phenotype of autosomal dominant lateral temporal lobe epilepsy related to reelin mutations. <i>Epilepsy and Behavior</i> , 2017, 68, 103-107. | 1.7 | 31 |
| 6 | Heterozygous Reelin Mutations Cause Autosomal-Dominant Lateral Temporal Epilepsy. <i>American Journal of Human Genetics</i> , 2015, 96, 992-1000. | 6.2 | 94 |
| 7 | 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. <i>Arthritis Research and Therapy</i> , 2015, 17, 93. | 3.5 | 43 |