Rui-Juan Lv

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7146645/publications.pdf

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15	321	1040056	996975
papers	citations	h-index	g-index
16 all docs	16 docs citations	16 times ranked	552 citing authors

#	Article	IF	CITATIONS
1	Status epilepticus-related etiology, incidence and mortality: A meta-analysis. Epilepsy Research, 2017, 136, 12-17.	1.6	79
2	Circular RNA: a new star in neurological diseases. International Journal of Neuroscience, 2017, 127, 726-734.	1.6	50
3	ASIC1a polymorphism is associated with temporal lobe epilepsy. Epilepsy Research, 2011, 96, 74-80.	1.6	36
4	The role of the microRNA-146a/complement factor H/interleukin- $1\hat{l}^2$ -mediated inflammatory loop circuit in the perpetuate inflammation of chronic temporal lobe epilepsy. DMM Disease Models and Mechanisms, 2018, 11, .	2.4	31
5	Seizure semiology: an important clinical clue to the diagnosis of autoimmune epilepsy. Annals of Clinical and Translational Neurology, 2018, 5, 208-215.	3.7	25
6	Correlation between tumor necrosis factor alpha mRNA and microRNA-155 expression in rat models and patients with temporal lobe epilepsy. Brain Research, 2018, 1700, 56-65.	2.2	21
7	A polymorphism in CALHM1 is associated with temporal lobe epilepsy. Epilepsy and Behavior, 2011, 20, 681-685.	1.7	16
8	Semi-quantitative FDG-PET Analysis Increases the Sensitivity Compared With Visual Analysis in the Diagnosis of Autoimmune Encephalitis. Frontiers in Neurology, 2019, 10, 576.	2.4	15
9	Clinical Characteristics and Long-Term Prognosis of Anti-LGI1 Encephalitis: A Single-Center Cohort Study in Beijing, China. Frontiers in Neurology, 2021, 12, 674368.	2.4	11
10	Intravenous methylprednisolone or immunoglobulin for anti-glutamic acid decarboxylase 65 antibody autoimmune encephalitis: which is better?. BMC Neuroscience, 2020, 21, 13.	1.9	10
11	Significance of MDR1 gene C3435T polymorphism in predicting childhood refractory epilepsy. Epilepsy Research, 2017, 132, 21-28.	1.6	8
12	A Chinese Family With Adult-Onset Leigh-Like Syndrome Caused by the Heteroplasmic m.10191T>C Mutation in the Mitochondrial MTND3 Gene. Frontiers in Neurology, 2019, 10, 347.	2.4	8
13	Recognition of seizure semiology and semiquantitative FDGâ€PET analysis of antiâ€LGI1 encephalitis. CNS Neuroscience and Therapeutics, 2021, 27, 1173-1181.	3.9	7
14	Clinical and genetic characteristics of type I sialidosis patients in mainland China. Annals of Clinical and Translational Neurology, 2020, 7, 911-923.	3.7	2
15	Seizure as the unique clinical manifestation of cerebral metastases in a 27-year-old man with non-small cell lung cancer. Neurological Sciences, 2018, 39, 805-807.	1.9	0