

Lata Vadlamudi

List of Publications by Year in descending order

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Version: 2024-02-01

26
papers

1,410
citations

471509

17
h-index

580821

25
g-index

27
all docs

27
docs citations

27
times ranked

2053
citing authors

#	ARTICLE	IF	CITATIONS
1	Human induced pluripotent stem cells generated from epilepsy patients for use as in vitro models for drug screening. <i>Stem Cell Research</i> , 2022, 60, 102673.	0.7	0
2	Developing a gene panel for pharmaco-resistant epilepsy: a review of epilepsy pharmacogenetics. <i>Pharmacogenomics</i> , 2021, 22, 225-234.	1.3	12
3	Generation of induced pluripotent stem cell lines from peripheral blood mononuclear cells of three drug resistant and three drug responsive epilepsy patients. <i>Stem Cell Research</i> , 2021, 56, 102564.	0.7	0
4	Evidence for type-specific DNA methylation patterns in epilepsy: a discordant monozygotic twin approach. <i>Epigenomics</i> , 2019, 11, 951-968.	2.1	19
5	Deciphering the role of epigenetics in self-limited epilepsy with centrotemporal spikes. <i>Epilepsy Research</i> , 2019, 156, 106163.	1.6	5
6	Genetic epilepsy with febrile seizures plus. <i>Neurology</i> , 2017, 89, 1210-1219.	1.1	112
7	Mutations in mammalian target of rapamycin regulator <i>DEPDC5</i> cause focal epilepsy with brain malformations. <i>Annals of Neurology</i> , 2014, 75, 782-787.	5.3	193
8	Genetics of epilepsy. <i>Neurology</i> , 2014, 83, 1042-1048.	1.1	61
9	Genetics of febrile seizure subtypes and syndromes: A twin study. <i>Epilepsy Research</i> , 2013, 105, 103-109.	1.6	36
10	Timing of De Novo Mutagenesis – A Twin Study of Sodium-Channel Mutations. <i>New England Journal of Medicine</i> , 2010, 363, 1335-1340.	27.0	100
11	Array-Based Gene Discovery with Three Unrelated Subjects Shows SCARB2/LIMP-2 Deficiency Causes Myoclonus Epilepsy and Glomerulosclerosis. <i>American Journal of Human Genetics</i> , 2008, 82, 673-684.	6.2	230
12	Gene expression analysis in absence epilepsy using a monozygotic twin design. <i>Epilepsia</i> , 2008, 49, 1546-1554.	5.1	24
13	Obstetric Events as a Risk Factor for Febrile Seizures: A Community-Based Twin Study. <i>Twin Research and Human Genetics</i> , 2008, 11, 634-640.	0.6	2
14	Replicated effects of sex and genotype on gene expression in human lymphoblastoid cell lines. <i>Human Molecular Genetics</i> , 2007, 16, 364-373.	2.9	25
15	Volumetric analysis of a specific language region – the planum temporale. <i>Journal of Clinical Neuroscience</i> , 2006, 13, 206-213.	1.5	11
16	Analyzing the Etiology of Benign Rolandic Epilepsy: A Multicenter Twin Collaboration. <i>Epilepsia</i> , 2006, 47, 550-555.	5.1	135
17	Severe myoclonic epilepsy of infancy (Dravet syndrome): Recognition and diagnosis in adults. <i>Neurology</i> , 2006, 67, 2224-2226.	1.1	153
18	Action myoclonus-renal failure syndrome: A cause for worsening tremor in young adults. <i>Neurology</i> , 2006, 67, 1310-1311.	1.1	13

#	ARTICLE	IF	CITATIONS
19	Epilepsy in twins. <i>Neurology</i> , 2004, 62, 1127-1133.	1.1	50
20	Is benign rolandic epilepsy genetically determined?. <i>Annals of Neurology</i> , 2004, 56, 129-132.	5.3	52
21	Factors underlying scalp-EEG interictal epileptiform discharges in intractable frontal lobe epilepsy. <i>Epileptic Disorders</i> , 2004, 6, 89-95.	1.3	22
22	Electroencephalographic findings in Kufs disease. <i>Clinical Neurophysiology</i> , 2003, 114, 1738-1743.	1.5	11
23	Genetics of temporal lobe epilepsy. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2003, 74, 1359-1361.	1.9	48
24	Multifocal myoclonus due to verapamil overdose. <i>Neurology</i> , 2002, 58, 984-984.	1.1	40
25	Cerebral arterial gas embolism by helium: An unusual case successfully treated with hyperbaric oxygen and lidocaine. <i>Annals of Emergency Medicine</i> , 2000, 35, 300-303.	0.6	27
26	Rasmussen's syndrome in a 54 year old female: more support for an adult variant. <i>Journal of Clinical Neuroscience</i> , 2000, 7, 154-156.	1.5	28