

Chao Gao

List of Publications by Year in descending order

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Version: 2024-02-01

13
papers

227
citations

1307594

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1058476

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all docs

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docs citations

21
times ranked

402
citing authors

#	ARTICLE	IF	CITATIONS
1	Mutations disrupting neuritogenesis genes confer risk for cerebral palsy. <i>Nature Genetics</i> , 2020, 52, 1046-1056.	21.4	96
2	Diagnostic Yields of Trio-WES Accompanied by CNVseq for Rare Neurodevelopmental Disorders. <i>Frontiers in Genetics</i> , 2019, 10, 485.	2.3	28
3	Variants of the OLIG2 Gene are Associated with Cerebral Palsy in Chinese Han Infants with Hypoxic-Ischemic Encephalopathy. <i>NeuroMolecular Medicine</i> , 2019, 21, 75-84.	3.4	24
4	Lithium Treatment Is Safe in Children With Intellectual Disability. <i>Frontiers in Molecular Neuroscience</i> , 2018, 11, 425.	2.9	18
5	Autophagy-Related Gene 7 Polymorphisms and Cerebral Palsy in Chinese Infants. <i>Frontiers in Cellular Neuroscience</i> , 2019, 13, 494.	3.7	14
6	Genome-Wide Alteration of 5-Hydroxymethylcytosine in Hypoxic-Ischemic Neonatal Rat Model of Cerebral Palsy. <i>Frontiers in Molecular Neuroscience</i> , 2019, 12, 214.	2.9	8
7	Tuina Massage Improves Cognitive Functions of Hypoxic-Ischemic Neonatal Rats by Regulating Genome-Wide DNA Hydroxymethylation Levels. <i>Evidence-based Complementary and Alternative Medicine</i> , 2019, 2019, 1-11.	1.2	7
8	KecNet: A Light Neural Network for Arrhythmia Classification Based on Knowledge Reinforcement. <i>Journal of Healthcare Engineering</i> , 2021, 2021, 1-10.	1.9	7
9	The GluN2B-Trp373 NMDA Receptor Variant is Associated with Autism-, Epilepsy-Related Phenotypes and Reduces NMDA Receptor Currents in Rats. <i>Neurochemical Research</i> , 2022, 47, 1588-1597.	3.3	7
10	Neuronal Nitric Oxide Synthase Knockdown Within Basolateral Amygdala Induces Autistic-Related Phenotypes and Decreases Excitatory Synaptic Transmission in Mice. <i>Frontiers in Neuroscience</i> , 2020, 14, 886.	2.8	4
11	Overexpression of mGluR7 in the Prefrontal Cortex Attenuates Autistic Behaviors in Mice. <i>Frontiers in Cellular Neuroscience</i> , 2021, 15, 689611.	3.7	4
12	The Association Study of IL-23R Polymorphisms With Cerebral Palsy in Chinese Population. <i>Frontiers in Neuroscience</i> , 2020, 14, 590098.	2.8	3
13	TEP1 is a risk gene for sporadic cerebral palsy. <i>Journal of Genetics and Genomics</i> , 2021, 48, 1134-1134.	3.9	1