

# Yujiao Fu

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1604337/publications.pdf>

Version: 2024-02-01

10  
papers

117  
citations

1684188

5  
h-index

1474206

9  
g-index

10  
all docs

10  
docs citations

10  
times ranked

120  
citing authors

#	ARTICLE	IF	CITATIONS
1	Clinical features of patients with game-induced seizures in the Chinese population. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2016, 41, 51-55.	2.0	56
2	X-box binding protein I splicing attenuates brain microvascular endothelial cell damage induced by oxygen-glucose deprivation through the activation of phosphoinositide 3-kinase/protein kinase B, extracellular signal-regulated kinases, and hypoxia-inducible factor-1 $\alpha$ /vascular endothelial growth factor signaling pathways. <i>Journal of Cellular Physiology</i> , 2019, 234, 9316-9327.	4.1	16
3	Construction and analysis of a dysregulated lncRNA-associated ceRNA network in a rat model of temporal lobe epilepsy. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2019, 69, 105-114.	2.0	15
4	CpG methylation signature defines human temporal lobe epilepsy and predicts drug-resistant. <i>CNS Neuroscience and Therapeutics</i> , 2020, 26, 1021-1030.	3.9	10
5	&lt;p&gt;Effects of AQP4 and KCNJ10 Gene Polymorphisms on Drug Resistance and Seizure Susceptibility in Chinese Han Patients with Focal Epilepsy&lt;/p&gt;. <i>Neuropsychiatric Disease and Treatment</i> , 2020, Volume 16, 119-129.	2.2	7
6	&lt;p&gt;Impaired Cognitive Abilities in Siblings of Patients with Temporal Lobe Epilepsy&lt;/p&gt;. <i>Neuropsychiatric Disease and Treatment</i> , 2020, Volume 16, 3071-3079.	2.2	5
7	Dynamic Change of Shanks Gene mRNA Expression and DNA Methylation in Epileptic Rat Model and Human Patients. <i>Molecular Neurobiology</i> , 2020, 57, 3712-3726.	4.0	4
8	Intraspinal <i>Sparganum mansoni</i> infection with the extraction of a live adult worm. <i>Neurology: Clinical Practice</i> , 2019, 9, 472-474.	1.6	2
9	A novel compound heterozygous EPM2A mutation in a Chinese boy with Lafora disease. <i>Neurological Sciences</i> , 2020, 41, 2267-2270.	1.9	2
10	A case of organotin toxic encephalopathy with atypical imaging characteristic. <i>Neurological Sciences</i> , 2021, 42, 2579-2581.	1.9	0