

Shinsuke Ishigaki

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6967312/publications.pdf>

Version: 2024-02-01

10
papers

375
citations

932766

10
h-index

1372195

10
g-index

11
all docs

11
docs citations

11
times ranked

600
citing authors

#	ARTICLE	IF	CITATIONS
1	FUS regulates AMPA receptor function and FTL/ALS-associated behaviour via GluA1 mRNA stabilization. <i>Nature Communications</i> , 2015, 6, 7098.	5.8	129
2	Altered Tau Isoform Ratio Caused by Loss of FUS and SFPQ Function Leads to FTL-like Phenotypes. <i>Cell Reports</i> , 2017, 18, 1118-1131.	2.9	83
3	Lower Motor Neuron Involvement in TAR DNA-Binding Protein of 43 kDa-Related Frontotemporal Lobar Degeneration and Amyotrophic Lateral Sclerosis. <i>JAMA Neurology</i> , 2014, 71, 172.	4.5	33
4	Characteristic Features of FUS Inclusions in Spinal Motor Neurons of Sporadic Amyotrophic Lateral Sclerosis. <i>Journal of Neuropathology and Experimental Neurology</i> , 2020, 79, 370-377.	0.9	32
5	Aberrant interaction between FUS and SFPQ in neurons in a wide range of FTL spectrum diseases. <i>Brain</i> , 2020, 143, 2398-2405.	3.7	23
6	Marked Involvement of the Striatal Efferent System in TAR DNA-Binding Protein 43 kDa-Related Frontotemporal Lobar Degeneration and Amyotrophic Lateral Sclerosis. <i>Journal of Neuropathology and Experimental Neurology</i> , 2016, 75, 801-811.	0.9	17
7	Pathway from TDP-43-Related Pathology to Neuronal Dysfunction in Amyotrophic Lateral Sclerosis and Frontotemporal Lobar Degeneration. <i>International Journal of Molecular Sciences</i> , 2021, 22, 3843.	1.8	16
8	Motor neuron TDP-43 proteinopathy in progressive supranuclear palsy and corticobasal degeneration. <i>Brain</i> , 2022, 145, 2769-2784.	3.7	15
9	Actin-binding protein filamin-A drives tau aggregation and contributes to progressive supranuclear palsy pathology. <i>Science Advances</i> , 2022, 8, .	4.7	15
10	Pathologic Involvement of Glutamatergic Striatal Inputs From the Cortices in TAR DNA-Binding Protein 43 kDa-Related Frontotemporal Lobar Degeneration and Amyotrophic Lateral Sclerosis. <i>Journal of Neuropathology and Experimental Neurology</i> , 2017, 76, 759-768.	0.9	12