Luke McAlary

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3240259/publications.pdf

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23	896	17 h-index	23
papers	citations		g-index
33	33	33	1370 citing authors
all docs	docs citations	times ranked	

#	Article	IF	CITATIONS
1	CuATSM improves motor function and extends survival but is not tolerated at a high dose in SOD1G93A mice with a C57BL/6 background. Scientific Reports, 2021, 11, 19392.	1.6	9
2	The microglial NLRP3 inflammasome is activated by amyotrophic lateral sclerosis proteins. Glia, 2020, 68, 407-421.	2.5	133
3	Trajectory Taken by Dimeric Cu/Zn Superoxide Dismutase through the Protein Unfolding and Dissociation Landscape Is Modulated by Salt Bridge Formation. Analytical Chemistry, 2020, 92, 1702-1711.	3.2	9
4	The prion-like nature of amyotrophic lateral sclerosis. Progress in Molecular Biology and Translational Science, 2020, 175, 261-296.	0.9	14
5	Neurodegenerative disease-associated protein aggregates are poor inducers of the heat shock response in neuronal cells. Journal of Cell Science, 2020, 133, .	1.2	6
6	Ubiquitin Homeostasis Is Disrupted in TDP-43 and FUS Cell Models of ALS. IScience, 2020, 23, 101700.	1.9	28
7	Amyotrophic Lateral Sclerosis: Proteins, Proteostasis, Prions, and Promises. Frontiers in Cellular Neuroscience, 2020, 14, 581907.	1.8	25
8	Cerebrovascular amyloid Angiopathy in bioengineered vessels is reduced by high-density lipoprotein particles enriched in Apolipoprotein E. Molecular Neurodegeneration, 2020, 15, 23.	4.4	19
9	Tryptophan residue 32 in human Cu-Zn superoxide dismutase modulates prion-like propagation and strain selection. PLoS ONE, 2020, 15, e0227655.	1.1	22
10	Proteome Homeostasis Dysfunction: A Unifying Principle in ALS Pathogenesis. Trends in Neurosciences, 2020, 43, 274-284.	4.2	47
11	Thulium oxide nanoparticles as radioenhancers for the treatment of metastatic cutaneous squamous cell carcinoma. Physics in Medicine and Biology, 2020, 65, 215018.	1.6	6
12	Emerging Developments in Targeting Proteotoxicity in Neurodegenerative Diseases. CNS Drugs, 2019, 33, 883-904.	2.7	23
13	Prion-Like Propagation of Protein Misfolding and Aggregation in Amyotrophic Lateral Sclerosis. Frontiers in Molecular Neuroscience, 2019, 12, 262.	1.4	101
14	CuATSM Protects Against the <i>In Vitro</i> Cytotoxicity of Wild-Type-Like Copper–Zinc Superoxide Dismutase Mutants but not Mutants That Disrupt Metal Binding. ACS Chemical Neuroscience, 2019, 10, 1555-1564.	1.7	21
15	CNS-derived extracellular vesicles from superoxide dismutase 1 (SOD1)G93A ALS mice originate from astrocytes and neurons and carry misfolded SOD1. Journal of Biological Chemistry, 2019, 294, 3744-3759.	1.6	97
16	Strategies to promote the maturation of ALS-associated SOD1 mutants: small molecules return to the fold. Neural Regeneration Research, 2019, 14, 1511.	1.6	9
17	The cysteine-reactive small molecule ebselen facilitates effective SOD1 maturation. Nature Communications, 2018, 9, 1693.	5.8	71
18	TryptophanÂ32-mediated SOD1 aggregation is attenuated by pyrimidine-like compounds in living cells. Scientific Reports, 2018, 8, 15590.	1.6	32

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19	SOD1A4V aggregation alters ubiquitin homeostasis in a cell model of ALS. Journal of Cell Science, 2018, 131, .	1.2	39
20	Flow cytometric measurement of the cellular propagation of TDP-43 aggregation. Prion, 2017, 11, 195-204.	0.9	32
21	Susceptibility of Mutant SOD1 to Form a Destabilized Monomer Predicts Cellular Aggregation and Toxicity but Not In vitro Aggregation Propensity. Frontiers in Neuroscience, 2016, 10, 499.	1.4	75
22	Rapid flow cytometric measurement of protein inclusions and nuclear trafficking. Scientific Reports, 2016, 6, 31138.	1.6	31
23	Glutathionylation potentiates benign superoxide dismutase 1 variants to the toxic forms associated with amyotrophic lateral sclerosis. Scientific Reports, 2013, 3, 3275.	1.6	41