

Laura Pasetto

List of Publications by Year in descending order

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Version: 2024-02-01

17
papers

599
citations

840119

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h-index

887659

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docs citations

19
times ranked

1093
citing authors

#	ARTICLE	IF	CITATIONS
1	Mutant Copper-Zinc Superoxide Dismutase (SOD1) Induces Protein Secretion Pathway Alterations and Exosome Release in Astrocytes. <i>Journal of Biological Chemistry</i> , 2013, 288, 15699-15711.	1.6	216
2	New Insights on the Mechanisms of Disease Course Variability in ALS from Mutant SOD1 Mouse Models. <i>Brain Pathology</i> , 2016, 26, 237-247.	2.1	56
3	Role of Extracellular Vesicles in Amyotrophic Lateral Sclerosis. <i>Frontiers in Neuroscience</i> , 2018, 12, 574.	1.4	47
4	Targeting Extracellular Cyclophilin A Reduces Neuroinflammation and Extends Survival in a Mouse Model of Amyotrophic Lateral Sclerosis. <i>Journal of Neuroscience</i> , 2017, 37, 1413-1427.	1.7	42
5	Peptidylprolyl isomerase A governs TARDBP function and assembly in heterogeneous nuclear ribonucleoprotein complexes. <i>Brain</i> , 2015, 138, 974-991.	3.7	40
6	Lack of TNF α receptor type 2 protects motor neurons in a cellular model of amyotrophic lateral sclerosis and in mutant SOD1 mice but does not affect disease progression. <i>Journal of Neurochemistry</i> , 2015, 135, 109-124.	2.1	33
7	RNS60 exerts therapeutic effects in the SOD1 ALS mouse model through protective glia and peripheral nerve rescue. <i>Journal of Neuroinflammation</i> , 2018, 15, 65.	3.1	33
8	Decreased Levels of Foldase and Chaperone Proteins Are Associated with an Early-Onset Amyotrophic Lateral Sclerosis. <i>Frontiers in Molecular Neuroscience</i> , 2017, 10, 99.	1.4	30
9	Diagnostic and prognostic values of PBMC proteins in amyotrophic lateral sclerosis. <i>Neurobiology of Disease</i> , 2020, 139, 104815.	2.1	19
10	Decoding distinctive features of plasma extracellular vesicles in amyotrophic lateral sclerosis. <i>Molecular Neurodegeneration</i> , 2021, 16, 52.	4.4	19
11	Motor neuron degeneration, severe myopathy and TDP-43 increase in a transgenic pig model of SOD1-linked familial ALS. <i>Neurobiology of Disease</i> , 2019, 124, 263-275.	2.1	17
12	Micro-computed tomography for non-invasive evaluation of muscle atrophy in mouse models of disease. <i>PLoS ONE</i> , 2018, 13, e0198089.	1.1	13
13	Defective cyclophilin A induces TDP-43 proteinopathy: implications for amyotrophic lateral sclerosis and frontotemporal dementia. <i>Brain</i> , 2021, 144, 3710-3726.	3.7	13
14	A Novel HGF/SF Receptor (MET) Agonist Transiently Delays the Disease Progression in an Amyotrophic Lateral Sclerosis Mouse Model by Promoting Neuronal Survival and Dampening the Immune Dysregulation. <i>International Journal of Molecular Sciences</i> , 2020, 21, 8542.	1.8	8
15	Trabectedin and Lurbinectedin Extend Survival of Mice Bearing C26 Colon Adenocarcinoma, without Affecting Tumor Growth or Cachexia. <i>Cancers</i> , 2020, 12, 2312.	1.7	5
16	Contingent intramuscular boosting of P2XR7 axis improves motor function in transgenic ALS mice. <i>Cellular and Molecular Life Sciences</i> , 2022, 79, 7.	2.4	5
17	Cyclophilin A deficiency accelerates RML-induced prion disease. <i>Neurobiology of Disease</i> , 2019, 130, 104498.	2.1	2