

François Gros-Louis

List of Publications by Year in descending order

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Version: 2024-02-01

15
papers

1,295
citations

758635

12
h-index

996533

15
g-index

15
all docs

15
docs citations

15
times ranked

2825
citing authors

#	ARTICLE	IF	CITATIONS
1	Wild-type and mutant SOD1 share an aberrant conformation and a common pathogenic pathway in ALS. <i>Nature Neuroscience</i> , 2010, 13, 1396-1403.	7.1	600
2	Genetics of familial and sporadic amyotrophic lateral sclerosis. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2006, 1762, 956-972.	1.8	214
3	Intracerebroventricular infusion of monoclonal antibody or its derived Fab fragment against misfolded forms of SOD1 mutant delays mortality in a mouse model of ALS. <i>Journal of Neurochemistry</i> , 2010, 113, 1188-1199.	2.1	158
4	Misfolded SOD1 pathology in sporadic Amyotrophic Lateral Sclerosis. <i>Scientific Reports</i> , 2018, 8, 14223.	1.6	85
5	Chromogranin B P413L variant as risk factor and modifier of disease onset for amyotrophic lateral sclerosis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 21777-21782.	3.3	49
6	Early detection of structural abnormalities and cytoplasmic accumulation of TDP-43 in tissue-engineered skins derived from ALS patients. <i>Acta Neuropathologica Communications</i> , 2015, 3, 5.	2.4	47
7	Human Organ-Specific 3D Cancer Models Produced by the Stromal Self-Assembly Method of Tissue Engineering for the Study of Solid Tumors. <i>BioMed Research International</i> , 2020, 2020, 1-23.	0.9	28
8	Moyamoya Disease Susceptibility Gene <i><i>RNF213</i></i> Regulates Endothelial Barrier Function. <i>Stroke</i> , 2022, 53, 1263-1275.	1.0	26
9	Characterization of the phenotype with cognitive impairment and protein mislocalization in SCA34. <i>Neurology: Genetics</i> , 2020, 6, e403.	0.9	21
10	An Optimized Approach to Recover Secreted Proteins from Fibroblast Conditioned-Media for Secretomic Analysis. <i>Frontiers in Cellular Neuroscience</i> , 2016, 10, 70.	1.8	16
11	Potential skin involvement in ALS: revisiting Charcot's observation "a review of skin abnormalities in ALS. <i>Reviews in the Neurosciences</i> , 2017, 28, 551-572.	1.4	16
12	Sex-dependent effects of chromogranin B P413L allelic variant as disease modifier in amyotrophic lateral sclerosis. <i>Human Molecular Genetics</i> , 2016, 25, ddw304.	1.4	15
13	Cell Seeding on UV-treated 3D Polymeric Templates Allows for Cost-effective Production of Small-caliber Tissue-engineered Blood Vessels. <i>Biotechnology Journal</i> , 2019, 14, e1800306.	1.8	10
14	Biofabrication of a three dimensional human-based personalized neurofibroma model. <i>Biotechnology Journal</i> , 2021, 16, e2000250.	1.8	7
15	Diagnosis of idiopathic amyotrophic lateral sclerosis using Fourier-transform infrared spectroscopic analysis of patient-derived skin. <i>Analyst, The</i> , 2020, 145, 3678-3685.	1.7	3