FranÃ§ois Gros-Louis

List of Publications by Year
in descending order

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15	1,295	758635 12	996533 15
papers	citations	h-index	g-index

15	15	15	2825
all docs	docs citations	times ranked	citing authors

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