Françoise Piguet

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	AAVâ€delivered diacylglycerol kinase DGKk achieves longâ€ŧerm rescue of fragile X syndrome mouse model. EMBO Molecular Medicine, 2022, 14, e14649.	6.9	11
2	The Challenge of Gene Therapy for Neurological Diseases: Strategies and Tools to Achieve Efficient Delivery to the Central Nervous System. Human Gene Therapy, 2021, 32, 349-374.	2.7	21
3	Complete Correction of Brain and Spinal Cord Pathology in Metachromatic Leukodystrophy Mice. Frontiers in Molecular Neuroscience, 2021, 14, 677895.	2.9	10
4	Real-Time Monitoring of Exosome Enveloped-AAV Spreading by Endomicroscopy Approach: A New Tool for Gene Delivery in the Brain. Molecular Therapy - Methods and Clinical Development, 2019, 14, 237-251.	4.1	35
5	Rapid and Complete Reversal of Sensory Ataxia by Gene Therapy in a Novel Model of Friedreich Ataxia. Molecular Therapy, 2018, 26, 1940-1952.	8.2	86
6	Clinical Gene Therapy for Neurodegenerative Diseases: Past, Present, and Future. Human Gene Therapy, 2017, 28, 988-1003.	2.7	82
7	Niacinâ€mediated Tace activation ameliorates <scp>CMT</scp> neuropathies with focal hypermyelination. EMBO Molecular Medicine, 2016, 8, 1438-1454.	6.9	48
8	CYP46A1, the rate-limiting enzyme for cholesterol degradation, is neuroprotective in Huntington's disease. Brain, 2016, 139, 953-970.	7.6	135
9	Intracerebral Gene Therapy Using AAVrh.10-hARSA Recombinant Vector to Treat Patients with Early-Onset Forms of Metachromatic Leukodystrophy: Preclinical Feasibility and Safety Assessments in Nonhuman Primates. Human Gene Therapy Clinical Development, 2015, 26, 113-124.	3.1	68
10	Correction of Brain Oligodendrocytes by AAVrh.10 Intracerebral Gene Therapy in Metachromatic Leukodystrophy Mice. Human Gene Therapy, 2012, 23, 903-914.	2.7	73
11	Efficient intracerebral delivery of AAV5 vector encoding human ARSA in non-human primate. Human Molecular Cenetics, 2010, 19, 147-158	2.9	67