Vincent Picher-Martel

List of Publications by Year
in descending order

12 papers	365 citations	1163117 8 h-index	1199594 12 g-index
12 all docs	12 docs citations	12 times ranked	707 citing authors

#	Article	IF	Citations
1	From animal models to human disease: a genetic approach for personalized medicine in ALS. Acta Neuropathologica Communications, 2016, 4, 70.	5.2	115
2	Key role of UBQLN2 in pathogenesis of amyotrophic lateral sclerosis and frontotemporal dementia. Acta Neuropathologica Communications, 2019, 7, 103.	5.2	78
3	Ubiquilin-2 drives NF-Î¹B activity and cytosolic TDP-43 aggregation in neuronal cells. Molecular Brain, 2015, 8, 71.	2.6	54
4	Transmission of ALS pathogenesis by the cerebrospinal fluid. Acta Neuropathologica Communications, 2020, 8, 65.	5.2	30
5	Neuronal Expression of UBQLN2P497H Exacerbates TDP-43 Pathology in TDP-43G348C Mice through Interaction with Ubiquitin. Molecular Neurobiology, 2019, 56, 4680-4696.	4.0	23
6	The Occurrence of FUS Mutations in Pediatric Amyotrophic Lateral Sclerosis: A Case Report and Review of the Literature. Journal of Child Neurology, 2020, 35, 556-562.	1.4	23
7	Targeting TDP-43 Pathology Alleviates Cognitive and Motor Deficits Caused by Chronic Cerebral Hypoperfusion. Neurotherapeutics, 2021, 18, 1095-1112.	4.4	11
8	Current and Promising Therapies in Autosomal Recessive Ataxias. CNS and Neurological Disorders - Drug Targets, 2018, 17, 161-171.	1.4	9
9	Whole-exome sequencing identifies homozygous mutation in TTI2 in a child with primary microcephaly: a case report. BMC Neurology, 2020, 20, 58.	1.8	9
10	Case Report: Acute Necrotizing Encephalopathy Following COVID-19 Vaccine. Frontiers in Neurology, 2022, 13, 872734.	2.4	7
11	SMALED2 with BICD2 gene mutations: Report of two cases and portrayal of a classical phenotype. Neuromuscular Disorders, 2020, 30, 669-673.	0.6	3
12	CAPTURE ALS: the comprehensive analysis platform to understand, remedy and eliminate ALS. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2022, , 1-7.	1.7	3