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List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Altered cortical processing of sensory input in Huntington disease mouse models. Neurobiology of Disease, 2022, 169, 105740.	4.4	9
2	Regulation of hippocampal excitatory synapses by the Zdhhc5 palmitoyl acyltransferase. Journal of Cell Science, 2021, 134, .	2.0	13
3	Impaired Refinement of Kinematic Variability in Huntington Disease Mice on an Automated Home Cage Forelimb Motor Task. Journal of Neuroscience, 2021, 41, 8589-8602.	3.6	4
4	Endocannabinoid LTD in Accumbal D1 Neurons Mediates Reward-Seeking Behavior. IScience, 2020, 23, 100951.	4.1	27
5	Alterations in synaptic function and plasticity in Huntington disease. Journal of Neurochemistry, 2019, 150, 346-365.	3.9	90
6	Endocannabinoid-Specific Impairment in Synaptic Plasticity in Striatum of Huntington's Disease Mouse Model. Journal of Neuroscience, 2018, 38, 544-554.	3.6	28
7	Interacting Cannabinoid and Opioid Receptors in the Nucleus Accumbens Core Control Adolescent Social Play. Frontiers in Behavioral Neuroscience, 2016, 10, 211.	2.0	55
8	Influence of cortical synaptic input on striatal neuronal dendritic arborization and sensitivity to excitotoxicity in corticostriatal coculture. Journal of Neurophysiology, 2016, 116, 380-390.	1.8	7
9	An enhanced Q175 knock-in mouse model of Huntington disease with higher mutant huntingtin levels and accelerated disease phenotypes. Human Molecular Genetics, 2016, 25, 3654-3675.	2.9	85
10	Functional and structural deficits at accumbens synapses in a mouse model of Fragile X. Frontiers in Cellular Neuroscience, 2015, 9, 100.	3.7	42
11	Chronic blockade of extrasynaptic NMDA receptors ameliorates synaptic dysfunction and pro-death signaling in Huntington disease transgenic mice. Neurobiology of Disease, 2014, 62, 533-542.	4.4	74
12	Genetic rescue of CB1 receptors on medium spiny neurons prevents loss of excitatory striatal synapses but not motor impairment in HD mice. Neurobiology of Disease, 2014, 71, 140-150.	4.4	46
13	Mechanisms of synaptic dysfunction and excitotoxicity in Huntington's disease. Drug Discovery Today, 2014, 19, 990-996.	6.4	101
14	Uncoupling of the endocannabinoid signalling complex in a mouse model of fragile X syndrome. Nature Communications, 2012, 3, 1080.	12.8	234
15	Opposing Roles of Synaptic and Extrasynaptic NMDA Receptor Signaling in Cocultured Striatal and Cortical Neurons. Journal of Neuroscience, 2012, 32, 3992-4003.	3.6	121
16	Calpain and STriatal-Enriched protein tyrosine Phosphatase (STEP) activation contribute to extrasynaptic NMDA receptor localization in a Huntington's disease mouse model. Human Molecular Genetics, 2012, 21, 3739-3752.	2.9	75
17	Mitigation of augmented extrasynaptic NMDAR signaling and apoptosis in cortico-striatal co-cultures from Huntington's disease mice. Neurobiology of Disease, 2012, 48, 40-51.	4.4	74
18	Nutritional omega-3 deficiency abolishes endocannabinoid-mediated neuronal functions. Nature Neuroscience, 2011, 14, 345-350.	14.8	276