Maija L Castrén

List of Publications by Year in descending order

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ΜΑΠΑΙ CASTRÃON

#	Article	IF	CITATIONS
1	Generation of the Human Pluripotent Stem-Cell-Derived Astrocyte Model with Forebrain Identity. Brain Sciences, 2021, 11, 209.	2.3	10
2	Increased iron content in the heart of the Fmr1 knockout mouse. BioMetals, 2021, 34, 947-954.	4.1	5
3	Urokinase plasminogen activator mediates changes in human astrocytes modeling fragile X syndrome. Glia, 2021, 69, 2947-2962.	4.9	12
4	Elevated de novo protein synthesis in FMRP-deficient human neurons and its correction by metformin treatment. Molecular Autism, 2020, 11, 41.	4.9	23
5	Urine microRNA Profiling Displays miR-125a Dysregulation in Children with Fragile X Syndrome. Cells, 2020, 9, 289.	4.1	10
6	Integrative Analysis Identifies Key Molecular Signatures Underlying Neurodevelopmental Deficits in Fragile X Syndrome. Biological Psychiatry, 2020, 88, 500-511.	1.3	33
7	Modeling FXS with Mouse Neural Progenitors. Methods in Molecular Biology, 2019, 1942, 71-78.	0.9	Ο
8	Dysregulated Ca2+-Permeable AMPA Receptor Signaling in Neural Progenitors Modeling Fragile X Syndrome. Frontiers in Synaptic Neuroscience, 2019, 11, 2.	2.5	10
9	Astrocytes in Neuropathologies Affecting the Frontal Cortex. Frontiers in Cellular Neuroscience, 2019, 13, 44.	3.7	24
10	Functional changes of AMPA responses in human induced pluripotent stem cell–derived neural progenitors in fragile X syndrome. Science Signaling, 2018, 11, .	3.6	54
11	Increased Calcium Influx through L-type Calcium Channels in Human and Mouse Neural Progenitors Lacking Fragile X Mental Retardation Protein. Stem Cell Reports, 2018, 11, 1449-1461.	4.8	27
12	Metabotropic glutamate receptor 5 responses dictate differentiation of neural progenitors to NMDAâ€responsive cells in fragile X syndrome. Developmental Neurobiology, 2017, 77, 438-453.	3.0	38
13	Epileptic Electroencephalography Profile Associates with Attention Problems in Children with Fragile X Syndrome: Review and Case Series. Frontiers in Human Neuroscience, 2016, 10, 353.	2.0	13
14	Cortical neurogenesis in fragile X syndrome. Frontiers in Bioscience - Scholar, 2016, 8, 160-168.	2.1	17
15	Distinctive behavioral and cellular responses to fluoxetine in the mouse model for Fragile X syndrome. Frontiers in Cellular Neuroscience, 2014, 8, 150.	3.7	32
16	Tissue Plasminogen Activator Contributes to Alterations of Neuronal Migration and Activity-Dependent Responses in Fragile X Mice. Journal of Neuroscience, 2014, 34, 1916-1923.	3.6	22
17	BDNF in fragile X syndrome. Neuropharmacology, 2014, 76, 729-736.	4.1	60
18	Effect of glutamate receptor antagonists on migrating neural progenitor cells. European Journal of Neuroscience, 2013, 37, 1369-1382.	2.6	30

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19	Neural Stem Cells. Results and Problems in Cell Differentiation, 2012, 54, 33-40.	0.7	4
20	BDNF and TrkB in neuronal differentiation of Fmr1-knockout mouse. Neurobiology of Disease, 2011, 41, 469-480.	4.4	81
21	Aberrant differentiation of glutamatergic cells in neocortex of mouse model for fragile X syndrome. Neurobiology of Disease, 2009, 33, 250-259.	4.4	72
22	The Val66Met polymorphism in the BDNF gene is associated with epilepsy in fragile X syndrome. Epilepsy Research, 2009, 85, 114-117.	1.6	29
23	Neurotransmitter responsiveness during early maturation of neural progenitor cells. Differentiation, 2009, 77, 188-198.	1.9	27
24	Differentiation of Neuronal Cells in Fragile X Syndrome. Cell Cycle, 2006, 5, 1528-1530.	2.6	13
25	Altered differentiation of neural stem cells in fragile X syndrome. Proceedings of the National Academy of Sciences of the United States of America, 2005, 102, 17834-17839.	7.1	155
26	Rgs4 mRNA expression is decreased in the brain of Fmr1 knockout mouse. Molecular Brain Research, 2005, 133, 162-165.	2.3	11