

# ChloÃ© DelÃ©pine

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3284678/publications.pdf>

Version: 2024-02-01

9  
papers

190  
citations

1307366

7  
h-index

1474057

9  
g-index

11  
all docs

11  
docs citations

11  
times ranked

379  
citing authors

#	ARTICLE	IF	CITATIONS
1	Altered microtubule dynamics and vesicular transport in mouse and human MeCP2-deficient astrocytes. <i>Human Molecular Genetics</i> , 2016, 25, 146-157.	1.4	53
2	Astrocyte Transcriptome from the <i>Mecp2308-Truncated</i> Mouse Model of Rett Syndrome. <i>NeuroMolecular Medicine</i> , 2015, 17, 353-363.	1.8	31
3	Altered microtubule dynamics in <i>Mecp2</i> -deficient astrocytes. <i>Journal of Neuroscience Research</i> , 2012, 90, 990-998.	1.3	27
4	MeCP2 deficiency is associated with impaired microtubule stability. <i>FEBS Letters</i> , 2013, 587, 245-253.	1.3	23
5	Differential Expression and Regulation of Brain-Derived Neurotrophic Factor (BDNF) mRNA Isoforms in Brain Cells from <i>Mecp2308/y</i> Mouse Model. <i>Journal of Molecular Neuroscience</i> , 2015, 56, 758-767.	1.1	17
6	Mutations in the C-terminus of CDKL5: proceed with caution. <i>European Journal of Human Genetics</i> , 2014, 22, 270-272.	1.4	12
7	GSK3 $\gamma$ inhibitor CHIR 99021 modulates cerebral organoid development through dose-dependent regulation of apoptosis, proliferation, differentiation and migration. <i>PLoS ONE</i> , 2021, 16, e0251173.	1.1	12
8	HDAC inhibitor ameliorates behavioral deficits in <i>Mecp2308/y</i> mouse model of Rett syndrome. <i>Brain Research</i> , 2021, 1772, 147670.	1.1	8
9	Somatic mosaicism for a <i>FOXP1</i> mutation: diagnostic implication. <i>Clinical Genetics</i> , 2014, 85, 589-591.	1.0	7