Sabrina Oishi

List of Publications by Year in descending order

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Version: 2024-02-01

1478505 1588992 8 144 6 8 citations h-index g-index papers 9 9 9 319 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Partial Loss of USP9X Function Leads to a Male Neurodevelopmental and Behavioral Disorder Converging on Transforming Growth Factor \hat{l}^2 Signaling. Biological Psychiatry, 2020, 87, 100-112.	1.3	42
2	Investigating cortical features of Sotos syndrome using mice heterozygous for <i>Nsd1</i> . Genes, Brain and Behavior, 2020, 19, e12637.	2.2	16
3	Heterozygosity for Nuclear Factor One X in mice models features of Malan syndrome. EBioMedicine, 2019, 39, 388-400.	6.1	9
4	NFIX-Mediated Inhibition of Neuroblast Branching Regulates Migration Within the Adult Mouse Ventricular–Subventricular Zone. Cerebral Cortex, 2019, 29, 3590-3604.	2.9	10
5	Neurogenic differentiation by hippocampal neural stem and progenitor cells is biased by NFIX expression. Development (Cambridge), 2018, 145, .	2.5	29
6	A morphology independent approach for identifying dividing adult neural stem cells in the mouse hippocampus. Developmental Dynamics, 2018, 247, 194-200.	1.8	7
7	Usp9x-deficiency disrupts the morphological development of the postnatal hippocampal dentate gyrus. Scientific Reports, 2016, 6, 25783.	3.3	28
8	USP9X deletion elevates the density of oligodendrocytes within the postnatal dentate gyrus. Neurogenesis (Austin, Tex), 2016, 3, e1235524.	1.5	3