Constance Smith-Hicks

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/10650512/publications.pdf

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10 papers	250 citations	1478505 6 h-index	1372567 10 g-index
10 all docs	10 docs citations	10 times ranked	607 citing authors

#	Article	IF	Citations
1	SRF binding to SRE 6.9 in the Arc promoter is essential for LTD in cultured Purkinje cells. Nature Neuroscience, 2010, 13, 1082-1089.	14.8	72
2	De Novo Variants in MAPK8IP3 Cause Intellectual Disability with Variable Brain Anomalies. American Journal of Human Genetics, 2019, 104, 203-212.	6.2	44
3	Monogenic disorders that mimic the phenotype of Rett syndrome. Neurogenetics, 2018, 19, 41-47.	1.4	41
4	De Novo Missense Variants in FBXW11 Cause Diverse Developmental Phenotypes Including Brain, Eye, and Digit Anomalies. American Journal of Human Genetics, 2019, 105, 640-657.	6.2	31
5	Rett Syndrome and CDKL5 Deficiency Disorder: From Bench to Clinic. International Journal of Molecular Sciences, 2019, 20, 5098.	4.1	30
6	Bi-allelic variants in SPATA5L1 lead to intellectual disability, spastic-dystonic cerebral palsy, epilepsy, and hearing loss. American Journal of Human Genetics, 2021, 108, 2006-2016.	6.2	11
7	Heterozygous de novo variants in <scp>CSNK1G1</scp> are associated with syndromic developmental delay and autism spectrum disorder. Clinical Genetics, 2020, 98, 571-576.	2.0	10
8	Truncating variants in the SHANK1 gene are associated with a spectrum of neurodevelopmental disorders. Genetics in Medicine, 2021, 23, 1912-1921.	2.4	5
9	Delineating the genotypic and phenotypic spectrum of HECW2-related neurodevelopmental disorders. Journal of Medical Genetics, 2022, 59, 669-677.	3.2	5
10	mTORC1 functional assay revealsSZT2loss-of-function variants and a founder in-frame deletion. Brain, 2022, 145, 1939-1948.	7.6	1