Julian Curiel

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/11061960/publications.pdf

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7 papers	391 citations	1478505 6 h-index	1720034 7 g-index
8 all docs	8 docs citations	8 times ranked	916 citing authors

#	Article	IF	Citations
1	TUBB4A mutations result in both glial and neuronal degeneration in an H-ABC leukodystrophy mouse model. ELife, 2020, 9, .	6.0	15
2	Adulthood leukodystrophies. Nature Reviews Neurology, 2018, 14, 94-105.	10.1	119
3	Absence of Axoglial Paranodal Junctions in a Child With CNTNAP1 Mutations, Hypomyelination, and Arthrogryposis. Journal of Child Neurology, 2018, 33, 642-650.	1.4	11
4	Loss of CLOCK Results in Dysfunction of Brain Circuits Underlying Focal Epilepsy. Neuron, 2017, 96, 387-401.e6.	8.1	66
5	TUBB4A mutations result in specific neuronal and oligodendrocytic defects that closely match clinically distinct phenotypes. Human Molecular Genetics, 2017, 26, 4506-4518.	2.9	59
6	X-linked adrenoleukodystrophy in a chimpanzee due to an ABCD1 mutation reported in multiple unrelated humans. Molecular Genetics and Metabolism, 2017, 122, 130-133.	1.1	5
7	The emerging role of the tubulin code: From the tubulin molecule to neuronal function and disease. Cytoskeleton, 2016, 73, 521-550.	2.0	116