

# Julian Curiel

## List of Publications by Year in descending order

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

Version: 2024-02-01

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