Nathan D Kopp

List of Publications by Year
in descending order

8	154	1478280 6	1719901 7
papers	citations	h-index	g-index

8	8	8	327
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Human iPSC-Derived Neurons and Cerebral Organoids Establish Differential Effects of Germline NF1 Gene Mutations. Stem Cell Reports, 2020, 14, 541-550.	2.3	48
2	The anatomical distribution of genetic associations. Nucleic Acids Research, 2015, 43, 10804-10820.	6.5	37
3	Gtf2i and Gtf2ird1 mutation do not account for the full phenotypic effect of the Williams syndrome critical region in mouse models. Human Molecular Genetics, 2019, 28, 3443-3465.	1.4	23
4	Functions of Gtf2i and Gtf2ird1 in the developing brain: transcription, DNA binding and long-term behavioral consequences. Human Molecular Genetics, 2020, 29, 1498-1519.	1.4	18
5	Moving from capstones toward cornerstones: successes and challenges in applying systems biology to identify mechanisms of autism spectrum disorders. Frontiers in Genetics, 2015, 6, 301.	1.1	14
6	Exome sequencing of 85 Williamsâ€“Beuren syndrome cases rules out coding variation as a major contributor to remaining variance in social behavior. Molecular Genetics & Genomic Medicine, 2018, 6, 749-765.	0.6	9
7	Pathogenic paternally inherited NLGN4X deletion in a female with autism spectrum disorder: Clinical, cytogenetic, and molecular characterization. American Journal of Medical Genetics, Part A, 2021, 185, 894-900.	0.7	5
8	Expert Curation of Somatic FLT3 Variants By the Clingen Somatic Hematologic Cancer Taskforce (ClinGen HCT). Blood, 2021, 138, 4387-4387.	0.6	0