Nathan D Kopp

List of Publications by Citations

8 papers	84 citations	5 h-index	8 g-index
8 ext. papers	123 ext. citations	6.2 avg, IF	2.34 L-index

#	Paper	IF	Citations
8	The anatomical distribution of genetic associations. *Nucleic Acids Research*, **2015**, 43, 10804-20	20.1	26
7	Human iPSC-Derived Neurons and Cerebral Organoids Establish Differential Effects of Germline NF1 Gene Mutations. *Stem Cell Reports*, **2020**, 14, 541-550	8	21
6	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	4.5	12
5	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	5.6	11
4	Exome sequencing of 85 Williams-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	2.3	9
3	Functions of Gtf2i and Gtf2ird1 in the developing brain: transcription, DNA binding and long-term behavioral consequences. *Human Molecular Genetics*, **2020**, 29, 1498-1519	5.6	5
2	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	2.5	0
1	Expert Curation of Somatic FLT3 Variants By the Clingen Somatic Hematologic Cancer Taskforce (ClinGen HCT). *Blood*, **2021**, 138, 4387-4387	2.2	