

# Nathan D Kopp

## List of Publications by Citations

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**Version:** 2024-04-27

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

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. <i>Nucleic Acids Research</i> , <b>2015</b> , 43, 10804-20	20.1	26
7	Human iPSC-Derived Neurons and Cerebral Organoids Establish Differential Effects of Germline NF1 Gene Mutations. <i>Stem Cell Reports</i> , <b>2020</b> , 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. <i>Frontiers in Genetics</i> , <b>2015</b> , 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. <i>Human Molecular Genetics</i> , <b>2019</b> , 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. <i>Molecular Genetics &amp; Genomic Medicine</i> , <b>2018</b> , 6, 749-765	2.3	9
3	Functions of Gtf2i and Gtf2ird1 in the developing brain: transcription, DNA binding and long-term behavioral consequences. <i>Human Molecular Genetics</i> , <b>2020</b> , 29, 1498-1519	5.6	5
2	Pathogenic paternally inherited NLGN4X deletion in a female with autism spectrum disorder: Clinical, cytogenetic, and molecular characterization. <i>American Journal of Medical Genetics, Part A</i> , <b>2021</b> , 185, 894-900	2.5	0
1	Expert Curation of Somatic FLT3 Variants By the Clingen Somatic Hematologic Cancer Taskforce (ClinGen HCT). <i>Blood</i> , <b>2021</b> , 138, 4387-4387	2.2	