Yonit A Addissie

List of Publications by Year
in descending order

19	514	840776 11	940533 16
papers	citations	h-index	g-index

19	19	19	879
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	22q11.2 deletion syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2017, 173, 879-888.	1.2	103
2	Down syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2017, 173, 42-53.	1.2	75
3	Noonan syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2017, 173, 2323-2334.	1.2	68
4	Williamsâ€“Beuren syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2018, 176, 1128-1136.	1.2	55
5	Craniosynostosis and Noonan syndrome with KRAS mutations: Expanding the phenotype with a case report and review of the literature. American Journal of Medical Genetics, Part A, 2015, 167, 2657-2663.	1.2	38
6	Muenke syndrome: An international multicenter natural history study. American Journal of Medical Genetics, Part A, 2016, 170, 918-929.	1.2	37
7	Loss of function inROBO1is associated with tetralogy of Fallot and septal defects. Journal of Medical Genetics, 2017, 54, 825-829.	3.2	27
8	Prenatal exposure to pesticides and risk for holoprosencephaly: a case-control study. Environmental Health, 2020, 19, 65.	4.0	20
9	Rubinsteinâ€“Taybi syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2020, 182, 2939-2950.	1.2	16
10	Exome Sequencing and Congenital Heart Disease in Sub-Saharan Africa. Circulation Genomic and Precision Medicine, 2021, 14, e003108.	3.6	16
11	In-depth investigations of adolescents and adults with holoprosencephaly identify unique characteristics. Genetics in Medicine, 2018, 20, 14-23.	2.4	15
12	Turner syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2020, 182, 303-313.	1.2	15
13	Identifying environmental risk factors and <scp>geneâ€“environment</scp> interactions in holoprosencephaly. Birth Defects Research, 2021, 113, 63-76.	1.5	14
14	Executive Function and Adaptive Behavior in Muenke Syndrome. Journal of Pediatrics, 2015, 167, 428-434.	1.8	9
15	Tuberous sclerosis in a patient from Nigeria. American Journal of Medical Genetics, Part A, 2019, 179, 1423-1425.	1.2	4
16	Circle of Willis anomalies in Turner syndrome: Absent A1 segment of the anterior cerebral artery. Birth Defects Research, 2019, 111, 1584-1588.	1.5	2
17	Cover Image, Volume 173A, Number 4, April 2017. American Journal of Medical Genetics, Part A, 2017, 173, i.	1.2	0
18	Cover Image, Volume 173A, Number 9, September 2017. American Journal of Medical Genetics, Part A, 2017, 173, i.	1.2	0

#	Article	IF	Citations
19	Cover Image, Volume 176A, Number 5, May 2018. American Journal of Medical Genetics, Part A, 2018, 176, .	1.2	0