

Yonit A Addissie

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/883756/publications.pdf>

Version: 2024-02-01

19
papers

514
citations

840776

11
h-index

940533

16
g-index

19
all docs

19
docs citations

19
times ranked

879
citing authors

#	ARTICLE	IF	CITATIONS
1	Identifying environmental risk factors and <sc>gene"environment</sc> interactions in holoprosencephaly. Birth Defects Research, 2021, 113, 63-76.	1.5	14
2	Exome Sequencing and Congenital Heart Disease in Sub-Saharan Africa. Circulation Genomic and Precision Medicine, 2021, 14, e003108.	3.6	16
3	Turner syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2020, 182, 303-313.	1.2	15
4	Rubinstein"Taybi syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2020, 182, 2939-2950.	1.2	16
5	Prenatal exposure to pesticides and risk for holoprosencephaly: a case-control study. Environmental Health, 2020, 19, 65.	4.0	20
6	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
7	Tuberous sclerosis in a patient from Nigeria. American Journal of Medical Genetics, Part A, 2019, 179, 1423-1425.	1.2	4
8	Williams"Beuren syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2018, 176, 1128-1136.	1.2	55
9	In-depth investigations of adolescents and adults with holoprosencephaly identify unique characteristics. Genetics in Medicine, 2018, 20, 14-23.	2.4	15
10	Cover Image, Volume 176A, Number 5, May 2018. American Journal of Medical Genetics, Part A, 2018, 176, .	1.2	0
11	Loss of function in<i>ROBO1</i> is associated with tetralogy of Fallot and septal defects. Journal of Medical Genetics, 2017, 54, 825-829.	3.2	27
12	22q11.2 deletion syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2017, 173, 879-888.	1.2	103
13	Cover Image, Volume 173A, Number 4, April 2017. American Journal of Medical Genetics, Part A, 2017, 173, i.	1.2	0
14	Down syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2017, 173, 42-53.	1.2	75
15	Cover Image, Volume 173A, Number 9, September 2017. American Journal of Medical Genetics, Part A, 2017, 173, i.	1.2	0
16	Noonan syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2017, 173, 2323-2334.	1.2	68
17	Muenke syndrome: An international multicenter natural history study. American Journal of Medical Genetics, Part A, 2016, 170, 918-929.	1.2	37
18	Craniosynostosis and Noonan syndrome with <i>KRAS</i> 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

#	ARTICLE	IF	CITATIONS
19	Executive Function and Adaptive Behavior in Muenke Syndrome. Journal of Pediatrics, 2015, 167, 428-434.	1.8	9