Cedrik Tekendo-Ngongang

List of Publications by Year
in descending order

24 papers	389 citations	933447 10 h-index	839539 18 g-index
26 all docs	26 docs citations	26 times ranked	594 citing authors

#	Article	IF	Citations
1	Neural network classifiers for images of genetic conditions with cutaneous manifestations. Human Genetics and Genomics Advances, 2022, 3, 100053.	1.7	7
2	Neural Networks for Classification and Image Generation of Aging in Genetic Syndromes. Frontiers in Genetics, 2022, 13, 864092.	2.3	14
3	Exome Sequencing and Congenital Heart Disease in Sub-Saharan Africa. Circulation Genomic and Precision Medicine, 2021, 14, e003108.	3.6	16
4	Crossâ€cultural representations of conjoined twins. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2021, 187, 240-253.	1.6	5
5	Beyond Trinucleotide Repeat Expansion in Fragile X Syndrome: Rare Coding and Noncoding Variants in FMR1 and Associated Phenotypes. Genes, 2021, 12, 1669.	2.4	10
6	A Monoallelic Variant in REST Is Associated with Non-Syndromic Autosomal Dominant Hearing Impairment in a South African Family. Genes, 2021, 12, 1765.	2.4	5
7	Turner syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2020, 182, 303-313.	1.2	15
8	Rubinsteinâ€“Taybi syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2020, 182, 2939-2950.	1.2	16
9	Noonan syndrome on the African Continent. Birth Defects Research, 2020, 112, 718-724.	1.5	5
10	MECP2 duplication syndrome in a patient from Cameroon. American Journal of Medical Genetics, Part A, 2020, 182, 619-622.	1.2	4
11	Comorbidity of congenital heart defects and holoprosencephaly is likely genetically driven and geneâ€specific. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2020, 184, 154-158.	1.6	6
12	Novel heterozygous variants in KMT2D associated with holoprosencephaly. Clinical Genetics, 2019, 96, 266-270.	2.0	13
13	Noonan Syndrome in South Africa: Clinical and Molecular Profiles. Frontiers in Genetics, 2019, 10, 333.	2.3	13
14	Diversity and dysmorphology. Current Opinion in Pediatrics, 2019, 31, 702-707.	2.0	17
15	Cornelia de Lange syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2019, 179, 150-158.	1.2	40
16	Williamsâ€“Beuren syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2018, 176, 1128-1136.	1.2	55
17	Cover Image, Volume 176A, Number 5, May 2018. American Journal of Medical Genetics, Part A, 2018, 176, .	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	Noonan syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2017, 173, 2323-2334.	1.2	68
20	The 22q11.2 Deletion Syndrome in Congenital Heart Defects: Prevalence of Microdeletion Syndrome in Cameroon. Global Heart, 2017, 12, 115.	2.3	8
21	Challenges in Clinical Diagnosis of Williams-Beuren Syndrome in Sub-Saharan Africans: Case Reports from Cameroon. Molecular Syndromology, 2014, 5, 287-292.	0.8	20
22	Initiation of prenatal genetic diagnosis of chromosomal anomalies in Cameroon. International Journal of Gynecology and Obstetrics, 2012, 116, 174-175.	2.3	1
23	Initiation of a medical genetics service in sub-Saharan Africa: Experience of prenatal diagnosis in Cameroon. European Journal of Medical Genetics, 2011, 54, e399-e404.	1.3	40
24	Initiation of prenatal genetic diagnosis of sickle cell anaemia in Cameroon (subâ€Saharan Africa). Prenatal Diagnosis, 2011, 31, 1210-1212.	2.3	10