Andrew K Sobering

List of Publications by Year
in descending order

20 papers	201 citations	1683934 5 h-index	1125617 13 g-index
22 all docs	22 docs citations	22 times ranked	374 citing authors

#	Article	IF	Citations
1	Down syndrome in diverse populations. American Journal of Medical Genetics, Part A, 2017, 173, 42-53.	0.7	75
2	Expanding the genotypic and phenotypic spectrum in a diverse cohort of 104 individuals with Wiedemannâ€Steiner syndrome. American Journal of Medical Genetics, Part A, 2021, 185, 1649-1665.	0.7	34
3	Towards the optimal use of video recordings to support the flipped classroom in medical school basic sciences education. Medical Education Online, 2021, 26, 1841406.	1.1	28
4	Presentation and care of a family with Huntington disease in a resource-limited community. Journal of Clinical Movement Disorders, 2017, 4, 4.	2.2	10
5	Experiences with offering pro bono medical genetics services in the West Indies: Benefits to patients, physicians, and the community. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2020, 184, 1030-1041.	0.7	8
6	Genetic diagnosis of Down syndrome in an underserved community. American Journal of Medical Genetics, Part A, 2018, 176, 483-486.	0.7	6
7	Childhood-Onset Spinocerebellar Ataxia 3: Tongue Dystonia as an Early Manifestation. Tremor and Other Hyperkinetic Movements, 2019, 9, .	1.1	6
8	Quiz Discuss Compare: Using Audience Response Devices to Actively Engage Students. Medical Science Educator, 2015, 25, 299-302.	0.7	5
9	Variants in PHF8 cause a spectrum of X-linked neurodevelopmental disorders and facial dysmorphology. Human Genetics and Genomics Advances, 2022, 3, 100102.	1.0	5
10	Diagnosis of Spinocerebellar Ataxia in the West Indies. Tremor and Other Hyperkinetic Movements, 2018, 8, 567.	1.1	4
11	Identification of a de novo case of COL5A1â€related EhlersâĐanlos syndrome in an infant in the West Indies leading to improved targeted clinical care. Clinical Case Reports (discontinued), 2018, 6, 2256-2261.	0.2	3
12	A homozygous truncating NALCN variant in two Afroâ€Caribbean siblings with hypotonia and dolichocephaly. American Journal of Medical Genetics, Part A, 2020, 182, 1877-1880.	0.7	3
13	Diagnosis of Spinocerebellar Ataxia in the West Indies. Tremor and Other Hyperkinetic Movements, 2020, 8, 567.	1.1	3
14	Reevaluating Biochemistry and Nutrition Education Through Opinions of Clinicians and Educators. Medical Science Educator, 2019, 29, 189-197.	0.7	2
15	Application of exome sequencing to diagnose a novel presentation of the Cornelia de Lange syndrome in an Afroâ€Caribbean family. Molecular Genetics & Genomic Medicine, 2020, 8, e1318.	0.6	2
16	Childhood-Onset Spinocerebellar Ataxia 3: Tongue Dystonia as an Early Manifestation. Tremor and Other Hyperkinetic Movements, 2020, 9, .	1.1	2
17	Pseudo-ataxia due to Osteoid Osteoma. Tremor and Other Hyperkinetic Movements, 2020, 9, 631.	1.1	2
18	Sickle cell disease in Grenada: Quality of life and barriers to care. Molecular Genetics & Genomic Medicine, 2021, 9, e1567.	0.6	1

#	Article	IF	Citations
19	Pseudo-ataxia due to Osteoid Osteoma. Tremor and Other Hyperkinetic Movements, 2019, 9, 631.	1.1	1
20	A novel unbalanced translocation between chromosomes 5p and 18q leading to dysmorphology and global developmental delay. Molecular Genetics & Genomic Medicine, 2022, , e1900.	0.6	1