Alexander E Katz

List of Publications by Year
in descending order

10 papers	394 citations	1307594 7 h-index	1474206 9 g-index
10 all docs	10 docs citations	10 times ranked	797 citing authors

#	Article	IF	Citations
1	The ACMG SF v3.0 gene list increases returnable variant detection by 22% when compared with v2.0 in the ClinSeqÂcohort. Genetics in Medicine, 2022, 24, 736-743.	2.4	7
2	Pitfalls and challenges in genetic test interpretation: An exploration of genetic professionals experience with interpretation of results. Clinical Genetics, 2021, 99, 638-649.	2.0	15
3	Chromosome 1q21.2 and additional loci influence risk of spontaneous coronary artery dissection and myocardial infarction. Nature Communications, 2020, 11, 4432.	12.8	60
4	A Novel Recurrent COL5A1 Genetic Variant Is Associated With a Dysplasia-Associated Arterial Disease Exhibiting Dissections and Fibromuscular Dysplasia. Arteriosclerosis, Thrombosis, and Vascular Biology, 2020, 40, 2686-2699.	2.4	30
5	Common genetic susceptibility loci link PFAPA syndrome, BehÃ§etâ€™s disease, and recurrent aphthous stomatitis. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 14405-14411.	7.1	52
6	Management of Secondary Genomic Findings. American Journal of Human Genetics, 2020, 107, 3-14.	6.2	29
7	Variants in myelin regulatory factor (MYRF) cause autosomal dominant and syndromic nanophthalmos in humans and retinal degeneration in mice. PLoS Genetics, 2019, 15, e1008130.	3.5	50
8	PHACTR1 Is a Genetic Susceptibility Locus for Fibromuscular Dysplasia Supporting Its Complex Genetic Pattern of Inheritance. PLoS Genetics, 2016, 12, e1006367.	3.5	146
9	Abstract 15370: Genetic Study Identifies Common Variation in PHACTR1 to Associate With Fibromuscular Dysplasia (Best of Basic Science Abstract). Circulation, 2015, 132, .	1.6	5
10	Conservative management of an elderly patient with Eisenmenger syndrome. Journal of Cardiology Cases, 2013, 7, e114-e116.	0.5	0