

# Alexander E Katz

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/2254017/publications.pdf>

Version: 2024-02-01

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