

# Oyediran Akinrinade

## List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

21  
papers

427  
citations

11  
h-index

20  
g-index

25  
ext. papers

510  
ext. citations

4.3  
avg, IF

3.5  
L-index

#	Paper	IF	Citations
21	Whole genome sequencing delineates regulatory, copy number, and cryptic splice variants in early onset cardiomyopathy.. <i>Npj Genomic Medicine</i> , <b>2022</b> , 7, 18	6.2	0
20	Evolving Up-regulation of Biliary Fibrosis-Related Extracellular Matrix Molecules After Successful Portoenterostomy. <i>Hepatology Communications</i> , <b>2021</b> , 5, 1036-1050	6	5
19	Everolimus Rescues the Phenotype of Elastin Insufficiency in Patient Induced Pluripotent Stem Cell-Derived Vascular Smooth Muscle Cells. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , <b>2020</b> , 40, 1325-1339	9.4	9
18	Quality of whole genome sequencing from blood versus saliva derived DNA in cardiac patients. <i>BMC Medical Genomics</i> , <b>2020</b> , 13, 11	3.7	13
17	Machine Learning Identifies Clinical and Genetic Factors Associated With Anthracycline Cardiotoxicity in Pediatric Cancer Survivors. <i>JACC: CardioOncology</i> , <b>2020</b> , 2, 690-706	3.8	6
16	Relevance of Titin Missense and Non-Frameshifting Insertions/Deletions Variants in Dilated Cardiomyopathy. <i>Scientific Reports</i> , <b>2019</b> , 9, 4093	4.9	19
15	Targeted next generation sequencing reveals genetic defects underlying inherited retinal disease in Iranian families. <i>Molecular Vision</i> , <b>2019</b> , 25, 106-117	2.3	10
14	Return of genetic and genomic research findings: experience of a pediatric biorepository. <i>BMC Medical Genomics</i> , <b>2019</b> , 12, 173	3.7	10
13	Transcription factor GATA4 associates with mesenchymal-like gene expression in human hepatoblastoma cells. <i>Tumor Biology</i> , <b>2018</b> , 40, 1010428318785498	2.9	8
12	Accessory genetic content in <i>Campylobacter jejuni</i> ST21CC isolates from feces and blood. <i>International Journal of Medical Microbiology</i> , <b>2017</b> , 307, 233-240	3.7	10
11	GATA4 Regulates Blood-Testis Barrier Function and Lactate Metabolism in Mouse Sertoli Cells. <i>Endocrinology</i> , <b>2016</b> , 157, 2416-31	4.8	26
10	Loss of PPAR $\alpha$ in endothelial cells leads to impaired angiogenesis. <i>Journal of Cell Science</i> , <b>2016</b> , 129, 693-705	4.9	28
9	Germline copy number variation analysis in Finnish families with hereditary prostate cancer. <i>Prostate</i> , <b>2016</b> , 76, 316-24	4.2	11
8	Relevance of truncating titin mutations in dilated cardiomyopathy. <i>Clinical Genetics</i> , <b>2016</b> , 90, 49-54	4	34
7	Genetics and genotype-phenotype correlations in Finnish patients with dilated cardiomyopathy. <i>European Heart Journal</i> , <b>2015</b> , 36, 2327-37	9.5	101
6	Comparative genomics and genome biology of invasive <i>Campylobacter jejuni</i> . <i>Scientific Reports</i> , <b>2015</b> , 5, 17300	4.9	14
5	Prevalence of Titin Truncating Variants in General Population. <i>PLoS ONE</i> , <b>2015</b> , 10, e0145284	3.7	67

4	GATA4 is a key regulator of steroidogenesis and glycolysis in mouse Leydig cells. <i>Endocrinology</i> , <b>2015</b> , 156, 1860-72	4.8	27
3	copy number variation analysis in familial BRCA1/2-negative Finnish breast and ovarian cancer. <i>PLoS ONE</i> , <b>2013</b> , 8, e71802	3.7	25
2	Whole genome sequencing delineates regulatory and novel genic variants in childhood cardiomyopathy		3
1	Age and Sex Differences in the Genetics of Cardiomyopathy		1