Eddie Ip

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.

10	216	5	10
papers	citations	h-index	g-index
10	330 ext. citations	10.8	1.99
ext. papers		avg, IF	L-index

#	Paper	IF	Citations
10	Congenital Heart Disease Gene: a Curated Database for Congenital Heart Disease Genes <i>Circulation Genomic and Precision Medicine</i> , 2022 , 101161CIRCGEN121003539	5.2	
9	Whole genome sequencing in transposition of the great arteries and associations with clinically relevant heart, brain and laterality genes. <i>American Heart Journal</i> , 2021 , 244, 1-13	4.9	1
8	Functional genomics and gene-environment interaction highlight the complexity of congenital heart disease caused by Notch pathway variants. <i>Human Molecular Genetics</i> , 2020 , 29, 566-579	5.6	16
7	Heterozygous loss of WBP11 function causes multiple congenital defects in humans and mice. <i>Human Molecular Genetics</i> , 2020 , 29, 3662-3678	5.6	3
6	Spliceogen: an integrative, scalable tool for the discovery of splice-altering variants. <i>Bioinformatics</i> , 2019 , 35, 4405-4407	7.2	3
5	VPOT: A Customizable Variant Prioritization Ordering Tool for Annotated Variants. <i>Genomics, Proteomics and Bioinformatics,</i> 2019 , 17, 540-545	6.5	2
4	Identification of clinically actionable variants from genome sequencing of families with congenital heart disease. <i>Genetics in Medicine</i> , 2019 , 21, 1111-1120	8.1	25
3	Genetic burden and associations with adverse neurodevelopment in neonates with congenital heart disease. <i>American Heart Journal</i> , 2018 , 201, 33-39	4.9	15
2	A Screening Approach to Identify Clinically Actionable Variants Causing Congenital Heart Disease in Exome Data. <i>Circulation Genomic and Precision Medicine</i> , 2018 , 11, e001978	5.2	37
1	NAD Deficiency, Congenital Malformations, and Niacin Supplementation. <i>New England Journal of Medicine</i> , 2017 , 377, 544-552	59.2	114