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## List of Publications by Year in descending order

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7  
papers

208  
citations

1478505

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1720034

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docs citations

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281  
citing authors

#	ARTICLE	IF	CITATIONS
1	Common Genetic Variants Contribute to Risk of Transposition of the Great Arteries. <i>Circulation Research</i> , 2022, 130, 166-180.	4.5	15
2	Genome-wide association analyses identify new Brugada syndrome risk loci and highlight a new mechanism of sodium channel regulation in disease susceptibility. <i>Nature Genetics</i> , 2022, 54, 232-239.	21.4	55
3	Enhancing rare variant interpretation in inherited arrhythmias through quantitative analysis of consortium disease cohorts and population controls. <i>Genetics in Medicine</i> , 2021, 23, 47-58.	2.4	57
4	Biallelic loss-of-function variants in <i>PLD1</i> cause congenital right-sided cardiac valve defects and neonatal cardiomyopathy. <i>Journal of Clinical Investigation</i> , 2021, 131, .	8.2	16
5	Rare variants in <i>KDR</i> , encoding VEGF Receptor 2, are associated with tetralogy of Fallot. <i>Genetics in Medicine</i> , 2021, 23, 1952-1960.	2.4	7
6	<i>SCN5A</i> Mutation Type and a Genetic Risk Score Associate Variably With Brugada Syndrome Phenotype in <i>SCN5A</i> Families. <i>Circulation Genomic and Precision Medicine</i> , 2020, 13, e002911.	3.6	41
7	<i>GATA6</i> mutations: Characterization of two novel patients and a comprehensive overview of the <i>GATA6</i> genotypic and phenotypic spectrum. <i>American Journal of Medical Genetics, Part A</i> , 2019, 179, 1836-1845.	1.2	16