

# John Garcia

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/780697/publications.pdf>

Version: 2024-02-01

8  
papers

1,193  
citations

1684188

5  
h-index

1588992

8  
g-index

8  
all docs

8  
docs citations

8  
times ranked

2627  
citing authors

#	ARTICLE	IF	CITATIONS
1	Evaluation of gene validity for CPVT and short QT syndrome in sudden arrhythmic death. <i>European Heart Journal</i> , 2022, 43, 1500-1510.	2.2	57
2	Common Variants in <i>KCNE1</i> , <i>KCNH2</i> , and <i>SCN5A</i> May Impact Cardiac Arrhythmia Risk. <i>Circulation Genomic and Precision Medicine</i> , 2021, 14, e003206.	3.6	3
3	Clinical Implications of Pathogenic Germline Variants in Small Intestine Neuroendocrine Tumors (SI-NETs). <i>JCO Precision Oncology</i> , 2021, 5, 808-816.	3.0	7
4	<i>ATP1A3</i> Encoded Sodium-Potassium ATPase Subunit Alpha 3 D801N Variant Is Associated With Shortened QT Interval and Predisposition to Ventricular Fibrillation Preceded by Bradycardia. <i>Journal of the American Heart Association</i> , 2021, 10, e019887.	3.7	3
5	An International, Multicentered, Evidence-Based Reappraisal of Genes Reported to Cause Congenital Long QT Syndrome. <i>Circulation</i> , 2020, 141, 418-428.	1.6	238
6	ClinVar database of global familial hypercholesterolemia-associated DNA variants. <i>Human Mutation</i> , 2018, 39, 1631-1640.	2.5	84
7	Reappraisal of Reported Genes for Sudden Arrhythmic Death. <i>Circulation</i> , 2018, 138, 1195-1205.	1.6	271
8	Sherloc: a comprehensive refinement of the ACMG-AMP variant classification criteria. <i>Genetics in Medicine</i> , 2017, 19, 1105-1117.	2.4	530