

Sharon Cresci

List of Publications by Year in descending order

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Version: 2024-02-01

35
papers

1,565
citations

471509

17
h-index

395702

33
g-index

35
all docs

35
docs citations

35
times ranked

3716
citing authors

#	ARTICLE	IF	CITATIONS
1	Pulsus Alternans in Cardiogenic Shock Recapitulated in Single Cell Fluorescence Imaging of a Patient's Cardiomyocyte. <i>Circulation: Heart Failure</i> , 2022, 15, CIRCHEARTFAILURE121008855.	3.9	5
2	COVID-19: Understanding Inter-Individual Variability and Implications for Precision Medicine. <i>Mayo Clinic Proceedings</i> , 2021, 96, 446-463.	3.0	62
3	Genetically determined NLRP3 inflammasome activation associates with systemic inflammation and cardiovascular mortality. <i>European Heart Journal</i> , 2021, 42, 1742-1756.	2.2	63
4	Phenylephrine Provocation to Evaluate the Cause of Mitral Regurgitation in Patients With Obstructive Hypertrophic Cardiomyopathy. <i>Circulation: Cardiovascular Imaging</i> , 2021, 14, e012656.	2.6	1
5	Exploring experiences of hypertrophic cardiomyopathy diagnosis, treatment, and impacts on quality of life among middle-aged and older adults: An interview study. <i>Heart and Lung: Journal of Acute and Critical Care</i> , 2021, 50, 788-793.	1.6	2
6	Syndrome of Reversible Cardiogenic Shock and Left Ventricular Ballooning in Obstructive Hypertrophic Cardiomyopathy. <i>Journal of the American Heart Association</i> , 2021, 10, e021141.	3.7	9
7	Association of Factor V Leiden With Subsequent Atherothrombotic Events. <i>Circulation</i> , 2020, 142, 546-555.	1.6	11
8	<i>PPARA</i> Polymorphism Influences the Cardiovascular Benefit of Fenofibrate in Type 2 Diabetes: Findings From ACCORD-Lipid. <i>Diabetes</i> , 2020, 69, 771-783.	0.6	28
9	Recurrent Takotsubo Cardiomyopathy in a Patient With Hypertrophic Cardiomyopathy Leading to Cardiogenic Shock Requiring VA-ECMO. <i>JACC: Case Reports</i> , 2020, 2, 1014-1018.	0.6	4
10	Genetic Architecture of Circulating Very-Long-Chain (C24:0 and C22:0) Ceramide Concentrations. <i>Journal of Lipid and Atherosclerosis</i> , 2020, 9, 172.	3.5	10
11	A Genetic Locus on Chromosome 2q24 Predicting Peripheral Neuropathy Risk in Type 2 Diabetes: Results From the ACCORD and BARI 2D Studies. <i>Diabetes</i> , 2019, 68, 1649-1662.	0.6	22
12	Subsequent Event Risk in Individuals With Established Coronary Heart Disease. <i>Circulation Genomic and Precision Medicine</i> , 2019, 12, e002470.	3.6	17
13	Association between the EPHX2 p.Lys55Arg polymorphism and prognosis following an acute coronary syndrome. <i>Prostaglandins and Other Lipid Mediators</i> , 2018, 138, 15-22.	1.9	9
14	A novel genetic marker of decreased inflammation and improved survival after acute myocardial infarction. <i>Basic Research in Cardiology</i> , 2018, 113, 38.	5.9	58
15	Relations between lipoprotein(a) concentrations, LPA genetic variants, and the risk of mortality in patients with established coronary heart disease: a molecular and genetic association study. <i>Lancet Diabetes and Endocrinology</i> , 2017, 5, 534-543.	11.4	84
16	Change in Angina Symptom Status After Acute Myocardial Infarction and Its Association With Readmission Risk: An Analysis of the Translational Research Investigating Underlying Disparities in Acute Myocardial Infarction Patients' Health Status (TRIUMPH) Registry. <i>Journal of the American Heart Association</i> , 2016, 5, .	3.7	9
17	Association between diabetes mellitus and angina after acute myocardial infarction: analysis of the TRIUMPH prospective cohort study. <i>European Journal of Preventive Cardiology</i> , 2015, 22, 779-787.	1.8	15
18	Utility of a genetic risk score to predict recurrent cardiovascular events 1 year after an acute coronary syndrome: A pooled analysis of the RISCA, PRAXY, and TRIUMPH cohorts. <i>Atherosclerosis</i> , 2015, 242, 261-267.	0.8	21

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19	CYP450 pharmacogenomics: a cardiology perspective. <i>Personalized Medicine</i> , 2015, 12, 59-62.	1.5	0
20	Exome sequencing identifies rare LDLR and APOA5 alleles conferring risk for myocardial infarction. <i>Nature</i> , 2015, 518, 102-106.	27.8	581
21	CHRNA5 Variant Predicts Smoking Cessation in Patients With Acute Myocardial Infarction. <i>Nicotine and Tobacco Research</i> , 2014, 16, 1224-1231.	2.6	25
22	Cytochrome P450 Gene Variants, Race, and Mortality Among Clopidogrel-Treated Patients After Acute Myocardial Infarction. <i>Circulation: Cardiovascular Genetics</i> , 2014, 7, 277-286.	5.1	50
23	Adrenergic-Pathway Gene Variants Influence Beta-Blocker-Related Outcomes After Acute Coronary Syndrome in a Race-Specific Manner. <i>Journal of the American College of Cardiology</i> , 2012, 60, 898-907.	2.8	35
24	Factors influencing patient willingness to participate in genetic research after a myocardial infarction. <i>Genome Medicine</i> , 2011, 3, 39.	8.2	21
25	Investigation of 95 variants identified in a genome-wide study for association with mortality after acute coronary syndrome. <i>BMC Medical Genetics</i> , 2011, 12, 127.	2.1	14
26	Peroxisome Proliferator-Activated Receptor Pathway Gene Polymorphism Associated With Extent of Coronary Artery Disease in Patients With Type 2 Diabetes in the Bypass Angioplasty Revascularization Investigation 2 Diabetes Trial. <i>Circulation</i> , 2011, 124, 1426-1434.	1.6	28
27	A PPAR± Promoter Variant Impairs ERR-Dependent Transactivation and Decreases Mortality after Acute Coronary Ischemia in Patients with Diabetes. <i>PLoS ONE</i> , 2010, 5, e12584.	2.5	18
28	Clinical and Genetic Modifiers of Long-Term Survival in Heart Failure. <i>Journal of the American College of Cardiology</i> , 2009, 54, 432-444.	2.8	105
29	From SNPs to Functional Studies in Cardiovascular Pharmacogenomics. <i>Methods in Molecular Biology</i> , 2008, 448, 379-393.	0.9	3
30	PPAR Genomics and Pharmacogenomics: Implications for Cardiovascular Disease. <i>PPAR Research</i> , 2008, 2008, 1-11.	2.4	11
31	Interaction between <i>PPARA</i> genotype and ß ² -blocker treatment influences clinical outcomes following acute coronary syndromes. <i>Pharmacogenomics</i> , 2008, 9, 1403-1417.	1.3	16
32	Epithelial Neutrophil-Activating Peptide (ENA-78), Acute Coronary Syndrome Prognosis, and Modulatory Effect of Statins. <i>PLoS ONE</i> , 2008, 3, e3117.	2.5	32
33	Pharmacogenetics of the PPAR genes and cardiovascular disease. <i>Pharmacogenomics</i> , 2007, 8, 1581-1595.	1.3	7
34	ß ₂ -Adrenergic Receptor Genotype and Survival Among Patients Receiving ß ² -Blocker Therapy After an Acute Coronary Syndrome. <i>JAMA - Journal of the American Medical Association</i> , 2005, 294, 1526.	7.4	177
35	The PPAR genes, cardiovascular disease and the emergence of PPAR pharmacogenetics. <i>Expert Opinion on Pharmacotherapy</i> , 2005, 6, 2577-2591.	1.8	12