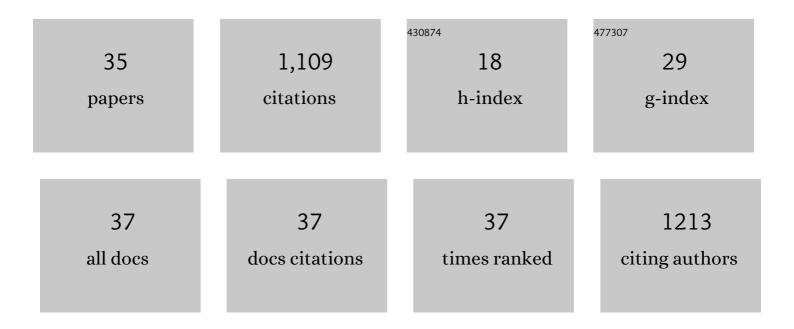
## Michelle S Parvatiyar

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Post-translational modification patterns on $\hat{l}^2$ -myosin heavy chain are altered in ischemic and nonischemic human hearts. ELife, 2022, 11, .	6.0	10
2	Sarcospanâ€deficient mice exhibit a heightened inflammatory phenotype under obesiogenic conditions. FASEB Journal, 2022, 36, .	0.5	0
3	Anomalous structural dynamics of minimally frustrated residues in cardiac troponin C triggers hypertrophic cardiomyopathy. Chemical Science, 2021, 12, 7308-7323.	7.4	7
4	Cardiovascular Injury Due to SARS-CoV-2. Current Clinical Microbiology Reports, 2021, 8, 167-177.	3.4	18
5	Essential roles of the dystrophin-glycoprotein complex in different cardiac pathologies. Advances in Medical Sciences, 2021, 66, 52-71.	2.1	11
6	Establishing a Role for Sarcospan as an Obesity‣usceptibility Gene in Mice. FASEB Journal, 2021, 35, .	0.5	0
7	A comprehensive guide to genetic variants and post-translational modifications of cardiac troponin C. Journal of Muscle Research and Cell Motility, 2020, 42, 323-342.	2.0	12
8	Sexual dimorphism in cardiac transcriptome associated with a troponin C murine model of hypertrophic cardiomyopathy. Physiological Reports, 2020, 8, e14396.	1.7	7
9	Meta-analysis of cardiomyopathy-associated variants in troponin genes identifies loci and intragenic hot spots that are associated with worse clinical outcomes. Journal of Molecular and Cellular Cardiology, 2020, 142, 118-125.	1.9	30
10	Beta-Myosin Heavy Chain Post-Translational Modifications in Failing and Non-Failing Human Hearts. Biophysical Journal, 2019, 116, 29a-30a.	0.5	0
11	The missing links within troponin. Archives of Biochemistry and Biophysics, 2019, 663, 95-100.	3.0	6
12	Stabilization of the cardiac sarcolemma by sarcospan rescues DMD-associated cardiomyopathy. JCI Insight, 2019, 4, .	5.0	18
13	Hypertrophic Cardiomyopathy Cardiac Troponin C Mutations Differentially Affect Slow Skeletal and Cardiac Muscle Regulation. Frontiers in Physiology, 2017, 8, 221.	2.8	16
14	Sarcospan Regulates Cardiac Isoproterenol Response and Prevents Duchenne Muscular Dystrophy–Associated Cardiomyopathy. Journal of the American Heart Association, 2015, 4, .	3.7	31
15	In Vivo Analysis of Troponin C Knock-In (A8V) Mice. Circulation: Cardiovascular Genetics, 2015, 8, 653-664.	5.1	32
16	Pathogenesis associated with a restrictive cardiomyopathy mutant in cardiac troponin T is due to reduced protein stability and greatly increased myofilament Ca2+ sensitivity. Biochimica Et Biophysica Acta - General Subjects, 2015, 1850, 365-372.	2.4	6
17	The Role of Sarcospan in Cardiac Sarcolemma Organization and Function. FASEB Journal, 2015, 29, 801.3.	0.5	0
18	A Mutation in TNNC1-encoded Cardiac Troponin C, TNNC1-A31S, Predisposes to Hypertrophic Cardiomyopathy and Ventricular Fibrillation. Journal of Biological Chemistry, 2012, 287, 31845-31855.	3.4	50

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#	Article	IF	CITATIONS
19	The Functional Properties of Human Slow Skeletal Troponin T Isoforms in Cardiac Muscle Regulation. Journal of Biological Chemistry, 2012, 287, 37362-37370.	3.4	21
20	Structural considerations for chromatin state models with transcription as a functional readout. FEBS Letters, 2012, 586, 3548-3554.	2.8	5
21	Cardiac Linker Histones Are Differentially Regulated Following Hypertrophic Stimuli. FASEB Journal, 2012, 26, 1127.9.	0.5	0
22	Fetal Cardiac Troponin Isoforms Rescue the Increased Ca2+ Sensitivity Produced by a Novel Double Deletion in Cardiac Troponin T Linked to Restrictive Cardiomyopathy. Biophysical Journal, 2011, 100, 114a-115a.	0.5	0
23	Functional Characterization of TNNC1 Rare Variants Identified in Dilated Cardiomyopathy. Journal of Biological Chemistry, 2011, 286, 34404-34412.	3.4	43
24	Strong Cross-bridges Potentiate the Ca2+ Affinity Changes Produced by Hypertrophic Cardiomyopathy Cardiac Troponin C Mutants in Myofilaments. Journal of Biological Chemistry, 2011, 286, 1005-1013.	3.4	28
25	Fetal Cardiac Troponin Isoforms Rescue the Increased Ca2+ Sensitivity Produced by a Novel Double Deletion in Cardiac Troponin T Linked to Restrictive Cardiomyopathy. Journal of Biological Chemistry, 2011, 286, 20901-20912.	3.4	19
26	Predicting Cardiomyopathic Phenotypes by Altering Ca2+ Affinity of Cardiac Troponin C. Journal of Biological Chemistry, 2010, 285, 27785-27797.	3.4	26
27	Cardiac Troponin Mutations and Restrictive Cardiomyopathy. Journal of Biomedicine and Biotechnology, 2010, 2010, 1-9.	3.0	54
28	Mutations in Troponin that cause HCM, DCM AND RCM: What can we learn about thin filament function?. Journal of Molecular and Cellular Cardiology, 2010, 48, 882-892.	1.9	176
29	Clinical and Functional Characterization of <i>TNNT2</i> Mutations Identified in Patients With Dilated Cardiomyopathy. Circulation: Cardiovascular Genetics, 2009, 2, 306-313.	5.1	95
30	A Functional and Structural Study of Troponin C Mutations Related to Hypertrophic Cardiomyopathy. Journal of Biological Chemistry, 2009, 284, 19090-19100.	3.4	83
31	Molecular and functional characterization of novel hypertrophic cardiomyopathy susceptibility mutations in TNNC1-encoded troponin C. Journal of Molecular and Cellular Cardiology, 2008, 45, 281-288.	1.9	111
32	Troponin and cardiomyopathy. Biochemical and Biophysical Research Communications, 2008, 369, 74-81.	2.1	36
33	A Troponin T Mutation That Causes Infantile Restrictive Cardiomyopathy Increases Ca2+ Sensitivity of Force Development and Impairs the Inhibitory Properties of Troponin. Journal of Biological Chemistry, 2008, 283, 2156-2166.	3.4	52
34	Transgenic mice expressing Na+-K+-ATPase in smooth muscle decreases blood pressure. American Journal of Physiology - Heart and Circulatory Physiology, 2007, 293, H1172-H1182.	3.2	39
35	Global Analysis of Cellular Factors and Responses Involved in Pseudomonas aeruginosa Resistance to Arsenite. Journal of Bacteriology, 2005, 187, 4853-4864.	2.2	67