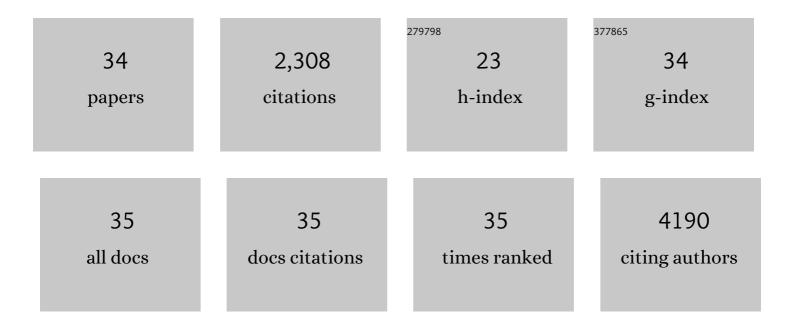
Yusu Gu

List of Publications by Year in descending order

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Yusu Cu

#	Article	IF	CITATIONS
1	Resident fibroblast lineages mediate pressure overload–induced cardiac fibrosis. Journal of Clinical Investigation, 2014, 124, 2921-2934.	8.2	497
2	Pericytes of Multiple Organs Do Not Behave as Mesenchymal Stem Cells InÂVivo. Cell Stem Cell, 2017, 20, 345-359.e5.	11.1	393
3	Nesprin 1 is critical for nuclear positioning and anchorage. Human Molecular Genetics, 2010, 19, 329-341.	2.9	131
4	Mouse and computational models link Mlc2v dephosphorylation to altered myosin kinetics in early cardiac disease. Journal of Clinical Investigation, 2012, 122, 1209-1221.	8.2	131
5	Loss-of-function mutations in co-chaperone BAC3 destabilize small HSPs and cause cardiomyopathy. Journal of Clinical Investigation, 2017, 127, 3189-3200.	8.2	107
6	Progressive Cardiac Dysfunction and Fibrosis in the Cardiomyopathic Hamster and Effects of Growth Hormone and Angiotensin-Converting Enzyme Inhibition. Circulation, 1999, 100, 1734-1743.	1.6	94
7	Cardiac-specific ablation of Cypher leads to a severe form of dilated cardiomyopathy with premature death. Human Molecular Genetics, 2009, 18, 701-713.	2.9	88
8	Infarct Fibroblasts Do Not Derive From Bone Marrow Lineages. Circulation Research, 2018, 122, 583-590.	4.5	65
9	Adipocyte-specific loss of PPARÎ ³ attenuates cardiac hypertrophy. JCI Insight, 2016, 1, e89908.	5.0	65
10	Reduction of myocardial ischaemia–reperfusion injury by inactivating oxidized phospholipids. Cardiovascular Research, 2019, 115, 179-189.	3.8	61
11	Impaired mitophagy facilitates mitochondrial damage in Danon disease. Journal of Molecular and Cellular Cardiology, 2017, 108, 86-94.	1.9	57
12	β1 Integrin Gene Excision in the Adult Murine Cardiac Myocyte Causes Defective Mechanical and Signaling Responses. American Journal of Pathology, 2012, 180, 952-962.	3.8	51
13	The Muscle Ankyrin Repeat Proteins CARP, Ankrd2, and DARP Are Not Essential for Normal Cardiac Development and Function at Basal Conditions and in Response to Pressure Overload. PLoS ONE, 2014, 9, e93638.	2.5	49
14	Systemic AAV9.LAMP2B injection reverses metabolic and physiologic multiorgan dysfunction in a murine model of Danon disease. Science Translational Medicine, 2020, 12, .	12.4	49
15	Mitochondrial Reprogramming Induced by CaMKIIδ Mediates Hypertrophy Decompensation. Circulation Research, 2015, 116, e28-39.	4.5	47
16	Suppression of Endothelial AGO1 Promotes Adipose Tissue Browning and Improves Metabolic Dysfunction. Circulation, 2020, 142, 365-379.	1.6	44
17	PRDM16 Is a Compact Myocardium-Enriched Transcription Factor Required to Maintain Compact Myocardial Cardiomyocyte Identity in Left Ventricle. Circulation, 2022, 145, 586-602.	1.6	44
18	Normalization of Naxos plakoglobin levels restores cardiac function in mice. Journal of Clinical Investigation, 2015, 125, 1708-1712.	8.2	39

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19	Combinatorial interactions of genetic variants in human cardiomyopathy. Nature Biomedical Engineering, 2019, 3, 147-157.	22.5	37
20	Luma is not essential for murine cardiac development and function. Cardiovascular Research, 2018, 114, 378-388.	3.8	35
21	Discoidin domain receptor 2 germline gene deletion leads to altered heart structure and function in the mouse. American Journal of Physiology - Heart and Circulatory Physiology, 2014, 307, H773-H781.	3.2	33
22	Desmosomal junctions are necessary for adult sinus node function. Cardiovascular Research, 2016, 111, 274-286.	3.8	33
23	Aortic pathology from protein kinase G activation is prevented by an antioxidant vitamin B12 analog. Nature Communications, 2019, 10, 3533.	12.8	30
24	Nebulette knockout mice have normal cardiac function, but show Z-line widening and up-regulation of cardiac stress markers. Cardiovascular Research, 2015, 107, 216-225.	3.8	27
25	A secretory pathway kinase regulates sarcoplasmic reticulum Ca2+ homeostasis and protects against heart failure. ELife, 2018, 7, .	6.0	22
26	Desmosomal COP9 regulates proteome degradation in arrhythmogenic right ventricular dysplasia/cardiomyopathy. Journal of Clinical Investigation, 2021, 131, .	8.2	18
27	Cardiolipin Remodeling Defects Impair Mitochondrial Architecture and Function in a Murine Model of Barth Syndrome Cardiomyopathy. Circulation: Heart Failure, 2021, 14, e008289.	3.9	17
28	Urocortin 2 Lowers Blood Pressure and Reduces Plasma Catecholamine Levels in Mice with Hyperadrenergic Activity. Endocrinology, 2010, 151, 4820-4829.	2.8	10
29	Selective Life‣ong Skeletal Myofiber—Targeted VEGF Gene Ablation Impairs Exercise Capacity in Adult Mice. Journal of Cellular Physiology, 2016, 231, 505-511.	4.1	8
30	Cardiomyocyte Expression of ZO-1 Is Essential for Normal Atrioventricular Conduction but Does Not Alter Ventricular Function. Circulation Research, 2020, 127, 284-297.	4.5	8
31	Constitutive protein kinase G activation exacerbates stressâ€induced cardiomyopathy. British Journal of Pharmacology, 2022, 179, 2413-2429.	5.4	7
32	Mediator complex proximal Tail subunit MED30 is critical for Mediator core stability and cardiomyocyte transcriptional network. PLoS Genetics, 2021, 17, e1009785.	3.5	4
33	Increased Echogenicity and Radiodense Foci on Echocardiogram and MicroCT in Murine Myocarditis. PLoS ONE, 2016, 11, e0159971.	2.5	4
34	Atypical ALPK2 kinase is not essential for cardiac development and function. American Journal of Physiology - Heart and Circulatory Physiology, 2020, 318, H1509-H1515.	3.2	3