

Wuqiang Zhu

List of Publications by Year in descending order

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39
papers

1,749
citations

394421

19
h-index

315739

38
g-index

40
all docs

40
docs citations

40
times ranked

2763
citing authors

#	ARTICLE	IF	CITATIONS
1	Optogenetic Control of Engrafted Human Induced Pluripotent Stem Cell-Derived Cardiomyocytes in Live Mice: A Proof-of-Concept Study. <i>Cells</i> , 2022, 11, 951.	4.1	2
2	Nanofiber capsules for minimally invasive sampling of biological specimens from gastrointestinal tract. <i>Acta Biomaterialia</i> , 2022, 146, 211-221.	8.3	5
3	Identification of metabolic pathways underlying FGF1 and CHIR99021-mediated cardioprotection. <i>IScience</i> , 2022, 25, 104447.	4.1	5
4	Turning back the clock: A concise viewpoint of cardiomyocyte cell cycle activation for myocardial regeneration and repair. <i>Journal of Molecular and Cellular Cardiology</i> , 2022, 170, 15-21.	1.9	4
5	Minimally Invasive Delivery of 3D Shape Recoverable Constructs with Ordered Structures for Tissue Repair. <i>ACS Biomaterials Science and Engineering</i> , 2021, 7, 2204-2211.	5.2	16
6	Circular RNAs and Cardiovascular Regeneration. <i>Frontiers in Cardiovascular Medicine</i> , 2021, 8, 672600.	2.4	5
7	Cyclin D2 Overexpression Enhances the Efficacy of Human Induced Pluripotent Stem Cell-Derived Cardiomyocytes for Myocardial Repair in a Swine Model of Myocardial Infarction. <i>Circulation</i> , 2021, 144, 210-228.	1.6	61
8	De novo Drug Delivery Modalities for Treating Damaged Hearts: Current Challenges and Emerging Solutions. <i>Frontiers in Cardiovascular Medicine</i> , 2021, 8, 742315.	2.4	2
9	Metabolic Profile in Neonatal Pig Hearts. <i>Frontiers in Cardiovascular Medicine</i> , 2021, 8, 763984.	2.4	3
10	N-cadherin overexpression enhances the reparative potency of human-induced pluripotent stem cell-derived cardiac myocytes in infarcted mouse hearts. <i>Cardiovascular Research</i> , 2020, 116, 671-685.	3.8	25
11	Targeting exosome-associated human antigen R attenuates fibrosis and inflammation in diabetic heart. <i>FASEB Journal</i> , 2020, 34, 2238-2251.	0.5	50
12	CHIR99021 and fibroblast growth factor 1 enhance the regenerative potency of human cardiac muscle patch after myocardial infarction in mice. <i>Journal of Molecular and Cellular Cardiology</i> , 2020, 141, 1-10.	1.9	40
13	Utilization of Human Induced Pluripotent Stem Cells for Cardiac Repair. <i>Frontiers in Cell and Developmental Biology</i> , 2020, 8, 36.	3.7	20
14	Fluorescent indicators for continuous and lineage-specific reporting of cell cycle phases in human pluripotent stem cells. <i>Biotechnology and Bioengineering</i> , 2020, 117, 2177-2186.	3.3	10
15	Myocardial protection by nanomaterials formulated with CHIR99021 and FGF1. <i>JCI Insight</i> , 2020, 5, .	5.0	15
16	Editorial: Nanotechnology in Cardiovascular Regenerative Medicine. <i>Frontiers in Bioengineering and Biotechnology</i> , 2020, 8, 608844.	4.1	5
17	Y-27632 preconditioning enhances transplantation of human-induced pluripotent stem cell-derived cardiomyocytes in myocardial infarction mice. <i>Cardiovascular Research</i> , 2019, 115, 343-356.	3.8	30
18	Cardiomyocytes from CCND2-overexpressing human induced-pluripotent stem cells repopulate the myocardial scar in mice: A 6-month study. <i>Journal of Molecular and Cellular Cardiology</i> , 2019, 137, 25-33.	1.9	19

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19	Enhancing the Engraftment of Human Induced Pluripotent Stem Cell-derived Cardiomyocytes via a Transient Inhibition of Rho Kinase Activity. <i>Journal of Visualized Experiments</i> , 2019, , .	0.3	4
20	Targeted expression of cyclin D2 ameliorates late stage anthracycline cardiotoxicity. <i>Cardiovascular Research</i> , 2019, 115, 960-965.	3.8	19
21	Optogenetics: Background, Methodological Advances and Potential Applications for Cardiovascular Research and Medicine. <i>Frontiers in Bioengineering and Biotechnology</i> , 2019, 7, 466.	4.1	57
22	CCND2 Overexpression Enhances the Regenerative Potency of Human Induced Pluripotent Stem Cell-Derived Cardiomyocytes. <i>Circulation Research</i> , 2018, 122, 88-96.	4.5	113
23	Large Cardiac Muscle Patches Engineered From Human Induced-Pluripotent Stem Cell-Derived Cardiac Cells Improve Recovery From Myocardial Infarction in Swine. <i>Circulation</i> , 2018, 137, 1712-1730.	1.6	332
24	VEGF nanoparticles repair the heart after myocardial infarction. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2018, 314, H278-H284.	3.2	101
25	Protein phosphatase 5 and the tumor suppressor p53 down-regulate each other's activities in mice. <i>Journal of Biological Chemistry</i> , 2018, 293, 18218-18229.	3.4	14
26	Spheroids of cardiomyocytes derived from human-induced pluripotent stem cells improve recovery from myocardial injury in mice. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2018, 315, H327-H339.	3.2	65
27	Can We Engineer a Human Cardiac Patch for Therapy?. <i>Circulation Research</i> , 2018, 123, 244-265.	4.5	121
28	Regenerative Potential of Neonatal Porcine Hearts. <i>Circulation</i> , 2018, 138, 2809-2816.	1.6	179
29	Meeting Report for the 2017 National Institutes of Health National Heart, Lung, and Blood Institute Progenitor Cell Biology Consortium. <i>Circulation Research</i> , 2017, 120, 1709-1712.	4.5	2
30	Biomarkers for monitoring chemotherapy-induced cardiotoxicity. <i>Critical Reviews in Clinical Laboratory Sciences</i> , 2017, 54, 87-101.	6.1	22
31	Pluripotent Stem Cell Derived Cardiac Cells for Myocardial Repair. <i>Journal of Visualized Experiments</i> , 2017, , .	0.3	9
32	Overcoming the Roadblocks to Cardiac Cell Therapy Using Tissue Engineering. <i>Journal of the American College of Cardiology</i> , 2017, 70, 766-775.	2.8	82
33	Cardiomyocyte proliferation prevents failure in pressure overload but not volume overload. <i>Journal of Clinical Investigation</i> , 2017, 127, 4285-4296.	8.2	31
34	³¹ P NMR 2D Mapping of Creatine Kinase Forward Flux Rate in Hearts with Postinfarction Left Ventricular Remodeling in Response to Cell Therapy. <i>PLoS ONE</i> , 2016, 11, e0162149.	2.5	4
35	Cyclin D2-mediated cardiomyocyte cell cycle activity reverses doxorubicin-induced cardiotoxicity. <i>FASEB Journal</i> , 2013, 27, 1105.26.	0.5	0
36	The pivotal role of p53 in doxorubicin-induced acute versus chronic cardiotoxicity. <i>FASEB Journal</i> , 2013, 27, 528.2.	0.5	0

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37	Cell-Cycle-Based Strategies to Drive Myocardial Repair. <i>Pediatric Cardiology</i> , 2009, 30, 710-715.	1.3	12
38	Acute Doxorubicin Cardiotoxicity Is Associated With p53-Induced Inhibition of the Mammalian Target of Rapamycin Pathway. <i>Circulation</i> , 2009, 119, 99-106.	1.6	190
39	A Mouse Model for Juvenile Doxorubicin-Induced Cardiac Dysfunction. <i>Pediatric Research</i> , 2008, 64, 488-494.	2.3	61