

David A Elliott

List of Publications by Year in descending order

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Version: 2024-02-01

69
papers

6,252
citations

109321

35
h-index

102487

66
g-index

74
all docs

74
docs citations

74
times ranked

8379
citing authors

#	ARTICLE	IF	CITATIONS
1	Isolation and characterization of human embryonic stem cell-derived heart field-specific cardiomyocytes unravels new insights into their transcriptional and electrophysiological profiles. <i>Cardiovascular Research</i> , 2022, 118, 828-843.	3.8	14
2	3D-cardiomics: A spatial transcriptional atlas of the mammalian heart. <i>Journal of Molecular and Cellular Cardiology</i> , 2022, 163, 20-32.	1.9	16
3	Effect and application of cryopreserved three-dimensional microcardiac spheroids in myocardial infarction therapy. <i>Clinical and Translational Medicine</i> , 2022, 12, e721.	4.0	5
4	The Australia and New Zealand Cardio-Oncology Registry: evaluation of chemotherapy-related cardiotoxicity in a national cohort of paediatric cancer patients. <i>Internal Medicine Journal</i> , 2021, 51, 229-234.	0.8	6
5	Optimal Detection of Cardiac Sequelae. <i>JACC: CardioOncology</i> , 2021, 3, 154-156.	4.0	0
6	BET inhibition blocks inflammation-induced cardiac dysfunction and SARS-CoV-2 infection. <i>Cell</i> , 2021, 184, 2167-2182.e22.	28.9	131
7	Sex-Specific Control of Human Heart Maturation by the Progesterone Receptor. <i>Circulation</i> , 2021, 143, 1614-1628.	1.6	42
8	Modelling Mitochondrial Disease in Human Pluripotent Stem Cells: What Have We Learned?. <i>International Journal of Molecular Sciences</i> , 2021, 22, 7730.	4.1	14
9	Alpha-protein kinase 3 (<i>ALPK3</i>) truncating variants are a cause of autosomal dominant hypertrophic cardiomyopathy. <i>European Heart Journal</i> , 2021, 42, 3063-3073.	2.2	51
10	Exercise cardiovascular magnetic resonance reveals reduced cardiac reserve in pediatric cancer survivors with impaired cardiopulmonary fitness. <i>Journal of Cardiovascular Magnetic Resonance</i> , 2020, 22, 64.	3.3	22
11	β -catenin drives distinct transcriptional networks in proliferative and non-proliferative cardiomyocytes. <i>Development (Cambridge)</i> , 2020, 147, .	2.5	24
12	Evaluating anthracycline cardiotoxicity associated single nucleotide polymorphisms in a paediatric cohort with early onset cardiomyopathy. <i>Cardio-Oncology</i> , 2020, 6, 5.	1.7	6
13	The role of cardiac transcription factor NKX2-5 in regulating the human cardiac miRNAome. <i>Scientific Reports</i> , 2019, 9, 15928.	3.3	3
14	Drug Screening in Human PSC-Cardiac Organoids Identifies Pro-proliferative Compounds Acting via the Mevalonate Pathway. <i>Cell Stem Cell</i> , 2019, 24, 895-907.e6.	11.1	199
15	Pediatric Anthracycline-Induced Cardiotoxicity: Mechanisms, Pharmacogenomics, and Pluripotent Stem Cell Modeling. <i>Clinical Pharmacology and Therapeutics</i> , 2019, 105, 614-624.	4.7	30
16	Stem cell topography splits growth and homeostatic functions in the fish gill. <i>ELife</i> , 2019, 8, .	6.0	16
17	NKX2-5 regulates human cardiomyogenesis via a HEY2 dependent transcriptional network. <i>Nature Communications</i> , 2018, 9, 1373.	12.8	77
18	Systematic review of pharmacogenomics and adverse drug reactions in paediatric oncology patients. <i>Pediatric Blood and Cancer</i> , 2018, 65, e26937.	1.5	13

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19	Isolation and characterization of ventricular-like cells derived from NKX2-5 and MLC2v double knock-in human pluripotent stem cells. <i>Biochemical and Biophysical Research Communications</i> , 2018, 495, 1278-1284.	2.1	9
20	3D aggregate culture improves metabolic maturation of human pluripotent stem cell derived cardiomyocytes. <i>Biotechnology and Bioengineering</i> , 2018, 115, 630-644.	3.3	108
21	Coculturing with endothelial cells promotes in vitro maturation and electrical coupling of human embryonic stem cell-derived cardiomyocytes. <i>Journal of Heart and Lung Transplantation</i> , 2017, 36, 684-693.	0.6	29
22	Development of a human cardiac organoid injury model reveals innate regenerative potential. <i>Development (Cambridge)</i> , 2017, 144, 1118-1127.	2.5	127
23	Genetic determinants of anthracycline cardiotoxicity – ready for the clinic?. <i>British Journal of Clinical Pharmacology</i> , 2017, 83, 1141-1142.	2.4	10
24	Chemotherapy-related cardiotoxicity: are Australian practitioners missing the point?. <i>Internal Medicine Journal</i> , 2017, 47, 1166-1172.	0.8	6
25	Biomarkers of Human Pluripotent Stem Cell-Derived Cardiac Lineages. <i>Trends in Molecular Medicine</i> , 2017, 23, 651-668.	6.7	21
26	Functional screening in human cardiac organoids reveals a metabolic mechanism for cardiomyocyte cell cycle arrest. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, E8372-E8381.	7.1	361
27	ALPK3-deficient cardiomyocytes generated from patient-derived induced pluripotent stem cells and mutant human embryonic stem cells display abnormal calcium handling and establish that ALPK3 deficiency underlies familial cardiomyopathy. <i>European Heart Journal</i> , 2016, 37, 2586-2590.	2.2	49
28	GAPTrap: A Simple Expression System for Pluripotent Stem Cells and Their Derivatives. <i>Stem Cell Reports</i> , 2016, 7, 518-526.	4.8	27
29	Differentiation of human embryonic stem cells to HOXA+ hemogenic vasculature that resembles the aorta-gonad-mesonephros. <i>Nature Biotechnology</i> , 2016, 34, 1168-1179.	17.5	150
30	CD13 and ROR2 Permit Isolation of Highly Enriched Cardiac Mesoderm from Differentiating Human Embryonic Stem Cells. <i>Stem Cell Reports</i> , 2016, 6, 95-108.	4.8	30
31	Magnetic Resonance Imaging of Iron Oxide-Labeled Human Embryonic Stem Cell-Derived Cardiac Progenitors. <i>Stem Cells Translational Medicine</i> , 2016, 5, 67-74.	3.3	23
32	Atrial-like cardiomyocytes from human pluripotent stem cells are a robust preclinical model for assessing atrial-selective pharmacology. <i>EMBO Molecular Medicine</i> , 2015, 7, 394-410.	6.9	310
33	Cardiac Repair With a Novel Population of Mesenchymal Stem Cells Resident in the Human Heart. <i>Stem Cells</i> , 2015, 33, 3100-3113.	3.2	53
34	Cardiomyocyte differentiation of pluripotent stem cells with SB203580 analogues correlates with Wnt pathway CK1 inhibition independent of p38 MAPK signaling. <i>Journal of Molecular and Cellular Cardiology</i> , 2015, 80, 56-70.	1.9	18
35	Multipotent Caudal Neural Progenitors Derived from Human Pluripotent Stem Cells That Give Rise to Lineages of the Central and Peripheral Nervous System. <i>Stem Cells</i> , 2015, 33, 1759-1770.	3.2	80
36	Comparing mouse and human pluripotent stem cell derived cardiac cells: Both systems have advantages for pharmacological and toxicological screening. <i>Journal of Pharmacological and Toxicological Methods</i> , 2015, 74, 17-25.	0.7	2

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37	Dual Reporter <i><i>MESP1mCherry/w-NKX2-5eGFP/w</i></i> hESCs Enable Studying Early Human Cardiac Differentiation. <i>Stem Cells</i> , 2015, 33, 56-67.	3.2	65
38	Differentiation of Human Pluripotent Stem Cells to Cardiomyocytes Under Defined Conditions. <i>Methods in Molecular Biology</i> , 2014, 1353, 163-180.	0.9	48
39	Controlling Expansion and Cardiomyogenic Differentiation of Human Pluripotent Stem Cells in Scalable Suspension Culture. <i>Stem Cell Reports</i> , 2014, 3, 1132-1146.	4.8	189
40	SIRPA, VCAM1 and CD34 identify discrete lineages during early human cardiovascular development. <i>Stem Cell Research</i> , 2014, 13, 172-179.	0.7	63
41	Strategies for rapidly mapping proviral integration sites and assessing cardiogenic potential of nascent human induced pluripotent stem cell clones. <i>Experimental Cell Research</i> , 2014, 327, 297-306.	2.6	13
42	FOXN1GFP/w Reporter hESCs Enable Identification of Integrin- β 24, HLA-DR, and EpCAM as Markers of Human PSC-Derived FOXN1+ Thymic Epithelial Progenitors. <i>Stem Cell Reports</i> , 2014, 2, 925-937.	4.8	42
43	Cellular Reprogramming. <i>Circulation: Heart Failure</i> , 2013, 6, 1102-1107.	3.9	2
44	Isogenic human pluripotent stem cell pairs reveal the role of a KCNH2 mutation in long-QT syndrome. <i>EMBO Journal</i> , 2013, 32, 3161-3175.	7.8	174
45	PGC-1 β and Reactive Oxygen Species Regulate Human Embryonic Stem Cell-Derived Cardiomyocyte Function. <i>Stem Cell Reports</i> , 2013, 1, 560-574.	4.8	59
46	The use of agarose microwells for scalable embryoid body formation and cardiac differentiation of human and murine pluripotent stem cells. <i>Biomaterials</i> , 2013, 34, 2463-2471.	11.4	131
47	Human Embryonic Stem Cell Derived Mesenchymal Progenitors Express Cardiac Markers but Do Not Form Contractile Cardiomyocytes. <i>PLoS ONE</i> , 2013, 8, e54524.	2.5	26
48	Congenital Asplenia in Mice and Humans with Mutations in a Pbx/Nrx2-5/p15 Module. <i>Developmental Cell</i> , 2012, 22, 913-926.	7.0	70
49	Differentiation of Human Embryonic Stem Cells and Induced Pluripotent Stem Cells to Cardiomyocytes. <i>Circulation Research</i> , 2012, 111, 344-358.	4.5	641
50	INS GFP/w human embryonic stem cells facilitate isolation of in vitro derived insulin-producing cells. <i>Diabetologia</i> , 2012, 55, 694-706.	6.3	113
51	Analysis of Mitochondrial Function and Localisation during Human Embryonic Stem Cell Differentiation In Vitro. <i>PLoS ONE</i> , 2012, 7, e52214.	2.5	37
52	NKX2-5eGFP/w hESCs for isolation of human cardiac progenitors and cardiomyocytes. <i>Nature Methods</i> , 2011, 8, 1037-1040.	19.0	384
53	SIRPA is a specific cell-surface marker for isolating cardiomyocytes derived from human pluripotent stem cells. <i>Nature Biotechnology</i> , 2011, 29, 1011-1018.	17.5	500
54	A Targeted <i><i>NKX2.1</i></i> Human Embryonic Stem Cell Reporter Line Enables Identification of Human Basal Forebrain Derivatives. <i>Stem Cells</i> , 2011, 29, 462-473.	3.2	99

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55	NK-2 Class Homeodomain Proteins. , 2010, , 569-597.		10
56	Elastomeric nanocomposites as cell delivery vehicles and cardiac support devices. <i>Soft Matter</i> , 2010, 6, 4715.	2.7	65
57	The GAL4 System. <i>Methods in Molecular Biology</i> , 2008, 420, 79-95.	0.9	120
58	The GAL4 System: A Versatile Toolkit for Gene Expression in <i>Drosophila</i> . <i>Cold Spring Harbor Protocols</i> , 2008, 2008, pdb.top49.	0.3	22
59	Time to mend a broken heart. <i>Stem Cell Research</i> , 2007, 1, 4-6.	0.7	2
60	A tyrosine-rich domain within homeodomain transcription factor Nkx2-5 is an essential element in the early cardiac transcriptional regulatory machinery. <i>Development (Cambridge)</i> , 2006, 133, 1311-1322.	2.5	28
61	Transcriptional regulation of the murine promoter by cardiac factors Nkx2-5, GATA4 and Tbx5. <i>Cardiovascular Research</i> , 2004, 64, 402-411.	3.8	91
62	Independent Regulation of Synaptic Size and Activity by the Anaphase-Promoting Complex. <i>Cell</i> , 2004, 119, 707-718.	28.9	214
63	Cardiac homeobox gene NKX2-5 mutations and congenital heart disease. <i>Journal of the American College of Cardiology</i> , 2003, 41, 2072-2076.	2.8	231
64	Cardiac T-box factor Tbx20 directly interacts with Nkx2-5, GATA4, and GATA5 in regulation of gene expression in the developing heart. <i>Developmental Biology</i> , 2003, 262, 206-224.	2.0	260
65	Developmental paradigms in heart disease: insights from tinman. <i>Annals of Medicine</i> , 2002, 34, 148-156.	3.8	39
66	Developmental paradigms in heart disease: insights from tinman. <i>Annals of Medicine</i> , 2002, 34, 148-156.	3.8	21
67	Homeodomain Factor Nkx2-5 in Heart Development and Disease. <i>Cold Spring Harbor Symposia on Quantitative Biology</i> , 2002, 67, 107-114.	1.1	67
68	Cardiac Septal and Valvular Dymorphogenesis in Mice Heterozygous for Mutations in the Homeobox Gene <i>Nkx2-5</i> . <i>Circulation Research</i> , 2000, 87, 888-895.	4.5	325
69	Transcriptional Control and Pattern Formation in the Developing Vertebrate Heart. , 1999, , 111-129.		16