List of Publications by Year in descending order

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	94433	102487
4,994	37	66
citations	h-index	g-index
123	123	7640
docs citations	times ranked	citing authors
	citations 123	4,99437citationsh-index123123

#	Article	IF	CITATIONS
1	CYP1A2 polymorphisms modify the association of habitual coffee consumption with appetite, macronutrient intake, and body mass index: results from an observational cohort and a cross-over randomized study. International Journal of Obesity, 2022, 46, 162-168.	3.4	10
2	Editorial: Obesity, metabolic phenotypes and COVID-19. Metabolism: Clinical and Experimental, 2022, 128, 155121.	3.4	20
3	Reconstituted HDL-apoE3 promotes endothelial cell migration through ID1 and its downstream kinases ERK1/2, AKT and p38 MAPK. Metabolism: Clinical and Experimental, 2022, 127, 154954.	3.4	12
4	A junctional cAMP compartment regulates rapid Ca2+ signaling in atrial myocytes. Journal of Molecular and Cellular Cardiology, 2022, 165, 141-157.	1.9	6
5	Genetically-Guided Medical Nutrition Therapy in Type 2 Diabetes Mellitus and Pre-diabetes: A Series of n-of-1 Superiority Trials. Frontiers in Nutrition, 2022, 9, 772243.	3.7	9
6	Darling: A Web Application for Detecting Disease-Related Biomedical Entity Associations with Literature Mining. Biomolecules, 2022, 12, 520.	4.0	9
7	The Challenge and Importance of Integrating Drug–Nutrient–Genome Interactions in Personalized Cardiovascular Healthcare. Journal of Personalized Medicine, 2022, 12, 513.	2.5	3
8	Nonalcoholic Fatty Liver Disease and Cardiovascular Disease: a Review of Shared Cardiometabolic Risk Factors. Hypertension, 2022, 79, 1319-1326.	2.7	50
9	Epitranscriptomic challenges and promises in metabolic diseases. Metabolism: Clinical and Experimental, 2022, 132, 155219.	3.4	9
10	Aberrant PLN-R14del Protein Interactions Intensify SERCA2a Inhibition, Driving Impaired Ca2+ Handling and Arrhythmogenesis. International Journal of Molecular Sciences, 2022, 23, 6947.	4.1	11
11	Structure–function analysis of naturally occurring apolipoprotein A-I L144R, A164S and L178P mutants provides insight on their role on HDL levels and cardiovascular risk. Cellular and Molecular Life Sciences, 2021, 78, 1523-1544.	5.4	8
12	Intracoronary Administration of Allogeneic Cardiosphere-Derived Cells Immediately Prior to Reperfusion in Pigs With Acute Myocardial Infarction Reduces Infarct Size and Attenuates Adverse Cardiac Remodeling. Journal of Cardiovascular Pharmacology and Therapeutics, 2021, 26, 88-99.	2.0	3
13	Advances in biological therapies for dyslipidemias and atherosclerosis. Metabolism: Clinical and Experimental, 2021, 116, 154461.	3.4	41
14	COVID-19 enters the expanding network of apolipoprotein E4-related pathologies. Redox Biology, 2021, 41, 101938.	9.0	31
15	Impaired Right Ventricular Calcium Cycling Is an Early Risk Factor in R14del-Phospholamban Arrhythmias. Journal of Personalized Medicine, 2021, 11, 502.	2.5	12
16	Genotype-guided dietary supplementation in precision nutrition. Nutrition Reviews, 2021, 79, 1225-1235.	5.8	10
17	Epitranscriptomics of cardiovascular diseases (Review). International Journal of Molecular Medicine, 2021, 49, .	4.0	9
18	Precision Medicine in Aortic Anastomosis: A Numerical and Experimental Study of a Novel Double-Sided Needle. Journal of Personalized Medicine, 2021, 11, 1385.	2.5	2

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19	Levosimendan prevents doxorubicin-induced cardiotoxicity in time- and dose-dependent manner: implications for inotropy. Cardiovascular Research, 2020, 116, 576-591.	3.8	32
20	The "Virtual Digital Twins―Concept in Precision Nutrition. Advances in Nutrition, 2020, 11, 1405-1413.	6.4	48
21	The Cardioprotective PKA-Mediated Hsp20 Phosphorylation Modulates Protein Associations Regulating Cytoskeletal Dynamics. International Journal of Molecular Sciences, 2020, 21, 9572.	4.1	9
22	Genome-wide transcriptomics identifies an early preclinical signature of prion infection. PLoS Pathogens, 2020, 16, e1008653.	4.7	40
23	Genome-wide transcriptomics identifies an early preclinical signature of prion infection. , 2020, 16, e1008653.		Ο
24	Genome-wide transcriptomics identifies an early preclinical signature of prion infection. , 2020, 16, e1008653.		0
25	Cardiosphere-Derived Cells Attenuate Inflammation, Preserve Systolic Function, and Prevent Adverse Remodeling in Rat Hearts With Experimental Autoimmune Myocarditis. Journal of Cardiovascular Pharmacology and Therapeutics, 2019, 24, 70-77.	2.0	19
26	Personalized Assessment of the Coronary Atherosclerotic Arteries by Intravascular Ultrasound Imaging: Hunting the Vulnerable Plaque. Journal of Personalized Medicine, 2019, 9, 8.	2.5	12
27	The Crocus sativus Compounds trans-Crocin 4 and trans-Crocetin Modulate the Amyloidogenic Pathway and Tau Misprocessing in Alzheimer Disease Neuronal Cell Culture Models. Frontiers in Neuroscience, 2019, 13, 249.	2.8	42
28	The future of apolipoprotein E mimetic peptides in the prevention of cardiovascular disease. Current Opinion in Lipidology, 2019, 30, 326-341.	2.7	10
29	Prevalence and cardiac phenotype of patients with aÂphospholamban mutation. Netherlands Heart Journal, 2019, 27, 64-69.	0.8	52
30	Regulation of BECN1-mediated autophagy by HSPB6: Insights from a human HSPB6 ^{S10F} mutant. Autophagy, 2018, 14, 80-97.	9.1	27
31	HAX-1 regulates SERCA2a oxidation and degradation. Journal of Molecular and Cellular Cardiology, 2018, 114, 220-233.	1.9	20
32	Current and Emerging Reconstituted HDL-apoA-I and HDL-apoE Approaches to Treat Atherosclerosis. Journal of Personalized Medicine, 2018, 8, 34.	2.5	23
33	The Histidine-Rich Calcium Binding Protein in Regulation of Cardiac Rhythmicity. Frontiers in Physiology, 2018, 9, 1379.	2.8	12
34	Glial responses during epileptogenesis in Mus musculus point to potential therapeutic targets. PLoS ONE, 2018, 13, e0201742.	2.5	24
35	HDL-apoA-I induces the expression of angiopoietin like 4 (ANGPTL4) in endothelial cells via a PI3K/AKT/FOXO1 signaling pathway. Metabolism: Clinical and Experimental, 2018, 87, 36-47.	3.4	21
36	The march of pluripotent stem cells in cardiovascular regenerative medicine. Stem Cell Research and Therapy, 2018, 9, 201.	5.5	32

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37	Beneficial Effects of Sideritis scardica and Cichorium spinosum against Amyloidogenic Pathway and Tau Misprocessing in Alzheimer's Disease Neuronal Cell Culture Models. Journal of Alzheimer's Disease, 2018, 64, 787-800.	2.6	12
38	Protein aggregation and neurodegeneration in prototypical neurodegenerative diseases: Examples of amyloidopathies, tauopathies and synucleinopathies. Progress in Neurobiology, 2017, 155, 171-193.	5.7	137
39	Muscle Lim Protein and myosin binding protein C form a complex regulating muscle differentiation. Biochimica Et Biophysica Acta - Molecular Cell Research, 2017, 1864, 2308-2321.	4.1	7
40	Phosphorylation of serine96 of histidine-rich calcium-binding protein by the Fam20C kinase functions to prevent cardiac arrhythmia. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, 9098-9103.	7.1	43
41	NADPH oxidases as drug targets and biomarkers in neurodegenerative diseases: What is the evidence?. Free Radical Biology and Medicine, 2017, 112, 387-396.	2.9	88
42	Impaired calcium homeostasis is associated with sudden cardiac death and arrhythmias in a genetic equivalent mouse model of the human HRC-Ser96Ala variant. Cardiovascular Research, 2017, 113, 1403-1417.	3.8	14
43	Strictly co-isogenic C57BL/6J- <i>Prnp</i> â^'/â^' mice: A rigorous resource for prion science. Journal of Experimental Medicine, 2016, 213, 313-327.	8.5	98
44	Ciliary neurotrophic factor upregulates follistatin and Pak1, causes overexpression of muscle differentiation related genes and downregulation of established atrophy mediators in skeletal muscle. Metabolism: Clinical and Experimental, 2016, 65, 915-925.	3.4	16
45	Strictly co-isogenic C57BL/6J-Prnpâ^'/â^'mice: A rigorous resource for prion science. Journal of Cell Biology, 2016, 212, 2126OIA42.	5.2	0
46	Genetic modifiers to the PLN L39X mutation in a patient with DCM and sustained ventricular tachycardia?. Global Cardiology Science & Practice, 2015, 2015, 29.	0.4	6
47	Correction of human phospholamban R14del mutation associated with cardiomyopathy using targeted nucleases and combination therapy. Nature Communications, 2015, 6, 6955.	12.8	155
48	Muscle LIM Protein: Master regulator of cardiac and skeletal muscle functions. Gene, 2015, 566, 1-7.	2.2	65
49	Cardioprotection by H2S engages a cGMP-dependent protein kinase G/phospholamban pathway. Cardiovascular Research, 2015, 106, 432-442.	3.8	72
50	Calcium/calmodulin-dependent protein kinase II (CaMKII) inhibition ameliorates arrhythmias elicited by junctin ablation under stress conditions. Heart Rhythm, 2015, 12, 1599-1610.	0.7	11
51	Prion Infections and Anti-PrP Antibodies Trigger Converging Neurotoxic Pathways. PLoS Pathogens, 2015, 11, e1004662.	4.7	76
52	Human G109E-inhibitor-1 impairs cardiac function and promotes arrhythmias. Journal of Molecular and Cellular Cardiology, 2015, 89, 349-359.	1.9	12
53	Outpatient management of chronic heart failure. Expert Opinion on Pharmacotherapy, 2015, 16, 17-41.	1.8	3
54	Natural Products as Sources of Potential Antiamyloidogenic Agents. GSTF Journal of Advances in Medical Research, 2015, 1, .	0.0	3

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55	Calcium Cycling Circuits in Cardiac Physiology and Pathophysiology. , 2015, , 205-215.		1
56	Pharmacogenomics in the Development and Characterization of Atheroprotective Drugs. Methods in Molecular Biology, 2014, 1175, 259-300.	0.9	5
57	MicroRNAs in heart failure: Small molecules with major impact. Global Cardiology Science & Practice, 2014, 2014, 30.	0.4	32
58	Forced swim test induces divergent global transcriptomic alterations in the hippocampus of high versus low novelty-seeker rats. Human Genomics, 2014, 8, 4.	2.9	8
59	Muscle lim protein isoform negatively regulates striated muscle actin dynamics and differentiation. FEBS Journal, 2014, 281, 3261-3279.	4.7	26
60	Reprogramming of the MicroRNA Transcriptome Mediates Resistance to Rapamycin. Journal of Biological Chemistry, 2013, 288, 6034-6044.	3.4	41
61	Pharmacogenetic considerations for late life depression therapy. Expert Opinion on Drug Metabolism and Toxicology, 2013, 9, 989-999.	3.3	9
62	Science and practice of arrhythmogenic cardiomyopathy: A paradigm shift. Global Cardiology Science & Practice, 2013, 2013, 8.	0.4	4
63	Identification of a Protein Phosphatase-1/Phospholamban Complex That Is Regulated by cAMP-Dependent Phosphorylation. PLoS ONE, 2013, 8, e80867.	2.5	13
64	Rapamycin Resistance Is Linked to Defective Regulation of Skp2. Cancer Research, 2012, 72, 1836-1843.	0.9	38
65	Molecular genetics made simple. Global Cardiology Science & Practice, 2012, 2012, 6.	0.4	6
66	The expanding role of epigenetics. Global Cardiology Science & Practice, 2012, 2012, 7.	0.4	12
67	Regulation of adverse remodelling by osteopontin in a genetic heart failure model. European Heart Journal, 2012, 33, 1954-1963.	2.2	80
68	Array-based pharmacogenomics of molecular-targeted therapies in oncology. Pharmacogenomics Journal, 2012, 12, 185-196.	2.0	13
69	Combining multiple hypothesis testing and affinity propagation clustering leads to accurate, robust and sample size independent classification on gene expression data. BMC Bioinformatics, 2012, 13, 270.	2.6	11
70	Histidine-rich calcium binding protein: The new regulator of sarcoplasmic reticulum calcium cycling. Journal of Molecular and Cellular Cardiology, 2011, 50, 43-49.	1.9	53
71	Investigating the Minimum Required Number of Genes for the Classification of Neuromuscular Disease Microarray Data. IEEE Transactions on Information Technology in Biomedicine, 2011, 15, 349-355.	3.2	6
72	Small Heat Shock Protein 20 Interacts With Protein Phosphatase-1 and Enhances Sarcoplasmic Reticulum Calcium Cycling. Circulation Research, 2011, 108, 1429-1438.	4.5	67

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73	SERCA2a superinhibition by human phospholamban triggers electrical and structural remodeling in mouse hearts. Physiological Genomics, 2011, 43, 357-364.	2.3	11
74	In Silico Dynamic Molecular Interaction Networks for the Discovery of New Therapeutic Targets. Current Pharmaceutical Design, 2010, 16, 2241-2251.	1.9	6
75	Pharmacogenetically Tailored Treatments for Heart Disease. Current Pharmaceutical Design, 2010, 16, 2194-2213.	1.9	6
76	Editorial [Hot Topic: Pharmacogenomics: Achievements, Challenges and Prospects, for Patients, Pharmaceutical Industries and Healthcare Systems (Guest Editor: Despina Sanoudou)]. Current Pharmaceutical Design, 2010, 16, 2182-2183.	1.9	3
77	Histopathologic and genetic alterations as predictors of response to treatment and survival in lung cancer: A review of published data. Critical Reviews in Oncology/Hematology, 2010, 75, 94-109.	4.4	38
78	Role of Esrrg in the fibrate-mediated regulation of lipid metabolism genes in human ApoA-I transgenic mice. Pharmacogenomics Journal, 2010, 10, 165-179.	2.0	16
79	A comparison of batch effect removal methods for enhancement of prediction performance using MAQC-II microarray gene expression data. Pharmacogenomics Journal, 2010, 10, 278-291.	2.0	249
80	Clinical Pharmacogenetics in Oncology: the Paradigm of Molecular Targeted Therapies. Current Pharmaceutical Design, 2010, 16, 2184-2193.	1.9	10
81	Regulation of ApoA-I Gene Expression and Prospects to Increase Plasma ApoA-I and HDL Levels. , 2010, , 15-24.		1
82	Muscle Lim Protein Interacts with Cofilin 2 and Regulates F-Actin Dynamics in Cardiac and Skeletal Muscle. Molecular and Cellular Biology, 2009, 29, 6046-6058.	2.3	51
83	T-tubule disorganization and defective excitation-contraction coupling in muscle fibers lacking myotubularin lipid phosphatase. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 18763-18768.	7.1	167
84	The Anti-apoptotic Protein HAX-1 Interacts with SERCA2 and Regulates Its Protein Levels to Promote Cell Survival. Molecular Biology of the Cell, 2009, 20, 306-318.	2.1	106
85	The role of SERCA2a/PLN complex, Ca2+ homeostasis, and anti-apoptotic proteins in determining cell fate. Pflugers Archiv European Journal of Physiology, 2009, 457, 687-700.	2.8	37
86	G.P.12.02 T-tubule disorganisation and defective excitation–contraction coupling in muscle fibres lacking myotubularin lipid phosphatase. Neuromuscular Disorders, 2009, 19, 635-636.	0.6	0
87	Genes and Gene–Environment Interactions in the Pathogenesis of Obesity and the Metabolic Syndrome. , 2009, , 11-39.		0
88	Investigating the minimum required number of genes for optimum classification of myopathy microarray data. , 2009, , .		0
89	Search for Potential Markers for Prostate Cancer Diagnosis, Prognosis and Treatment in Clinical Tissue Specimens Using Amine-Specific Isobaric Tagging (iTRAQ) with Two-Dimensional Liquid Chromatography and Tandem Mass Spectrometry. Journal of Proteome Research, 2008, 7, 3146-3158.	3.7	92
90	The Ser96Ala variant in histidine-rich calcium-binding protein is associated with life-threatening ventricular arrhythmias in idiopathic dilated cardiomyopathy. European Heart Journal, 2008, 29, 2514-2525.	2.2	48

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91	Histidine-rich Ca-binding protein interacts with sarcoplasmic reticulum Ca-ATPase. American Journal of Physiology - Heart and Circulatory Physiology, 2007, 293, H1581-H1589.	3.2	75
92	A Dominant Negative Form of the Transcription Factor c-Jun Affects Genes That Have Opposing Effects on Lipid Homeostasis in Mice. Journal of Biological Chemistry, 2007, 282, 19556-19564.	3.4	23
93	C.P.1.10 Molecular mechanisms underlying X-linked myotubular myopathy. Neuromuscular Disorders, 2007, 17, 836-837.	0.6	0
94	Phospholamban Interacts with HAX-1, a Mitochondrial Protein with Anti-apoptotic Function. Journal of Molecular Biology, 2007, 367, 65-79.	4.2	85
95	Defective Ribosomal Protein Gene Expression Alters Transcription, Translation, Apoptosis, and Oncogenic Pathways in Diamond-Blackfan Anemia. Stem Cells, 2006, 24, 2034-2044.	3.2	75
96	Skeletal muscle repair in a mouse model of nemaline myopathy. Human Molecular Genetics, 2006, 15, 2603-2612.	2.9	44
97	Genetics of Obesity and Diabetes. , 2006, , 39-67.		0
98	Interferon-α/β-mediated innate immune mechanisms in dermatomyositis. Annals of Neurology, 2005, 57, 664-678.	5.3	530
99	Variations in gene expression among different types of human skeletal muscle. Muscle and Nerve, 2005, 32, 483-491.	2.2	28
100	The influence of muscle type and dystrophin deficiency on murine expression profiles. Mammalian Genome, 2005, 16, 739-748.	2.2	35
101	Array lessons from the heart: focus on the genome and transcriptome of cardiomyopathies. Physiological Genomics, 2005, 21, 131-143.	2.3	34
102	Evidence by molecular profiling for a placental origin of infantile hemangioma. Proceedings of the National Academy of Sciences of the United States of America, 2005, 102, 19097-19102.	7.1	170
103	Side Population cells isolated from different tissues share transcriptome signatures and express tissue-specific markers. Experimental Cell Research, 2005, 303, 360-374.	2.6	45
104	Defective Ribosomal Protein Gene Expression Alters Transcription, Translation and Oncogenic Pathways in Diamond-Blackfan Anemia Blood, 2005, 106, 3546-3546.	1.4	5
105	Transcriptional profile of postmortem skeletal muscle. Physiological Genomics, 2004, 16, 222-228.	2.3	38
106	Expression profiling and identification of novel genes involved in myogenic differentiation. FASEB Journal, 2004, 18, 1-23.	0.5	157
107	Molecular classification of nemaline myopathies: "nontyping―specimens exhibit unique patterns of gene expression. Neurobiology of Disease, 2004, 15, 590-600.	4.4	13
108	Gene Expression Changes in Bone Marrow Cells from Diamond-Blackfan Anemia Patients Blood, 2004, 104, 720-720.	1.4	8

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109	Gene expression profiling of Duchenne muscular dystrophy skeletal muscle. Neurogenetics, 2003, 4, 163-171.	1.4	82
110	Reproducibility of gene expression across generations of Affymetrix microarrays. BMC Bioinformatics, 2003, 4, 27.	2.6	67
111	Expression profiling reveals altered satellite cell numbers and glycolytic enzyme transcription in nemaline myopathy muscle. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 4666-4671.	7.1	68
112	Structural Genomic Abnormalities of Chromosomes 9 and 18 in Myxopapillary Ependymomas. Journal of Neuropathology and Experimental Neurology, 2003, 62, 927-935.	1.7	15
113	Gene expression comparison of biopsies from Duchenne muscular dystrophy (DMD) and normal skeletal muscle. Proceedings of the National Academy of Sciences of the United States of America, 2002, 99, 15000-15005.	7.1	312
114	Clinical and genetic heterogeneity in nemaline myopathy – a disease of skeletal muscle thin filaments. Trends in Molecular Medicine, 2001, 7, 362-368.	6.7	145
115	Interstitial colocalization of two cervid satellite DNAs involved in the genesis of the Indian muntjac karyotype. Chromosome Research, 2000, 8, 363-373.	2.2	39
116	Analysis of pilocytic astrocytoma by comparative genomic hybridization. British Journal of Cancer, 2000, 82, 1218-1222.	6.4	96
117	Acquired Rearrangement of an Amplified Epidermal Growth Factor Receptor (EGFR) Gene in a Human Glioblastoma Xenograft. Journal of Neuropathology and Experimental Neurology, 1999, 58, 697-701.	1.7	12
118	Chromosome specific comparative genome hybridisation for determining the origin of intrachromosomal duplications Journal of Medical Genetics, 1998, 35, 37-41.	3.2	23
119	Neuropharmacogenetics of Major Depression: Has the Time Come to Take both Sexes into Account?. , 0, , .		0
120	Continuous glucose monitoring and hypoglycaemia events: unmet needs. Diabetologia, 0, , .	6.3	1