Jian-Yuan Zhao

List of Publications by Year in descending order

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361296 345118 1,544 45 20 36 citations h-index g-index papers 47 47 47 1736 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Biallelic DNAH9 mutations are identified in Chinese patients with defective left–right patterning and cilia-related complex congenital heart disease. Human Genetics, 2022, 141, 1339-1353.	1.8	7
2	Fructose-1,6-bisphosphate prevents pregnancy loss by inducing decidual COX-2 ⁺ macrophage differentiation. Science Advances, 2022, 8, eabj2488.	4.7	22
3	A defective lysophosphatidic acid-autophagy axis increases miscarriage risk by restricting decidual macrophage residence. Autophagy, 2022, 18, 2459-2480.	4.3	26
4	Gestational Leucylation Suppresses Embryonic Tâ€Box Transcription Factor 5 Signal and Causes Congenital Heart Disease. Advanced Science, 2022, 9, e2201034.	5.6	38
5	Acox2 is a regulator of lysine crotonylation that mediates hepatic metabolic homeostasis in mice. Cell Death and Disease, 2022, 13, 279.	2.7	12
6	Methylene-bridge tryptophan fatty acylation regulates PI3K-AKT signaling and glucose uptake. Cell Reports, 2022, 38, 110509.	2.9	5
7	A proteogenomic analysis of clear cell renal cell carcinoma in a Chinese population. Nature Communications, 2022, 13, 2052.	5.8	48
8	Nicotinamide Mononucleotide Alleviates Cardiomyopathy Phenotypes Caused by Short-Chain Enoyl-Coa Hydratase 1 Deficiency. JACC Basic To Translational Science, 2022, 7, 348-362.	1.9	32
9	Master microRNA-222 regulates cardiac microRNA maturation and triggers Tetralogy of Fallot. Signal Transduction and Targeted Therapy, 2022, 7, .	7.1	2
10	Increased expression of tribbles homolog 3 predicts poor prognosis and correlates with tumor immunity in clear cell renal cell carcinoma: a bioinformatics study. Bioengineered, 2022, 13, 14000-14012.	1.4	6
11	Rapamycin prevents spontaneous abortion by triggering decidual stromal cell autophagy-mediated NK cell residence. Autophagy, 2021, 17, 2511-2527.	4.3	65
12	Two sides of NNMT in alcoholic and non-alcoholic fatty liver development. Journal of Hepatology, 2021, 74, 1250-1253.	1.8	11
13	Ketogenic diets inhibit mitochondrial biogenesis and induce cardiac fibrosis. Signal Transduction and Targeted Therapy, 2021, 6, 54.	7.1	91
14	SINO Syndrome Causative KIDINS220/ARMS Gene Regulates Adipocyte Differentiation. Frontiers in Cell and Developmental Biology, 2021, 9, 619475.	1.8	5
15	Low chorionic villous succinate accumulation associates with recurrent spontaneous abortion risk. Nature Communications, 2021, 12, 3428.	5.8	76
16	Nuclear dihydroxyacetone phosphate signals nutrient sufficiency and cell cycle phase to global histone acetylation. Nature Metabolism, 2021, 3, 859-875.	5.1	23
17	Mitochondrial STAT5A promotes metabolic remodeling and the Warburg effect by inactivating the pyruvate dehydrogenase complex. Cell Death and Disease, 2021, 12, 634.	2.7	13
18	High maternal blood lipid levels during early pregnancy are associated with increased risk of congenital heart disease in offspring. Acta Obstetricia Et Gynecologica Scandinavica, 2021, 100, 1806-1813.	1.3	9

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19	Transmission of a Novel Imprinting Center Deletion Associated With Prader–Willi Syndrome Through Three Generations of a Chinese Family: Case Presentation, Differential Diagnosis, and a Lesson Worth Thinking About. Frontiers in Genetics, 2021, 12, 630650.	1.1	1
20	Calcineurin inactivation inhibits pyruvate dehydrogenase complex activity and induces the Warburg effect. Oncogene, 2021, 40, 6692-6702.	2.6	3
21	Homocysteine inhibits pro-insulin receptor cleavage and causes insulin resistance via protein cysteine-homocysteinylation. Cell Reports, 2021, 37, 109821.	2.9	104
22	Adenylate cyclaseâ€activating polypeptide 1 gene methylation predicts prognosis and the immune microenvironment of bladder cancer. Clinical and Translational Medicine, 2021, 11, e597.	1.7	1
23	Inactivation of the AMPK–GATA3–ECHS1 Pathway Induces Fatty Acid Synthesis That Promotes Clear Cell Renal Cell Carcinoma Growth. Cancer Research, 2020, 80, 319-333.	0.4	90
24	Inhibiting MARSs reduces hyperhomocysteinemiaâ€associated neural tube and congenital heart defects. EMBO Molecular Medicine, 2020, 12, e9469.	3.3	21
25	APC/CCDH1 synchronizes ribose-5-phosphate levels and DNA synthesis to cell cycle progression. Nature Communications, 2019, 10, 2502.	5.8	44
26	Bi-allelic Mutations in TTC21A Induce Asthenoteratospermia in Humans and Mice. American Journal of Human Genetics, 2019, 104, 738-748.	2.6	103
27	Elevated CD36 expression correlates with increased visceral adipose tissue and predicts poor prognosis in ccRCC patients. Journal of Cancer, 2019, 10, 4522-4531.	1.2	29
28	The Prognostic Value of Programmed Death-Ligand 1 in a Chinese Cohort With Clear Cell Renal Cell Carcinoma. Frontiers in Oncology, 2019, 9, 879.	1.3	6
29	Susceptibility to congenital heart defects associated with a polymorphism in TBX2 3′ untranslated region in the Han Chinese population. Pediatric Research, 2019, 85, 378-383.	1.1	4
30	Sensing and Transmitting Intracellular Amino Acid Signals through Reversible Lysine Aminoacylations. Cell Metabolism, 2018, 27, 151-166.e6.	7.2	97
31	High expression of F2RL3 correlates with aggressive features and poor survival in clear cell renal cell carcinoma. Journal of Cancer, 2018, 9, 3400-3406.	1.2	1
32	Colonic Lysine Homocysteinylation Induced by High-Fat Diet Suppresses DNA Damage Repair. Cell Reports, 2018, 25, 398-412.e6.	2.9	70
33	Elevated H3K79 homocysteinylation causes abnormal gene expression during neural development and subsequent neural tube defects. Nature Communications, 2018, 9, 3436.	5.8	56
34	PD-L1 expression in Xp11.2 translocation renal cell carcinoma: Indicator of tumor aggressiveness. Scientific Reports, 2017, 7, 2074.	1.6	21
35	Lower Circulating Folate Induced by a Fidgetin Intronic Variant Is Associated With Reduced Congenital Heart Disease Susceptibility. Circulation, 2017, 135, 1733-1748.	1.6	50
36	A TBX5 3â€2UTR variant increases the risk of congenital heart disease in the Han Chinese population. Cell Discovery, 2017, 3, 17026.	3.1	23

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37	Functional variants of the 5-methyltetrahydrofolate-homocysteine methyltransferase gene significantly increase susceptibility to prostate cancer: Results from an ethnic Han Chinese population. Scientific Reports, 2016, 6, 36264.	1.6	12
38	MTHFR c.677C>T Inhibits Cell Proliferation and Decreases Prostate Cancer Susceptibility in the Han Chinese Population in Shanghai. Scientific Reports, 2016, 6, 36290.	1.6	7
39	The emergence of intracellular metabolite signaling networks. IUBMB Life, 2016, 68, 871-872.	1.5	0
40	NADP+-IDH Mutations Promote Hypersuccinylation that Impairs Mitochondria Respiration and Induces Apoptosis Resistance. Molecular Cell, 2015, 60, 661-675.	4.5	175
41	A Genetic Variant in Vitamin B12 Metabolic Genes That Reduces the Risk of Congenital Heart Disease in Han Chinese Populations. PLoS ONE, 2014, 9, e88332.	1.1	10
42	Genetic variants reducing MTR gene expression increase the risk of congenital heart disease in Han Chinese populations. European Heart Journal, 2014, 35, 733-742.	1.0	31
43	A functional variant in the cystathionine \hat{l}^2 -synthase gene promoter significantly reduces congenital heart disease susceptibility in a Han Chinese population. Cell Research, 2013, 23, 242-253.	5.7	27
44	Functional Variant in Methionine Synthase Reductase Intron-1 Significantly Increases the Risk of Congenital Heart Disease in the Han Chinese Population. Circulation, 2012, 125, 482-490.	1.6	57
45	Genetic Polymorphisms of the TYMS Gene Are Not Associated with Congenital Cardiac Septal Defects in a Han Chinese Population. PLoS ONE, 2012, 7, e31644.	1.1	9