

Li Xie

List of Publications by Year in descending order

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Version: 2024-02-01

26
papers

296
citations

1039406

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940134

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27
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27
docs citations

27
times ranked

413
citing authors

#	ARTICLE	IF	CITATIONS
1	Case Report: A Young Man With Giant Pericardial Synovial Sarcoma. <i>Frontiers in Cardiovascular Medicine</i> , 2022, 9, 829328.	1.1	2
2	Whole-Exome Sequencing Identifies a Novel Variant (c.1538T > C) of TNNI3K in Arrhythmogenic Right Ventricular Cardiomyopathy. <i>Frontiers in Cardiovascular Medicine</i> , 2022, 9, 843837.	1.1	3
3	Surgery for Primary Cardiac Tumors in Children: Successful Management of Large Fibromas. <i>Frontiers in Cardiovascular Medicine</i> , 2022, 9, 808394.	1.1	4
4	FOXP3 Contributes to TMZ Resistance, Prognosis, and Immune Infiltration in GBM from a Novel Pyroptosis-Associated Risk Signature. <i>Disease Markers</i> , 2022, 2022, 1-21.	0.6	6
5	Hand-sewn expanded polytetrafluoroethylene valved conduit for right ventricular outflow tract reconstruction. <i>Journal of Central South University (Medical Sciences)</i> , 2022, 47, 94-100.	0.1	1
6	Clinical exome sequencing reveals a mutation in <i>PDHA1</i> in Leigh syndrome: A case of a Chinese boy with lethal neuropathy. <i>Molecular Genetics & Genomic Medicine</i> , 2021, 9, e1651.	0.6	2
7	Comprehensive analysis of lncRNA biomarkers in kidney renal clear cell carcinoma by lncRNA-mediated ceRNA network. <i>PLoS ONE</i> , 2021, 16, e0252452.	1.1	4
8	The roles of GTPase-activating proteins in regulated cell death and tumor immunity. <i>Journal of Hematology and Oncology</i> , 2021, 14, 171.	6.9	17
9	Identification and Integrate Analysis of Key Biomarkers for Diagnosis and Prognosis of Non-Small Cell Lung Cancer Based on Bioinformatics Analysis. <i>Technology in Cancer Research and Treatment</i> , 2021, 20, 153303382110602.	0.8	17
10	Close interactions between lncRNAs, lipid metabolism and ferroptosis in cancer. <i>International Journal of Biological Sciences</i> , 2021, 17, 4493-4513.	2.6	29
11	Establishment of a Dihydrofolate Reductase Gene Knock-In Zebrafish Strain to Aid Preliminary Analysis of Congenital Heart Disease Mechanisms. <i>Frontiers in Cardiovascular Medicine</i> , 2021, 8, 763851.	1.1	3
12	Modified Maxillary-Swing Approach for Resection of Primary Malignancies in the Pterygopalatine Fossa. <i>Frontiers in Oncology</i> , 2020, 10, 530381.	1.3	2
13	Partial Resection of a Huge Left Ventricle Cardiac Fibroma in an Asymptomatic Child. <i>Annals of Thoracic Surgery</i> , 2019, 108, e393-e395.	0.7	3
14	In-Continuity Neck Dissection: Long-Term Oncological Outcomes in Squamous Cell Carcinoma of the Buccal Mucosa. <i>Journal of Oral and Maxillofacial Surgery</i> , 2018, 76, 1107-1116.	0.5	0
15	Repression of Noxa by Bmi1 contributes to deguelin-induced apoptosis in non-small cell lung cancer cells. <i>Journal of Cellular and Molecular Medicine</i> , 2018, 22, 6213-6227.	1.6	29
16	Deguelin attenuates non-small cell lung cancer cell metastasis through inhibiting the CtsZ/FAK signaling pathway. <i>Cellular Signalling</i> , 2018, 50, 131-141.	1.7	40
17	Targeting MCL-1 sensitizes human esophageal squamous cell carcinoma cells to cisplatin-induced apoptosis. <i>BMC Cancer</i> , 2017, 17, 449.	1.1	42
18	A clinical and molecular analysis of a patient with Emanuel syndrome. <i>Molecular Medicine Reports</i> , 2017, 15, 1348-1352.	1.1	5

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19	19p13.2 Microdeletion including NF1 associated with overgrowth and intellectual disability suggestive of Malan syndrome. <i>Molecular Cytogenetics</i> , 2016, 9, 71.	0.4	6
20	Facial lymph node involvement as a prognostic factor for patient survival in oral cavity squamous cell carcinoma. <i>Tumor Biology</i> , 2016, 37, 3489-3496.	0.8	8
21	Anomalous origin of the right pulmonary artery from the ascending aorta: results of direct implantation surgical repair in 6 infants. <i>Journal of Cardiothoracic Surgery</i> , 2015, 10, 97.	0.4	8
22	Duplication of 10q22.3-q23.3 encompassing <i>BMPR1A</i> and <i>NGR3</i> associated with congenital heart disease, microcephaly, and mild intellectual disability. <i>American Journal of Medical Genetics, Part A</i> , 2015, 167, 3174-3179.	0.7	12
23	Whole-exome sequencing identifies Y1495X of <i>SCN5A</i> to be associated with familial conduction disease and sudden death. <i>Scientific Reports</i> , 2015, 4, 5616.	1.6	26
24	Rare De Novo Copy Number Variants in Patients with Congenital Pulmonary Atresia. <i>PLoS ONE</i> , 2014, 9, e96471.	1.1	19
25	Echocardiographic Assessment and Guidance in Minimally Invasive Surgical Device Closure of Perimembranous Ventricular Septal Defects. <i>Heart Surgery Forum</i> , 2014, 17, 206.	0.2	7
26	Two novel mutations of the LDL receptor gene associated with familial hypercholesterolemia in a Chinese family. <i>Chinese Medical Journal</i> , 2007, 120, 1694-9.	0.9	1