Yi Huang

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

31	802	15	28
papers	citations	h-index	g-index
35	1,170 ext. citations	7.3	4.46
ext. papers		avg, IF	L-index

#	Paper	IF	Citations
31	Apatinib promotes autophagy and apoptosis through VEGFR2/STAT3/BCL-2 signaling in osteosarcoma. <i>Cell Death and Disease</i> , 2017 , 8, e3015	9.8	131
30	Knockdown of long non-coding RNA HOTAIR increases miR-454-3p by targeting Stat3 and Atg12 to inhibit chondrosarcoma growth. <i>Cell Death and Disease</i> , 2017 , 8, e2605	9.8	76
29	PD-1 axis expression in musculoskeletal tumors and antitumor effect of nivolumab in osteosarcoma model of humanized mouse. <i>Journal of Hematology and Oncology</i> , 2018 , 11, 16	22.4	68
28	Tumor-associated macrophages promote lung metastasis and induce epithelial-mesenchymal transition in osteosarcoma by activating the COX-2/STAT3 axis. <i>Cancer Letters</i> , 2019 , 440-441, 116-125	9.9	68
27	miR-16-5p inhibits chordoma cell proliferation, invasion and metastasis by targeting Smad3. <i>Cell Death and Disease</i> , 2018 , 9, 680	9.8	55
26	Apatinib inhibits migration and invasion as well as PD-L1 expression in osteosarcoma by targeting STAT3. <i>Biochemical and Biophysical Research Communications</i> , 2018 , 495, 1695-1701	3.4	54
25	Bone marrow mesenchymal stem cell-derived exosomal miR-206 inhibits osteosarcoma progression by targeting TRA2B. <i>Cancer Letters</i> , 2020 , 490, 54-65	9.9	41
24	Immunotherapy for osteosarcoma: Fundamental mechanism, rationale, and recent breakthroughs. <i>Cancer Letters</i> , 2021 , 500, 1-10	9.9	41
23	Osteosarcoma cell intrinsic PD-L2 signals promote invasion and metastasis via the RhoA-ROCK-LIMK2 and autophagy pathways. <i>Cell Death and Disease</i> , 2019 , 10, 261	9.8	34
22	Apatinib plus camrelizumab (anti-PD1 therapy, SHR-1210) for advanced osteosarcoma (APFAO) progressing after chemotherapy: a single-arm, open-label, phase 2 trial 2020 , 8,		34
21	BMPR2 promotes invasion and metastasis via the RhoA-ROCK-LIMK2 pathway in human osteosarcoma cells. <i>Oncotarget</i> , 2017 , 8, 58625-58641	3.3	20
20	Identification of Potential Therapeutic Targets and Immune Cell Infiltration Characteristics in Osteosarcoma Using Bioinformatics Strategy. <i>Frontiers in Oncology</i> , 2020 , 10, 1628	5.3	20
19	Exosomal PD-L1 and N-cadherin predict pulmonary metastasis progression for osteosarcoma patients. <i>Journal of Nanobiotechnology</i> , 2020 , 18, 151	9.4	19
18	BMPR2 and HIF1- overexpression in resected osteosarcoma correlates with distant metastasis and patient survival. <i>Chinese Journal of Cancer Research: Official Journal of China Anti-Cancer Association, Beijing Institute for Cancer Research,</i> 2017 , 29, 447-454	3.8	18
17	Bortezomib induces apoptosis and suppresses cell growth and metastasis by inactivation of Stat3 signaling in chondrosarcoma. <i>International Journal of Oncology</i> , 2017 , 50, 477-486	4.4	16
16	The role of tumor-associated macrophages in osteosarcoma progression - therapeutic implications. <i>Cellular Oncology (Dordrecht)</i> , 2021 , 44, 525-539	7.2	15
15	Knockdown of HMGA2 regulates the level of autophagy via interactions between MSI2 and Beclin1 to inhibit NF1-associated malignant peripheral nerve sheath tumour growth. <i>Journal of Experimental and Clinical Cancer Research.</i> 2019 . 38. 185	12.8	14

LIST OF PUBLICATIONS

14	Induction of the mesenchymal to epithelial transition by demethylation-activated microRNA-125b is involved in the anti-migration/invasion effects of arsenic trioxide on human chondrosarcoma. Journal of Experimental and Clinical Cancer Research, 2016, 35, 129	12.8	14
13	LncRNA CASC15 is Upregulated in Osteosarcoma Plasma Exosomes and CASC15 Knockdown Inhibits Osteosarcoma Progression by Regulating miR-338-3p/RAB14 Axis. <i>OncoTargets and Therapy</i> , 2020 , 13, 12055-12066	4.4	13
12	Macrophages-derived exosomal lncRNA LIFR-AS1 promotes osteosarcoma cell progression via miR-29a/NFIA axis. <i>Cancer Cell International</i> , 2021 , 21, 192	6.4	13
11	Development of a prognostic gene signature based on an immunogenomic infiltration analysis of osteosarcoma. <i>Journal of Cellular and Molecular Medicine</i> , 2020 , 24, 11230-11242	5.6	12
10	Macrophages reduce the sensitivity of osteosarcoma to neoadjuvant chemotherapy drugs by secreting Interleukin-1 beta. <i>Cancer Letters</i> , 2020 , 480, 4-14	9.9	9
9	miR-100-5p Inhibits Malignant Behavior of Chordoma Cells by Targeting IGF1R. <i>Cancer Management and Research</i> , 2020 , 12, 4129-4137	3.6	6
8	circUSP34 accelerates osteosarcoma malignant progression by sponging miR-16-5p. <i>Cancer Science</i> , 2021 , 113, 120	6.9	3
7	Apatinib plus camrelizumab (SHR-1210) for unresectable high-grade osteosarcoma (APFAO) progressing after chemotherapy: A prospective, open label, phase II trial <i>Journal of Clinical Oncology</i> , 2019 , 37, 11013-11013	2.2	2
6	Chloroquine suppresses proliferation and invasion and induces apoptosis of osteosarcoma cells associated with inhibition of phosphorylation of STAT3. <i>Aging</i> , 2021 , 13, 17901-17913	5.6	2
5	PI3K inhibitor impairs tumor progression and enhances sensitivity to anlotinib in anlotinib-resistant osteosarcoma <i>Cancer Letters</i> , 2022 , 536, 215660	9.9	2
4	Immune-Related LncRNAs Affect the Prognosis of Osteosarcoma, Which Are Related to the Tumor Immune Microenvironment. <i>Frontiers in Cell and Developmental Biology</i> , 2021 , 9, 731311	5.7	O
3	Development of a Nomogram for Predicting the Efficacy of Preoperative Chemotherapy in Osteosarcoma. <i>International Journal of General Medicine</i> , 2021 , 14, 4819-4827	2.3	O
2	Quality of life and Q-TWiST were not adversely affected in Ewing sarcoma patients treated with combined anlotinib, irinotecan, and vincristine: (Peking University People Hospital Ewing sarcoma trial-02, PKUPH-EWS-02) <i>Medicine (United States)</i> , 2021 , 100, e28078	1.8	0
1	Anlotinib, vincristine, and irinotecan for advanced Ewing sarcoma after failure of standard multimodal therapy: A multicenter, two-cohort, phase Ib/II trial (NCT03416517). 2019 , 5, 118-118		