Xiao Wang

List of Publications by Year in descending order

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623734 501196 39 874 14 28 h-index citations g-index papers 42 42 42 1320 all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	Crystal structure of the YTH domain of YTHDF2 reveals mechanism for recognition of N6-methyladenosine. Cell Research, 2014, 24, 1493-1496.	12.0	266
2	Selective inhibition of CDK4/6: A safe and effective strategy for developing anticancer drugs. Acta Pharmaceutica Sinica B, 2021, 11, 30-54.	12.0	66
3	Synthesis of 2 <i>H</i> â€Indazolo[2,1â€ <i>b</i>]Phthalazineâ€1,6,11(13 <i>H</i>)â€Trione Derivatives Using We Cyanuric Chloride under Solventâ€Free Condition. Journal of the Chinese Chemical Society, 2010, 57, 1341-1345.	et 1.4	47
4	Access to Paediatric Essential Medicines: A Survey of Prices, Availability, Affordability and Price Components in Shaanxi Province, China. PLoS ONE, 2014, 9, e90365.	2.5	38
5	DKK1 inhibits breast cancer cell migration and invasion through suppression of \hat{l}^2 -catenin/MMP7 signaling pathway. Cancer Cell International, 2019, 19, 168.	4.1	38
6	Synergistic cytotoxic effects of arsenite and tetrandrine in human breast cancer cell line MCF-7. International Journal of Oncology, 2017, 51, 587-598.	3.3	33
7	A specific gut microbiota and metabolomic profiles shifts related to antidiabetic action: The similar and complementary antidiabetic properties of type 3 resistant starch from Canna edulis and metformin. Pharmacological Research, 2020, 159, 104985.	7.1	33
8	Antitumor activity of arsenite in combination with tetrandrine against human breast cancer cell line MDA-MB-231 in vitro and in vivo. Cancer Cell International, 2018, 18, 113.	4.1	31
9	Transdermal microemulsion drug delivery system for impairing male reproductive toxicity and enhancing efficacy of Tripterygium Wilfordii Hook f. Fìtoterapìâ, 2012, 83, 690-698.	2.2	26
10	Inhibition of PIKfyve using YM201636 suppresses the growth of liver cancer via the induction of autophagy. Oncology Reports, 2019, 41, 1971-1979.	2.6	24
11	EGF promotes <i>DKK1</i> transcription in hepatocellular carcinoma by enhancing the phosphorylation and acetylation of histone H3. Science Signaling, 2020, 13, .	3.6	21
12	Deciphering the mechanism of Fang-Ji-Di-Huang-Decoction in ameliorating psoriasis-like skin inflammation via the inhibition of IL-23/Th17Âcell axis. Journal of Ethnopharmacology, 2021, 281, 114571.	4.1	18
13	Protective Effects of Fisetin on Hepatic Ischemia-reperfusion Injury Through Alleviation of Apoptosis and Oxidative Stress. Archives of Medical Research, 2021, 52, 163-173.	3.3	17
14	Reusable melamine trisulfonic acid-catalyzed three-component synthesis of 7-alkyl-6H,7H-naphtho[1′,2′:5,6]pyrano[3,2-c]chromen-6-ones. Monatshefte FÃ⅓r Chemie, 2011, 142, 163-	-1 ¹ 6 ⁸ 7.	15
15	Wnt5b/Ryk-mediated membrane trafficking of P2X3 receptors contributes to bone cancer pain. Experimental Neurology, 2020, 334, 113482.	4.1	15
16	Silica Chloride Catalysed Oneâ€pot Synthesis of 13â€Arylâ€indeno[1,2â€ <i>b</i>]naphtha[1,2â€e]pyranâ€12(13 <i>H</i>)â€ones under Solventâ€free Conditions of the Chinese Chemical Society, 2010, 57, 738-741.	s 1Jø urnal	14
17	Cyanuric chloride-catalyzed synthesis of <i>N</i> -sulfonyl imines. Journal of Sulfur Chemistry, 2010, 31, 509-513.	2.0	14
18	PKM2-Induced the Phosphorylation of Histone H3 Contributes to EGF-Mediated PD-L1 Transcription in HCC. Frontiers in Pharmacology, 2020, 11, 577108.	3.5	14

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19	Preparative separation of quaternary ammonium alkaloids from Caulis Mahoniae by conventional and pH-zone-refining counter-current chromatography. RSC Advances, 2016, 6, 83343-83349.	3.6	13
20	Preparative separation of alkaloids from Litsea cubeba using combined applications of pH-zone-refining and high-speed counter-current chromatography. RSC Advances, 2015, 5, 75831-75837.	3.6	12
21	P300â€dependent acetylation of histone H3 is required for epidermal growth factor receptorâ€mediated highâ€mobility group protein A2 transcription in hepatocellular carcinoma. Cancer Science, 2021, 112, 679-690.	3.9	12
22	New Monoterpenoid Indoles with Osteoclast Activities from Gelsemium elegans. Molecules, 2021, 26, 7457.	3.8	10
23	Zr(HSO ₄) ₄ â€catalyzed oneâ€pot threeâ€component synthesis of 7â€alkylâ€6 <i>H</i> ,7 <i>H</i> ,8€naphtho[1′,2′:5,6]pyrano[3,2â€ <i>C</i>)]chromenâ€6â€ones. Journal of IChemistry, 2011, 48, 1379-1382.	He t esocyc	lic9
24	Discovery of novel and selective CDK4/6 inhibitors by pharmacophore and structure-based virtual screening. Future Medicinal Chemistry, 2020, 12, 1121-1136.	2.3	9
25	Development and clinical advancement of small molecules for exÂvivo expansion of hematopoietic stem cell. Acta Pharmaceutica Sinica B, 2022, 12, 2808-2831.	12.0	9
26	Molecular Iodine: A Versatile Catalyst for the Synthesis of 2H-indazolo[2,1-b]phthalazine-1,6,11(13H)-trione Derivatives in Ethanol. E-Journal of Chemistry, 2011, 8, 1000-1005.	0.5	8
27	Poly[4-diacetoxyiodo] Styrene–Promoted Thiocyanation of Aromatic Ethers, Anilines, and Indoles. Phosphorus, Sulfur and Silicon and the Related Elements, 2011, 186, 304-310.	1.6	8
28	One-pot synthesis of 1-aryl-1 <i>H</i> ,3 <i>H</i> -thiazolo[3,4- <i>a</i>]benzimidazoles using magnetite-linked sulfonic acid as catalyst. Phosphorus, Sulfur and Silicon and the Related Elements, 2014, 189, 1851-1857.	1.6	8
29	Oneâ€Pot Threeâ€Component Synthesis of 6â€Bromoquinolines and 6â€lodoquinolines. Journal of the Chinese Chemical Society, 2010, 57, 616-621.	1.4	7
30	P ₂ O ₅ /SiO ₂ as a New, Efficient and Reusable Catalyst for Preparation of 4,4′-Epoxydicoumarins Under Solvent-free Conditions. E-Journal of Chemistry, 2011, 8, 1626-1631.	0.5	7
31	Release profile of insulin from pH-sensitive hydrogel and its hypoglycemic effect by oral administration. Journal of Biomaterials Science, Polymer Edition, 2016, 27, 86-96.	3.5	7
32	A Study on the Mechanism of Lavender in the Treatment of Insomnia Based on Network Pharmacology. Combinatorial Chemistry and High Throughput Screening, 2020, 23, 419-432.	1.1	5
33	Discovery of novel and orally bioavailable CDK 4/6 inhibitors with high kinome selectivity, low toxicity and long-acting stability for the treatment of multiple myeloma. European Journal of Medicinal Chemistry, 2022, 228, 114024.	5.5	4
34	Discovery of Novel Phosphoinositide-3-Kinase α Inhibitors with High Selectivity, Excellent Bioavailability, and Long-Acting Efficacy for Gastric Cancer. Journal of Medicinal Chemistry, 2022, 65, 9873-9892.	6.4	4
35	Characterization of Formononetin Sulfonation in SULT1A3 Overexpressing HKE293 Cells: Involvement of Multidrug Resistance-Associated Protein 4 in Excretion of Sulfate. Frontiers in Pharmacology, 2020, 11, 614756.	3.5	3
36	Suppression of epidermal growth factor receptorâ€mediated βâ€catenin nuclear accumulation enhances the antiâ€tumor activity of phosphoinositide 3â€kinase inhibitor in breast cancer. Cell Biology International, 2019, 43, 931-939.	3.0	2

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37	Fabrication of poly(<i>t</i> -butyl betaine carboxylate)-based nanoparticles and study on their <i>in vivo</i> biosecurity. Journal of Biomaterials Science, Polymer Edition, 2021, 32, 2387-2401.	3.5	2
38	Fabrication and characterization of glutathioneâ€responsive nanoparticles from the disulfide bondâ€bridged block copolymer. Polymers for Advanced Technologies, 2022, 33, 180-188.	3.2	1
39	Exploration of the active components and pharmacological mechanism of Compound Longmaining for the treatment of myocardial infarction. Frontiers in Bioscience, 2021, 26, 813.	2.1	1