Jiaming Li

List of Publications by Year
in descending order

28	320	1040056	888059
papers	citations	9	17
		h-index	g-index

29	29	29	500
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Discovery of novel brain-penetrant GluN2B NMDAR antagonists via pharmacophore-merging strategy as anti-stroke therapeutic agents. European Journal of Medicinal Chemistry, 2022, 227, 113876.	5.5	7
2	Design, synthesis and biological evaluation of indoline derivatives as multifunctional agents for the treatment of ischemic stroke. Medicinal Chemistry Research, 2022, 31, 805-818.	2.4	1
3	Hybrids of aurantiamide acetate and isopropylated genipin as potential antiâ€inflammatory agents: The design, synthesis, and biological evaluation. Chemical Biology and Drug Design, 2021, 97, 797-808.	3.2	3
4	Design, synthesis and biological evaluation of novel benzoxaborole derivatives as potent PDE4 inhibitors for topical treatment of atopic dermatitis. European Journal of Medicinal Chemistry, 2021, 213, 113171.	5.5	20
5	Synthesis and Anti-proliferative Activity of Indole-2-amide Derivatives as Cyclooxygenase-2/5-lipoxygenase (COX-2/5-LOX) Dual Inhibitors. Chinese Journal of Organic Chemistry, 2021, 41, 1631.	1.3	3
6	Discovery of talmapimod analogues as polypharmacological anti-inflammatory agents. Journal of Enzyme Inhibition and Medicinal Chemistry, 2020, 35, 187-198.	5.2	9
7	Structurally novel PI3KÎ´/Î³ dual inhibitors characterized by a seven-membered spirocyclic spacer: The SARs investigation and PK evaluation. European Journal of Medicinal Chemistry, 2020, 191, 112143.	5.5	9
8	A conjugated mTOR/MEK bifunctional inhibitor as potential polypharmacological anticancer agent: the prototype compound discovery. Medicinal Chemistry Research, 2020, 29, 519-527.	2.4	4
9	Design, synthesis, biological evaluation and docking study of novel indole-2-amide as anti-inflammatory agents with dual inhibition of COX and 5-LOX. European Journal of Medicinal Chemistry, 2019, 180, 41-50.	5.5	54
10	Design, synthesis and biological evaluation of novel desloratadine derivatives with anti-inflammatory and H1 antagonize activities. Bioorganic and Medicinal Chemistry Letters, 2019, 29, 126712.	2.2	2
11	Conformationally restricted quinazolone derivatives as PI3KÎ´-selective inhibitors: the design, synthesis and biological evaluation. MedChemComm, 2019, 10, 413-420.	3.4	5
12	The synthesis and biological evaluation of novel gardenamide A derivatives as multifunctional neuroprotective agents. MedChemComm, 2019, 10, 1180-1186.	3.4	2
13	Design, synthesis, and biological evaluation of tetrahydroisoquinoline-based diaryl urea derivatives for suppressing VEGFR-2 signaling. Anti-Cancer Drugs, 2019, 30, 508-516.	1.4	2
14	Design, synthesis, and evaluation of genipin derivatives for the treatment of Alzheimer's Disease. Chemical Biology and Drug Design, 2019, 93, 110-122.	3.2	13
15	Synthesis and Biological Evaluation of Novel Phenylpropenoyl-amino Acid Derivatives. Chinese Journal of Organic Chemistry, 2019, 39, 1953.	1.3	2
16	Design, Synthesis and Evaluation of Novel Tacrine-3-n-butylphthalide Hybrids as Multifunctional Cholinesterase Inhibitors. Chinese Journal of Organic Chemistry, 2019, 39, 3505.	1.3	0
17	Design, synthesis chalcone derivatives as AdipoR agonist for type 2 diabetes. Chemical Biology and Drug Design, 2018, 92, 1525-1536.	3.2	11
18	Discovery of traditional Chinese medicine monomers and theirÂsynthetic intermediates, analogs or derivatives for battling P-gp-mediated multi-drug resistance. European Journal of Medicinal Chemistry, 2018, 159, 381-392.	5.5	38

#	Article	IF	Citations
19	Design, Synthesis, and Biological Evaluation of Novel PDE-4 Inhibitors. Chinese Journal of Organic Chemistry, 2018, 38, 478.	1.3	0
20	Design, synthesis, and biological evaluation of novel tetrahydroisoquinoline derivatives as potential antitumor candidate. Chemical Biology and Drug Design, 2017, 89, 443-455.	3.2	16
21	Design, Synthesis and Pharmacological Evaluation of Novel Piperlongumine derivatives as Potential Antiplatelet Aggregation Candidate. Chemical Biology and Drug Design, 2016, 87, 833-840.	3.2	9
22	The salicylanilide derivatives inhibit signal transducer and activator of transcription 3 pathways in A549 lung cancer cells. Anti-Cancer Drugs, 2016, 27, 41-47.	1.4	6
23	Synthesis and Evaluation of Paeonol Derivatives as Potential Multifunctional Agents for the Treatment of Alzheimerâ€™s Disease. Molecules, 2015, 20, 1304-1318.	3.8	31
24	Synthesis and Evaluation of Salicylanilide Derivatives as Potential Epidermal Growth Factor Receptor Inhibitors. Chemical Biology and Drug Design, 2015, 85, 280-289.	3.2	4
25	Design, Synthesis and Biological Activity of 1,2-Benzothiazine Derivatives as Potential Anticancer Agents. Chinese Journal of Organic Chemistry, 2014, 34, 2040.	1.3	4
26	Design, Synthesis, and Biological Evaluation ofN-Aryl-salicylamide Derivatives as Potential Antitumor Agents. Chinese Journal of Organic Chemistry, 2013, 33, 1026.	1.3	1
27	Gambogenic acid induced mitochondrial-dependent apoptosis and referred to Phospho-Erk1/2 and Phospho-p38 MAPK in human hepatoma HepG2 cells. Environmental Toxicology and Pharmacology, 2012, 33, 181-190.	4.0	62
28	A Practical Method to Stereospecifically Synthesize transâ€Stilbene Derivatives. Chinese Journal of Chemistry, 2011, 29, 1423-1428.	4.9	2