

Ping Xu

List of Publications by Year in descending order

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39
papers

515
citations

623734
14
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752698
20
g-index

41
all docs

41
docs citations

41
times ranked

634
citing authors

#	ARTICLE	IF	CITATIONS
1	The state of the art of PROTAC technologies for drug discovery. <i>European Journal of Medicinal Chemistry</i> , 2022, 235, 114290.	5.5	45
2	Research progress of MEK1/2 inhibitors and degraders in the treatment of cancer. <i>European Journal of Medicinal Chemistry</i> , 2021, 218, 113386.	5.5	29
3	A Smart Fluorescent Probe for NO Detection and Application in Myocardial Fibrosis Imaging. <i>Analytical Chemistry</i> , 2020, 92, 5064-5072.	6.5	28
4	Design and synthesis of a novel class of furan-based molecules as potential 20S proteasome inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007, 17, 1102-1106.	2.2	25
5	Substituted 3-Benzylcoumarins as Allosteric MEK1 Inhibitors: Design, Synthesis and Biological Evaluation as Antiviral Agents. <i>Molecules</i> , 2013, 18, 6057-6091.	3.8	24
6	Copper-catalyzed Aerobic Dehydrogenation of C ₆ H ₅ C ₂ H ₃ to C=C Bonds in the Synthesis of Pyridazinones. <i>European Journal of Organic Chemistry</i> , 2013, 2013, 6130-6136.	2.4	23
7	Discovery of 3-benzyl-1,3-benzoxazine-2,4-dione analogues as allosteric mitogen-activated kinase kinase (MEK) inhibitors and anti-enterovirus 71 (EV71) agents. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 3472-3482.	3.0	23
8	Rational Design, Green Synthesis, and Initial Evaluation of a Series of Full-color Tunable Fluorescent Dyes Enabled by the Copper-catalyzed N-arylation of 6-phenyl Pyridazinones and Their Application in Cell Imaging. <i>Chemistry - A European Journal</i> , 2013, 19, 13774-13782.	3.3	22
9	A Copper-catalyzed Aerobic Cascade Dehydrogenative Dehalogenative Reaction. <i>Advanced Synthesis and Catalysis</i> , 2013, 355, 1284-1290.	4.3	20
10	2-Substituted-thio- N -(4-substituted-thiazol-1 H -imidazol-2-yl)acetamides as BACE1 inhibitors: Synthesis, biological evaluation and Docking studies. <i>European Journal of Medicinal Chemistry</i> , 2017, 137, 462-475.	5.5	19
11	Development of novel proteasome inhibitors based on phthalazinone scaffold. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 2801-2805.	2.2	16
12	Relay Catalysis by a Multifunctional Cu Catalyst in a Tandem Dehydro-/Dehalogenation Sequence along with N-Arylation. <i>Organic Letters</i> , 2013, 15, 2770-2773.	4.6	15
13	4-Oxo-1,4-dihydro-quinoline-3-carboxamides as BACE-1 inhibitors: Synthesis, biological evaluation and docking studies. <i>European Journal of Medicinal Chemistry</i> , 2014, 79, 413-421.	5.5	15
14	Novel CADD-based peptidyl vinyl ester derivatives as potential proteasome inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 2198-2202.	2.2	14
15	The Discovery of Novel Secretase Inhibitors: Pharmacophore Modeling, Virtual Screening, and Docking Studies. <i>Chemical Biology and Drug Design</i> , 2012, 79, 972-980.	3.2	14
16	Ruthenium-catalyzed Switchable N ² -H/C ² -H Alkenylation of 6-phenyl(dihydro)pyridazinones with Alkynes. <i>European Journal of Organic Chemistry</i> , 2014, 2014, 6863-6867.	2.4	14
17	Capillary electrophoresis for screening of 20S proteasome inhibitors. <i>Analytical Biochemistry</i> , 2009, 394, 62-67.	2.4	13
18	Synthesis, <i>in vitro</i> Biological Evaluation and Molecular Docking Studies of Benzimidamides as Potential BACE1 Inhibitors. <i>Chemical Biology and Drug Design</i> , 2012, 80, 775-780.	3.2	12

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19	Rational Design of Fluorescent Phthalazinone Derivatives for One- and Two-Photon Imaging. Chemistry - A European Journal, 2016, 22, 12363-12370.	3.3	12
20	Discovery of carbazole derivatives as novel allosteric MEK inhibitors by pharmacophore modeling and virtual screening. European Journal of Medicinal Chemistry, 2019, 178, 802-817.	5.5	12
21	Hybrids of MEK inhibitor and NO donor as multitarget antitumor drugs. European Journal of Medicinal Chemistry, 2020, 196, 112271.	5.5	12
22	Regulation of enterovirus 2A protease-associated viral IRES activities by the cell's ERK signaling cascade: Implicating ERK as an efficiently antiviral target. Antiviral Research, 2017, 143, 13-21.	4.1	11
23	Revelation of the dynamic progression of hypoxia-reoxygenation injury by visualization of the lysosomal hydrogen peroxide. Biochemical and Biophysical Research Communications, 2017, 486, 904-908.	2.1	11
24	Structure-Activity Study of Nitazoxanide Derivatives as Novel STAT3 Pathway Inhibitors. ACS Medicinal Chemistry Letters, 2021, 12, 696-703.	2.8	9
25	Synthesis and Biological Evaluations of 3-Benzothiazol-2-yl Coumarin Derivatives as MEK1 Inhibitors. Letters in Drug Design and Discovery, 2013, 10, 727-732.	0.7	9
26	Template Synthesis of Peptidomimetics Composed of Aspartic Acid Moiety by Ugi Four-Component Condensation Reaction. Synthetic Communications, 2005, 35, 1881-1888.	2.1	8
27	Synthesis, Bioactivity, Docking and Molecular Dynamics Studies of Furan-Based Peptides as 20S Proteasome Inhibitors. ChemMedChem, 2015, 10, 498-510.	3.2	8
28	Discovery of a potent and highly specific β 2 proteasome inhibitor from a library of copper complexes. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 5780-5784.	2.2	8
29	Design, synthesis and biological evaluation of cobalt(II)-Schiff base complexes as ATP-noncompetitive MEK1 inhibitors. Journal of Inorganic Biochemistry, 2019, 195, 174-181.	3.5	8
30	Design, synthesis, and biological evaluation of 4-aminopyrimidine or 4,6-diaminopyrimidine derivatives as beta amyloid cleaving enzyme-1 inhibitors. Chemical Biology and Drug Design, 2019, 93, 926-933.	3.2	7
31	Synthesis of β -Secretase Inhibitors Containing a Hydroxyethylene Dipeptide Isostere. Synthetic Communications, 2007, 37, 9-24.	2.1	6
32	Substituted 3-benzylcoumarins 13 and 14 suppress enterovirus A71 replication by impairing viral 2Apro dependent IRES-driven translation. Antiviral Research, 2018, 160, 10-16.	4.1	4
33	Nitazoxanide and related thiazolides induce cell death in cancer cells by targeting the 20S proteasome with novel binding modes. Biochemical Pharmacology, 2022, 197, 114913.	4.4	4
34	Rational Design of a Near-Infrared Fluorescent Probe Based on a Pyridazinone Scaffold. European Journal of Organic Chemistry, 2017, 2017, 3274-3281.	2.4	3
35	Early Diagnosis of Cerebral Ischemia Reperfusion Injury and Revelation of Its Regional Development by a H_{3R} Receptor-Directed Probe. ACS Sensors, 2021, 6, 1330-1338.	7.8	3
36	Induction of Apoptosis in Cancer Cells by Glutathione Transferase Inhibitor Mediated Hydrophobic Tagging Molecules. ACS Medicinal Chemistry Letters, 2021, 12, 720-725.	2.8	3

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37	Synthesis and preliminary evaluation of peptidomimetic inhibitors of human β^2 -secretase. European Journal of Medicinal Chemistry, 2010, 45, 2089-2094.	5.5	2
38	Discovery of a cobalt complex with high MEK1 binding affinity. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 2221-2224.	2.2	2
39	Design and synthesis of tripeptidyl furylketones as selective inhibitors against the β^5 subunit of human 20S proteasome. European Journal of Medicinal Chemistry, 2020, 192, 112160.	5.5	2