

Ping Xu

List of Publications by Year in descending order

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39
papers

515
citations

623188

14
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752256

20
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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.	2.6	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.	2.6	29
3	A Smart Fluorescent Probe for NO Detection and Application in Myocardial Fibrosis Imaging. <i>Analytical Chemistry</i> , 2020, 92, 5064-5072.	3.2	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.	1.0	25
5	Substituted 3-Benzylcoumarins as Allosteric MEK1 Inhibitors: Design, Synthesis and Biological Evaluation as Antiviral Agents. <i>Molecules</i> , 2013, 18, 6057-6091.	1.7	24
6	Copper-catalyzed Aerobic Dehydrogenation of C=C to C=C Bonds in the Synthesis of Pyridazinones. <i>European Journal of Organic Chemistry</i> , 2013, 2013, 6130-6136.	1.2	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.	1.4	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.	1.7	22
9	A Copper-catalyzed Aerobic Cascade Dehydrogenative Dehalogenative Reaction. <i>Advanced Synthesis and Catalysis</i> , 2013, 355, 1284-1290.	2.1	20
10	2-Substituted-thio-N-(4-substituted-thiazol/1H-imidazol-2-yl)acetamides as BACE1 inhibitors: Synthesis, biological evaluation and docking studies. <i>European Journal of Medicinal Chemistry</i> , 2017, 137, 462-475.	2.6	19
11	Development of novel proteasome inhibitors based on phthalazinone scaffold. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 2801-2805.	1.0	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.	2.4	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.	2.6	15
14	Novel CADD-based peptidyl vinyl ester derivatives as potential proteasome inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 2198-2202.	1.0	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.	1.5	14
16	Ruthenium-catalyzed Switchable N-H/C-H Alkenylation of 6-phenyl(dihydro)pyridazinones with Alkynes. <i>European Journal of Organic Chemistry</i> , 2014, 2014, 6863-6867.	1.2	14
17	Capillary electrophoresis for screening of 20S proteasome inhibitors. <i>Analytical Biochemistry</i> , 2009, 394, 62-67.	1.1	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.	1.5	12

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19	Rational Design of Fluorescent Phthalazinone Derivatives for One- and Two-Photon Imaging. <i>Chemistry - A European Journal</i> , 2016, 22, 12363-12370.	1.7	12
20	Discovery of carbazole derivatives as novel allosteric MEK inhibitors by pharmacophore modeling and virtual screening. <i>European Journal of Medicinal Chemistry</i> , 2019, 178, 802-817.	2.6	12
21	Hybrids of MEK inhibitor and NO donor as multitarget antitumor drugs. <i>European Journal of Medicinal Chemistry</i> , 2020, 196, 112271.	2.6	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. <i>Antiviral Research</i> , 2017, 143, 13-21.	1.9	11
23	Revelation of the dynamic progression of hypoxia-reoxygenation injury by visualization of the lysosomal hydrogen peroxide. <i>Biochemical and Biophysical Research Communications</i> , 2017, 486, 904-908.	1.0	11
24	Structure-Activity Study of Nitazoxanide Derivatives as Novel STAT3 Pathway Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 696-703.	1.3	9
25	Synthesis and Biological Evaluations of 3-Benzothiazol-2-yl Coumarin Derivatives as MEK1 Inhibitors. <i>Letters in Drug Design and Discovery</i> , 2013, 10, 727-732.	0.4	9
26	Template Synthesis of Peptidomimetics Composed of Aspartic Acid Moiety by Ugi Four-Component Condensation Reaction. <i>Synthetic Communications</i> , 2005, 35, 1881-1888.	1.1	8
27	Synthesis, Bioactivity, Docking and Molecular Dynamics Studies of Furan-Based Peptides as 20S Proteasome Inhibitors. <i>ChemMedChem</i> , 2015, 10, 498-510.	1.6	8
28	Discovery of a potent and highly specific β 2 proteasome inhibitor from a library of copper complexes. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 5780-5784.	1.0	8
29	Design, synthesis and biological evaluation of cobalt(II)-Schiff base complexes as ATP-noncompetitive MEK1 inhibitors. <i>Journal of Inorganic Biochemistry</i> , 2019, 195, 174-181.	1.5	8
30	Design, synthesis, and biological evaluation of 4-aminopyrimidine or 4,6-diaminopyrimidine derivatives as beta amyloid cleaving enzyme-1 inhibitors. <i>Chemical Biology and Drug Design</i> , 2019, 93, 926-933.	1.5	7
31	Synthesis of β -Secretase Inhibitors Containing a Hydroxyethylene Dipeptide Isostere. <i>Synthetic Communications</i> , 2007, 37, 9-24.	1.1	6
32	Substituted 3-benzylcoumarins 13 and 14 suppress enterovirus A71 replication by impairing viral 2Apro dependent IRES-driven translation. <i>Antiviral Research</i> , 2018, 160, 10-16.	1.9	4
33	Nitazoxanide and related thiazolides induce cell death in cancer cells by targeting the 20S proteasome with novel binding modes. <i>Biochemical Pharmacology</i> , 2022, 197, 114913.	2.0	4
34	Rational Design of a Near-Infrared Fluorescent Probe Based on a Pyridazinone Scaffold. <i>European Journal of Organic Chemistry</i> , 2017, 2017, 3274-3281.	1.2	3
35	Early Diagnosis of Cerebral Ischemia Reperfusion Injury and Revelation of Its Regional Development by a H_{3R} Receptor-Directed Probe. <i>ACS Sensors</i> , 2021, 6, 1330-1338.	4.0	3
36	Induction of Apoptosis in Cancer Cells by Glutathione Transferase Inhibitor Mediated Hydrophobic Tagging Molecules. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 720-725.	1.3	3

#	ARTICLE	IF	CITATIONS
37	Synthesis and preliminary evaluation of peptidomimetic inhibitors of human \hat{I}^2 -secretase. European Journal of Medicinal Chemistry, 2010, 45, 2089-2094.	2.6	2
38	Discovery of a cobalt complex with high MEK1 binding affinity. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 2221-2224.	1.0	2
39	Design and synthesis of tripeptidyl furylketones as selective inhibitors against the \hat{I}^{25} subunit of human 20S proteasome. European Journal of Medicinal Chemistry, 2020, 192, 112160.	2.6	2