

Zi-Ning Cui

List of Publications by Year in descending order

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26
papers

358
citations

759233

12
h-index

839539

18
g-index

26
all docs

26
docs citations

26
times ranked

359
citing authors

#	ARTICLE	IF	CITATIONS
1	Synthesis and biological evaluation of 2,5-disubstituted furan derivatives as P-glycoprotein inhibitors for Doxorubicin resistance in MCF-7/ADR cell. <i>European Journal of Medicinal Chemistry</i> , 2018, 151, 546-556.	5.5	32
2	Synthesis and bioactivity of pyrazole and triazole derivatives as potential PDE4 inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 3632-3635.	2.2	31
3	Synthesis and fungicidal activity of novel pyrazole derivatives containing 5-Phenyl-2-Furan. <i>Bioorganic and Medicinal Chemistry</i> , 2019, 27, 115048.	3.0	27
4	Novel <i>S</i> -Thiazol-2-yl-furan-2-carbothioate Derivatives as Potential T3SS Inhibitors Against <i>Xanthomonas oryzae</i> on Rice. <i>Journal of Agricultural and Food Chemistry</i> , 2019, 67, 11867-11876.	5.2	27
5	Design, synthesis and biological evaluation of 2,4-disubstituted oxazole derivatives as potential PDE4 inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 1852-1859.	3.0	21
6	Design, synthesis and bioactivity study on 5-phenylfuran derivatives as potent reversal agents against P-glycoprotein-mediated multidrug resistance in MCF-7/ADR cell. <i>European Journal of Medicinal Chemistry</i> , 2021, 216, 113336.	5.5	21
7	Synthesis and bioactivity of thiazolidin-2-cyanamide derivatives against type III secretion system of <i>Xanthomonas oryzae</i> on rice. <i>Pesticide Biochemistry and Physiology</i> , 2018, 149, 89-97.	3.6	20
8	Synthesis and biological evaluation of 1,3,4-thiadiazole derivatives as type III secretion system inhibitors against <i>Xanthomonas oryzae</i> . <i>Pesticide Biochemistry and Physiology</i> , 2019, 160, 87-94.	3.6	19
9	Diversity-oriented synthesis of fluoroalkylated amines <i>via</i> the palladium-catalyzed divergent fluoroalkylation of 1,3-dienes. <i>Chemical Communications</i> , 2022, 58, 5614-5617.	4.1	15
10	Design, synthesis and fungicidal activity of novel 2-substituted aminocycloalkylsulfonamides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 271-276.	2.2	13
11	2,5-Disubstituted furan derivatives containing 1,3,4-thiadiazole moiety as potent β -glucosidase and <i>E. coli</i> β -glucuronidase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2021, 216, 113322.	5.5	13
12	Small Molecule Inhibitors Specifically Targeting the Type III Secretion System of <i>Xanthomonas oryzae</i> on Rice. <i>International Journal of Molecular Sciences</i> , 2019, 20, 971.	4.1	12
13	Synthesis, Fungicidal Activity, and Structure Activity Relationship of β -Acylaminocycloalkylsulfonamides against <i>Botrytis cinerea</i> . <i>Scientific Reports</i> , 2017, 7, 42096.	3.3	11
14	Tetrahydroquinoline and tetrahydroisoquinoline derivatives as potential selective PDE4B inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018, 28, 3271-3275.	2.2	11
15	Thiazolidin-2-cyanamides derivatives as novel potent <i>Escherichia coli</i> β -glucuronidase inhibitors and their structure-activity relationships. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020, 35, 1736-1742.	5.2	11
16	Synthesis and bioactivity of phenyl substituted furan and oxazole carboxylic acid derivatives as potential PDE4 inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2020, 207, 112795.	5.5	10
17	Synthesis and bioactivity of 3,5-dimethylpyrazole derivatives as potential PDE4 inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018, 28, 3276-3280.	2.2	9
18	Discovery of Ethyl 2-Nitro-3-Arylacrylates Molecules as T3SS Inhibitor Reducing the Virulence of Plant Pathogenic Bacteria <i>Xanthomonas</i> . <i>Frontiers in Microbiology</i> , 2019, 10, 1874.	3.5	8

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19	Synthesis and bioactivity of 1,3-thiazolidine-2-thione derivatives against type III secretion system of <i>Xanthomonas oryzae</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2019, 27, 3364-3371.	3.0	8
20	TEMPO-Mediated Synthesis of <i>N</i> -(Fluoroalkyl)imidazolones via Reaction of Imidazoles with Iodo-fluoroacetate. <i>Advanced Synthesis and Catalysis</i> , 2020, 362, 269-276.	4.3	8
21	Rational design of phenyl thiophene (pyridine) derivatives that overcome P-glycoprotein mediated MDR in MCF-7/ADR cell. <i>Bioorganic Chemistry</i> , 2021, 114, 105075.	4.1	8
22	Rational design of conformationally constrained oxazolidinone-fused 1,2,3,4-tetrahydroisoquinoline derivatives as potential PDE4 inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 5709-5717.	3.0	7
23	Structure-aided optimization of 3-O- β -chacotriosyl ursolic acid as novel H5N1 entry inhibitors with high selective index. <i>Bioorganic and Medicinal Chemistry</i> , 2019, 27, 4048-4058.	3.0	7
24	Synthesis and biological evaluation of acylated oligorhamnoside derivatives structurally related to mezzettiaside-6 with cytotoxic activity. <i>Organic and Biomolecular Chemistry</i> , 2016, 14, 6691-6702.	2.8	6
25	Discovery of a series of 5-phenyl-2-furan derivatives containing 1,3-thiazole moiety as potent <i>Escherichia coli</i> β -glucuronidase inhibitors. <i>Bioorganic Chemistry</i> , 2021, 116, 105306.	4.1	2
26	Sulfite-Promoted 1 H Fluoroalkyl Sulfuration of Imidazoheterocycles with Bromofluoroacetate and Elemental Sulfur. <i>Synthesis</i> , 2020, 52, 2541-2550.	2.3	1