

Yuanyuan Qian

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9140832/publications.pdf>

Version: 2024-02-01

10
papers

103
citations

1478505

6
h-index

1474206

9
g-index

10
all docs

10
docs citations

10
times ranked

132
citing authors

#	ARTICLE	IF	CITATIONS
1	Exploration of the Structural Space in 4(3 <i>H</i>)-Quinazolinone Antibacterials. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 5287-5296.	6.4	28
2	Structure-Activity Relationship for the Oxadiazole Class of Antibacterials. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 322-326.	2.8	18
3	An Atypical ABC Transporter Is Involved in Antifungal Resistance and Host Interactions in the Pathogenic Fungus <i>Cryptococcus neoformans</i> . <i>MBio</i> , 2022, 13, .	4.1	16
4	Discovery of Potent and Selective Agonists of μ Opioid Receptor by Revisiting the "Message-Address" Concept. <i>ACS Medicinal Chemistry Letters</i> , 2016, 7, 391-396.	2.8	13
5	Discovery of a Highly Selective and Potent μ Opioid Receptor Agonist from <i>N</i> -Cyclopropylmethyl-7-phenyl-6,14-endoethanotetrahydronorthebaines with Reduced Central Nervous System (CNS) Side Effects Navigated by the Message-Address Concept. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 11054-11070.	6.4	12
6	Cinnamionitrile Adjuvants Restore Susceptibility to β -Lactams against Methicillin-Resistant <i>Staphylococcus aureus</i> . <i>ACS Medicinal Chemistry Letters</i> , 2019, 10, 1148-1153.	2.8	10
7	Design, Synthesis, and Structure-Activity Relationship Exploration of Alkyl/Phenylalkyl Piperidine Analogues as Novel Highly Potent and Selective μ Opioid Receptor Agonists. <i>ACS Chemical Neuroscience</i> , 2021, 12, 285-299.	3.5	3
8	Susceptibility of Methicillin-Resistant <i>Staphylococcus aureus</i> to Five Quinazolinone Antibacterials. <i>Antimicrobial Agents and Chemotherapy</i> , 2019, 64, .	3.2	2
9	Discovery, Structure-Activity Relationship, and Mechanistic Studies of 1-((3 <i>R</i> ,4 <i>S</i>)-3-((Dimethylamino)methyl)-4-hydroxy-4-(3-methoxyphenyl)piperidin-1-yl)-2-(2,4,5-trifluorophenyl)ethan-1-one as a Novel Potent Analgesic. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 9458-9483.	6.0	1
10	Structure-Activity Relationship for the Picolinamide Antibacterials that Selectively Target <i>Clostridioides difficile</i> . <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 991-995.	2.8	0