

Shinmee Mah

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/2135828/publications.pdf>

Version: 2024-02-01

9
papers

210
citations

1307594

7
h-index

1474206

9
g-index

10
all docs

10
docs citations

10
times ranked

393
citing authors

#	ARTICLE	IF	CITATIONS
1	Catalyst Controlled Divergent C4/C8 Site-Selective C-H Arylation of Isoquinolones. <i>Organic Letters</i> , 2015, 17, 3864-3867.	4.6	66
2	Discovery of EGF Receptor Inhibitors That Are Selective for the d746â€‹/b>750/T790M/C797S Mutant through Structureâ€‹Based de Novo Design. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 7634-7638.	13.8	58
3	Systematic Computational Design and Identification of Low Picomolar Inhibitors of Aurora Kinase A. <i>Journal of Chemical Information and Modeling</i> , 2018, 58, 700-709.	5.4	20
4	Identification of 4-Phenoxyquinoline Based Inhibitors for L1196M Mutant of Anaplastic Lymphoma Kinase by Structure-Based Design. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 9205-9221.	6.4	18
5	HS-173 as a novel inducer of RIP3-dependent necroptosis in lung cancer. <i>Cancer Letters</i> , 2019, 444, 94-104.	7.2	16
6	Discovery of fluorescent 3-heteroaryl coumarin derivatives as novel inhibitors of anaplastic lymphoma kinase. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 186-194.	2.8	12
7	Discovery of Low Micromolar Dual Inhibitors for Wild Type and L1196M Mutant of Anaplastic Lymphoma Kinase through Structure-Based Virtual Screening. <i>Journal of Chemical Information and Modeling</i> , 2016, 56, 802-810.	5.4	9
8	Discovery of EGF Receptor Inhibitors That Are Selective for the d746â€‹/b>750/T790M/C797S Mutant through Structureâ€‹Based de Novo Design. <i>Angewandte Chemie</i> , 2017, 129, 7742-7746.	2.0	7
9	Kinase and GPCR polypharmacological approach for the identification of efficient anticancer medicines. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 8402-8413.	2.8	4