

Xiaobao Yang

List of Publications by Year in descending order

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Version: 2024-02-01

18
papers

1,021
citations

516215

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839053

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all docs

19
docs citations

19
times ranked

1431
citing authors

#	ARTICLE	IF	CITATIONS
1	Proteolysis Targeting Chimeras (PROTACs) of Anaplastic Lymphoma Kinase (ALK). <i>European Journal of Medicinal Chemistry</i> , 2018, 151, 304-314.	2.6	165
2	Discovery of a first-in-class EZH2 selective degrader. <i>Nature Chemical Biology</i> , 2020, 16, 214-222.	3.9	148
3	The First Structure-Activity Relationship Studies for Designer Receptors Exclusively Activated by Designer Drugs. <i>ACS Chemical Neuroscience</i> , 2015, 6, 476-484.	1.7	128
4	Distinct cortical and striatal actions of a β^2 -arrestin-biased dopamine D2 receptor ligand reveal unique antipsychotic-like properties. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, E8178-E8186.	3.3	117
5	Discovery of SIAIS178 as an Effective BCR-ABL Degradator by Recruiting Von Hippel-Lindau (VHL) E3 Ubiquitin Ligase. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 9281-9298.	2.9	79
6	Effective degradation of EGFR L858R+T790M mutant proteins by CRBN-based PROTACs through both proteasome and autophagy/lysosome degradation systems. <i>European Journal of Medicinal Chemistry</i> , 2021, 218, 113328.	2.6	55
7	Development of a Brigatinib degrader (SIAIS117) as a potential treatment for ALK positive cancer resistance. <i>European Journal of Medicinal Chemistry</i> , 2020, 193, 112190.	2.6	50
8	Chemoselective Synthesis of Lenalidomide-Based PROTAC Library Using Alkylation Reaction. <i>Organic Letters</i> , 2019, 21, 3838-3841.	2.4	48
9	Structure-Activity Relationship Studies for Enhancer of Zeste Homologue 2 (EZH2) and Enhancer of Zeste Homologue 1 (EZH1) Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 7617-7633.	2.9	46
10	ROCK1 mechano-signaling dependency of human malignancies driven by TEAD/YAP activation. <i>Nature Communications</i> , 2022, 13, 703.	5.8	31
11	Distinct CDK6 complexes determine tumor cell response to CDK4/6 inhibitors and degraders. <i>Nature Cancer</i> , 2021, 2, 429-443.	5.7	29
12	Discovery of Potent and Selective Allosteric Inhibitors of Protein Arginine Methyltransferase 3 (PRMT3). <i>Journal of Medicinal Chemistry</i> , 2018, 61, 1204-1217.	2.9	27
13	Structure-based discovery of SIAIS001 as an oral bioavailability ALK degrader constructed from Alectinib. <i>European Journal of Medicinal Chemistry</i> , 2021, 217, 113335.	2.6	26
14	Discovery of a Brigatinib Degradator SIAIS164018 with Destroying Metastasis-Related Oncoproteins and a Reshuffling Kinome Profile. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 9152-9165.	2.9	23
15	Discovery of novel BCR-ABL PROTACs based on the cereblon E3 ligase design, synthesis, and biological evaluation. <i>European Journal of Medicinal Chemistry</i> , 2021, 223, 113645.	2.6	23
16	Construction of an IMiD-based azide library as a kit for PROTAC research. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 166-170.	1.5	21
17	Development of an MDM2 Degradator for Treatment of Acute Leukemias. <i>Blood</i> , 2021, 138, 1866-1866.	0.6	3
18	Abstract 41: Tumor resistance to CDK4/6 inhibitors and degraders determined by the expression state of CDK6. , 2021, , .		0