Xiaobao Yang

List of Publications by Year
in descending order

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18	1,021	516215 16	839053 18
papers	citations	h-index	g-index

19	19	19	1431
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	ROCK1 mechano-signaling dependency of human malignancies driven by TEAD/YAP activation. Nature Communications, 2022, 13, 703.	5.8	31
2	Construction of an IMiD-based azide library as a kit for PROTAC research. Organic and Biomolecular Chemistry, 2021, 19, 166-170.	1.5	21
3	Distinct CDK6 complexes determine tumor cell response to CDK4/6 inhibitors and degraders. Nature Cancer, 2021, 2, 429-443.	5.7	29
4	Structure-based discovery of SIAIS001 as an oral bioavailability ALK degrader constructed from Alectinib. European Journal of Medicinal Chemistry, 2021, 217, 113335.	2.6	26
5	Effective degradation of EGFRL858R+T790M mutant proteins by CRBN-based PROTACs through both proteosome and autophagy/lysosome degradation systems. European Journal of Medicinal Chemistry, 2021, 218, 113328.	2.6	55
6	Discovery of a Brigatinib Degrader SIAIS164018 with Destroying Metastasis-Related Oncoproteins and a Reshuffling Kinome Profile. Journal of Medicinal Chemistry, 2021, 64, 9152-9165.	2.9	23
7	Abstract 41: Tumor resistance to CDK4/6 inhibitors and degraders determined by the expression state of CDK6. , 2021, , .		0
8	Discovery of novel BCR-ABL PROTACs based on the cereblon E3 ligase design, synthesis, and biological evaluation. European Journal of Medicinal Chemistry, 2021, 223, 113645.	2.6	23
9	Development of an MDM2 Degrader for Treatment of Acute Leukemias. Blood, 2021, 138, 1866-1866.	0.6	3
10	Discovery of a first-in-class EZH2 selective degrader. Nature Chemical Biology, 2020, 16, 214-222.	3.9	148
11	Development of a Brigatinib degrader (SIAIS117) as a potential treatment for ALK positive cancer resistance. European Journal of Medicinal Chemistry, 2020, 193, 112190.	2.6	50
12	Discovery of SIAIS178 as an Effective BCR-ABL Degrader by Recruiting Von Hippelâ€“Lindau (VHL) E3 Ubiquitin Ligase. Journal of Medicinal Chemistry, 2019, 62, 9281-9298.	2.9	79
13	Chemoselective Synthesis of Lenalidomide-Based PROTAC Library Using Alkylation Reaction. Organic Letters, 2019, 21, 3838-3841.	2.4	48
14	Proteolysis Targeting Chimeras (PROTACs) of Anaplastic Lymphoma Kinase (ALK). European Journal of Medicinal Chemistry, 2018, 151, 304-314.	2.6	165
15	Discovery of Potent and Selective Allosteric Inhibitors of Protein Arginine Methyltransferase 3 (PRMT3). Journal of Medicinal Chemistry, 2018, 61, 1204-1217.	2.9	27
16	Distinct cortical and striatal actions of a Î²-arrestinâ€“biased dopamine D2 receptor ligand reveal unique antipsychotic-like properties. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, E8178-E8186.	3.3	117
17	Structureâ€“Activity Relationship Studies for Enhancer of Zeste Homologue 2 (EZH2) and Enhancer of Zeste Homologue 1 (EZH1) Inhibitors. Journal of Medicinal Chemistry, 2016, 59, 7617-7633.	2.9	46
18	The First Structureâ€“Activity Relationship Studies for Designer Receptors Exclusively Activated by Designer Drugs. ACS Chemical Neuroscience, 2015, 6, 476-484.	1.7	128