

Naixia Zhang

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7903778/publications.pdf>

Version: 2024-02-01

22
papers

573
citations

840585

11
h-index

713332

21
g-index

22
all docs

22
docs citations

22
times ranked

944
citing authors

#	ARTICLE	IF	CITATIONS
1	Rpn1 provides adjacent receptor sites for substrate binding and deubiquitination by the proteasome. <i>Science</i> , 2016, 351, .	6.0	234
2	Small-Molecule Targeting of E3 Ligase Adaptor SPOP in Kidney Cancer. <i>Cancer Cell</i> , 2016, 30, 474-484.	7.7	74
3	Discovery and biological evaluation of vinylsulfonamide derivatives as highly potent, covalent TEAD autopalmitylation inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2019, 184, 111767.	2.6	55
4	A Dynamic View of ATP-coupled Functioning Cycle of Hsp90 N-terminal Domain. <i>Scientific Reports</i> , 2015, 5, 9542.	1.6	29
5	Metabolomic investigation of regional brain tissue dysfunctions induced by global cerebral ischemia. <i>BMC Neuroscience</i> , 2016, 17, 25.	0.8	23
6	Tetramerized Sesquiterpenoid Ainsliatetramers A and B from <i>Ainsliaea fragrans</i> and Their Cytotoxic Activities. <i>Organic Letters</i> , 2019, 21, 8211-8214.	2.4	21
7	The Intervention Effects of Acupuncture on Fatigue Induced by Exhaustive Physical Exercises: A Metabolomics Investigation. <i>Evidence-based Complementary and Alternative Medicine</i> , 2015, 2015, 1-11.	0.5	20
8	USP28 and USP25 are downregulated by Vismodegib in vitro and in colorectal cancer cell lines. <i>FEBS Journal</i> , 2021, 288, 1325-1342.	2.2	17
9	Exploration of Fragment Binding Poses Leading to Efficient Discovery of Highly Potent and Orally Effective Inhibitors of FABP4 for Anti-inflammation. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 4090-4106.	2.9	15
10	Applications of Solution NMR in Drug Discovery. <i>Molecules</i> , 2021, 26, 576.	1.7	12
11	Allosteric Regulation of Hsp90's Activity by Small Molecules Targeting the Middle Domain of the Chaperone. <i>IScience</i> , 2020, 23, 100857.	1.9	11
12	Structural and functional studies of USP20 ZnF-UBP domain by NMR. <i>Protein Science</i> , 2019, 28, 1606-1619.	3.1	10
13	The N-terminal ubiquitin-binding region of ubiquitin-specific protease 28 modulates its deubiquitination function: NMR structural and mechanistic insights. <i>Biochemical Journal</i> , 2015, 471, 155-165.	1.7	8
14	Design, synthesis, and biological evaluation of tetrahydroquinolin derivatives as potent inhibitors of CBP bromodomain. <i>Bioorganic Chemistry</i> , 2020, 101, 103991.	2.0	7
15	Kynurenine derivative 3-HAA is an agonist ligand for transcription factor YY1. <i>Journal of Hematology and Oncology</i> , 2021, 14, 153.	6.9	7
16	Structural and Functional Investigations of the N-Terminal Ubiquitin Binding Region of Usp25. <i>Biophysical Journal</i> , 2017, 112, 2099-2108.	0.2	6
17	¹ H, ¹³ C and ¹⁵ N backbone and side-chain resonance assignments of the ZnF-UBP domain of USP20/VDU2. <i>Biomolecular NMR Assignments</i> , 2017, 11, 91-93.	0.4	6
18	Structure-based drug optimization and biological evaluation of tetrahydroquinolin derivatives as selective and potent CBP bromodomain inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020, 30, 127480.	1.0	6

#	ARTICLE	IF	CITATIONS
19	Aha1 Exhibits Distinctive Dynamics Behavior and Chaperone-Like Activity. <i>Molecules</i> , 2021, 26, 1943.	1.7	6
20	Suppression of asparagine synthetase enhances the antitumor potency of ART and artemalogue SOMCL-14-221 in non-small cell lung cancer. <i>Cancer Letters</i> , 2020, 475, 22-33.	3.2	5
21	Ribonucleotide reductase holoenzyme inhibitor COH29 interacts with deubiquitinase ubiquitin-specific protease 2 and downregulates its substrate protein cyclin D1. <i>FASEB Journal</i> , 2022, 36, e22329.	0.2	1
22	Expression, purification and characterization of the second DUSP domain of deubiquitinase USP20/VDU2. <i>Protein Expression and Purification</i> , 2021, 181, 105836.	0.6	0