

Naixia Zhang

List of Publications by Citations

Source: <https://exaly.com/author-pdf/7903778/naixia-zhang-publications-by-citations.pdf>

Version: 2024-04-27

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

22
papers

340
citations

7
h-index

18
g-index

22
ext. papers

469
ext. citations

7.2
avg, IF

3.01
L-index

#	Paper	IF	Citations
22	Rpn1 provides adjacent receptor sites for substrate binding and deubiquitination by the proteasome. <i>Science</i> , 2016 , 351,	33.3	166
21	Small-Molecule Targeting of E3 Ligase Adaptor SPOP in Kidney Cancer. <i>Cancer Cell</i> , 2016 , 30, 474-484	24.3	51
20	Discovery and biological evaluation of vinylsulfonamide derivatives as highly potent, covalent TEAD autopalmitylation inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2019 , 184, 111767	6.8	22
19	A dynamic view of ATP-coupled functioning cycle of Hsp90 N-terminal domain. <i>Scientific Reports</i> , 2015 , 5, 9542	4.9	18
18	Metabolomic investigation of regional brain tissue dysfunctions induced by global cerebral ischemia. <i>BMC Neuroscience</i> , 2016 , 17, 25	3.2	13
17	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, 508302	2.3	11
16	Tetramerized Sesquiterpenoid Ainsliatetramers A and B from and Their Cytotoxic Activities. <i>Organic Letters</i> , 2019 , 21, 8211-8214	6.2	9
15	USP28 and USP25 are downregulated by Vismodegib in vitro and in colorectal cancer cell lines. <i>FEBS Journal</i> , 2021 , 288, 1325-1342	5.7	7
14	H, C and N backbone and side-chain resonance assignments of the ZnF-UBP domain of USP20/VDU2. <i>Biomolecular NMR Assignments</i> , 2017 , 11, 91-93	0.7	5
13	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	8.3	5
12	Allosteric Regulation of Hsp90 α Activity by Small Molecules Targeting the Middle Domain of the Chaperone. <i>IScience</i> , 2020 , 23, 100857	6.1	5
11	Structural and Functional Investigations of the N-Terminal Ubiquitin Binding Region of Usp25. <i>Biophysical Journal</i> , 2017 , 112, 2099-2108	2.9	4
10	Design, synthesis, and biological evaluation of tetrahydroquinolin derivatives as potent inhibitors of CBP bromodomain. <i>Bioorganic Chemistry</i> , 2020 , 101, 103991	5.1	4
9	Structural and functional studies of USP20 ZnF-UBP domain by NMR. <i>Protein Science</i> , 2019 , 28, 1606-1610	10.3	4
8	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	3.8	4
7	Applications of Solution NMR in Drug Discovery. <i>Molecules</i> , 2021 , 26,	4.8	4
6	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	9.9	3

5	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	2.9	3
4	Aha1 Exhibits Distinctive Dynamics Behavior and Chaperone-Like Activity. <i>Molecules</i> , 2021 , 26,	4.8	1
3	Kynurenine derivative 3-HAA is an agonist ligand for transcription factor YY1. <i>Journal of Hematology and Oncology</i> , 2021 , 14, 153	22.4	1
2	Expression, purification and characterization of the second DUSP domain of deubiquitinase USP20/VDU2. <i>Protein Expression and Purification</i> , 2021 , 181, 105836	2	
1	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	0.9	