

Xiaoyu Zhang

List of Publications by Year in Descending Order

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

38
papers

1,149
citations

19
h-index

33
g-index

44
ext. papers

1,557
ext. citations

9.8
avg, IF

4.7
L-index

#	Paper	IF	Citations
38	Expanding the landscape of E3 ligases for targeted protein degradation. <i>Current Research in Chemical Biology</i> , 2022 , 2, 100020		1
37	DCAF11 Supports Targeted Protein Degradation by Electrophilic Proteolysis-Targeting Chimeras. <i>Journal of the American Chemical Society</i> , 2021 , 143, 5141-5149	16.4	23
36	Chemical Inhibition of ENL/AF9 YEATS Domains in Acute Leukemia. <i>ACS Central Science</i> , 2021 , 7, 815-830	6.8	12
35	SPIN4 Is a Principal Endogenous Substrate of the E3 Ubiquitin Ligase DCAF16. <i>Biochemistry</i> , 2021 , 60, 637-642	3.2	3
34	High-Throughput Enzyme Assay for Screening Inhibitors of the ZDHHC3/7/20 Acyltransferases. <i>ACS Chemical Biology</i> , 2021 , 16, 1318-1324	4.9	0
33	NMT1 and NMT2 are lysine myristoyltransferases regulating the ARF6 GTPase cycle. <i>Nature Communications</i> , 2020 , 11, 1067	17.4	28
32	A Chemical Proteomic Probe for the Mitochondrial Pyruvate Carrier Complex. <i>Angewandte Chemie</i> , 2020 , 132, 3924-3927	3.6	
31	A Chemical Proteomic Probe for the Mitochondrial Pyruvate Carrier Complex. <i>Angewandte Chemie - International Edition</i> , 2020 , 59, 3896-3899	16.4	6
30	An Activity-Guided Map of Electrophile-Cysteine Interactions in Primary Human T Cells. <i>Cell</i> , 2020 , 182, 1009-1026.e29	56.2	57
29	Chemical Proteomics for Expanding the Druggability of Human Disease. <i>ChemBioChem</i> , 2020 , 21, 3319-3330	3.8	1
28	SIRT2 and Lysine Fatty Acylation Regulate the Activity of RalB and Cell Migration. <i>ACS Chemical Biology</i> , 2019 , 14, 2014-2023	4.9	19
27	Electrophilic PROTACs that degrade nuclear proteins by engaging DCAF16. <i>Nature Chemical Biology</i> , 2019 , 15, 737-746	11.7	154
26	Loss of Sirtuin 1 Alters the Secretome of Breast Cancer Cells by Impairing Lysosomal Integrity. <i>Developmental Cell</i> , 2019 , 49, 393-408.e7	10.2	66
25	A Small-Molecule SIRT2 Inhibitor That Promotes K-Ras4a Lysine Fatty-Acylation. <i>ChemMedChem</i> , 2019 , 14, 744-748	3.7	25
24	HDAC11 regulates type I interferon signaling through defatty-acylation of SHMT2. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019 , 116, 5487-5492	11.5	79
23	Comparative Nucleotide-Dependent Interactome Analysis Reveals Shared and Differential Properties of KRas4a and KRas4b. <i>ACS Central Science</i> , 2018 , 4, 71-80	16.8	11
22	Protein Lipidation: Occurrence, Mechanisms, Biological Functions, and Enabling Technologies. <i>Chemical Reviews</i> , 2018 , 118, 919-988	68.1	166

21	Direct Comparison of SIRT2 Inhibitors: Potency, Specificity, Activity-Dependent Inhibition, and On-Target Anticancer Activities. <i>ChemMedChem</i> , 2018 , 13, 1890-1894	3.7	28
20	HPLC-Based Enzyme Assays for Sirtuins. <i>Methods in Molecular Biology</i> , 2018 , 1813, 225-234	1.4	2
19	SIRT6 regulates Ras-related protein R-Ras2 by lysine defatty-acylation. <i>ELife</i> , 2017 , 6,	8.9	45
18	SIRT2 and lysine fatty acylation regulate the transforming activity of K-Ras4a. <i>ELife</i> , 2017 , 6,	8.9	45
17	Identifying the functional contribution of the defatty-acylase activity of SIRT6. <i>Nature Chemical Biology</i> , 2016 , 12, 614-20	11.7	68
16	Lysine fatty acylation promotes lysosomal targeting of TNF- α . <i>Scientific Reports</i> , 2016 , 6, 24371	4.9	24
15	SIRT7 Is Activated by DNA and Deacetylates Histone H3 in the Chromatin Context. <i>ACS Chemical Biology</i> , 2016 , 11, 742-7	4.9	41
14	SIRT2 Reverses 4-Oxononanoyl Lysine Modification on Histones. <i>Journal of the American Chemical Society</i> , 2016 , 138, 12304-7	16.4	51
13	Thiomyrystoyl peptides as cell-permeable Sirt6 inhibitors. <i>Organic and Biomolecular Chemistry</i> , 2014 , 12, 7498-502	3.9	59
12	Improving the NQO1-inducing activities of phenolic acids from radix <i>Salvia miltiorrhiza</i> : a methylation strategy. <i>Chemical Biology and Drug Design</i> , 2011 , 78, 558-66	2.9	3
11	A deuterium-labelling mass spectrometry-tandem diode-array detector screening method for rapid discovery of naturally occurring electrophiles. <i>Analytical and Bioanalytical Chemistry</i> , 2011 , 400, 3463-71	4.4	4
10	Terpenoids from <i>Tripterygium wilfordii</i> . <i>Phytochemistry</i> , 2011 , 72, 1482-7	4	32
9	Characterization of aromatase binding agents from the dichloromethane extract of <i>Corydalis yanhusuo</i> using ultrafiltration and liquid chromatography tandem mass spectrometry. <i>Molecules</i> , 2010 , 15, 3556-66	4.8	14
8	A new fluorescein isothiocyanate-based screening method for the rapid discovery of electrophilic compounds. <i>Analytical Methods</i> , 2010 , 2, 1472	3.2	2
7	PYDDT, a novel phase 2 enzymes inducer, activates Keap1-Nrf2 pathway via depleting the cellular level of glutathione. <i>Toxicology Letters</i> , 2010 , 199, 93-101	4.4	17
6	2-(penta-1,3-dienyl)-5-(3,4-dihydroxybut-1-enyl)thiophene, a novel NQO1 inducing agent from <i>Echinops grijisii</i> Hance. <i>Molecules</i> , 2010 , 15, 5273-81	4.8	11
5	Characterization of bioactive thiophenes from the dichloromethane extract of <i>Echinops grijisii</i> as Michael addition acceptors. <i>Analytical and Bioanalytical Chemistry</i> , 2010 , 397, 1975-84	4.4	12
4	A secoiridoid with quinone reductase inducing activity from <i>Cortex fraxini</i> . <i>Phytotherapy Research</i> , 2010 , 81, 834-7	3.2	8

3	Three lignans and one coumarinolignoid with quinone reductase activity from <i>Eurycorymbus cavaleriei</i> . <i>Phytotherapy Research</i> , 2009 , 80, 320-6	3.2	5
2	Seven new benzeneacetic acid derivatives and their quinone reductase activity from <i>Eurycorymbus cavaleriei</i> . <i>Phytochemistry Letters</i> , 2009 , 2, 152-158	1.9	6
1	Characterization of chemopreventive agents from the dichloromethane extract of <i>Eurycorymbus cavaleriei</i> by liquid chromatography-ion trap mass spectrometry. <i>Journal of Chromatography A</i> , 2009 , 1216, 4859-67	4.5	20