

# Xiang Li

## List of Publications by Year in descending order

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Version: 2024-02-01

48  
papers

2,433  
citations

236912

25  
h-index

233409

45  
g-index

53  
all docs

53  
docs citations

53  
times ranked

3099  
citing authors

#	ARTICLE	IF	CITATIONS
1	YEATS Domains as Novel Epigenetic Readers: Structures, Functions, and Inhibitor Development. <i>ACS Chemical Biology</i> , 2023, 18, 994-1013.	3.4	21
2	Roles of Negatively Charged Histone Lysine Acylations in Regulating Nucleosome Structure and Dynamics. <i>Frontiers in Molecular Biosciences</i> , 2022, 9, 899013.	3.5	4
3	Lysine succinylation on non-histone chromosomal protein HMG-17 (HMGN2) regulates nucleosomal DNA accessibility by disrupting the HMGN2â€“nucleosome association. <i>RSC Chemical Biology</i> , 2021, 2, 1257-1262.	4.1	4
4	A tri-functional amino acid enables mapping of binding sites for posttranslational-modification-mediated protein-protein interactions. <i>Molecular Cell</i> , 2021, 81, 2669-2681.e9.	9.7	39
5	AtHDA6 functions as an H3K18ac eraser to maintain pericentromeric CHG methylation in <i>Arabidopsis thaliana</i> . <i>Nucleic Acids Research</i> , 2021, 49, 9755-9767.	14.5	6
6	Integrative Chemical Biology Approaches to Deciphering the Histone Code: A Problem-Driven Journey. <i>Accounts of Chemical Research</i> , 2021, 54, 3734-3747.	15.6	17
7	Protocol for the preparation of site-specific succinylated histone mimics to investigate the impact on nucleosome dynamics. <i>STAR Protocols</i> , 2021, 2, 100604.	1.2	0
8	Concise solid-phase synthesis enables derivatisation of YEATS domain cyclopeptide inhibitors for improved cellular uptake. <i>Bioorganic and Medicinal Chemistry</i> , 2021, 45, 116342.	3.0	9
9	Chemoproteomic approach for mapping binding sites of post-translational-modification-mediated proteinâ€“protein interactions. <i>Trends in Biochemical Sciences</i> , 2021, 46, 1030-1031.	7.5	0
10	A bifunctional amino acid to study proteinâ€“protein interactions. <i>RSC Advances</i> , 2020, 10, 42076-42083.	3.6	8
11	Biomimetic Î±-selective ribosylation enables two-step modular synthesis of biologically important ADP-ribosylated peptides. <i>Nature Communications</i> , 2020, 11, 5600.	12.8	13
12	Semisynthesis of site-specifically succinylated histone reveals that succinylation regulates nucleosome unwrapping rate and DNA accessibility. <i>Nucleic Acids Research</i> , 2020, 48, 9538-9549.	14.5	34
13	Selective Targeting of AF9 YEATS Domain by Cyclopeptide Inhibitors with Preorganized Conformation. <i>Journal of the American Chemical Society</i> , 2020, 142, 21450-21459.	13.7	25
14	Rational Design of Reversible Redox Shuttle for Highly Efficient Light-Driven Microswimmer. <i>ACS Nano</i> , 2020, 14, 3272-3280.	14.6	25
15	Editorial overview: Recent advance in chemical genetics and chemical epigenetics. <i>Current Opinion in Chemical Biology</i> , 2019, 51, A1-A3.	6.1	0
16	Glutarylation of Histone H4 Lysine 91 Regulates Chromatin Dynamics. <i>Molecular Cell</i> , 2019, 76, 660-675.e9.	9.7	112
17	Chemical Proteomic Profiling of Bromodomains Enables the Wide-Spectrum Evaluation of Bromodomain Inhibitors in Living Cells. <i>Journal of the American Chemical Society</i> , 2019, 141, 11497-11505.	13.7	21
18	Thermodynamic insights into an interaction between ACYL-CoAâ€“BINDING PROTEIN2 and LYSOPHOSPHOLIPASE2 in <i>Arabidopsis</i> . <i>Journal of Biological Chemistry</i> , 2019, 294, 6214-6226.	3.4	24

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19	Interrogating Interactions and Modifications of Histones in Live Cells. <i>Cell Chemical Biology</i> , 2018, 25, 1-3.	5.2	48
20	Site-Specific Installation of Succinyl Lysine Analog into Histones Reveals the Effect of H2BK34 Succinylation on Nucleosome Dynamics. <i>Cell Chemical Biology</i> , 2018, 25, 166-174.e7.	5.2	42
21	Structure-guided development of YEATS domain inhibitors by targeting $\pi$ - $\pi$ stacking. <i>Nature Chemical Biology</i> , 2018, 14, 1140-1149.	8.0	76
22	DNA-Encoded Dynamic Chemical Library and Its Applications in Ligand Discovery. <i>Journal of the American Chemical Society</i> , 2018, 140, 15859-15867.	13.7	83
23	A chemical reporter facilitates the detection and identification of lysine HMGylation on histones. <i>Chemical Science</i> , 2018, 9, 7797-7801.	7.4	11
24	Metabolic Labeling of Pseudaminic Acid-Containing Glycans on Bacterial Surfaces. <i>ACS Chemical Biology</i> , 2018, 13, 3030-3037.	3.4	41
25	Peptide-based approaches to identify and characterize proteins that recognize histone post-translational modifications. <i>Chinese Chemical Letters</i> , 2018, 29, 1051-1057.	9.0	11
26	Genetically Encoded Photoaffinity Histone Marks. <i>Journal of the American Chemical Society</i> , 2017, 139, 6522-6525.	13.7	55
27	Histone Ketoamide Adduction by 4-Oxo-2-nonenal Is a Reversible Posttranslational Modification Regulated by Sirt2. <i>ACS Chemical Biology</i> , 2017, 12, 47-51.	3.4	24
28	Crystal structure of the thioesterification conformation of <i>Bacillus subtilis</i> $\alpha$ -succinylbenzoyl-CoA synthetase reveals a distinct substrate-binding mode. <i>Journal of Biological Chemistry</i> , 2017, 292, 12296-12310.	3.4	6
29	Photo-lysine captures proteins that bind lysine post-translational modifications. <i>Nature Chemical Biology</i> , 2016, 12, 70-72.	8.0	77
30	Integrative Chemical Biology Approaches for Identification and Characterization of $\alpha$ -Erasers for Fatty Acid-Acylated Lysine Residues within Proteins. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 1149-1152.	13.8	62
31	Developing diazirine-based chemical probes to identify histone modification $\alpha$ -readers <sup>TM</sup> and $\alpha$ -erasers <sup>TM</sup> . <i>Chemical Science</i> , 2015, 6, 1011-1017.	7.4	56
32	Chemical proteomics approaches to examine novel histone posttranslational modifications. <i>Current Opinion in Chemical Biology</i> , 2015, 24, 80-90.	6.1	22
33	Identification of $\alpha$ -erasers <sup>TM</sup> for lysine crotonylated histone marks using a chemical proteomics approach. <i>ELife</i> , 2014, 3, .	6.0	237
34	A Chemical Probe for Lysine Malonylation. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 4883-4886.	13.8	64
35	Examining post-translational modification-mediated protein-protein interactions using a chemical proteomics approach. <i>Protein Science</i> , 2013, 22, 287-295.	7.6	33
36	Abstract: A Chemical Probe for Lysine Malonylation ( <i>Angew. Chem.</i> 18/2013). <i>Angewandte Chemie</i> , 2013, 125, 5056-5056.	2.0	0

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37	Dense Chromatin Activates Polycomb Repressive Complex 2 to Regulate H3 Lysine 27 Methylation. <i>Science</i> , 2012, 337, 971-975.	12.6	240
38	Quantitative Chemical Proteomics Approach To Identify Post-translational Modification-Mediated Protein-Protein Interactions. <i>Journal of the American Chemical Society</i> , 2012, 134, 1982-1985.	13.7	114
39	A Synthetic Chloride Channel Restores Chloride Conductance in Human Cystic Fibrosis Epithelial Cells. <i>PLoS ONE</i> , 2012, 7, e34694.	2.5	64
40	Examining the Mechanism of Action of a Kinesin Inhibitor Using Stable Isotope Labeled Inhibitors for Cross-Linking (SILIC). <i>Journal of the American Chemical Society</i> , 2011, 133, 12386-12389.	13.7	11
41	An Optical Switch for a Motor Protein. <i>ChemBioChem</i> , 2011, 12, 2265-2266.	2.6	0
42	Approach to Profile Proteins That Recognize Post-Translationally Modified Histone "Tails". <i>Journal of the American Chemical Society</i> , 2010, 132, 2504-2505.	13.7	46
43	Synthetic Chloride Channel Regulates Cell Membrane Potentials and Voltage-Gated Calcium Channels. <i>Journal of the American Chemical Society</i> , 2009, 131, 13676-13680.	13.7	90
44	$\hat{\pm}$ -Aminoxy Acids: New Possibilities from Foldamers to Anion Receptors and Channels. <i>Accounts of Chemical Research</i> , 2008, 41, 1428-1438.	15.6	183
45	A Small Synthetic Molecule Forms Chloride Channels to Mediate Chloride Transport across Cell Membranes. <i>Journal of the American Chemical Society</i> , 2007, 129, 7264-7265.	13.7	106
46	Peptides of aminoxy acids as foldamers. <i>Chemical Communications</i> , 2006, , 3367.	4.1	103
47	A Cyclic Hexapeptide Comprising Alternating $\hat{\pm}$ -Aminoxy and $\hat{\pm}$ -Amino Acids is a Selective Chloride Ion Receptor. <i>Chemistry - A European Journal</i> , 2005, 11, 3005-3009.	3.3	30
48	Enantioselective Recognition of Carboxylates: A Receptor Derived from $\hat{\pm}$ -Aminoxy Acids Functions as a Chiral Shift Reagent for Carboxylic Acids. <i>Journal of the American Chemical Society</i> , 2005, 127, 7996-7997.	13.7	117