

Edward A Lemke

List of Publications by Year in descending order

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Version: 2024-02-01

104
papers

10,945
citations

43973

48
h-index

33814

99
g-index

117
all docs

117
docs citations

117
times ranked

12645
citing authors

#	ARTICLE	IF	CITATIONS
1	Molecular Anatomy of a Trafficking Organelle. <i>Cell</i> , 2006, 127, 831-846.	13.5	1,985
2	A near-infrared fluorophore for live-cell super-resolution microscopy of cellular proteins. <i>Nature Chemistry</i> , 2013, 5, 132-139.	6.6	779
3	Interplay of $\hat{\pm}$ -synuclein binding and conformational switching probed by single-molecule fluorescence. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 5645-5650.	3.3	379
4	In situ structural analysis of the human nuclear pore complex. <i>Nature</i> , 2015, 526, 140-143.	13.7	361
5	Precision and accuracy of single-molecule FRET measurements—a multi-laboratory benchmark study. <i>Nature Methods</i> , 2018, 15, 669-676.	9.0	350
6	Amino Acids for Diels–Alder Reactions in Living Cells. <i>Angewandte Chemie - International Edition</i> , 2012, 51, 4166-4170.	7.2	298
7	A natively unfolded yeast prion monomer adopts an ensemble of collapsed and rapidly fluctuating structures. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2007, 104, 2649-2654.	3.3	296
8	Cell type–specific nuclear pores: a case in point for context–dependent stoichiometry of molecular machines. <i>Molecular Systems Biology</i> , 2013, 9, 648.	3.2	277
9	Genetically Encoded Copper–Free Click Chemistry. <i>Angewandte Chemie - International Edition</i> , 2011, 50, 3878-3881.	7.2	269
10	Fourier ring correlation as a resolution criterion for super-resolution microscopy. <i>Journal of Structural Biology</i> , 2013, 183, 363-367.	1.3	269
11	Single-molecule biophysics: at the interface of biology, physics and chemistry. <i>Journal of the Royal Society Interface</i> , 2008, 5, 15-45.	1.5	263
12	Plasticity of an Ultrafast Interaction between Nucleoporins and Nuclear Transport Receptors. <i>Cell</i> , 2015, 163, 734-745.	13.5	255
13	Minimal Tags for Rapid Dual–Color Live–Cell Labeling and Super–Resolution Microscopy. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 2245-2249.	7.2	254
14	Control of protein phosphorylation with a genetically encoded photocaged amino acid. <i>Nature Chemical Biology</i> , 2007, 3, 769-772.	3.9	208
15	Decoupling of size and shape fluctuations in heteropolymeric sequences reconciles discrepancies in SAXS vs. FRET measurements. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, E6342-E6351.	3.3	195
16	Genetic Incorporation of a Small, Environmentally Sensitive, Fluorescent Probe into Proteins in <i>Saccharomyces cerevisiae</i> . <i>Journal of the American Chemical Society</i> , 2009, 131, 12921-12923.	6.6	183
17	A General and Efficient Method for the Site-Specific Dual-Labeling of Proteins for Single Molecule Fluorescence Resonance Energy Transfer. <i>Journal of the American Chemical Society</i> , 2008, 130, 17664-17665.	6.6	159
18	Associating HIV-1 envelope glycoprotein structures with states on the \hat{A} virus observed by smFRET. <i>Nature</i> , 2019, 568, 415-419.	13.7	156

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19	Genetic Encoding of a Bicyclo[6.1.0]nonyne- ϵ -Charged Amino Acid Enables Fast Cellular Protein Imaging by Metal-Free Ligation. <i>ChemBioChem</i> , 2012, 13, 2094-2099.	1.3	153
20	FRET-based dynamic structural biology: Challenges, perspectives and an appeal for open-science practices. <i>ELife</i> , 2021, 10, .	2.8	152
21	Designer membraneless organelles enable codon reassignment of selected mRNAs in eukaryotes. <i>Science</i> , 2019, 363, .	6.0	129
22	Visualizing a one-way protein encounter complex by ultrafast single-molecule mixing. <i>Nature Methods</i> , 2011, 8, 239-241.	9.0	128
23	Labeling proteins on live mammalian cells using click chemistry. <i>Nature Protocols</i> , 2015, 10, 780-791.	5.5	127
24	Direct Visualization of the Conformational Dynamics of Single Influenza Hemagglutinin Trimers. <i>Cell</i> , 2018, 174, 926-937.e12.	13.5	118
25	Debugging Eukaryotic Genetic Code Expansion for Site-Specific Click-Paint Super-Resolution Microscopy. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 16172-16176.	7.2	117
26	Conserved features of intermediates in amyloid assembly determine their benign or toxic states. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 11172-11177.	3.3	115
27	Click Strategies for Single-Molecule Protein Fluorescence. <i>Journal of the American Chemical Society</i> , 2012, 134, 5187-5195.	6.6	106
28	Principles for designing fluorescent sensors and reporters. <i>Nature Chemical Biology</i> , 2011, 7, 480-483.	3.9	97
29	Single Molecule Study of the Intrinsically Disordered FG-Repeat Nucleoporin 153. <i>Biophysical Journal</i> , 2011, 101, 1710-1719.	0.2	97
30	PED in 2021: a major update of the protein ensemble database for intrinsically disordered proteins. <i>Nucleic Acids Research</i> , 2021, 49, D404-D411.	6.5	95
31	The liquid state of FG-nucleoporins mimics permeability barrier properties of nuclear pore complexes. <i>Journal of Cell Biology</i> , 2020, 219, .	2.3	93
32	Monomeric Huntingtin Exon 1 Has Similar Overall Structural Features for Wild-Type and Pathological Polyglutamine Lengths. <i>Journal of the American Chemical Society</i> , 2017, 139, 14456-14469.	6.6	87
33	The Multiple Faces of Disordered Nucleoporins. <i>Journal of Molecular Biology</i> , 2016, 428, 2011-2024.	2.0	82
34	Floppy but not sloppy: Interaction mechanism of FG-nucleoporins and nuclear transport receptors. <i>Seminars in Cell and Developmental Biology</i> , 2017, 68, 34-41.	2.3	81
35	Facilitated aggregation of FG nucleoporins under molecular crowding conditions. <i>EMBO Reports</i> , 2013, 14, 178-183.	2.0	78
36	Visualization of Synaptic Vesicle Movement in Intact Synaptic Boutons Using Fluorescence Fluctuation Spectroscopy. <i>Biophysical Journal</i> , 2005, 89, 2091-2102.	0.2	76

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37	Direct single-molecule observation of a protein living in two opposed native structures. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 10153-10158.	3.3	72
38	Kirkwood's "Buff Approach Rescues Overcollapse of a Disordered Protein in Canonical Protein Force Fields. Journal of Physical Chemistry B, 2015, 119, 7975-7984.	1.2	70
39	Genetic code expansion enabled site-specific dual-color protein labeling: superresolution microscopy and beyond. Current Opinion in Chemical Biology, 2015, 28, 164-173.	2.8	65
40	Genetic code expansion for multiprotein complex engineering. Nature Methods, 2016, 13, 997-1000.	9.0	63
41	Microfluidic Device for Single-Molecule Experiments with Enhanced Photostability. Journal of the American Chemical Society, 2009, 131, 13610-13612.	6.6	61
42	Highly Stable <i>trans</i> -Cyclooctene Amino Acids for Live-Cell Labeling. Chemistry - A European Journal, 2015, 21, 12266-12270.	1.7	58
43	New Red-Emitting Tetrazine-Phenoxazine Fluorogenic Labels for Live-Cell Intracellular Bioorthogonal Labeling Schemes. Chemistry - A European Journal, 2016, 22, 8972-8979.	1.7	58
44	Bio-orthogonal Red and Far-Red Fluorogenic Probes for Wash-Free Live-Cell and Super-resolution Microscopy. ACS Central Science, 2021, 7, 1561-1571.	5.3	57
45	A Versatile Tool for Live-Cell Imaging and Super-Resolution Nanoscopy Studies of HIV-1 Env Distribution and Mobility. Cell Chemical Biology, 2017, 24, 635-645.e5.	2.5	55
46	Orthogonal spin labeling using click chemistry for in vitro and in vivo applications. Journal of Magnetic Resonance, 2017, 275, 38-45.	1.2	54
47	Continuous throughput and long-term observation of single-molecule FRET without immobilization. Nature Methods, 2014, 11, 297-300.	9.0	53
48	Cargo transport through the nuclear pore complex at a glance. Journal of Cell Science, 2021, 134, .	1.2	53
49	Schnelle, zweifarbige Proteinmarkierung an lebenden Zellen für die hochauflösende Mikroskopie. Angewandte Chemie, 2014, 126, 2278-2282.	1.6	51
50	Single Synaptic Vesicle Tracking in Individual Hippocampal Boutons at Rest and during Synaptic Activity. Journal of Neuroscience, 2005, 25, 11034-11044.	1.7	49
51	Large-Scale Conformational Dynamics Control H5N1 Influenza Polymerase PB2 Binding to Importin β . Journal of the American Chemical Society, 2015, 137, 15122-15134.	6.6	49
52	A new family of bioorthogonally applicable fluorogenic labels. Organic and Biomolecular Chemistry, 2013, 11, 3297.	1.5	46
53	The Exploding Genetic Code. ChemBioChem, 2014, 15, 1691-1694.	1.3	44
54	Origin of Orthogonality of Strain-Promoted Click Reactions. Chemistry - A European Journal, 2015, 21, 12431-12435.	1.7	44

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55	Physics of the nuclear pore complex: Theory, modeling and experiment. <i>Physics Reports</i> , 2021, 921, 1-53.	10.3	44
56	Shining a Light on Phase Separation in the Cell. <i>Cell</i> , 2017, 168, 11-13.	13.5	42
57	Two Differential Binding Mechanisms of FG-Nucleoporins and Nuclear Transport Receptors. <i>Cell Reports</i> , 2018, 22, 3660-3671.	2.9	41
58	Hydrophilic <i>trans</i> -Cyclooctenylated Noncanonical Amino Acids for Fast Intracellular Protein Labeling. <i>ChemBioChem</i> , 2016, 17, 1518-1524.	1.3	39
59	The CD27L and CTP1L Endolysins Targeting Clostridia Contain a Built-in Trigger and Release Factor. <i>PLoS Pathogens</i> , 2014, 10, e1004228.	2.1	37
60	Mapping Multivalency and Differential Affinities within Large Intrinsically Disordered Protein Complexes with Segmental Motion Analysis. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 7364-7367.	7.2	37
61	New Generation of Bioorthogonally Applicable Fluorogenic Dyes with Visible Excitations and Large Stokes Shifts. <i>Bioconjugate Chemistry</i> , 2014, 25, 1370-1374.	1.8	34
62	Labeling of virus components for advanced, quantitative imaging analyses. <i>FEBS Letters</i> , 2016, 590, 1896-1914.	1.3	34
63	Intramolecular three-colour single pair FRET of intrinsically disordered proteins with increased dynamic range. <i>Molecular BioSystems</i> , 2012, 8, 2531.	2.9	32
64	Molecular determinants of large cargo transport into the nucleus. <i>ELife</i> , 2020, 9, .	2.8	31
65	Genetically Encoded Click Chemistry for Single-Molecule FRET of Proteins. <i>Methods in Cell Biology</i> , 2013, 113, 169-187.	0.5	30
66	Application of Noncanonical Amino Acids for Protein Labeling in a Genomically Recoded <i>Escherichia coli</i> . <i>ACS Synthetic Biology</i> , 2017, 6, 233-255.	1.9	29
67	Architecture of TAF11/TAF13/TBP complex suggests novel regulation properties of general transcription factor TFIID. <i>ELife</i> , 2017, 6, .	2.8	29
68	Dual film-like organelles enable spatial separation of orthogonal eukaryotic translation. <i>Cell</i> , 2021, 184, 4886-4903.e21.	13.5	28
69	Comment on "Innovative scattering analysis shows that hydrophobic disordered proteins are expanded in water". <i>Science</i> , 2018, 361, .	6.0	27
70	Site-Specific Labeling of Proteins for Single-Molecule FRET Measurements Using Genetically Encoded Ketone Functionalities. <i>Methods in Molecular Biology</i> , 2011, 751, 3-15.	0.4	26
71	Super-resolution Microscopy of Clickable Amino Acids Reveals the Effects of Fluorescent Protein Tagging on Protein Assemblies. <i>ACS Nano</i> , 2015, 9, 11034-11041.	7.3	26
72	Identification and mutational studies of conserved amino acids in the outer membrane receptor protein, FepA, which affect transport but not binding of ferric-enterobactin in <i>Escherichia coli</i> . <i>BioMetals</i> , 2003, 16, 507-518.	1.8	25

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73	Synthetic biomolecular condensates to engineer eukaryotic cells. <i>Current Opinion in Chemical Biology</i> , 2021, 64, 174-181.	2.8	25
74	Single-molecule FRET and crosslinking studies in structural biology enabled by noncanonical amino acids. <i>Current Opinion in Structural Biology</i> , 2015, 32, 66-73.	2.6	24
75	Sampling Long- versus Short-Range Interactions Defines the Ability of Force Fields To Reproduce the Dynamics of Intrinsically Disordered Proteins. <i>Journal of Chemical Theory and Computation</i> , 2017, 13, 3964-3974.	2.3	22
76	Nanoscale devices for linkerless long-term single-molecule observation. <i>Current Opinion in Biotechnology</i> , 2016, 39, 105-112.	3.3	21
77	Bisazide Cyanine Dyes as Fluorogenic Probes for Bis-Cyclooctynylated Peptide Tags and as Fluorogenic Cross-Linkers of Cyclooctynylated Proteins. <i>Bioconjugate Chemistry</i> , 2017, 28, 1552-1559.	1.8	20
78	Bistetrazine-Cyanines as Double-Clicking Fluorogenic Two-Point Binder or Crosslinker Probes. <i>Chemistry - A European Journal</i> , 2018, 24, 8841-8847.	1.7	19
79	Mechanism-Dependent Modulation of Ultrafast Interfacial Water Dynamics in Intrinsically Disordered Protein Complexes. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 4720-4724.	7.2	19
80	What precision-protein-tuning and nano-resolved single molecule sciences can do for each other. <i>BioEssays</i> , 2013, 35, 65-74.	1.2	16
81	Beyond the Transport Function of Import Receptors: What's All the FUS about?. <i>Cell</i> , 2018, 173, 549-553.	13.5	14
82	Synthesis and Evaluation of Novel Ring-Strained Noncanonical Amino Acids for Residue-Specific Bioorthogonal Reactions in Living Cells. <i>Chemistry - A European Journal</i> , 2021, 27, 6094-6099.	1.7	14
83	Unnatural Amino Acid Mutagenesis Reveals Dimerization As a Negative Regulatory Mechanism of VHR's Phosphatase Activity. <i>ACS Chemical Biology</i> , 2014, 9, 1451-1459.	1.6	12
84	Palladium-unleashed proteins: gentle aldehyde decaging for site-selective protein modification. <i>Chemical Communications</i> , 2018, 54, 1501-1504.	2.2	12
85	Precision Control of Cellular Pathways with Light. <i>ChemBioChem</i> , 2010, 11, 1825-1827.	1.3	11
86	Verbesserte Erweiterung des eukaryotischen genetischen Codes für seitenspezifische, hochauflösende Click-PAINT-Mikroskopie. <i>Angewandte Chemie</i> , 2016, 128, 16406-16410.	1.6	11
87	Raising the ribosomal repertoire. <i>Nature Chemistry</i> , 2020, 12, 503-504.	6.6	10
88	Comparative analysis of the coordinated motion of Hsp70s from different organelles observed by single-molecule three-color FRET. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2021, 118, .	3.3	10
89	Multifunctionality of F-rich nucleoporins. <i>Biochemical Society Transactions</i> , 2020, 48, 2603-2614.	1.6	10
90	Detektion von Mehrbindigkeit und differenziellen Affinitäten in großen, intrinsisch ungeordneten Proteinen mithilfe von Segmentbewegungsanalyse. <i>Angewandte Chemie</i> , 2014, 126, 7492-7496.	1.6	7

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91	Synthesis of Azido-Glycans for Chemical Glycomodification of Proteins. <i>European Journal of Organic Chemistry</i> , 2018, 2018, 4296-4305.	1.2	7
92	Inducible Genetic Code Expansion in Eukaryotes. <i>ChemBioChem</i> , 2020, 21, 3216-3219.	1.3	7
93	Role of Solvent Compatibility in the Phase Behavior of Binary Solutions of Weakly Associating Multivalent Polymers. <i>Biomacromolecules</i> , 2022, 23, 349-364.	2.6	7
94	Condensed, Microtubule-coating Thin Organelles for Orthogonal Translation in Mammalian Cells. <i>Journal of Molecular Biology</i> , 2022, 434, 167454.	2.0	6
95	There is plenty of room in protein-RNA condensates. <i>Biophysical Journal</i> , 2021, 120, 1121-1122.	0.2	5
96	MultiBacTAG-Genetic Code Expansion Using the Baculovirus Expression System in Sf21 Cells. <i>Methods in Molecular Biology</i> , 2018, 1728, 297-311.	0.4	4
97	Fluorogenic Tetrazine-Siliconrhodamine Probe for the Labeling of Noncanonical Amino Acid Tagged Proteins. <i>Methods in Molecular Biology</i> , 2018, 1728, 337-363.	0.4	2
98	Mechanismusabhängige Regulation der ultraschnellen Dynamik von Wasser an Grenzflächen in Komplexen mit intrinsisch ungeordneten Proteinen. <i>Angewandte Chemie</i> , 2019, 131, 4769-4774.	1.6	2
99	Phase Separation Comes of Age: From Phenomenology to Single Molecules. <i>Molecular Cell</i> , 2019, 74, 413-415.	4.5	2
100	Probing Differential Binding Mechanisms of Phenylalanine-Glycine-Rich Nucleoporins by Single-Molecule FRET. <i>Methods in Enzymology</i> , 2018, 611, 327-346.	0.4	1
101	Faces, facets, and functions of biomolecular condensates driven by multivalent proteins and nucleic acids. <i>Biophysical Journal</i> , 2021, 120, E1-E4.	0.2	1
102	When two become one: Integrating FRET and EPR into one structural model. <i>Biophysical Journal</i> , 2021, 120, 4637-4638.	0.2	1
103	Frontispiece: Highly Stable trans-Cyclooctene Amino Acids for Live-Cell Labeling. <i>Chemistry - A European Journal</i> , 2015, 21, n/a-n/a.	1.7	0
104	Titelbild: Verbesserte Erweiterung des eukaryotischen genetischen Codes für seitenspezifische, hochauflösende Click-PAINT-Mikroskopie (<i>Angew. Chem.</i> 52/2016). <i>Angewandte Chemie</i> , 2016, 128, 16163-16163.	1.6	0