

Stephan A Sieber

List of Publications by Year in descending order

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Version: 2024-02-01

130
papers

4,767
citations

101543
36
h-index

128289
60
g-index

143
all docs

143
docs citations

143
times ranked

5564
citing authors

#	ARTICLE	IF	CITATIONS
1	Chemical proteomics approaches for identifying the cellular targets of natural products. <i>Natural Product Reports</i> , 2016, 33, 681-708.	10.3	295
2	The microstructure and micromechanics of the tendonâ€“bone insertion. <i>Nature Materials</i> , 2017, 16, 664-670.	27.5	250
3	Electrophilic natural products and their biological targets. <i>Natural Product Reports</i> , 2012, 29, 659.	10.3	232
4	Proteomic profiling of metalloprotease activities with cocktails of active-site probes. , 2006, 2, 274-281.		224
5	β -Lactones as Specific Inhibitors of ClpP Attenuate the Production of Extracellular Virulence Factors of <i>< i>Staphylococcus aureus</i> . <i>Journal of the American Chemical Society</i> , 2008, 130, 14400-14401.	13.7	177
6	β -Lactones as Privileged Structures for the Activeâ€“Site Labeling of Versatile Bacterial Enzyme Classes. <i>Angewandte Chemie - International Edition</i> , 2008, 47, 4600-4603.	13.8	168
7	Thinking Outside the Boxâ€”Novel Antibacterials To Tackle the Resistance Crisis. <i>Angewandte Chemie - International Edition</i> , 2018, 57, 14440-14475.	13.8	129
8	AAA+ chaperones and acyldepsipeptides activate the ClpP protease via conformational control. <i>Nature Communications</i> , 2015, 6, 6320.	12.8	110
9	Phenyl Esters Are Potent Inhibitors of Caseinolytic Protease P and Reveal a Stereogenic Switch for Deoligomerization. <i>Journal of the American Chemical Society</i> , 2015, 137, 8475-8483.	13.7	89
10	A Whole Proteome Inventory of Background Photocrosslinker Binding. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 1396-1401.	13.8	87
11	Vibralactone as a Tool to Study the Activity and Structure of the ClpP1P2 Complex from <i>< i>Listeria monocytogenes</i> . <i>Angewandte Chemie - International Edition</i> , 2011, 50, 11001-11004.	13.8	80
12	Repurposing human kinase inhibitors to create an antibiotic active against drug-resistant <i>Staphylococcus aureus</i> , persisters and biofilms. <i>Nature Chemistry</i> , 2020, 12, 145-158.	13.6	78
13	Towards synthetic cells using peptide-based reaction compartments. <i>Nature Communications</i> , 2018, 9, 3862.	12.8	75
14	Duocarmycin Analogues Target Aldehyde Dehydrogenaseâ€“1 in Lung Cancer Cells. <i>Angewandte Chemie - International Edition</i> , 2012, 51, 2874-2877.	13.8	72
15	A Conformational Switch Underlies ClpP Protease Function. <i>Angewandte Chemie - International Edition</i> , 2011, 50, 5749-5752.	13.8	69
16	Cryo-EM structure of the ClpXP protein degradation machinery. <i>Nature Structural and Molecular Biology</i> , 2019, 26, 946-954.	8.2	68
17	The Heat Shock Response in Yeast Maintains Protein Homeostasis by Chaperoning and Replenishing Proteins. <i>Cell Reports</i> , 2019, 29, 4593-4607.e8.	6.4	67
18	Analytical platforms for activity-based protein profiling ? exploiting the versatility of chemistry for functional proteomics. <i>Chemical Communications</i> , 2006, , 2311.	4.1	64

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19	Activity-based Probes for Studying the Activity of Flavin-dependent Oxidases and for the Protein Target Profiling of Monoamine Oxidase Inhibitors. <i>Angewandte Chemie - International Edition</i> , 2012, 51, 7035-7040.	13.8	63
20	Promysalin Elicits Species-Selective Inhibition of <i>< i> Pseudomonas aeruginosa </i></i> by Targeting Succinate Dehydrogenase. <i>Journal of the American Chemical Society</i> , 2018, 140, 1774-1782.	13.7	63
21	Insights into Structural Network Responsible for Oligomerization and Activity of Bacterial Virulence Regulator Caseinolytic Protease P (ClpP) Protein. <i>Journal of Biological Chemistry</i> , 2012, 287, 9484-9494.	3.4	62
22	Microarray Platform for Profiling Enzyme Activities in Complex Proteomes. <i>Journal of the American Chemical Society</i> , 2004, 126, 15640-15641.	13.7	61
23	Structural and functional insights into caseinolytic proteases reveal an unprecedented regulation principle of their catalytic triad. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, 11302-11307.	7.1	60
24	Surface topology affects wetting behavior of <i>Bacillus subtilis</i> biofilms. <i>Npj Biofilms and Microbiomes</i> , 2017, 3, 11.	6.4	55
25	An Antibacterial β -Lactone Kills <i>Mycobacterium tuberculosis</i> by Disrupting Mycolic Acid Biosynthesis. <i>Angewandte Chemie - International Edition</i> , 2018, 57, 348-353.	13.8	55
26	The Mechanism of Caseinolytic Protease (ClpP) Inhibition. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 3009-3014.	13.8	53
27	Mining the cellular inventory of pyridoxal phosphate-dependent enzymes with functionalized cofactor mimics. <i>Nature Chemistry</i> , 2018, 10, 1234-1245.	13.6	51
28	Making a Long Journey Short: Alkyne Functionalization of Natural Product Scaffolds. <i>Chemistry - A European Journal</i> , 2016, 22, 4666-4678.	3.3	50
29	A cyanobacterial serine protease of <i>< i> Plasmodium falciparum </i></i> is targeted to the apicoplast and plays an important role in its growth and development. <i>Molecular Microbiology</i> , 2010, 77, 873-890.	2.5	48
30	Pretubulysin derived probes as novel tools for monitoring the microtubule network via activity-based protein profiling and fluorescence microscopy. <i>Molecular BioSystems</i> , 2012, 8, 2067.	2.9	48
31	Development and characterization of improved β -lactone-based anti-virulence drugs targeting ClpP. <i>Bioorganic and Medicinal Chemistry</i> , 2012, 20, 583-591.	3.0	47
32	Disruption of Oligomerization and Dehydroalanine Formation as Mechanisms for ClpP Protease Inhibition. <i>Journal of the American Chemical Society</i> , 2014, 136, 1360-1366.	13.7	47
33	Covalent Mucin Coatings Form Stable Anti-Biofouling Layers on a Broad Range of Medical Polymer Materials. <i>Advanced Materials Interfaces</i> , 2020, 7, 1902069.	3.7	43
34	Reversible Inhibitors Arrest ClpP in a Defined Conformational State that Can Be Revoked by ClpX Association. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 15892-15896.	13.8	42
35	Alkynol natural products target ALDH2 in cancer cells by irreversible binding to the active site. <i>Chemical Communications</i> , 2015, 51, 15784-15787.	4.1	42
36	Self-Assembled Palladium and Platinum Coordination Cages: Photophysical Studies and Anticancer Activity. <i>European Journal of Inorganic Chemistry</i> , 2016, 2016, 5189-5196.	2.0	40

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37	<scp>ECE</scp>2 regulates neurogenesis and neuronal migration during human cortical development. <i>EMBO Reports</i> , 2020, 21, e48204.	4.5	40
38	FICD activity and AMPylation remodelling modulate human neurogenesis. <i>Nature Communications</i> , 2020, 11, 517.	12.8	39
39	Electrophilic reactivities of cyclic enones and $\text{I}^{\pm}, \text{I}^2$ -unsaturated lactones. <i>Chemical Science</i> , 2021, 12, 4850-4865.	7.4	38
40	A chemical compound inhibiting the Aha1-Hsp90 chaperone complex. <i>Journal of Biological Chemistry</i> , 2017, 292, 17073-17083.	3.4	37
41	The Cytotoxic Natural Product Vioprolide...A Targets Nucleolar Protein 14, Which Is Essential for Ribosome Biogenesis. <i>Angewandte Chemie - International Edition</i> , 2020, 59, 1595-1600.	13.8	37
42	A I^2 -Based Antivulence Drug Ameliorates <i>S. aureus</i> Skin Infections in Mice. <i>ChemMedChem</i> , 2014, 9, 710-713.	3.2	35
43	Barrel-shaped ClpP Proteases Display Attenuated Cleavage Specificities. <i>ACS Chemical Biology</i> , 2016, 11, 389-399.	3.4	35
44	A Chemical Disruptor of the ClpX Chaperone Complex Attenuates the Virulence of Multidrug-Resistant <i>S. aureus</i> . <i>Angewandte Chemie - International Edition</i> , 2017, 56, 15746-15750.	13.8	34
45	Selective Activation of Human Caseinolytic Protease-P (ClpP). <i>Angewandte Chemie - International Edition</i> , 2018, 57, 14602-14607.	13.8	34
46	Influence of wing-tip substituents and reaction conditions on the structure, properties and cytotoxicity of Ag(<i>scp</i>) I and Au(<i>scp</i>) I bis(NHC) complexes. <i>Dalton Transactions</i> , 2017, 46, 2722-2735.	3.3	33
47	Polyamide/PEG Blends as Biocompatible Biomaterials for the Convenient Regulation of Cell Adhesion and Growth. <i>Macromolecular Rapid Communications</i> , 2019, 40, e1900091.	3.9	33
48	A network of chaperones prevents and detects failures in membrane protein lipid bilayer integration. <i>Nature Communications</i> , 2019, 10, 672.	12.8	33
49	Structure and Mechanism of the Caseinolytic Protease ClpP1/2 Heterocomplex from <i>Listeria monocytogenes</i> . <i>Angewandte Chemie - International Edition</i> , 2015, 54, 3598-3602.	13.8	32
50	Chemical Probes Unravel an Antimicrobial Defense Response Triggered by Binding of the Human Opioid Dynorphin to a Bacterial Sensor Kinase. <i>Journal of the American Chemical Society</i> , 2017, 139, 6152-6159.	13.7	32
51	Chemical Cross-Linking Enables Drafting ClpXP Proximity Maps and Taking Snapshots of In Situ Interaction Networks. <i>Cell Chemical Biology</i> , 2019, 26, 48-59.e7.	5.2	31
52	Rugulactone and its Analogues Exert Antibacterial Effects through Multiple Mechanisms Including Inhibition of Thiamine Biosynthesis. <i>ChemBioChem</i> , 2012, 13, 1439-1446.	2.6	28
53	I^{\pm} -Methylene- I^3 -butyrolactones attenuate <i>S. aureus</i> virulence by inhibition of transcriptional regulation. <i>Chemical Science</i> , 2014, 5, 1158.	7.4	27
54	Protein Reactivity of Natural Product-Derived I^3 -Butyrolactones. <i>Biochemistry</i> , 2011, 50, 910-916.	2.5	26

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55	Structural, Biochemical, and Computational Studies Reveal the Mechanism of Selective Aldehyde Dehydrogenase 1A1 Inhibition by Cytotoxic Duocarmycin Analogues. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 13550-13554.		13.8	25
56	Effective GTP-Replacing FtsZ Inhibitors and Antibacterial Mechanism of Action. <i>ACS Chemical Biology</i> , 2015, 10, 834-843.		3.4	25
57	Quantitative Map of β -Lactone-Induced Virulence Regulation. <i>Journal of Proteome Research</i> , 2017, 16, 1180-1192.		3.7	25
58	Topographical alterations render bacterial biofilms susceptible to chemical and mechanical stress. <i>Biomaterials Science</i> , 2019, 7, 220-232.		5.4	25
59	Insights into ClpXP proteolysis: heterooligomerization and partial deactivation enhance chaperone affinity and substrate turnover in <i>Listeria monocytogenes</i> . <i>Chemical Science</i> , 2017, 8, 1592-1600.		7.4	24
60	A Subfamily of Bacterial Ribokinases Utilizes a Hemithioacetal for Pyridoxal Phosphate Salvage. <i>Journal of the American Chemical Society</i> , 2014, 136, 4992-4999.		13.7	21
61	Eine Gesamtproteom-basierte Auflistung der Hintergrundbinder von Photovernetzern. <i>Angewandte Chemie</i> , 2017, 129, 1417-1422.		2.0	21
62	Design and synthesis of tailored human caseinolytic protease P inhibitors. <i>Chemical Communications</i> , 2018, 54, 9833-9836.		4.1	21
63	A Pronucleotide Probe for Live-Cell Imaging of Protein AMPylation. <i>ChemBioChem</i> , 2020, 21, 1285-1287.		2.6	21
64	Global Inventory of ClpP- and ClpX-Regulated Proteins in <i>< i>Staphylococcus aureus</i></i> . <i>Journal of Proteome Research</i> , 2021, 20, 867-879.		3.7	21
65	Chemical Phosphoproteomics Sheds New Light on the Targets and Modes of Action of AKT Inhibitors. <i>ACS Chemical Biology</i> , 2021, 16, 631-641.		3.4	21
66	Extracellular LGALS3BP regulates neural progenitor position and relates to human cortical complexity. <i>Nature Communications</i> , 2021, 12, 6298.		12.8	21
67	Synthesis of (\pm)-Spongiolactone Enabling Discovery of a More Potent Derivative. <i>Chemistry - A European Journal</i> , 2015, 21, 1425-1428.		3.3	20
68	An amino acid domino effect orchestrates ClpP's conformational states. <i>Current Opinion in Chemical Biology</i> , 2017, 40, 102-110.		6.1	20
69	Synthesis of ramariolide natural products and discovery of their targets in mycobacteria. <i>Chemical Communications</i> , 2017, 53, 107-110.		4.1	19
70	$\tilde{\text{O}}$ ber bisherige Denkweisen hinaus – neue Wirkstoffe zur $\tilde{\text{O}}$ berwindung der Antibiotika-Krise. <i>Angewandte Chemie</i> , 2018, 130, 14642-14682.		2.0	18
71	Fimbrolide Natural Products Disrupt Bioluminescence of <i>< i>Vibrio</i></i> By Targeting Autoinducer Biosynthesis and Luciferase Activity. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 1187-1191.		13.8	16
72	Broad Spectrum Antibiotic Xanthocillin X Effectively Kills <i>< i>Acinetobacter baumannii</i></i> <i>< i>via</i></i> Dysregulation of Heme Biosynthesis. <i>ACS Central Science</i> , 2021, 7, 488-498.		11.3	16

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73	Chemical Probe To Monitor the Parkinsonism-Associated Protein DJ-1 in Live Cells. <i>ACS Chemical Biology</i> , 2018, 13, 2016-2019.	3.4	15
74	From Young to Old: AMPylation Hits the Brain. <i>Cell Chemical Biology</i> , 2020, 27, 773-779.	5.2	15
75	Substrate Profiling of Mitochondrial Caseinolytic Protease P via a Site-Specific Photocrosslinking Approach. <i>Angewandte Chemie - International Edition</i> , 2022, 61, .	13.8	15
76	A Mass Spectrometry Platform for a Streamlined Investigation of Proteasome Integrity, Posttranslational Modifications, and Inhibitor Binding. <i>Chemistry and Biology</i> , 2015, 22, 404-411.	6.0	14
77	Natural Product-Inspired Aminoepoxybenzoquinones Kill Members of the Gram-Negative Pathogen <i>< i>Salmonella</i></i> by Attenuating Cellular Stress Response. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 14852-14857.	13.8	14
78	A strategy for dual inhibition of the proteasome and fatty acid synthase with belactosin C-orlistat hybrids. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 2901-2916.	3.0	14
79	Tailored Pyridoxal Probes Unravel Novel Cofactor-Dependent Targets and Antibiotic Hits in Critical Bacterial Pathogens. <i>Angewandte Chemie - International Edition</i> , 2022, 61, .	13.8	14
80	Customizing Functionalized Cofactor Mimics to Study the Human Pyridoxal 5'-Phosphate-Binding Proteome. <i>Cell Chemical Biology</i> , 2019, 26, 1461-1468.e7.	5.2	13
81	The Natural Product Elegaphenone Potentiates Antibiotic Effects against <i>< i>Pseudomonas aeruginosa</i></i> . <i>Angewandte Chemie - International Edition</i> , 2019, 58, 8581-8584.	13.8	13
82	Neocarzilin A Is a Potent Inhibitor of Cancer Cell Motility Targeting VAT-1 Controlled Pathways. <i>ACS Central Science</i> , 2019, 5, 1170-1178.	11.3	12
83	Profiling withanolide A for therapeutic targets in neurodegenerative diseases. <i>Bioorganic and Medicinal Chemistry</i> , 2019, 27, 2508-2520.	3.0	11
84	Biochemical and Proteomic Studies of Human Pyridoxal 5'-Phosphate-Binding Protein (PLPBP). <i>ACS Chemical Biology</i> , 2020, 15, 254-261.	3.4	11
85	Structure and Function of an Elongation Factor P Subfamily in Actinobacteria. <i>Cell Reports</i> , 2020, 30, 4332-4342.e5.	6.4	11
86	Quantitative chemoproteomic profiling reveals multiple target interactions of spongiolactone derivatives in leukemia cells. <i>Chemical Communications</i> , 2017, 53, 12818-12821.	4.1	10
87	Dual Inhibitor of <i>< i>Staphylococcus aureus</i></i> Virulence and Biofilm Attenuates Expression of Major Toxins and Adhesins. <i>Biochemistry</i> , 2018, 57, 1814-1820.	2.5	10
88	Tailored Peptide Phenyl Esters Block ClpXP Proteolysis by an Unusual Breakdown into a Heptamer-Hexamer Assembly. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 7127-7132.	13.8	10
89	Targeting the endoplasmic reticulum-mitochondria interface sensitizes leukemia cells to cytostatics. <i>Haematologica</i> , 2019, 104, 546-555.	3.5	10
90	Total Synthesis of the Cyclic Depsipeptide Vioprolide-D via its (<i>< i>Z</i></i>)Diastereoisomer. <i>Angewandte Chemie - International Edition</i> , 2020, 59, 12357-12361.	13.8	10

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91	A tailored phosphoaspartate probe unravels CprR as a response regulator in <i>< i>Pseudomonas aeruginosa</i></i> interkingdom signaling. <i>Chemical Science</i> , 2021, 12, 4763-4770.	7.4	10
92	Functionalised Cofactor Mimics for Interactome Discovery and Beyond. <i>Angewandte Chemie - International Edition</i> , 2022, , .	13.8	10
93	Fluorescent palladium(<i>< scp>ii</scp></i>) and platinum(<i>< scp>ii</scp></i>) NHC/1,2,3-triazole complexes: antiproliferative activity and selectivity against cancer cells. <i>Dalton Transactions</i> , 2021, 50, 2158-2166.	3.3	9
94	Activity-Based Protein Profiling in Bacteria. <i>Methods in Molecular Biology</i> , 2017, 1491, 57-74.	0.9	8
95	Degrasyn exhibits antibiotic activity against multi-resistant <i>< i>Staphylococcus aureus</i></i> by modifying several essential cysteines. <i>Chemical Communications</i> , 2020, 56, 2929-2932.	4.1	8
96	Natürliche Fimbrolide inhibieren Autoinduktorbiosynthese und Luziferaseaktivität und unterdrücken damit die Biolumineszenz in <i>< i>Vibrio</i></i> . <i>Angewandte Chemie</i> , 2016, 128, 1203-1207.	2.0	7
97	A Chemical Proteomic Analysis of Illudinâ€¢-Interacting Proteins. <i>Chemistry - A European Journal</i> , 2019, 25, 12644-12651.	3.3	7
98	MS-Based <i>< i>in Situ</i></i> Proteomics Reveals AMPylation of Host Proteins during Bacterial Infection. <i>ACS Infectious Diseases</i> , 2020, 6, 3277-3289.	3.8	7
99	Comparative Target Analysis of Chlorinated Biphenyl Antimicrobials Highlights MenG as a Molecular Target of Triclocarban. <i>Applied and Environmental Microbiology</i> , 2020, 86, .	3.1	7
100	Targeting the ER-Mitochondrial Interface of Cell Death Sensitizes Leukemia Cells Towards Cytostatics. <i>Blood</i> , 2016, 128, 2319-2319.	1.4	7
101	Self-Assembled Palladium and Platinum Coordination Cages: Photophysical Studies and Anticancer Activity. <i>European Journal of Inorganic Chemistry</i> , 2016, 2016, 5181-5181.	2.0	6
102	An Aromatic Hydroxyamide Attenuates Multiresistant <i>< i>Staphylococcus aureus</i></i> Toxin Expression. <i>Chemistry - A European Journal</i> , 2016, 22, 1622-1630.	3.3	6
103	Transcriptomic Profiling Suggests That Promysalin Alters the Metabolic Flux, Motility, and Iron Regulation in <i>< i>Pseudomonas putida</i></i> KT2440. <i>ACS Infectious Diseases</i> , 2018, 4, 1179-1187.	3.8	6
104	Eukaryotic catecholamine hormones influence the chemotactic control of <i>< i>Vibrio campbellii</i></i> by binding to the coupling protein CheW. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2022, 119, e2118227119.	7.1	6
105	Azidobupramine, an Antidepressant-Derived Bifunctional Neurotransmitter Transporter Ligand Allowing Covalent Labeling and Attachment of Fluorophores. <i>PLoS ONE</i> , 2016, 11, e0148608.	2.5	5
106	Hydantoin analogs inhibit the fully assembled ClpXP protease without affecting the individual peptidase and chaperone domains. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 7124-7127.	2.8	5
107	Acyldepsipeptide Probes Facilitate Specific Detection of Caseinolytic Proteaseâ€¢-P Independent of Its Oligomeric and Activity State. <i>ChemBioChem</i> , 2020, 21, 235-240.	2.6	5
108	Synthetic post-translational modifications of elongation factor P using the ligase EpmA. <i>FEBS Journal</i> , 2021, 288, 663-677.	4.7	5

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109	Total synthesis and mechanism of action of the antibiotic armeniaspirol A. <i>Chemical Science</i> , 2021, 12, 16023-16034.	7.4	5
110	Der zytotoxische Naturstoff Vioprolidâ€...A interagiert mit dem fÃ¼r die Ribosomenâ€Biogenese essentiellen nukleolÃ¤ren Protein 14. <i>Angewandte Chemie</i> , 2020, 132, 1611-1617.	2.0	4
111	Totalsynthese des cyclischen Dipeptids Vioprolidâ€...D Ã¼ber sein (Z)â€Diastereomer. <i>Angewandte Chemie</i> , 2020, 132, 12456-12460.	2.0	4
112	Inactivity of Peptidase ClpP Causes Primary Accumulation of Mitochondrial Disaggregase ClpX with Its Interacting Nucleoid Proteins, and of mtDNA. <i>Cells</i> , 2021, 10, 3354.	4.1	4
113	Ein antibakterielles β -Lacton bekämpft <i>Mycobacterium tuberculosis</i> durch Infiltration der Mykolsäurebiosynthese. <i>Angewandte Chemie</i> , 2018, 130, 354-359.	2.0	3
114	Selektive Aktivierung der humanen caseinolytischen Proteaseâ€...P (ClpP). <i>Angewandte Chemie</i> , 2018, 130, 14811-14816.	2.0	3
115	Broadâ€range metalloprotease profiling in plants uncovers immunity provided by defenceâ€related metalloenzyme. <i>New Phytologist</i> , 2022, 235, 1287-1301.	7.3	3
116	Mechanistic analysis of aliphatic β -lactones in <i>Vibrio harveyi</i> reveals a quorum sensing independent mode of action. <i>Chemical Communications</i> , 2016, 52, 11971-11974.	4.1	2
117	Verringerung der Virulenz von multiresistentem <i>Staphylococcus aureus</i> mithilfe eines chemischen Disruptors des ClpXâ€Chaperonâ€Komplexes. <i>Angewandte Chemie</i> , 2017, 129, 15952-15957.	2.0	2
118	Der Naturstoff Elegaphenon verstärkt antibiotische Effekte gegen <i>Pseudomonas aeruginosa</i> . <i>Angewandte Chemie</i> , 2019, 131, 8670-8674.	2.0	2
119	Tailored Cofactor Traps for the <i>in Situ</i> Detection of Hemithioacetal-Forming Pyridoxal Kinases. <i>ACS Chemical Biology</i> , 2020, 15, 3227-3234.	3.4	2
120	Tranylcypromine specificity for monoamine oxidase is limited by promiscuous protein labelling and lysosomal trapping. <i>RSC Chemical Biology</i> , 2020, 1, 209-213.	4.1	2
121	Substrate profiling of mitochondrial caseinolytic protease P via a siteâ€specific photocrosslinking approach. <i>Angewandte Chemie</i> , 0, . . .	2.0	2
122	Tailored Pyridoxal Probes Unravel Novel Cofactorâ€Dependent Targets and Antibiotic Hits in Critical Bacterial Pathogens. <i>Angewandte Chemie</i> , 0, . . .	2.0	2
123	Functionalised Cofactor Mimics for Interactome Discovery and Beyond. <i>Angewandte Chemie</i> , 0, . . .	2.0	2
124	<i>Listeria monocytogenes</i> utilizes the ClpP1/2 proteolytic machinery for fine-tuned substrate degradation at elevated temperatures. <i>RSC Chemical Biology</i> , 0, . . .	4.1	2
125	In Vesiculo Synthesis of Peptide Membrane Precursors for Autonomous Vesicle Growth. <i>Journal of Visualized Experiments</i> , 2019, . . .	0.3	1
126	Bifunctional Duocarmycin Analogues as Inhibitors of Protein Tyrosine Kinases. <i>Journal of Natural Products</i> , 2019, 82, 16-26.	3.0	1

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127	Small molecule inhibitors of the mitochondrial ClpXP protease possess cytostatic potential and re-sensitize chemo-resistant cancers. <i>Scientific Reports</i> , 2021, 11, 11185.	3.3	1
128	Knockout for malaria. <i>Nature Chemistry</i> , 2014, 6, 93-94.	13.6	0
129	Frontispiece: An Aromatic Hydroxyamide Attenuates Multiresistant <i>< i>Staphylococcus aureus</i></i> Toxin Expression. <i>Chemistry - A European Journal</i> , 2016, 22, .	3.3	0
130	Blockade der ClpXP-vermittelten Proteolyse mit markiert geschnittenen Peptid-Phenylestern durch den ungewöhnlichen Zerfall in eine Heptamer-Hexamer-Anordnung. <i>Angewandte Chemie</i> , 2019, 131, 7201-7206.	2.0	0