

# Herbert Waldmann

## List of Publications by Year in descending order

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667  
papers

41,599  
citations

2538

96  
h-index

6282

158  
g-index

862  
all docs

862  
docs citations

862  
times ranked

30930  
citing authors

#	ARTICLE	IF	CITATIONS
1	An Acylation Cycle Regulates Localization and Activity of Palmitoylated Ras Isoforms. <i>Science</i> , 2005, 307, 1746-1752.	6.0	761
2	Fluorogenic probes for live-cell imaging of the cytoskeleton. <i>Nature Methods</i> , 2014, 11, 731-733.	9.0	705
3	Activation of the Raf-MEK-ERK pathway is required for neutrophil extracellular trap formation. <i>Nature Chemical Biology</i> , 2011, 7, 75-77.	3.9	649
4	The Pictet-Spengler Reaction in Nature and in Organic Chemistry. <i>Angewandte Chemie - International Edition</i> , 2011, 50, 8538-8564.	7.2	581
5	Chemical Strategies for Generating Protein Biochips. <i>Angewandte Chemie - International Edition</i> , 2008, 47, 9618-9647.	7.2	551
6	Small molecule inhibition of the KRAS-PDE1 interaction impairs oncogenic KRAS signalling. <i>Nature</i> , 2013, 497, 638-642.	13.7	551
7	Charting biologically relevant chemical space: A structural classification of natural products (SCONP). <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005, 102, 17272-17277.	3.3	534
8	Highly enantioselective synthesis and cellular evaluation of spirooxindoles inspired by natural products. <i>Nature Chemistry</i> , 2010, 2, 735-740.	6.6	531
9	Biology-Oriented Synthesis. <i>Angewandte Chemie - International Edition</i> , 2011, 50, 10800-10826.	7.2	438
10	A functional screen implicates microRNA-138-dependent regulation of the depalmitoylation enzyme APT1 in dendritic spine morphogenesis. <i>Nature Cell Biology</i> , 2009, 11, 705-716.	4.6	437
11	Catalytic Enantioselective 1,3-Dipolar Cycloadditions of Azomethine Ylides for Biology-Oriented Synthesis. <i>Accounts of Chemical Research</i> , 2014, 47, 1296-1310.	7.6	418
12	Inhibitors of Protein Tyrosine Phosphatases: Next-Generation Drugs?. <i>Angewandte Chemie - International Edition</i> , 2005, 44, 3814-3839.	7.2	406
13	From Protein Domains to Drug Candidates – Natural Products as Guiding Principles in the Design and Synthesis of Compound Libraries. <i>Angewandte Chemie - International Edition</i> , 2002, 41, 2878.	7.2	397
14	The Palmitoylation Machinery Is a Spatially Organizing System for Peripheral Membrane Proteins. <i>Cell</i> , 2010, 141, 458-471.	13.5	393
15	Target Identification for Small Bioactive Molecules: Finding the Needle in the Haystack. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 2744-2792.	7.2	393
16	Small-molecule inhibition of APT1 affects Ras localization and signaling. <i>Nature Chemical Biology</i> , 2010, 6, 449-456.	3.9	353
17	The Scaffold Tree – Visualization of the Scaffold Universe by Hierarchical Scaffold Classification. <i>Journal of Chemical Information and Modeling</i> , 2007, 47, 47-58.	2.5	322
18	Charting, Navigating, and Populating Natural Product Chemical Space for Drug Discovery. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 5989-6001.	2.9	317

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19	Asymmetric Hetero Diels-Alder Reactions. <i>Synthesis</i> , 1994, 1994, 535-551.	1.2	307
20	Discovery of protein phosphatase inhibitor classes by biology-oriented synthesis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2006, 103, 10606-10611.	3.3	288
21	General Enantioselective C <sup>α</sup> H Activation with Efficiently Tunable Cyclopentadienyl Ligands. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 2429-2434.	7.2	287
22	Protecting Group Strategies in Organic Synthesis. <i>Angewandte Chemie International Edition in English</i> , 1996, 35, 2056-2083.	4.4	285
23	New Modalities for Challenging Targets in Drug Discovery. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 10294-10323.	7.2	275
24	Synthesis of Natural Product Inspired Compound Collections. <i>Angewandte Chemie - International Edition</i> , 2009, 48, 3224-3242.	7.2	272
25	Identification of Thiazolidinones Spiro-Fused to Indolinones as Potent and Selective Inhibitors of the <i>Mycobacterium tuberculosis</i> Protein Tyrosine Phosphatase... <i>B. Angewandte Chemie - International Edition</i> , 2010, 49, 5902-5905.	7.2	261
26	Development of Tau Aggregation Inhibitors for Alzheimer's Disease. <i>Angewandte Chemie - International Edition</i> , 2009, 48, 1740-1752.	7.2	249
27	Natural-product-derived fragments for fragment-based ligand discovery. <i>Nature Chemistry</i> , 2013, 5, 21-28.	6.6	249
28	Arl2-GTP and Arl3-GTP regulate a GDI-like transport system for farnesylated cargo. <i>Nature Chemical Biology</i> , 2011, 7, 942-949.	3.9	231
29	Bioactivity-Guided Navigation of Chemical Space. <i>Accounts of Chemical Research</i> , 2010, 43, 1103-1114.	7.6	229
30	Rabbit muscle aldolase as a catalyst in organic synthesis. <i>Journal of the American Chemical Society</i> , 1989, 111, 627-635.	6.6	223
31	Interactive exploration of chemical space with Scaffold Hunter. <i>Nature Chemical Biology</i> , 2009, 5, 581-583.	3.9	207
32	Biology-Oriented Synthesis: Harnessing the Power of Evolution. <i>Journal of the American Chemical Society</i> , 2014, 136, 11853-11859.	6.6	207
33	Enzymic Protecting Group Techniques. <i>Chemical Reviews</i> , 1994, 94, 911-937.	23.0	199
34	The Allyl Group as Mildly and Selectively Removable Carboxy-Protecting Group for the Synthesis of Labile O-Glycopeptides. <i>Angewandte Chemie International Edition in English</i> , 1984, 23, 71-72.	4.4	198
35	Compound library development guided by protein structure similarity clustering and natural product structure. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2004, 101, 16721-16726.	3.3	198
36	The Asymmetric Hetero-Diels-Alder Reaction in the Syntheses of Biologically Relevant Compounds. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 11146-11157.	7.2	194

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37	Structure of Rab GDP-Dissociation Inhibitor in Complex with Prenylated YPT1 GTPase. <i>Science</i> , 2003, 302, 646-650.	6.0	193
38	Small-molecule modulation of Ras signaling. <i>Nature Chemical Biology</i> , 2014, 10, 613-622.	3.9	191
39	Staudinger Ligation: A New Immobilization Strategy for the Preparation of Small-Molecule Arrays. <i>Angewandte Chemie - International Edition</i> , 2003, 42, 5830-5834.	7.2	186
40	Photochemical Surface Patterning by the Thiol-Ene Reaction. <i>Angewandte Chemie - International Edition</i> , 2008, 47, 4421-4424.	7.2	179
41	Therapeutic intervention based on protein prenylation and associated modifications. <i>Nature Chemical Biology</i> , 2006, 2, 518-528.	3.9	176
42	Bioorganic synthesis of lipid-modified proteins for the study of signal transduction. <i>Nature</i> , 2000, 403, 223-226.	13.7	174
43	A framework for identification of actionable cancer genome dependencies in small cell lung cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 17034-17039.	3.3	167
44	( $\hat{\omega}$ )-Englerin A is a Potent and Selective Activator of TRPC4 and TRPC5 Calcium Channels. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 3787-3791.	7.2	161
45	Analysis of the eukaryotic prenylome by isoprenoid affinity tagging. <i>Nature Chemical Biology</i> , 2009, 5, 227-235.	3.9	160
46	Ras <sup>WT</sup> A Molecular Switch Involved in Tumor Formation. <i>Angewandte Chemie - International Edition</i> , 2000, 39, 4192-4214.	7.2	158
47	Bioorthogonal Chemistry for Site-Specific Labeling and Surface Immobilization of Proteins. <i>Accounts of Chemical Research</i> , 2011, 44, 762-773.	7.6	156
48	Biology-inspired synthesis of compound libraries. <i>Cellular and Molecular Life Sciences</i> , 2008, 65, 1186-1201.	2.4	150
49	Diels-Alder Ligation and Surface Immobilization of Proteins. <i>Angewandte Chemie - International Edition</i> , 2006, 45, 296-301.	7.2	149
50	Membrane-Mediated Induction and Sorting of K-Ras Microdomain Signaling Platforms. <i>Journal of the American Chemical Society</i> , 2011, 133, 880-887.	6.6	147
51	Rhodanine-Based Tau Aggregation Inhibitors in Cell Models of Tauopathy. <i>Angewandte Chemie - International Edition</i> , 2007, 46, 9215-9219.	7.2	145
52	The therapeutic potential of phosphatase inhibitors. <i>Current Opinion in Chemical Biology</i> , 2009, 13, 272-283.	2.8	144
53	Protein structure similarity clustering and natural product structure as guiding principles in drug discovery. <i>Drug Discovery Today</i> , 2005, 10, 471-483.	3.2	143
54	Lipidated Ras and Rab Peptides and Proteins <sup>WT</sup> Synthesis, Structure, and Function. <i>Angewandte Chemie - International Edition</i> , 2006, 45, 6622-6646.	7.2	137

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55	Identification of pyrazolopyridazinones as PDE $\epsilon$ inhibitors. Nature Communications, 2016, 7, 11360.	5.8	137
56	Enzymatic Protecting Group Techniques. Chemical Reviews, 2001, 101, 3367-3396.	23.0	136
57	Site-Selective Protein Immobilization by Staudinger Ligation. Angewandte Chemie - International Edition, 2006, 45, 1408-1412.	7.2	136
58	Organic Synthesis and Biological Signal Transduction. Angewandte Chemie - International Edition, 1998, 37, 688-749.	7.2	135
59	Principle and design of pseudo-natural products. Nature Chemistry, 2020, 12, 227-235.	6.6	134
60	Allyl esters as carboxy protecting groups in the synthesis of O-glycopeptides. Journal of Organic Chemistry, 1989, 54, 751-756.	1.7	133
61	N-Ras Forms Dimers at POPC Membranes. Biophysical Journal, 2012, 103, 1585-1593.	0.2	133
62	Solid-Phase Synthesis of Dysidiolide-Derived Protein Phosphatase Inhibitors. Journal of the American Chemical Society, 2002, 124, 13171-13178.	6.6	131
63	Natural Products Are Biologically Validated Starting Points in Structural Space for Compound Library Development: Solid-Phase Synthesis of Dysidiolide-Derived Phosphatase Inhibitors This research was supported by the Fonds der Chemischen Industrie and the Bayer AG.. Angewandte Chemie - International Edition, 2002, 41, 307.	7.2	131
64	S-Acylation and Plasma Membrane Targeting of the Farnesylated Carboxyl-Terminal Peptide of N-ras in Mammalian Fibroblasts. Biochemistry, 1997, 36, 13102-13109.	1.2	129
65	Bioactivity-guided mapping and navigation of chemical space. Nature Chemical Biology, 2009, 5, 585-592.	3.9	129
66	Asymmetric tandem Mannich-Michael reactions of amino acid ester imines with Danishefsky's diene. Journal of Organic Chemistry, 1992, 57, 4444-4451.	1.7	127
67	3-Substituted indolizine-1-carbonitrile derivatives as phosphatase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2006, 16, 59-63.	1.0	127
68	Von Proteindomänen zu Wirkstoffkandidaten – Naturstoffe als Leitstrukturen für das Design und die Synthese von Substanzbibliotheken. Angewandte Chemie, 2002, 114, 3002.	1.6	126
69	Visualizing Association of N-Ras in Lipid Microdomains: Influence of Domain Structure and Interfacial Adsorption. Journal of the American Chemical Society, 2006, 128, 192-201.	6.6	125
70	Membrane targeting mechanism of Rab GTPases elucidated by semisynthetic protein probes. Nature Chemical Biology, 2010, 6, 534-540.	3.9	119
71	Amino Acid Esters: Versatile Chiral Auxiliary Groups for the Asymmetric Synthesis of Nitrogen Heterocycles. Synlett, 1995, 1995, 133-141.	1.0	118
72	Impairment of prostate cancer cell growth by a selective and reversible lysine-specific demethylase 1 inhibitor. International Journal of Cancer, 2012, 131, 2704-2709.	2.3	118

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73	Total Synthesis and Biological Evaluation of the Nakijiquinones. <i>Journal of the American Chemical Society</i> , 2001, 123, 11586-11593.	6.6	117
74	General Enantioselective C-H Activation with Efficiently Tunable Cyclopentadienyl Ligands. <i>Angewandte Chemie</i> , 2017, 129, 2469-2474.	1.6	117
75	Structure of Rab Escort Protein-1 in Complex with Rab Geranylgeranyltransferase. <i>Molecular Cell</i> , 2003, 11, 483-494.	4.5	116
76	Natural product-inspired cascade synthesis yields modulators of centrosome integrity. <i>Nature Chemical Biology</i> , 2012, 8, 179-184.	3.9	116
77	Highly Enantioselective Catalytic [6+3] Cycloadditions of Azomethine Ylides. <i>Angewandte Chemie - International Edition</i> , 2012, 51, 9512-9516.	7.2	115
78	A crowdsourcing evaluation of the NIH chemical probes. <i>Nature Chemical Biology</i> , 2009, 5, 441-447.	3.9	111
79	Catalytic Enantioselective Synthesis of Functionalized Tropanes Reveals Novel Inhibitors of Hedgehog Signaling. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 12892-12896.	7.2	111
80	Cheminformatic Analysis of Natural Products and their Chemical Space. <i>Chimia</i> , 2007, 61, 355-360.	0.3	109
81	Choline-releasing glycerophosphodiesterase EDI3 drives tumor cell migration and metastasis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 8155-8160.	3.3	109
82	Influence of the Lipidation Motif on the Partitioning and Association of N-Ras in Model Membrane Subdomains. <i>Journal of the American Chemical Society</i> , 2009, 131, 1557-1564.	6.6	108
83	Enantioselective Formal C(sp <sup>3</sup> )-H Bond Activation in the Synthesis of Bioactive Spiropyrazolone Derivatives. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 307-311.	7.2	108
84	N,N-Phthaloylamino Acids as Chiral Auxiliaries in Asymmetric Mannich-Type Reactions. <i>Angewandte Chemie - International Edition</i> , 1999, 38, 184-187.	7.2	107
85	Revealing conformational substates of lipidated N-Ras protein by pressure modulation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 460-465.	3.3	106
86	Structure of doubly prenylated Ypt1:GDI complex and the mechanism of GDI-mediated Rab recycling. <i>EMBO Journal</i> , 2006, 25, 13-23.	3.5	103
87	Small-Molecule Inhibitors of Islet Amyloid Polypeptide Fibril Formation. <i>Angewandte Chemie - International Edition</i> , 2008, 47, 4679-4682.	7.2	103
88	Applications of Protein Biochips in Biomedical and Biotechnological Research. <i>Angewandte Chemie - International Edition</i> , 2009, 48, 7744-7751.	7.2	103
89	Natural Product Synthesis on Polymeric Supports-Synthesis and Biological Evaluation of an Indolactam Library. <i>Angewandte Chemie - International Edition</i> , 1999, 38, 2902-2906.	7.2	102
90	The cholesterol membrane anchor of the Hedgehog protein confers stable membrane association to lipid-modified proteins. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2004, 101, 8531-8536.	3.3	102

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91	Identification and Structure of Small-Molecule Stabilizers of 14-3-3 Protein-Protein Interactions. <i>Angewandte Chemie - International Edition</i> , 2010, 49, 4129-4132.	7.2	102
92	Asymmetric Synthesis of Natural Product Inspired Tricyclic Benzopyrones by an Organocatalyzed Annulation Reaction. <i>Angewandte Chemie - International Edition</i> , 2008, 47, 6869-6872.	7.2	101
93	Synthesis of analogs of 1,3-dihydroxyacetone phosphate and glyceraldehyde 3-phosphate for use in studies of fructose-1,6-diphosphate aldolase. <i>Journal of Organic Chemistry</i> , 1988, 53, 3457-3465.	1.7	100
94	Interaction analysis of prenylated Rab GTPase with Rab escort protein and GDP dissociation inhibitor explains the need for both regulators. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2007, 104, 12294-12299.	3.3	99
95	Discovery of a New Class of Inhibitors of <i>Mycobacterium tuberculosis</i> Protein Tyrosine Phosphatase...B by Biology-Oriented Synthesis. <i>Angewandte Chemie - International Edition</i> , 2008, 47, 5973-5977.	7.2	98
96	Identification of Acyl Protein Thioesterases...1 and 2 as the Cellular Targets of the Ras Signaling Modulators Palmostatin...B and M. <i>Angewandte Chemie - International Edition</i> , 2011, 50, 9838-9842.	7.2	98
97	Synthesis and Structure-Activity Correlation of Natural-Product Inspired Cyclodepsipeptides Stabilizing F-Actin. <i>Journal of the American Chemical Society</i> , 2010, 132, 3063-3077.	6.6	97
98	Inhibition of Glucose Transporters and Glutaminase Synergistically Impairs Tumor Cell Growth. <i>Cell Chemical Biology</i> , 2019, 26, 1214-1228.e25.	2.5	97
99	An Oxidation-Labile Traceless Linker for Solid-Phase Synthesis. <i>Angewandte Chemie - International Edition</i> , 1999, 38, 1073-1077.	7.2	96
100	Ligand-Directed Divergent Synthesis of Carbo- and Heterocyclic Ring Systems. <i>Angewandte Chemie - International Edition</i> , 2018, 57, 5212-5226.	7.2	95
101	Allylester als selektiv abspaltbare Carboxyschutzgruppen in der Peptid- und N-Glycopeptidsynthese. <i>Liebigs Annalen Der Chemie</i> , 1983, 1983, 1712-1725.	0.8	94
102	Oriented Immobilization of Farnesylated Proteins by the Thiol-Ene Reaction. <i>Angewandte Chemie - International Edition</i> , 2010, 49, 1252-1257.	7.2	93
103	Small-Molecule Target Engagement in Cells. <i>Cell Chemical Biology</i> , 2016, 23, 435-441.	2.5	93
104	C-H Bond Activation for the Synthesis of Heterocyclic Atropisomers Yields Hedgehog Pathway Inhibitors. <i>Angewandte Chemie - International Edition</i> , 2018, 57, 14250-14254.	7.2	93
105	Biology-oriented synthesis of a natural-product inspired oxepane collection yields a small-molecule activator of the Wnt-pathway. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, 6805-6810.	3.3	91
106	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.	6.6	89
107	Identification and Specificity Profiling of Protein Prenyltransferase Inhibitors Using New Fluorescent Phosphoisoprenoids. <i>Journal of the American Chemical Society</i> , 2006, 128, 2822-2835.	6.6	88
108	Chemoenzymatic Synthesis of N-RasLipopeptides. <i>Journal of the American Chemical Society</i> , 1998, 120, 6889-6902.	6.6	87

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109	Lipid Modifications of a Ras Peptide Exhibit Altered Packing and Mobility versus Host Membrane as Detected by <sup>2</sup> H Solid-State NMR. <i>Journal of the American Chemical Society</i> , 2005, 127, 12263-12272.	6.6	86
110	Silver catalyzed cascade synthesis of alkaloid ring systems: concise total synthesis of faspaplysin, homofaspaplysin C and analogues. <i>Chemical Communications</i> , 2010, 46, 4622.	2.2	86
111	Membrane binding of lipidated Ras peptides and proteins – The structural point of view. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 2009, 1788, 273-288.	1.4	84
112	Orthogonal ring-closing alkyne and olefin metathesis for the synthesis of small GTPase-targeting bicyclic peptides. <i>Nature Communications</i> , 2016, 7, 11300.	5.8	84
113	Chromopyrones are pseudo natural product glucose uptake inhibitors targeting glucose transporters GLUT-1 and -3. <i>Nature Chemistry</i> , 2018, 10, 1103-1111.	6.6	84
114	Diels–Alder Ligation of Peptides and Proteins. <i>Chemistry - A European Journal</i> , 2006, 12, 6095-6109.	1.7	82
115	Synthesis of Functional Ras Lipoproteins and Fluorescent Derivatives. <i>Journal of the American Chemical Society</i> , 2001, 123, 1023-1035.	6.6	80
116	A Microarray Strategy for Mapping the Substrate Specificity of Protein Tyrosine Phosphatase. <i>Angewandte Chemie - International Edition</i> , 2007, 46, 7700-7703.	7.2	80
117	Asymmetric synthesis of bicyclic amino acid derivatives by Aza–Diels–Alder reactions in aqueous solution. <i>Liebigs Annalen Der Chemie</i> , 1991, 1991, 1045-1048.	0.8	79
118	Direct Targeting of Rab–GTPase–Effector Interactions. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 2498-2503.	7.2	79
119	Chiral Titanium Alkoxides as Catalysts for the Enantioselective Reduction of Ketones with Boranes. <i>Angewandte Chemie International Edition in English</i> , 1995, 34, 2005-2006.	4.4	78
120	Natural Product Guided Compound Library Development. <i>Current Medicinal Chemistry</i> , 2002, 9, 2129-2145.	1.2	78
121	Sulindac-Derived Ras Pathway Inhibitors Target the Ras–Raf Interaction and Downstream Effectors in the Ras Pathway. <i>Angewandte Chemie - International Edition</i> , 2004, 43, 454-458.	7.2	78
122	Brunsvicamides A–C: Sponge-Related Cyanobacterial Peptides with Mycobacterium tuberculosis Protein Tyrosine Phosphatase Inhibitory Activity. <i>Journal of Medicinal Chemistry</i> , 2006, 49, 4871-4878.	2.9	78
123	Protein Biochips: Oriented Surface Immobilization of Proteins. <i>Macromolecular Chemistry and Physics</i> , 2010, 211, 136-144.	1.1	78
124	A PDE1–Ras Inhibitor Chemotype with up to Seven H–Bonds and Picomolar Affinity that Prevents Efficient Inhibitor Release by Arl2. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 2423-2428.	7.2	78
125	Solid phase synthesis of cyclic peptides by oxidative cyclative cleavage of an aryl hydrazide linker – synthesis of stylostatin 1. <i>Tetrahedron Letters</i> , 2001, 42, 5677-5680.	0.7	77
126	Natural Product Derived Receptor Tyrosine Kinase Inhibitors: Identification of IGF1R, Tie-2, and VEGFR-3 Inhibitors. <i>Angewandte Chemie - International Edition</i> , 2002, 41, 1174-1178.	7.2	77



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127	Asymmetric Solid-Phase Synthesis of 6,6-Spiroketal. <i>Angewandte Chemie - International Edition</i> , 2004, 43, 3195-3199.	7.2	77
128	Picomolar, selective, and subtype-specific small-molecule inhibition of TRPC1/4/5 channels. <i>Journal of Biological Chemistry</i> , 2017, 292, 8158-8173.	1.6	77
129	Total Synthesis and Biological Evaluation of (âˆ“)â€Englerinâ€…A and B: Synthesis of Analogues with Improved Activity Profile. <i>Angewandte Chemie - International Edition</i> , 2011, 50, 3998-4002.	7.2	76
130	Development of a Natural-Product-Derived Chemical Toolbox for Modulation of Protein Function. <i>Chemical Reviews</i> , 2014, 114, 4621-4639.	23.0	76
131	Programmable enantioselective one-pot synthesis of molecules with eight stereocenters. <i>Nature Chemical Biology</i> , 2012, 8, 428-430.	3.9	75
132	Membrane Insertion of a Lipidated Ras Peptide Studied by FTIR, Solid-State NMR, and Neutron Diffraction Spectroscopyâ€. <i>Journal of the American Chemical Society</i> , 2003, 125, 4070-4079.	6.6	74
133	An Enzyme-Labile Linker Group for Organic Syntheses on Solid Supports. <i>Angewandte Chemie - International Edition</i> , 1998, 37, 1143-1146.	7.2	73
134	Manumycin A and Its Analogues Are Irreversible Inhibitors of Neutral Sphingomyelinase. <i>ChemBioChem</i> , 2001, 2, 141-143.	1.3	73
135	Synthesis of the Rheb and Kâ€Ras4B GTPases. <i>Angewandte Chemie - International Edition</i> , 2010, 49, 6090-6095.	7.2	73
136	Pseudo Natural Productsâ€”Chemical Evolution of Natural Product Structure. <i>Angewandte Chemie - International Edition</i> , 2021, 60, 15705-15723.	7.2	73
137	(S)-Proline benzyl ester as chiral auxiliary in Lewis acid catalyzed asymmetric Diels-Alder reactions. <i>Journal of Organic Chemistry</i> , 1988, 53, 6133-6136.	1.7	72
138	Discovery of Neuritogenic Compound Classes Inspired by Natural Products. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 9576-9581.	7.2	72
139	An Enantioselective Inverseâ€Electronâ€Demand Imino Dielsâ€Alder Reaction. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 2134-2137.	7.2	72
140	Development of Natural Product-Derived Receptor Tyrosine Kinase Inhibitors Based on Conservation of Protein Domain Fold. <i>Journal of Medicinal Chemistry</i> , 2003, 46, 2917-2931.	2.9	71
141	Total Synthesis and Biological Evaluation of the Protein Phosphatase 2A Inhibitor Cytostatin and Analogues. <i>Chemistry - A European Journal</i> , 2004, 10, 2759-2780.	1.7	71
142	Design of compound libraries based on natural product scaffolds and protein structure similarity clustering (PSSC). <i>Molecular BioSystems</i> , 2005, 1, 36.	2.9	71
143	Design, Synthesis, and Phenotypic Profiling of Pyranoâ€Furoâ€Pyridone Pseudo Natural Products. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 14715-14723.	7.2	71
144	Aldolase-catalyzed synthesis of complex C8 and C9 monosaccharides.. <i>Tetrahedron Letters</i> , 1986, 27, 5807-5810.	0.7	70

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145	Chemoenzymatic Synthesis of a Characteristic Phosphorylated and Glycosylated Peptide Fragment of the Large Subunit of Mammalian RNA Polymerase II. <i>Journal of the American Chemical Society</i> , 1997, 119, 6702-6710.	6.6	70
146	The Autodepalmitoylating Activity of APT Maintains the Spatial Organization of Palmitoylated Membrane Proteins. <i>Biophysical Journal</i> , 2014, 106, 93-105.	0.2	70
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