

Herbert Waldmann

List of Publications by Citations

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754
papers

34,966
citations

91
h-index

147
g-index

862
ext. papers

38,368
ext. citations

8
avg, IF

7.43
L-index

#	Paper	IF	Citations
754	An acylation cycle regulates localization and activity of palmitoylated Ras isoforms. <i>Science</i> , 2005 , 307, 1746-52	33.3	679
753	Fluorogenic probes for live-cell imaging of the cytoskeleton. <i>Nature Methods</i> , 2014 , 11, 731-3	21.6	507
752	Chemical strategies for generating protein biochips. <i>Angewandte Chemie - International Edition</i> , 2008 , 47, 9618-47	16.4	507
751	The Pictet-Spengler reaction in nature and in organic chemistry. <i>Angewandte Chemie - International Edition</i> , 2011 , 50, 8538-64	16.4	492
750	Highly enantioselective synthesis and cellular evaluation of spirooxindoles inspired by natural products. <i>Nature Chemistry</i> , 2010 , 2, 735-40	17.6	492
749	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, 17272-7	11.5	489
748	Small molecule inhibition of the KRAS-PDE α interaction impairs oncogenic KRAS signalling. <i>Nature</i> , 2013 , 497, 638-42	50.4	460
747	Activation of the Raf-MEK-ERK pathway is required for neutrophil extracellular trap formation. <i>Nature Chemical Biology</i> , 2011 , 7, 75-7	11.7	431
746	A functional screen implicates microRNA-138-dependent regulation of the depalmitoylation enzyme APT1 in dendritic spine morphogenesis. <i>Nature Cell Biology</i> , 2009 , 11, 705-16	23.4	396
745	Biology-oriented synthesis. <i>Angewandte Chemie - International Edition</i> , 2011 , 50, 10800-26	16.4	376
744	Inhibitors of protein tyrosine phosphatases: next-generation drugs?. <i>Angewandte Chemie - International Edition</i> , 2005 , 44, 3814-39	16.4	371
743	Catalytic enantioselective 1,3-dipolar cycloadditions of azomethine ylides for biology-oriented synthesis. <i>Accounts of Chemical Research</i> , 2014 , 47, 1296-310	24.3	346
742	Target identification for small bioactive molecules: finding the needle in the haystack. <i>Angewandte Chemie - International Edition</i> , 2013 , 52, 2744-92	16.4	339
741	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, 2879-90	16.4	338
740	The palmitoylation machinery is a spatially organizing system for peripheral membrane proteins. <i>Cell</i> , 2010 , 141, 458-71	56.2	326
739	The scaffold tree--visualization of the scaffold universe by hierarchical scaffold classification. <i>Journal of Chemical Information and Modeling</i> , 2007 , 47, 47-58	6.1	288
738	Small-molecule inhibition of APT1 affects Ras localization and signaling. <i>Nature Chemical Biology</i> , 2010 , 6, 449-56	11.7	287

737	Asymmetric Hetero Diels-Alder Reactions. <i>Synthesis</i> , 1994 , 1994, 535-551	2.9	279
736	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, 10606-11	11.5	271
735	Charting, navigating, and populating natural product chemical space for drug discovery. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 5989-6001	8.3	266
734	Synthesis of natural product inspired compound collections. <i>Angewandte Chemie - International Edition</i> , 2009 , 48, 3224-42	16.4	253
733	Protecting Group Strategies in Organic Synthesis. <i>Angewandte Chemie International Edition in English</i> , 1996 , 35, 2056-2083		235
732	Identification of thiazolidinones spiro-fused to indolin-2-ones as potent and selective inhibitors of the Mycobacterium tuberculosis protein tyrosine phosphatase B. <i>Angewandte Chemie - International Edition</i> , 2010 , 49, 5902-5	16.4	226
731	General Enantioselective C-H Activation with Efficiently Tunable Cyclopentadienyl Ligands. <i>Angewandte Chemie - International Edition</i> , 2017 , 56, 2429-2434	16.4	223
730	Development of tau aggregation inhibitors for Alzheimer's disease. <i>Angewandte Chemie - International Edition</i> , 2009 , 48, 1740-52	16.4	211
729	Natural-product-derived fragments for fragment-based ligand discovery. <i>Nature Chemistry</i> , 2013 , 5, 21-8	17.6	207
728	Bioactivity-guided navigation of chemical space. <i>Accounts of Chemical Research</i> , 2010 , 43, 1103-14	24.3	202
727	Arl2-GTP and Arl3-GTP regulate a GDI-like transport system for farnesylated cargo. <i>Nature Chemical Biology</i> , 2011 , 7, 942-9	11.7	199
726	Rabbit muscle aldolase as a catalyst in organic synthesis. <i>Journal of the American Chemical Society</i> , 1989 , 111, 627-635	16.4	195
725	New Modalities for Challenging Targets in Drug Discovery. <i>Angewandte Chemie - International Edition</i> , 2017 , 56, 10294-10323	16.4	193
724	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, 16721-6	11.5	191
723	Structure of Rab GDP-dissociation inhibitor in complex with prenylated YPT1 GTPase. <i>Science</i> , 2003 , 302, 646-50	33.3	187
722	Interactive exploration of chemical space with Scaffold Hunter. <i>Nature Chemical Biology</i> , 2009 , 5, 581-3	11.7	180
721	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		176
720	Biology-oriented synthesis: harnessing the power of evolution. <i>Journal of the American Chemical Society</i> , 2014 , 136, 11853-9	16.4	171

7 ¹⁹	Photochemical surface patterning by the thiol-ene reaction. <i>Angewandte Chemie - International Edition</i> , 2008 , 47, 4421-4	16.4	169
7 ¹⁸	Staudinger ligation: a new immobilization strategy for the preparation of small-molecule arrays. <i>Angewandte Chemie - International Edition</i> , 2003 , 42, 5830-4	16.4	166
7 ¹⁷	Small-molecule modulation of Ras signaling. <i>Nature Chemical Biology</i> , 2014 , 10, 613-22	11.7	161
7 ¹⁶	Bioorganic synthesis of lipid-modified proteins for the study of signal transduction. <i>Nature</i> , 2000 , 403, 223-6	50.4	160
7 ¹⁵	Enzymic Protecting Group Techniques. <i>Chemical Reviews</i> , 1994 , 94, 911-937	68.1	160
7 ¹⁴	The asymmetric hetero-Diels-Alder reaction in the syntheses of biologically relevant compounds. <i>Angewandte Chemie - International Edition</i> , 2014 , 53, 11146-57	16.4	159
7 ¹³	Therapeutic intervention based on protein prenylation and associated modifications. <i>Nature Chemical Biology</i> , 2006 , 2, 518-28	11.7	155
7 ¹²	Analysis of the eukaryotic prenylome by isoprenoid affinity tagging. <i>Nature Chemical Biology</i> , 2009 , 5, 227-35	11.7	145
7 ¹¹	Diels-Alder ligation and surface immobilization of proteins. <i>Angewandte Chemie - International Edition</i> , 2005 , 45, 296-301	16.4	143
7 ¹⁰	Bioorthogonal chemistry for site-specific labeling and surface immobilization of proteins. <i>Accounts of Chemical Research</i> , 2011 , 44, 762-73	24.3	141
7 ⁰⁹	Biology-inspired synthesis of compound libraries. <i>Cellular and Molecular Life Sciences</i> , 2008 , 65, 1186-2010.3	10.3	140
7 ⁰⁸	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, 17034-9	11.5	138
7 ⁰⁷	(-)-Englerin A is a potent and selective activator of TRPC4 and TRPC5 calcium channels. <i>Angewandte Chemie - International Edition</i> , 2015 , 54, 3787-91	16.4	136
7 ⁰⁶	The therapeutic potential of phosphatase inhibitors. <i>Current Opinion in Chemical Biology</i> , 2009 , 13, 272-83	8.7	135
7 ⁰⁵	Ras-A Molecular Switch Involved in Tumor Formation. <i>Angewandte Chemie - International Edition</i> , 2000 , 39, 4192-4214	16.4	135
7 ⁰⁴	Die Pictet-Spengler-Reaktion in der Natur und der organischen Chemie. <i>Angewandte Chemie</i> , 2011 , 123, 8692-8719	3.6	134
7 ⁰³	Protein structure similarity clustering and natural product structure as guiding principles in drug discovery. <i>Drug Discovery Today</i> , 2005 , 10, 471-83	8.8	134
7 ⁰²	Site-selective protein immobilization by Staudinger ligation. <i>Angewandte Chemie - International Edition</i> , 2006 , 45, 1408-12	16.4	129

701	Lipidated ras and rab peptides and proteins--synthesis, structure, and function. <i>Angewandte Chemie - International Edition</i> , 2006 , 45, 6622-46	16.4	124
700	Rhodanine-based tau aggregation inhibitors in cell models of tauopathy. <i>Angewandte Chemie - International Edition</i> , 2007 , 46, 9215-9	16.4	123
699	Enzymatic protecting group techniques. <i>Chemical Reviews</i> , 2001 , 101, 3367-96	68.1	123
698	Organic Synthesis and Biological Signal Transduction. <i>Angewandte Chemie - International Edition</i> , 1998 , 37, 688-749	16.4	121
697	Bioactivity-guided mapping and navigation of chemical space. <i>Nature Chemical Biology</i> , 2009 , 5, 585-92	11.7	120
696	Solid-phase synthesis of dysidiolide-derived protein phosphatase inhibitors. <i>Journal of the American Chemical Society</i> , 2002 , 124, 13171-8	16.4	120
695	S-Acylation and plasma membrane targeting of the farnesylated carboxyl-terminal peptide of N-ras in mammalian fibroblasts. <i>Biochemistry</i> , 1997 , 36, 13102-9	3.2	117
694	Visualizing association of N-ras in lipid microdomains: influence of domain structure and interfacial adsorption. <i>Journal of the American Chemical Society</i> , 2006 , 128, 192-201	16.4	117
693	Biologie-orientierte Synthese (BIOS). <i>Angewandte Chemie</i> , 2011 , 123, 10990-11018	3.6	116
692	Membrane-mediated induction and sorting of K-Ras microdomain signaling platforms. <i>Journal of the American Chemical Society</i> , 2011 , 133, 880-7	16.4	116
691	Allyl esters as carboxy protecting groups in the synthesis of O-glycopeptides. <i>Journal of Organic Chemistry</i> , 1989 , 54, 751-756	4.2	115
690	N-Ras forms dimers at POPC membranes. <i>Biophysical Journal</i> , 2012 , 103, 1585-93	2.9	113
689	Asymmetric tandem Mannich-Michael reactions of amino acid ester imines with Danishefsky's diene. <i>Journal of Organic Chemistry</i> , 1992 , 57, 4444-4451	4.2	111
688	General Enantioselective C-H Activation with Efficiently Tunable Cyclopentadienyl Ligands. <i>Angewandte Chemie</i> , 2017 , 129, 2469-2474	3.6	109
687	Natural products are biologically validated starting points in structural space for compound library development: solid-phase synthesis of dysidiolide-derived phosphatase inhibitors. <i>Angewandte Chemie - International Edition</i> , 2002 , 41, 307-11	16.4	109
686	Natural product-inspired cascade synthesis yields modulators of centrosome integrity. <i>Nature Chemical Biology</i> , 2011 , 8, 179-84	11.7	108
685	Amino Acid Esters: Versatile Chiral Auxiliary Groups for the Asymmetric Synthesis of Nitrogen Heterocycles. <i>Synlett</i> , 1995 , 1995, 133-141	2.2	108
684	3-Substituted indolizine-1-carbonitrile derivatives as phosphatase inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006 , 16, 59-63	2.9	107

683	Identification of pyrazolopyridazinones as PDE inhibitors. <i>Nature Communications</i> , 2016 , 7, 11360	17.4	106
682	Total synthesis and biological evaluation of the nakijiquinones. <i>Journal of the American Chemical Society</i> , 2001 , 123, 11586-93	16.4	105
681	Structure of Rab escort protein-1 in complex with Rab geranylgeranyltransferase. <i>Molecular Cell</i> , 2003 , 11, 483-94	17.6	101
680	Highly enantioselective catalytic [6+3] cycloadditions of azomethine ylides. <i>Angewandte Chemie - International Edition</i> , 2012 , 51, 9512-6	16.4	100
679	Impairment of prostate cancer cell growth by a selective and reversible lysine-specific demethylase 1 inhibitor. <i>International Journal of Cancer</i> , 2012 , 131, 2704-9	7.5	98
678	Membrane targeting mechanism of Rab GTPases elucidated by semisynthetic protein probes. <i>Nature Chemical Biology</i> , 2010 , 6, 534-40	11.7	98
677	Von Proteindomänen zu Wirkstoffkandidaten – Naturstoffe als Leitstrukturen für das Design und die Synthese von Substanzbibliotheken. <i>Angewandte Chemie</i> , 2002 , 114, 3002	3.6	98
676	A crowdsourcing evaluation of the NIH chemical probes. <i>Nature Chemical Biology</i> , 2009 , 5, 441-7	11.7	97
675	Small-molecule inhibitors of islet amyloid polypeptide fibril formation. <i>Angewandte Chemie - International Edition</i> , 2008 , 47, 4679-82	16.4	97
674	Die Synthese von naturstoffinspirierten Verbindungsbibliotheken. <i>Angewandte Chemie</i> , 2009 , 121, 3272-3290	3.2	96
673	Catalytic enantioselective synthesis of functionalized tropanes reveals novel inhibitors of hedgehog signaling. <i>Angewandte Chemie - International Edition</i> , 2013 , 52, 12892-6	16.4	95
672	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, 1557-64	16.4	94
671	Applications of protein biochips in biomedical and biotechnological research. <i>Angewandte Chemie - International Edition</i> , 2009 , 48, 7744-51	16.4	93
670	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, 460-5	11.5	93
669	Asymmetric synthesis of natural product inspired tricyclic benzopyrones by an organocatalyzed annulation reaction. <i>Angewandte Chemie - International Edition</i> , 2008 , 47, 6869-72	16.4	93
668	Cheminformatic Analysis of Natural Products and their Chemical Space. <i>Chimia</i> , 2007 , 61, 355-360	1.3	93
667	N,N-Phthaloylamino Acids as Chiral Auxiliaries in Asymmetric Mannich-Type Reactions. <i>Angewandte Chemie - International Edition</i> , 1999 , 38, 184-187	16.4	93
666	Natural Product Synthesis on Polymeric Supports-Synthesis and Biological Evaluation of an Indolactam Library. <i>Angewandte Chemie - International Edition</i> , 1999 , 38, 2902-2906	16.4	93

665	Identification and structure of small-molecule stabilizers of 14-3-3 protein-protein interactions. <i>Angewandte Chemie - International Edition</i> , 2010 , 49, 4129-32	16.4	92
664	Structure of doubly prenylated Ypt1:GDI complex and the mechanism of GDI-mediated Rab recycling. <i>EMBO Journal</i> , 2006 , 25, 13-23	13	92
663	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, 12294-9	11.5	91
662	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, 9838-42	16.4	90
661	Discovery of a new class of inhibitors of Mycobacterium tuberculosis protein tyrosine phosphatase B by biology-oriented synthesis. <i>Angewandte Chemie - International Edition</i> , 2008 , 47, 5973-7	16.4	90
660	Synthesis and structure-activity correlation of natural-product inspired cyclodepsipeptides stabilizing F-actin. <i>Journal of the American Chemical Society</i> , 2010 , 132, 3063-77	16.4	88
659	Oriented immobilization of farnesylated proteins by the thiol-ene reaction. <i>Angewandte Chemie - International Edition</i> , 2010 , 49, 1252-7	16.4	87
658	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, 8531-6	11.5	86
657	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	4.2	85
656	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, 6805-10	11.5	84
655	Die Allylgruppe als mild und selektiv abspaltbare Carboxy-Schutzgruppe zur Synthese empfindlicher O-Glycopeptide. <i>Angewandte Chemie</i> , 1984 , 96, 49-50	3.6	84
654	Lipid modifications of a Ras peptide exhibit altered packing and mobility versus host membrane as detected by 2H solid-state NMR. <i>Journal of the American Chemical Society</i> , 2005 , 127, 12263-72	16.4	82
653	Allylester als selektiv abspaltbare Carboxyschutzgruppen in der Peptid- und N-Glycopeptidsynthese. <i>Liebigs Annalen Der Chemie</i> , 1983 , 1983, 1712-1725		81
652	Orthogonal ring-closing alkyne and olefin metathesis for the synthesis of small GTPase-targeting bicyclic peptides. <i>Nature Communications</i> , 2016 , 7, 11300	17.4	79
651	BIOS: Similarity-based design of natural product derived compound collections. <i>Chemistry Central Journal</i> , 2008 , 2,		78
650	Silver catalyzed cascade synthesis of alkaloid ring systems: concise total synthesis of fascaplysin, homofascaplysin C and analogues. <i>Chemical Communications</i> , 2010 , 46, 4622-4	5.8	77
649	Diels-Alder ligation of peptides and proteins. <i>Chemistry - A European Journal</i> , 2006 , 12, 6095-109	4.8	77
648	Small-Molecule Target Engagement in Cells. <i>Cell Chemical Biology</i> , 2016 , 23, 435-41	8.2	77

- 647 Natural product guided compound library development. *Current Medicinal Chemistry*, **2002**, 9, 2129-45 4.3 76
- 646 Chemoenzymatic Synthesis of N-RasLipopeptides. *Journal of the American Chemical Society*, **1998**, 120, 6889-6902 16.4 76
- 645 A microarray strategy for mapping the substrate specificity of protein tyrosine phosphatase. *Angewandte Chemie - International Edition*, **2007**, 46, 7700-3 16.4 75
- 644 Synthesis of functional Ras lipoproteins and fluorescent derivatives. *Journal of the American Chemical Society*, **2001**, 123, 1023-35 16.4 75
- 643 An oxidation-labile traceless linker for solid-phase synthesis. *Angewandte Chemie - International Edition*, **1999**, 38, 1073-7 16.4 75
- 642 Protein Biochips: Oriented Surface Immobilization of Proteins. *Macromolecular Chemistry and Physics*, **2010**, 211, 136-144 2.6 74
- 641 Enantioselective Formal C(sp³)-H Bond Activation in the Synthesis of Bioactive Spiropyrazolone Derivatives. *Angewandte Chemie - International Edition*, **2019**, 58, 307-311 16.4 74
- 640 Asymmetric solid-phase synthesis of 6,6-spiroketal. *Angewandte Chemie - International Edition*, **2004**, 43, 3195-9 16.4 73
- 639 Membrane binding of lipidated Ras peptides and proteins--the structural point of view. *Biochimica Et Biophysica Acta - Biomembranes*, **2009**, 1788, 273-88 3.8 72
- 638 Membrane insertion of a lipidated ras peptide studied by FTIR, solid-state NMR, and neutron diffraction spectroscopy. *Journal of the American Chemical Society*, **2003**, 125, 4070-9 16.4 71
- 637 Development of a natural-product-derived chemical toolbox for modulation of protein function. *Chemical Reviews*, **2014**, 114, 4621-39 68.1 70
- 636 Programmable enantioselective one-pot synthesis of molecules with eight stereocenters. *Nature Chemical Biology*, **2012**, 8, 428-30 11.7 70
- 635 Phenyl Esters Are Potent Inhibitors of Caseinolytic Protease P and Reveal a Stereogenic Switch for Deoligomerization. *Journal of the American Chemical Society*, **2015**, 137, 8475-83 16.4 69
- 634 Choline-releasing glycerophosphodiesterase EDI3 drives tumor cell migration and metastasis. *Proceedings of the National Academy of Sciences of the United States of America*, **2012**, 109, 8155-60 11.5 69
- 633 Identification and specificity profiling of protein prenyltransferase inhibitors using new fluorescent phosphoisoprenoids. *Journal of the American Chemical Society*, **2006**, 128, 2822-35 16.4 69
- 632 Sulindac-derived Ras pathway inhibitors target the Ras-Raf interaction and downstream effectors in the Ras pathway. *Angewandte Chemie - International Edition*, **2004**, 43, 454-8 16.4 69
- 631 Inhibitoren der Proteintyrosinphosphatasen: Kandidaten für zukünftige Wirkstoffe?. *Angewandte Chemie*, **2005**, 117, 3880-3906 3.6 69
- 630 Solid phase synthesis of cyclic peptides by oxidative cyclative cleavage of an aryl hydrazide linker. Synthesis of stylostatin 1. *Tetrahedron Letters*, **2001**, 42, 5677-5680 2 69

629	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		69
628	Discovery of neuritogenic compound classes inspired by natural products. <i>Angewandte Chemie - International Edition</i> , 2013 , 52, 9576-81	16.4	68
627	Synthesis of the Rheb and K-Ras4B GTPases. <i>Angewandte Chemie - International Edition</i> , 2010 , 49, 6090-516.4	16.4	68
626	Development of natural product-derived receptor tyrosine kinase inhibitors based on conservation of protein domain fold. <i>Journal of Medicinal Chemistry</i> , 2003 , 46, 2917-31	8.3	68
625	Manumycin A and its analogues are irreversible inhibitors of neutral sphingomyelinase. <i>ChemBioChem</i> , 2001 , 2, 141-3	3.8	68
624	Total synthesis and biological evaluation of (-)-englerin A and B: synthesis of analogues with improved activity profile. <i>Angewandte Chemie - International Edition</i> , 2011 , 50, 3998-4002	16.4	67
623	Natural product derived receptor tyrosine kinase inhibitors: identification of IGF1R, Tie-2, and VEGFR-3 inhibitors. <i>Angewandte Chemie - International Edition</i> , 2002 , 41, 1174-8	16.4	67
622	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		67
621	Principle and design of pseudo-natural products. <i>Nature Chemistry</i> , 2020 , 12, 227-235	17.6	67
620	Chemical biology--identification of small molecule modulators of cellular activity by natural product inspired synthesis. <i>Chemical Society Reviews</i> , 2008 , 37, 1361-74	58.5	66
619	Brunsvicamides A-C: sponge-related cyanobacterial peptides with Mycobacterium tuberculosis protein tyrosine phosphatase inhibitory activity. <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 4871-8	8.3	66
618	C-H Bond Activation for the Synthesis of Heterocyclic Atropisomers Yields Hedgehog Pathway Inhibitors. <i>Angewandte Chemie - International Edition</i> , 2018 , 57, 14250-14254	16.4	66
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615	Total synthesis and biological evaluation of the protein phosphatase 2A inhibitor cytostatin and analogues. <i>Chemistry - A European Journal</i> , 2004 , 10, 2759-80	4.8	64
614	Protein structure similarity clustering (PSSC) and natural product structure as inspiration sources for drug development and chemical genomics. <i>Current Opinion in Chemical Biology</i> , 2005 , 9, 232-9	9.7	64
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- 609 A highly efficient strategy for modification of proteins at the C terminus. *Angewandte Chemie - International Edition*, **2010**, 49, 9417-21 16.4 62
- 608 Total synthesis of chondramide C and its binding mode to F-actin. *Angewandte Chemie - International Edition*, **2008**, 47, 6473-7 16.4 62
- 607 The lipidated membrane anchor of full length N-Ras protein shows an extensive dynamics as revealed by solid-state NMR spectroscopy. *Journal of the American Chemical Society*, **2006**, 128, 13840-6 16.4 62
- 606 Natural Product-Guided Synthesis of a Spiroacetal Collection Reveals Modulators of Tubulin Cytoskeleton Integrity. *European Journal of Organic Chemistry*, **2005**, 2005, 4773-4788 3.2 62
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- 598 An Enzyme-Labile Linker Group for Organic Syntheses on Solid Supports. *Angewandte Chemie - International Edition*, **1998**, 37, 1143-1146 16.4 59
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