



Complete List of Publications

Number of publications: > 700

Number of citations: > 34.000

H-Index: 92

H. Kunz, H. Waldmann	1,3-Dithian-2-yl-methylester als Zweistufenschutzgruppe für die Carboxylfunktion bei der Peptidsynthese	Angew. Chem.	1983	95	47
H. Waldmann, H. Kunz	Allylester als selektiv abspaltbare Carboxylschutzgruppen in der Peptid- und N-Glycopeptidsynthese	Liebigs Ann. Chem.	1983		1712
H. Kunz, H. Waldmann	Die Allylgruppe als selektiv abspaltbare Carboxyl- Schutzgruppe zur Synthese empfindlicher O-Glycopeptide	Angew. Chem.	1984	96	49
H. Kunz, H. Waldmann	Synthesis of the Glycopeptide Partial Sequence A ⁸⁰ -A ⁸⁴ of Human Fibroblast Interferon	Helv. Chim. Acta	1985	68	283
H. Kunz, H. Waldmann	Directed Stereoselective Synthesis of α - and β -N-Acetyl-Neuraminic Acid-Galactose Disaccharides Using 2-Chloro- and 2-Fluoro Derivatives of Neuraminic Acid Allyl Ester	J. Chem. Soc. Chem. Commun.	1985		638
H. Kunz, H. Waldmann, C. Unverzagt	The Allyl Ester as a Temporary Protecting Group for the β -Carboxy Function of Aspartic Acid	Int. J. Peptide Protein Res.	1985	26	493
H. Kunz, H. Waldmann	Aufbau disaccharidischer N-Glycopeptide - Synthese der Verknüpfungsregion der Transmembran-Neuraminidase eines Influenza Virus	Angew. Chem.	1985	97	885
H. Kunz, H. Waldmann, C. Unverzagt	Synthesis of Glycopeptides Using the Allyl Ester and the Allyloxycarbonyl Group as Protecting Functions	Peptides 1986, Ed. D. Theodoropoulos, W. de Gruyter, Berlin	1986		615
M. D. Bednarski, H. Waldmann, G. M. Whitesides	Aldolase-Catalyzed Synthesis of Complex C8 and C9 Monosaccharides	Tetrahedron Lett.	1986	27	5807
H. Waldmann, D. Gyax, M. D. Bednarski, W. R. Shangraw, G. M. Whitesides	The Enzymic Utilization of Sucrose in the Synthesis of Amylose and Derivatives of Amylose, Using Phosphorylases	Carbohydr. REs.	1986	157	C4-C7
A. Akiyama, M. D. Bednarski, M.-J. Kim, E. S. Simon, H. Waldmann, G. M.	Enzymes in Organic Synthesis	Chem. Brit.	1987	23	645



Whitesides

H. Waldmann, G. M. Whitesides	Enzymes in Organic Synthesis	Ullmanns Encyclopedia of Industrial Chemistry, 5th. Ed., Vol A9, Verlag Chemie, Weinheim	1987		341
N. Bischofberger, H. Waldmann, T. Saito, E. S. Simon, W. Lees, M. D. Bednarski, G. M. Whitesides	Synthesis of Analogues of 1,3-Dihydroxyacetone Phosphate and Glyceraldehyde-3- Phosphate for Use in Studies of Fructose-1,6-Diphosphate Aldolase	J. Org. Chem.	1988	53	3457
A. Akiyama, M. Bednarski, M.-J. Kim, E. S. Simon, H. Waldmann, G. M. Whitesides	Enzymes in Organic Synthesis	CHEMTECH	1988		627
H. Waldmann, H. Kunz	1,3-Dithian-2-yl-methyl Esters as Carboxy Protecting Groups in the Synthesis of N-Glycopeptides	J. Org. Chem.	1988	53	4172
H. Kunz, H. Waldmann, U. Klinkhammer	The Allyl Ester as Carboxy Protecting Group in the Stereoselective Construction of Neuraminic Acid Galactose	Helv. Chim. Acta	1988	71	1868
H. Waldmann	Aminosäuremethylester als chirale Auxiliare in Aza-Diels-Alder-Reaktionen in wässriger Lösung	Angew. Chemie	1988	100	307
H. Waldmann	The Use of Penicillin Acylase for Selective N-Terminal Deprotection in Peptide Synthesis	Tetrahedron Lett.	1988	29	1131
H. Waldmann	S)-Proline Benzyl Ester as Chiral Auxiliary in Lewis Acid Catalyzed Diels-Alder Reactions	J. Org. Chem.	1988	53	6133
H. Waldmann	Der Phenylacetyl-(PhAc)-Rest als enzymatisch ablösbare Schutzgruppe für Peptide und Kohlenhydrate: Selektive Schutzgruppenabspaltungen mit Penicillin Acylase	Liebigs Ann. Chem.	1988		1175
H. Waldmann	The Use of Penicillin Acylase for Selective N-Terminal Deprotection in Peptide Synthesis	Peptides 1988, W. de Gruyter, Berlin	1989		272
H. Kunz, S. Friedrich-Bochnitschek, H. Waldmann und C. Unverzagt	The Allyloxycarbonyl (Aloc) Group and Allyl Esters as Protecting Functions in the Synthesis of Peptides and Glycopeptides	Chemistry of Peptides and Proteins, Vol. 4, S. 119, Attempto Verlag, Tübingen	1989		



H. Kunz, H. Waldmann, J. März	Synthese von N-Glycopeptid-Partialstrukturen der Verknüpfungsregion sowohl der Transmembran-Neuraminidase eines Influenza-Virus als auch des Faktors B des menschlichen Komplementsystems	Liebigs Ann. Chem.	1989		45
S. Friedrich-Bochnitschek, H. Waldmann, H. Kunz	Allyl Esters as Carboxy Protecting Groups in the Synthesis of O-Glycopeptides	J. Org. Chem.	1989	54	751
M. Bednarski, E. S. Simon, N. Bischofberger, W.-D. Fessner, M.-J. Kim, W. Lees, T. Saito, H. Waldmann, G. M. Whitesides	Rabbit Muscle Aldolase as a Catalyst in Organic Synthesis	J. Am. Chem. Soc.	1989	111	627
H. Waldmann	Asymmetrische Hetero-Diels-Alder-Reaktionen in wässriger Lösung unter Verwendung von Aminosäureestern als chiralen Auxiliaren	Liebigs Ann. Chem.	1989		231
H. Waldmann	A New Access to Chiral 2-Furylcarbinols by Enantioselective Hydrolysis with Penicillin Acylase	Tetrahedron Lett.	1989	30	3057
H. Waldmann, M. Dräger	On the Enhancement of Stereoselection by Cooperation Between Chiral Auxiliaries. Asymmetric Diels-Alder Reactions with Fumaric Acid Bis((S)-Proline Benzyl Ester) Amide	Tetrahedron Lett.	1989	30	4227
H. Kunz, C. Unverzagt, B. Dombó, W. Kosch, H. Waldmann	Synthesis of Glycopeptide Partial Structures of Virus Coat Glycoproteins	Chemistry of Peptides and Proteins, Vol. 5	1989		
H. Kunz, P. Wernig, M. Schilling, J. März, C. Unverzagt, S. Birnbach, U. Lang, H. Waldmann	Synthetic Tumor-Associated Glycopeptide Antigens	Environmental Health Perspectives	1990	88	247-249
H. Waldmann, M. Braun, M. Dräger	Aminosäureester als chirale Hilfsgruppen in Lewis-Säure-katalysierten Umsetzungen elektronenreicher Siloxydiene mit Iminen	Angew. Chem.	1990	102	1445
H. Waldmann, J. März, H. Kunz	Synthesis of N-Acetylglucosaminylasparagine-Glycotri- and pentapeptides by Selective C- and N-terminal Elongation of the Peptide Chain	Carbohydr. Res.	1990	196	75-93



K. von dem Bruch, H. Waldmann, H. Kunz	Glycopeptide - Chemische Synthese biologischer Informationsträger und molekularer Erkennungsstrukturen	Forsch. der Universität Mainz	1990		57
H. Waldmann, G. M. Whitesides	Enzymes in Organic Synthesis	Enzymes in Industry" (W. Gerhartz, Hrsg.), Verlag Chemie, Weinheim	1990		141
M. Schultz, H. Waldmann, W. Vogt, H. Kunz	Stereospecific C-C-Bond Formation with Rabbit Muscle Aldolase - A Chemoenzymatic Synthesis of (+)-Exo-Brevicomine	Tetrahedron Lett.	1990	31	867
P. Braun, H. Waldmann, W. Vogt, H. Kunz	Selective Enzymatic Removal of Protecting Functions: n-Heptyl Esters as Carboxy Protecting Groups in Peptide Synthesis	SYNLETT	1990		105
H. Waldmann	Aminosäureester als chirale Hilfsgruppen in Lewis-Säure-katalysierten Diels-Alder-Reaktionen	Liebigs Ann. Chem.	1990		671
H. Waldmann, M. Dräger	Thermische Diels-Alder-Reaktionen mit N-(2-Alkenoyl)-(S)-prolinestern als chiralen Dienophilen in organischen und wässrigen Reaktionsmedien	Liebigs Ann. Chem.	1990		681
H. Waldmann	Proline Benzyl Ester as Chiral Auxiliary in Barbier-Type Reactions in Aqueous Solution	SYNLETT	1990		627
M. Schultz, H. Waldmann, H. Kunz, W. Vogt	Chemoenzymatische "Chiral-Pool"-Synthese von (+)-exo-Brevicomine aus Kohlenhydraten mit Fructose-1,6-diphosphat-Aldolase	Liebigs Ann. Chem.	1990		1017
H. Waldmann	Prolin-benzylester als chirale Hilfsgruppe in asymmetrischen 1,3-dipolaren Cycloadditionen mit Nitriloxiden	Liebigs Ann. Chem.	1990		1013
H. Kunz, H. Waldmann	Protecting Groups	Comprehensive Organic Chemistry, Pergamon Press, London	1991		631
P. Braun, H. Waldmann, W. Vogt, H. Kunz,	Selektive enzymatische Schutzgruppenabspaltungen: Der n-Heptylester als Carboxyl-schutzgruppe in der Peptidsynthese	Liebigs Ann. Chem.	1991		165
H. Waldmann, M. Braun	Amino Acid Esters as Chiral Auxiliaries in Cycloaddition Reactions	Gazz. Chim. Ital	1991	121	277
H. Waldmann	Enzymatic Protecting Group Techniques	Kontakte (Merck)	1991	2	33



H. Waldmann	Aminosäureester als chirale Hilfsgruppen	GIT Fachzeitschrift für das Laboratorium	1991	35	593
H. Waldmann, M. Braun	Asymmetric Synthesis of Bicyclic Amino Acid Derivatives by Aza-Diels-Alder-Reactions in Aqueous Solution	Liebigs Ann. Chem.	1991		1045
H. Waldmann, M. Braun, M. Dräger	Asymmetric Aza-Diels-Alder-Reactions of Amino Acid Ester Imines with Brassards Diene	Tetrahedron Asymmetry	1991	2	1991
H. Waldmann	Aminosäureester als chirale Hilfsgruppen bei Barbier-Reaktionen in wässriger Lösung	Liebigs Ann. Chem.	1991		1317
H. Waldmann	LiClO ₄ in Ether - ein ungewöhnliches Lösungsmittel	Angew. Chem.	1991	103	1335
H. Waldmann, M. Braun, M. Weymann, M. Gewehr	An Approach to the Skeleton of Yohimbine-Type Alkaloids via Mannich-Michael Reactions with Indolyethyl-Imines	Synlett	1991		881
H. Waldmann, P. Braun, H. Kunz,	New Enzymatic Protecting Group Techniques for the Construction of Peptides and Glycopeptides	Biomed. Biochim. Acta	1991	50	243-248
P. Braun, H. Kunz, H. Waldmann	New Enzymatic Protecting Group Techniques for the Construction of Peptides and Glycopeptides	Peptides: Chemistry and Biology (Proceedings of the 12th American Peptide Symposium, Cambridge 1991)" (J. A. Smith und J. E. Rivier, Hrsg.), Escom, Leiden	1992		527
P. Braun, H. Waldmann, H. Kunz	Selective Enzymatic Removal of Protecting Functions: Heptyl Esters as Carboxy Protecting Groups in Glycopeptide Synthesis	Synlett	1992		39
H. Waldmann, M. Braun	Amino Acid Ester Imines as Chiral Auxiliaries in Tandem Mannich-Michael Reactions with Danishefskys Diene	J. Org. Chem.	1992	57	5444
H. Waldmann, A. Heuser, P. Braun, M. Schultz	Neue enzymatische Methoden für die Kohlenhydratchemie	GIT Fachzeitschrift für das Laboratorium	1992	36	908
H. Waldmann, A. Heuser, P. Braun, H. Kunz	New Enzymatic Protecting Group Techniques for Peptide and Glycopeptide Chemistry	Indian J. Chem.	1992	31B	799
H. Waldmann, A.	New Enzymatic Methods for the	Microbial Reagents in	1992		113



Heuser, P. Braun, M. Schultz, H. Kunz	Selective Functionalization of Carbohydrate Derivatives	of Organic Synthesis", NATO ASI Series, Kluwer, Dordrecht			
H. Waldmann, P. Braun, H. Kunz	New Enzymatic Protecting Group Techniques for the Construction of Peptides and Glycopeptides	Chemistry of Peptides and Proteins	1993	5/6 (Pt. A)	227-235
H. Waldmann	Amino Acid Esters as Chiral Auxiliary Groups I and II	Kontakte (Merck) 1993 (1), 58 und 1993 (2), 34			
H. Waldmann, A. Reidel	Enzymatic Protecting Group Techniques	J. Prakt. Chem.	1993	335	109-127
H. Waldmann, M. Braun, M. Weymann, M. Gewehr	Asymmetric Synthesis of Indolo[2,3-a]quinolizidin-2-ones - Congeners to Yohimbine-Type Alkaloids	Tetrahedron	1993	49	397
R. Lock, H. Waldmann	Construction of Tetracyclic Indole Bases via Aza Diels-Alder Reactions of Indolylethylimines with Brassard's Diene	Nat. Prod. Lett.	1993	2	49
H. Waldmann, G. Schmidt, M. Jansen, J. Geb	Asymmetric Pictet-Spengler Reactions Employing Amino Acid Esters as Mediators of Selectivity	Tetrahedron Lett.	1993	34	5867
P. Braun, H. Waldmann, H. Kunz	Chemoenzymatic Synthesis of O-Glycopeptides Carrying the Tumor Associated T _N -Antigen Structure	Bioorg. Med. Chem.	1993	1	197
H. Waldmann, A. Heuser, A. Reidel	Selective Enzymatic Deprotection of Hydroxy- and Amino Groups in Carbohydrates and Nucleosides	Synlett	1994		65
H. Waldmann, E. Bläser, M. Jansen, H.-P. Letschert	Asymmetrische Synthese hochsubstituierter Pyrrolidine durch 1,3-dipolare Cycloaddition von Azomethinyliden an N-Acryloylprolinbenzylester	Angew. Chem.	1994	106	717
H. Waldmann, R. Lock	Construction of Tetracyclic Indole Bases via Aza Diels-Alder Reactions of Indolylethylimines with Brassard's Diene	Liebigs Ann. Chem.	1994		511
H. Waldmann, A. Heuser	Acetylerase from Orange Peel as Biocatalyst for the Chemo- and Regioselective Deprotection of Carbohydrates	Bioorg. Med. Chem.	1994	2	477-482
H. Waldmann, D. Sebastian	Enzymatic Protecting Group Techniques	Chem. Rev.	1994	94	911-937
H. Waldmann	Asymmetric Hetero Diels-Alder Reactions	Synthesis	1994		535
T. Pohl, E. Nägele, H. Waldmann	Biocatalysts as Chemo- and Regioselecting Tools in Organic Synthesis	Catalysis Today	1994	22	407



H. Waldmann, G. Schmidt, M. Jansen, J. Geb	Asymmetric Steering of the Pictet-Spengler Reaction by Means of Amino Acid Esters as Chiral Auxiliary Groups	Tetrahedron	1994	50	11865
H. Waldmann, G. Böhm, U. Schmid,	O-Glycosidsynthesen unter neutralen Bedingungen in konzentrierten Lösungen von LiClO ₄ in organischen Solvenzien	Angew. Chem.	1994	106	2024
H. Waldmann, M. Weigerding, C. Dreisbach, C. Wandrey	C ₂ -Symmetric Bicyclic Diols as Chiral Ligands in the Titanate-Catalyzed Enantioselective Addition of Alkylzinc Reagents to Aldehydes	Helv. Chim. Acta	1994	77	2111
G. Giffels, C. Dreisbach, U. Kragl, M. Weigerding, H. Waldmann, C. Wandrey	Chirale Titanalkoxide als Katalysatoren zur enantioselektiven Reduktion von Ketonen mit Boranen	Angew. Chem.	1995	107	2165
H. Waldmann, E. Nägele	Synthesis of the Palmitoylated and Farnesylated C-Terminal Lipohexapeptide of the Human N-Ras Protein by Employing an Enzymatically Removable Urethane Protecting Group	Angew. Chem. / Angew. Chemie Int. Ed.	1995	107/34	2425-2428/2259-2262
H. Waldmann, G. Schmidt, H. Henke, M. Burkard	Asymmetrische Steuerung der Pictet-Spengler-Reaktion unter Verwendung von N,N-Phthaloylaminosäuren als chirale Hilfsgruppen	Angew. Chem.	1995	107	2608
H. Waldmann, E. Bläser, M. Jansen, H.-P. Letschert	Asymmetric Steering of 1,3-Dipolar Cycloaddition Reactions by Means of Proline Esters as Chiral Auxiliary Groups	Chem. Eur. J.	1995	1	150
M. Nettekoven, M. Psiorz, H. Waldmann	Synthesis of Enantiomerically Pure 4-Alkylsubstituted Tryptophan Derivatives by a Combination of Organometallic Reactions with Enantioselective Enzymatic Transformations	Tetrahedron Lett.	1995	36	1425
T. Pohl, H. Waldmann	Enhancement of the Enantioselectivity of Penicillin G-Acylase from E. coli by "Substrate Tuning"	Tetrahedron Lett.	1995	36	2963
G. Böhm, H. Waldmann	Synthesis of Glycosides of Fucose under Neutral Conditions in Solutions of LiClO ₄ in Organic Solvents	Tetrahedron Lett.	1995	36	3843
H. Waldmann	Asymmetric Synthesis of Nitrogen Heterocycles Employing Amino Acid Esters as Chiral Auxiliary Groups	Synlett	1995		133
H. Waldmann	Enzymatische Synthese von	"45 Jahre Fonds der	1995		133



	Peptidkonjugaten - Werkzeuge zum Studium der biologischen Signaltransduktion	Chemischen Industrie“			
A. Flohr, H. Waldmann	LiClO ₄ and Organic Solvents - A Powerful Combination	J. Prakt. Chem.	1995	337	609
H. Waldmann, K. Drauz, Hrsg.	Enzymatic Protecting Group Techniques" H. Waldmann in "Enzyme Catalysis in Organic Synthesis: A Comprehensive Handbook	Verlag Chemie, Weinheim	1995		
M. Schelhaas, S. Glomsda, M. Hänslers, H.-D. Jakubke, H. Waldmann	Enzymatische Synthese von Peptiden und Ras-Lipopeptiden unter Verwendung des Cholinesters als löslichkeitsvermittelnder Schutz- und Aktivierungsgruppe	Angew. Chem. / Angew. Chem. Int. Ed.	1996	108/3 5	82-85 / 106-109
M. Schelhaas, H. Waldmann	Schutzgruppenstrategien in der Organischen Synthese	Angew. Chem. / Angew. Chem. Int. Ed.	1996	108 / 35	2192-2219 / 2056-2083
T. Pohl, H. Waldmann	Enzymatische Synthese eines charakteristischen phosphorylierten und glycosylierten Peptidfragments der großen katalytischen Untereinheit der RNA-Polymerase II	Angew. Chem. / Angew. Chem. Int. Ed.	1996	108 / 35	1829 / 1729-1722
G. Schmidt, H. Waldmann, H. Henke, M. Burkard	Asymmetric Steering of the Pictet-Spengler Reaction by Means of N-Protected Amino Acids as Chiral Auxiliary Groups	Chem. Eur. J.	1996	2	1566-1571
G. Böhm, H. Waldmann	O-Glycoside Synthesis Under Neutral Conditions in Concentrated Solutions of LiClO ₄ in Organic Solvents Employing Benzyl-Protected Glycosyl Donors	Liebigs Ann. Chem.	1996		613
G. Böhm, H. Waldmann	O-Glycoside Synthesis Under Neutral Conditions in Concentrated Solutions of LiClO ₄ in Organic Solvents Employing Acyl-Protected Glycosyl Donors	Liebigs Ann. Chem.	1996		621
R. Lock, H. Waldmann	Asymmetric Synthesis of Highly Functionalized Tetracyclic Indole Bases Embodying the Basic Skeleton of Yohimbine- and Reserpine Type Alkaloids	Tetrahedron Lett.	1996		2753
G. Müller, H. Waldmann	An Enzyme Initiated Domino Hydroxylation Oxidation Carbo Diels-Alder Reaction Cascade	Tetrahedron Lett.	1996		3833
U. Schmid, H. Waldmann	Activation of Glycosyl Phosphates by in situ Conversion to Glycosyl Iodides under Neutral Conditions in Concentrated Solutions of LiClO ₄ in	Tetrahedron Lett.	1996		3837



Organic Solvents

H. Waldmann, A. Heuser, S. Schulze	Selective Enzymatic Removal of Protecting Groups: The Phenylacetamide as Amino Protecting Group in Phosphopeptide Synthesis	Tetrahedron Lett.	1996	37	8725-8728
H. Waldmann, E. Nägele, M. Schelhaas, D. Sebastian	Enzymatic Synthesis of Peptide Conjugates - Tools for the Study of Signal Transduction Processes	in „Peptides - Chemistry, Structure and Biology“ (Hrsg.: P. T. P. Kaumaya und R. Hodges), Mayflower, Kingswinford	1996		397
K. Drauz, H. Waldmann, B. Sauerbrei	Biocatalysis and Enzyme-Analogous Processes	in „Applied Homogeneous Catalysis with Organometallic Compounds“ (Hrsg. B. Cornils und W. A. Hermann), VCH	1996		768
H. Waldmann, A. Reidel	Die Phenylacetamido(PhAc)-Gruppe - Die erste enzymatisch abspaltbare Aminoschutz-gruppe für die Deblockierung von Oligonucleotiden in Lösung und an der festen Phase	Angew. Chem. / Angew. Chem. Int. Ed.	1997	109 / 36	642-644/ 647-649
T. Zelinski, H. Waldmann	Quervernetzte Enzymkristalle (CLECs): Effiziente und stabile Biokatalysatoren für die präparative organische Chemie	Angew. Chem. / Angew. Chem. Int. Ed.	1997	109 / 36	746-748/ 722-724
H. Waldmann, K. Hinterding, P. Herrlich, H. J. Rahmsdorf, A. Knebe	Selective Inhibition of Receptor Tyrosine Kinases by Synthetic Analogs of Aeropylsinin	Angew. Chem.	1997	109	1553-1555
H. Waldmann, M. Schelhaas, E. Nägele, J. Kuhlmann, A. Wittinghofer, H. Schroeder, J. R. Silvius	Chemoenzymatic Synthesis of Fluorescent N-Ras Lipopeptides and their Use in Membrane Localization Studies in vivo	Angew. Chem./Angew. Chem. Int. Ed.	1997	109/ 36	2324 2238-2241
T. Pohl, H. Waldmann	Chemoenzymatic Synthesis of a Characteristic Phosphorylated and Glycosylated Fragment of the Large Subunit of Mammalian RNA Polymerase II	J. Am. Chem. Soc.	1997	119	6702-6710
H. Waldmann, S. Gabold	Chemoenzymatic Synthesis of Nucleopeptides	J. Chem. Soc. Chem. Comm.	1997		1861
R. Lock, H. Waldmann	Enantioselective Construction of Highly Functionalized Indoloquinolizines - Congeners to Polycyclic Indole Alkaloids	Chem. Eur. J.	1997	3	143-151



H. Schroeder, R. Leventis, S. Rex, M. Schelhaas, E. Nägele, H. Waldmann, J. R. Silvius	S-Acylation and Plasma Membrane Targeting of the Farnesylated Carboxyl-terminal Peptide of N-Ras in Mammalian Fibroblasts	Biochemistry	1997	36	13102-13109
P. Stöber, M. Schelhaas, E. Nägele, P. Hagenbuch, J. Rétey, H. Waldmann,	Synthesis of Characteristic Liopeptides of the Human N-Ras Protein and their Evaluation as Possible Inhibitors of Protein Farnesyl Transferase	Bioorg. Med. Chem.	1997	5	75
T. Kappes, H. Waldmann	Enzymatic Synthesis of Peptide Conjugates - Tools for the Study of Biological Signal Transduction	Liebigs Ann. Chem.	1997		808-813
D. Sebastian, H. Waldmann	Chemoenzymatic Synthesis of a Characteristic Phosphopeptide Fragment of the human c-Raf-Kinase	Tetrahedron Lett.	1997	38	2927-2930
S. Kirschbaum, H. Waldmann	Construction of the Tricyclic Benzoquinolizine Ring System by Combination of a Tandem Mannich-Michael Reaction with a Heck Reaction	Tetrahedron Lett.	1997	38	2829-2832
D. Sebastian, A. Heuser, S. Schulze, H. Waldmann	Selective Enzymatic Deprotection of Phosphopeptides - Chemoenzymatic Synthesis of a Characteristic Phosphopeptide Fragment of the Raf-1 Kinase	Synthesis	1997		1098
T. Kappes, H. Waldmann	The Tetrabenzylglucosyloxycarbonyl(BGloc) Group - An Enzyme Labile Carbohydrate Derived Urethane Blocking Group	Carbohydr. Res.	1997	305	341
U. Schmid, H. Waldmann	O-Glycoside Synthesis with Glycosyl Iodides under Neutral Conditions in 1M LiClO ₄ in CH ₂ Cl ₂	Liebigs Ann./Recueil	1997		2573-2577
B. Sauerbrei, T. Kappes, H. Waldmann	Enzymatic Synthesis of Peptide Conjugates - Tools for the Study of Biological Signal Transduction	Top. Curr. Chem.	1997	186	66-86
T. Zelinski, H. Waldmann	Cross-Linked Enzyme Crystals (CLECs) - Powerful Biocatalysts for Synthetic Chemistry	J. Prakt. Chem./Chem.-Ztg	1997	339	394-396
K. Hinterding, D. Alonso-Díaz, H. Waldmann	Organic Synthesis and Biological Signal Transduction	Angew. Chem. / Angew. Chem. Int. Ed.	1998	110 / 37	716-780 / 688-749
B. Sauerbrei, V. Jungmann, H. Waldmann	Entwicklung einer enzymaliblen Ankergruppe für organische Synthesen an polymeren Trägern	Angew. Chem. / Angew. Chem. Int. Ed.	1998	110 / 37	1187-1190 / 1143-1146



K. Hinterding, P. Hagenbuch, J. Rétey, H. Waldmann	Synthese und in vitro Evaluierung des Farnesyltransferase-Inhibitors Pepticcinnamin E	Angew. Chem. / Angew. Chem. Int. Ed.	1998	110 / 37	1298-1301/ 1236-1239
E. Nägele, M. Schelhaas, N. Kuder, H. Waldmann	Chemoenzymatic Synthesis of N-Ras Lipoptides	J. Am. Chem. Soc.	1998	120	6889-6902
T. Schmittberger, A. Cotté, H. Waldmann	Synthesis of Characteristic Lipoptides of Lipid Modified Proteins Employing the Allyl Ester as Protecting Group	J. Chem. Soc. Chem. Commun.	1998		937-938
H. Schene, H. Waldmann	Direct Synthesis of 2-Deoxy- and 2,6- Dideoxy Glycosides under Neutral Conditions in LiClO ₄ /Ether Mixtures	J. Chem. Soc. Chem. Commun.	1998		2759-2760
U. Schmid, H. Waldmann	Synthesis of Fucosyl Saccharides Under Neutral Conditions in Solutions of Lithium Perchlorate in Dichloromethane	Chem. Eur. J.	1998	4	494-501
G. H. Müller, A. Lang, D. R. Seithel, H. Waldmann	An Enzyme-Initiated Hydroxylation- Oxidation-Carbo-Diels-Alder Domino Reacion	Chem. Eur. J.	1998	4	2513-2522
S. Kirschbaum, H. Waldmann	A Three Step Access to the Tricyclic Benzoquinolizine Ring System	J. Org. Chem.	1998	63	4936-4946
V. Jungmann, H. Waldmann	An Enzymatic Protecting Group Strategy for the Synthesis of Nucleopeptides	Tetrahedron Lett.	1998	39	1139-1142
U. Schlede, M. Nazaré, H. Waldmann,	Efficient Enantioselective Synthesis of a β -Hydroxyepoxide Building Block for the Construction of Macrocyclic Natural Products	Tetrahedron Lett.	1998	39	1143-1144
D. Brohm, H. Waldmann	Stereoselective Synthesis of the Core Structure of the Protein Phosphatase Inhibitor Dysidiolide	Tetrahedron Lett.	1998		3995-3998
H. Schene, H. Waldmann	Activation of Glycosyl Phosphites under Neutral Conditions in Solutions of Metal Perchlorates in Organic Solvents	Eur. J. Org. Chem.	1998		1227-1230
K. Hinterding, A. Knebel, P. Herrlich, H. Waldmann	Synthesis and Biological Evaluation of Aeropylsinin Analogues: A new Class of Receptor Tyrosine Kinase Inhibitors	Bioorg. Med. Chem.	1998	6	1153-1162
T. Schmittberger, H. Waldmann	Organic Synthesis and Biological Signal Transduction	Synlett	1998		574-584
T. Pathak, H. Waldmann	Enzymes and Protecting Groups	Curr. Op. Chem. Biol.	1998	2	112-120



U. Schmid, H. Waldmann	LiClO ₄ and Organic Solvents - Unusual Reaction Media	in „Organic Synthesis Highlights III“ (Hrsg.: J. Mulzer, H. Waldmann), Wiley-VCH	1998		205-210
D. J. Owen, K. Alexandrov, E. Rostkova, A. J. Scheidig, R. S. Goody, H. Waldmann	Chemoenzymatic Synthesis of Fluorescent Rab 7 Proteins; Tools for Studying Vesicular Trafficking in Cells	Angew. Chem. / Angew. Chem. Int. Ed.	1999	111 / 38	570-573 / 509-512
R. Müller, H. Goesmann, H. Waldmann	<i>N,N</i> -Phthaloylamino Acids as Chiral Auxiliaries in Asymmetric Mannich-Type Reactions	Angew. Chem. / Angew. Chem. Int. Ed.	1999	111 / 38	166-169 / 184-187
J. Sander, H. Waldmann	Chemoenzymatic Synthesis of a Characteristic Glycophosphopeptide from the Transactivation Domain of Serum Response Factor	Angew. Chem. / Angew. Chem. Int. Ed.	1999	111 / 38	1337-1339 / 1250-1252
F. Stieber, U. Grether, H. Waldmann	An Oxidation-Labile Traceless Linker for Solid-Phase Synthesis	Angew. Chem. / Angew. Chem. Int. Ed.	1999	111 / 38	1142-1145 / 1073-1077
B. Meseguer, D. Alonso-Díaz, N. Griebenow, T. Herget, H. Waldmann	Natural Product Synthesis on Polymeric Supports – Synthesis and Biological Evaluation of an Indolactam Library	Angew. Chem. / Angew. Chem. Int. Ed.	1999	111 / 38	3083-3087 / 2902-2906
P. Stahl, H. Waldmann	Asymmetric Synthesis of the Nakijiquinones - Selective Inhibitors of the Her-2/Neu Protooncogene	Angew. Chem. / Angew. Chem. Int. Ed.	1999	111 / 38	3935-3938 / 3710-3713
K. Hinterding, P. Hagenbuch, J. Rétey, H. Waldmann	Synthesis and in vitro Evaluation of the Farnesyltransferase Inhibitor Peptidocinnamin E	Chem. Eur. J.	1999	5	227-236
S. Flohr, V. Jungmann, H. Waldmann	Chemoenzymatic Synthesis of Nucleopeptides	Chem. Eur. J.	1999	5	669-681
A. Cotté, B. Bader, J. Kuhlmann, A. Wittinghofer, H. Waldmann	Synthesis of the N-Terminal Lipohexapeptide of Human G _α -Protein and Fluorescent Labeled Analogs for Biological Studies	Chem. Eur. J.	1999	5	922-936
M. Schelhaas, E. Nägele, N. Kuder, H. B. Bader,	Chemoenzymatic Synthesis of Biotinylated Ras Peptides and their Use in Membrane Binding Studies of Lipidated Model Proteins by Surface	Chem. Eur. J.	1999	5	1239-1252



J. Kuhlmann, A. Wittinghofer, H. Waldmann	Plasmon Resonance				
F. Eisele, D. J. Owen, H. Waldmann	Peptide Conjugates as Tools for the Study of Biological Signal Transduction	Bioorg. Med. Chem.	1999	7	193-224
T. Schmittberger, H. Waldmann	Synthesis of Palmitoylated and Prenylated C-Terminal Lipopeptides of the human R- and N-Ras Proteins	Bioorg. Med. Chem.	1999	7	749-762
E. Bläser, P. Kolar, D. Fenske, H. Goesmann, H. Waldmann	Asymmetric Steering of Oxa Diels-Alder Reactions with Silyloxydienes Employing Proline Esters as Chiral Auxiliary Groups	Eur. J. Org. Chem.	1999	1	329-333
H. Schene, H. Waldmann	Synthesis of Deoxy Glycosides Under Neutral Conditions in LiClO ₄ /Solvent Mixtures	Synthesis	1999	S1	1411-1422
G. H. Müller, H. Waldmann	The Phenyl Hydrazone as an Enzyme- Labile Protecting Group - Oxidative Cleavage with Mushroom Tyrosinase	Tetrahedron Lett.	1999	40	3549-3552
K. Kuhn, H. Waldmann	Synthesis of Characteristic Palmitoylated Lipopeptides from Human Y ₁ Receptor by a Combination of Enzyme-Labile and Pd(0)-Sensitive Protecting Groups	Tetrahedron Lett.	1999	40	6369-6372
I. Rubio, U. Wittig, C. Meyer, R. Heinze, D. Kadereit, H. Waldmann, J. Downward R. Wetzker	Farnesylation of Ras is important for the interaction with phosphoinositide 3- kinase γ	Eur. J. Biochem.	1999	266	70-82
T. Pathak, H. Waldmann	Enzymatic Protecting Group Techniques in Organic Synthesis	in <i>Stereoselective Biocatalysis</i> (Hrsg.: R. Patel), Marcel Dekker, New York	1999		775-797
B. Bader, K. Kuhn, D. J. Owen, H. Waldmann, A. Wittinghofer, J. Kuhlmann	Bioorganic synthesis of lipid-modified proteins for the study of signal transduction	Nature	2000	403	223-226
M. Nazaré, H. Waldmann	Synthesis of the (9S,18R)-seco acid of the leukocyte adhesion inhibitor Cyclamenol A	Angew. Chem. / Angew. Chem. Int. Ed.	2000	112 / 39	1171-1174 / 1125-1128
R. Machauer, H. Waldmann	Synthesis of the N-Terminal N- Myristoylated and S-Palmitoylated Undetrigintapeptide of Endothelial NO-	Angew. Chem. / Angew. Chem. Int. Ed.	2000	112 / 39	1503-1507 / 1449-1453



Synthetase

U. Grether, H. Waldmann	An Enzyme-Labile Safety Catch Linker for Combinatorial Synthesis on Soluble Polymeric Support	Angew. Chem. / Angew. Chem. Int. Ed.	2000	112 / 39	1688-1691 / 1629-1632
H. Waldmann, A. Wittinghofer	Ras, a Molecular Switch Involved in Tumor Formation	Angew. Chem. / Angew. Chem. Int. Ed.	2000	112 / 39	4360-4383 / 4192-4214
J. Sander, H. Waldmann	Enzymatic Protecting Group Techniques for Glyco- and Phosphopeptide Chemistry - Synthesis of a Glycophosphopeptide from Human Serum Response Factor	Chem. Eur. J.	2000	6	1564-1577
R. Müller, H. Röttele, H. Henke, H. Waldmann	Asymmetric Steering of the Mannich Reaction with Phthaloyl Amino Acids	Chem. Eur. J.	2000	6	2032-2043
B. Meseguer, D. Alonso-Díaz, N. Griebenow, T. Herget, H. Waldmann	Solid Phase Synthesis and Biological Evaluation of a Teleocidin Library – Discovery of a Selective PKC δ Down Regulator	Chem. Eur. J.	2000	6	3943-3957
A. G. Gum, T. Kappes-Roth, H. Waldmann	Enzyme-Labile Protecting Groups in Peptide Synthesis: Development of Glucose- and Galactose Derived Urethanes	Chem. Eur. J.	2000	6	3714-3721
N.H. Thomä, A. Iakovenko, D. Owen, A.S. Scheidig, H. Waldmann, R.S. Goody, K. Alexandrov	Phosphoisoprenoid Specificity of Rab Geranylgeranyltransferase II	Biochemistry	2000	39	12043- 12052
D. Kadereit, J. Kuhlmann, H. Waldmann	Linking the Fields – The Interplay of Organic Synthesis, Biophysical Chemistry and Cell Biology in the Chemical Biology of Protein Lipidation	ChemBioChem	2000	1	144-169
D. Kadereit, H. Waldmann	Synthesis of Characteristic H-Ras Lipopeptides Employing Nobel Metal-, Acid- and Reduction Labile Blocking Groups	ChemBioChem.	2000	3	200-203
N. Kuder, T. Zelinski, T. Pathak, O. Seitz, H. Waldmann	Synthesis of a Triply Phosphorylated Pentapeptide from Human τ -Protein	Bioorg. Med. Chem.	2000	8	2433-2439
M. Nazaré, H. Waldmann	Enantioselective Synthesis of the Leukocyte Adhesion Inhibitor Cyclamenol A Seco Acid	Tetrahedron Lett.	2000	41	625-628
J. Spieler,	Synthesis of Chiral Amino Alcohols	Eur. J. Org. Chem.	2000	3	391-399



O. Huttenloch, H. Waldmann	Embodying the Bispidine Framework and their Application as Ligands in Enantioselectively Catalyzed Additions to C=O and C=C Groups				
T. Kappes-Roth, H. Waldmann	Chemoenzymatic Synthesis of a Biotin-Labeled Glycophosphonopeptide of the c-Myc Oncoprotein	J. Chem. Soc., Perkin Trans	2000	16	2579-2581
H. Waldmann, M. Thutewohl	Ras-Farnesyltransferase-Inhibitors as Promising Anti-Tumor Drugs	Top. Curr. Chem.	2000	211	118-130
D. Kadereit, J. Kuhlmann, H. Waldmann	Organic Synthesis and Cell Biology	in <i>Stimulating Concepts in Chemistry</i> (Hrsg.: M. Shibaski, J.F. Stoddart, F. Vögtle), Wiley-VCH	2000		369-382
D. Kadereit, H. Waldmann	Chemoenzymatic Synthesis of Lipidated Peptides	Chemical Monthly	2000	131	571-584
H. Waldmann	Chemoenzymatic synthesis of lipidated peptide and protein conjugates: Tools for the study of biological signal transduction	In: "Peptides for the New Millennium", (G.B. Fields, J.T. Tam, G. Barany eds.), Kluwer Academic Publishers, Dordrecht, Netherlands	2000		555-557
H. Waldmann	Bioorganische Synthese lipidierter Peptide und Proteine zum Studium der biologischen Signaltransduktion	Jahrbuch der Max-Planck-Gesellschaft", (Max-Planck-Gesellschaft, Ed.) Vandenhoeck & Ruprecht, Göttingen	2000		348-353
F. Eisele, J. Kuhlmann, H. Waldmann	Synthesis and Membrane-Binding Properties of a Characteristic Lipopeptide from the Membrane-Anchoring Domain of Influenza Virus A Hemagglutinin	Angew. Chem. / Angew. Chem. Int. Ed.	2001	113 / 40	382-386 / 369-373
D. Huster, K. Kuhn, D. Kadereit, H. Waldmann, K. Arnold	¹ H High-Resolution Magic Angle Spinning NMR Spectroscopy for the Investigation of a Ras Lipopeptide in a Lipid Membrane	Angew. Chem. / Angew. Chem. Int. Ed.	2001	113 / 40	1083-1085 / 1056-1058
K. Kuhn, D. J. Owen, B. Bader, A. Wittinghofer, J. Kuhlmann, H. Waldmann	Synthesis of Functional Ras Lipoproteins and Fluorescent Derivatives	J. Am. Chem. Soc.	2001	123	1023-1035



P. Stahl, L. Kissau, R. Mazitschek, A. Huwe, P. Furet, A. Giannis, H. Waldmann	Total Synthesis and Biological Evaluation of the Nakijiquinones	J. Am. Chem. Soc.	2001	123	11586-11593
D. Kadereit, H. Waldmann	Enzymatic Protecting Group Techniques	Chem. Rev.	2001	101	3367-3396
O. Huttenloch, J. Spieler, H. Waldmann	Chiral Bicyclic Phosphoramidites – A New Class of Ligands for Asymmetric Catalysis	Chem. Eur. J.	2001	7	671-675
D. Kadereit, P. Deck, I. Heinemann, H. Waldmann	Acid-labile Protecting Groups for the Synthesis of Lipidated Peptides	Chem. Eur. J.	2001	7	1184-1193
U. Grether, H. Waldmann	An Enzyme-labile Safety Catch Linker for Synthesis on a Soluble Polymeric Support	Chem. Eur. J.	2001	7	959-971
R. Machauer, H. Waldmann	Synthesis of Lipidated eNOS Peptides by Combining Enzymatic, Noble Metal- and Acid Mediated Protecting Group Techniques with Solid Phase Peptide Synthesis and Fragment Condensation in Solution	Chem. Eur. J.	2001	7	2940-2956
M. Nazaré, H. Waldmann	Enantiospecific Synthesis of the (9 <i>S</i> , 18 <i>R</i>)-Diastereomer of the Leukocyte Adhesion Inhibitor Cyclamenol A	Chem. Eur. J.	2001	7	3363-3376
C. Arenz, M. Thutewohl, O. Block, H. Waldmann, H.-J. Altenbach, A. Giannis	Manumycin A and its Analogues are Irreversible Inhibitors of Neutral Sphingomyelinase	ChemBioChem	2001	2	141-143
H. Waldmann, M. Famulok	Chemistry meets Biology	ChemBioChem	2001	2	3-6
O. Seitz, I. Heinemann, A. Mattes, H. Waldmann	Synthetic Peptide Conjugates – Tailor-Made Probes for the Biology of Protein Modification and Protein Processing	Tetrahedron	2001	57	2247-2277
D. A. Jeyaraj, H. Waldmann	Synthesis of Nucleopeptides by an Enzyme Labile Urethane Protecting Group	Tetrahedron Lett.	2001	42	835-837
C. Rosenbaum, H. Waldmann	Solid Phase Synthesis of Cyclic Peptides by Oxidative Cyclative Cleavage of an Aryl Hydrazide Linker – Synthesis of Stylostatin 1	Tetrahedron Lett.	2001	42	5677-5680
R. Reents, D. A. Jeyaraj, H. Waldmann	Biocatalysis in Polymer-Supported Synthesis: Enzyme-labile Linker Groups	Adv. Synth. Catal.	2001	343	501-513
M. Völkert,	The Chemical Biology of Ras Lipidation	Bio. Chem.	2001	382	1133-1145



M. Wagner,
C. Peters,
H. Waldmann

N. H. Thomä, A. Iakovenko, A. Kalinin, H. Waldmann, R. S. Goody, K. Alexandrov	Allosteric Regulation of Substrate Binding and Product Release in Geranylgeranyltransferase Type II	Biochemistry	2001	40	268-274
I. Heinemann, C. Katzka, H. Waldmann	„Sane in the membrane“ – Wie gelangen lipidierete Ras-Proteine in die Plasmamembran?	BIOforum	2001	5	324-327
A. Rak, R. Reents, O. Pylypenko, A. Niculae, V. Sidorovitch, N. H. Thomä, H. Waldmann, I. Schlichting, R. S. Goody, K. Alexandrov	Crystallization and Preliminary X-ray Diffraction Analysis of the Rab Escort Protein-1 in Complex with Rab geranylgeranyltransferase	J. Struct. Biol.	2001	136	158-161
P. Stahl, L. Kissau, R. Mazitschek, A. Giannis, H. Waldmann	Natural Product Derived Receptor Tyrosine Kinase Inhibitors: Identification of IGF1R, Tie-2 and VEGFR-3 Inhibitors	Angew. Chem. / Angew. Chem. Int. Ed.	2002	114 / 41	1222-1226 / 1174-1178
D. Brohm, S. Metzger, A. Bhargava, O. Müller, F. Lieb, H. Waldmann	Natural Products are Biologically Validated Starting Points in Structural Space for Compound Library Development: Solid Phase Synthesis of Dysidiolide-Derived Phosphatase Inhibitors	Angew. Chem. / Angew. Chem. Int. Ed.	2002	114 / 41	319-323 / 307-311
L. Bialy, H. Waldmann	Synthesis of the Protein Phosphatase 2A Inhibitor (4S,5S,6S,10S,11S,12S)-Cytostatin	Angew. Chem. / Angew. Chem. Int. Ed.	2002	114 / 41	1819-1822 / 1748-1751
J. Kuhlmann, A. Tebbe, M. Wagner, K. Uwai, M. Völkert, H. Waldmann	Photoactivatable Synthetic Ras-Proteins: Baits for the Identification of Plasma-membrane Bound Binding Partners of Ras	Angew. Chem. / Angew. Chem. Int. Ed.	2002	114 / 41	2655-2658 / 2546-2550
R. P. Breinbauer, I. R. Vetter, H. Waldmann	From Protein Domains to Drug Candidates – Natural Products as Guiding Principles in the Design and Synthesis of Compound Libraries	Angew. Chem. / Angew. Chem. Int. Ed.	2002	114 / 41	3002-3015 / 2878-2890
M. Thutewohl, L. Kissau, B. Popkirova, I.-M. Karaguni, T. Nowak, M. Bate, J. Kuhlmann, O. Müller, H. Waldmann	Solid-Phase Synthesis and Biological Evaluation of a Peptidocinnamin E Library	Angew. Chem. Int. Ed.	2002	41	3616-3620
F. Stieber, R. Mazitschek,	Traceless Solid-Phase Synthesis of 2-Aminothiazoles: Receptor Tyrosine	Angew. Chem. Int. Ed.	2002	41	4757-4761



N. Soric, A. Giannis, H. Waldmann	Kinase Inhibitors with Dual Selectivity for Tie-2 and VEGFR-2					
K. Alexandrov, I. Heinemann, R. S. Goody, T. Durek, H. Waldmann	Intein-Mediated Synthesis of Geranylgeranylated Rab7 Protein <i>In Vitro</i>	J. Am. Chem. Soc.	2002	124	5648-5649	
B. Ludolph, F. Eisele, H. Waldmann	Solid Phase Synthesis of Lipidated Peptides	J. Am. Chem. Soc.	2002	124	5954-5955	
D. Brohm, N. Philippe, S. Metzger, A. Bhargava, O. Müller, F. Lieb, H. Waldmann	Solid Phase Synthesis of Dysidiolide-Derived Protein Phosphatase Inhibitors	J. Am. Chem. Soc.	2002	124	13171-13178	
O. Huttenloch, E. Laxman, H. Waldmann	Combinatorial Development of Chiral Phosphoramidite-Ligands for Enantioselective Conjugate Addition Reactions	Chem. Comm.	2002		673-675	
F. Stieber, H. Waldmann	Development of new acid-functionalized resins for combinatorial synthesis on solid supports	Chem. Comm.	2002	16	1748-1749	
D.A. Jeyaraj, H. Prinz, H. Waldmann	Synthesis of Nucleopeptides by Employing an Enzyme Labile Urethane Protecting Group	Chem. Eur. J.	2002	8	1879-1887	
F. Eisele, J. Kuhlmann, H. Waldmann	Synthesis and Membrane Binding Properties of a Lipopeptide Fragment from Influenza Virus A Hemagglutinin	Chem. Eur. J.	2002	15	3363-3376	
O. Huttenloch, E. Laxman, H. Waldmann	Solid-Phase Development of Chiral Phosphoramidite Ligands for Enantioselective Conjugate Addition Reactions	Chem. Eur. J.	2002	8	4767-4780	
M. Völkert, S. Koul, G. H. Müller, M. Lehnig, H. Waldmann	Phenylhydrazide as an Enzyme-Labile Protecting Group in Peptide Synthesis	J. Org. Chem.	2002	67	6902-6910	
B. Ludolph, F. Eisele, H. Waldmann	Solution-and Solid-Phase Synthesis of the Polybasic Lipid-Modified C-Termini of Rho A and K-Ras 4B	ChemBioChem	2002	9	901-904	
L. Bialy, M. Lopez-Canet, H. Waldmann	Determination of the Relative Configuration of the C-2-C-1'-Fragment of Cytostatin	Synthesis (Special issue dedicated to Dieter Seebach)	2002	14	2096-2104	
R. Reents, D.A. Jeyaraj, H. Waldmann	Enzymatically cleavable linker groups in polymer supported synthesis	Drug Discovery Today	2002	7	71-76	
M. Scheck, H. Waldmann	Chemoenzymatic synthesis of the C ₁₅ -C ₂₃ unit of Leptomycin B	Can. J. Chem.	2002	80	571-576	



D. Kadereit, R. Reents, D. A. Jeyaraj, H. Waldmann	Introduction and Removal of Protecting Groups	Enzyme Catalysis in Organic Synthesis, Second Edition, Ed. by K. Drauz and H. Waldmann, Wiley-VCH, Weinheim	2002	<i>I-III</i>	1333-1417
T. Pathak, H. Waldmann	Enzymatically Cleavable Protecting Groups	In: <i>Houben-Weyl „Methods of Organic Chemistry“</i> , vol E22a (L. Moroder, M. Goodman, eds., Thieme-Verlag Stuttgart-New York	2002		295-314
C. Schultz, H. Gröger, C. Dinkel, K. Drauz, H. Waldmann	Biocatalysis and Enzyme-Analogous Processes	in „Applied Homogeneous Catalysis with Organometallic Compounds“, Second Edition, Ed. by B. Cornils und W. A. Hermann), Wiley-VCH	2002	2	872-911
R. P. Breinbauer, M. Manger, M. Scheck, H. Waldmann	Natural Product Guided Library Development	Compound Current Medicinal Chemistry	2002	9	2129-2145
H. Waldmann, R. P. Breinbauer	Nature provides the answer - Natural Product Structures as Guiding Principle in Combinatorial Chemistry	Screening	2002	3	46-48
C. Peters, M. Wagner, M. Völkert, H. Waldmann	Bridging the Gap Between Cell Biology and Organic Chemistry	Naturwissenschaften	2002	89	381-390
A. Rak, O. Pylypenko, T. Durek, A. Watzke, S. Kushnir, L. Brunsveld, H. Waldmann, R. S. Goody, K. Alexandrov	Structure of Rab GDP-Dissociation Inhibitor in Complex with Prenylated YPT1 GTPase	Science	2003	302	646-650
O. Pylypenko, A. Rak, R. Reents, A. Niculae, V. Sidorovitch, M.-D. Cioca, E. Bessolitsyna, N. H. Thomä, H. Waldmann, I. Schlichting, R. S.	Structure of Rab Escort Protein-1 in Complex with Geranylgeranyltransferase”,	Mol. Cell	2003	11	483-494



Goody, K. Alexandrov

A. Fürstner, F. Feyen, H. Prinz, H. Waldmann	Total Synthesis and Reassessment of the Phosphatase Inhibitory Activity of the Antitumor Agent TMC-69-6H	Angew. Chem. / Angew. Chem. Int. Ed.	2003	115 / 42	5519-5522 / 5361-5364
M. Köhn, R. Wacker, C. Peters, H. Schröder, L. Soullère, R. Breinbauer, C. M. Niemeyer, H. Waldmann	Staudinger-Ligation: A New Immobilization Strategy for the Preparation of Small Molecule Arrays	Angew. Chem. / Angew. Chem. Int. Ed.	2003	115 / 42	6010-6014 / 5830-5834
D. Huster, A. Vogel, C. Katzka, H. A. Scheidt, H. Binder, S. Dante, T. Gutberlet, O. Zschörnig, H. Waldmann, K. Arnold	Membrane Insertion of a Lipidated Ras Peptide Studied by FTIR, Solid-State NMR, and Neutron Diffraction Spectroscopy	J. Am. Chem. Soc.	2003	125	4070-4079
M. Völkert, K. Uwai, A. Tebbe, B. Popkirova, M. Wagner, J. Kuhlmann, H. Waldmann	Synthesis and Biological Activity of Photoactivatable N-Ras Peptides and – Proteins“	J. Am. Chem. Soc.	2003	125	12749- 12758
L. Bialy, H. Waldmann	Synthesis and Biological Evaluation of Cytostatin Analogues	Chem. Commun.	2003	15	1872-1873
C. Rosenbaum, C. Katzka, A. Marzinzik, H. Waldmann	Traceless Fischer indole synthesis on the solid phase	Chem. Commun.	2003	15	1822-1823
F. Stieber, U. Grether, H. Waldmann	Development of the Traceless Phenylhydrazide-Linker for Solid-Phase Synthesis	Chem. Eur. J.	2003	9	3270-3281
F. Stieber, U. Grether, R. Mazitschek, N. Soric, A. Giannis, H. Waldmann	Multi-step Solid-Phase Synthesis of an Antibiotic and Receptor Tyrosine Kinase Inhibitors using the Traceless Phenylhydrazide Linker	Chem. Eur. J.	2003	9	3282-3291
B. Ludolph, H. Waldmann	The Synthesis of Acid- and Base-labile Lipopeptides on Solid Support	Chem. Eur. J.	2003	9	3683-3691
C. Peters, H. Waldmann	Solid-Phase Synthesis of Peptide Esters Employing the Hydrazide Linker	J. Org. Chem.	2003	15	6053-6055
J. M. Gerdes, H. Waldmann	Direct Mass Spectrometric Monitoring of Solid Phase Organic Syntheses	J. Comb. Chem.	2003	5	814-820
L. Kissau, P. Stahl, R. Mazitschek, A. Giannis,	Development of Natural Product Derived Receptor Tyrosine Kinase Inhibitors Based on Conservation of Protein Domain Fold	J. Med. Chem.	2003	46	2917-2931



H. Waldmann

M. Thutewohl, H. Waldmann	Solid Phase Synthesis of a Peptidocinnamin E-Library	Bioorg. Med. Chem.	2003	11 / 12	2591-2615
M. Thutewohl, L. Kissau, B. Popkurova, I.-M. Karaguni, T. Nowak, M. Bate, J. Kuhlmann, O. Müller, H. Waldmann	Identification of Mono- and Bisubstrate Inhibitors of Protein Farnesyltransferase and Inducers of Apoptosis from a Peptidocinnamin E Library	Bioorg. Med. Chem.	2003	11 / 12	2617-2626
H. Waldmann	At the Crossroads of Chemistry and Biology	Bioorg. Med. Chem.	2003	11	3045-3051
M. A. Koch, R. P. Breinbauer, H. Waldmann	Protein Structure Similarity as Guiding Principle for Combinatorial Library Design	Biol. Chem.	2003	384	1265-1272
S. Sommer, R. P. Breinbauer, H. Waldmann	Polymer-Supported Synthesis of Non- Oligomeric Natural Products	Organic Synthesis Highlights V, G. Schmalz, T. Wirth (Eds), Wiley-VCH	2003		395-408
H. Waldmann	Natürlich kombinatorisch – naturstoffgetriebene Wirkstoffentwicklung	Nachrichten aus der Chemie	2003	2	126-131
R. P. Breinbauer, I. R. Vetter, H. Waldmann	From Protein Domains to Drug Candidates – Natural Products as Guiding Principles in Compound Library Design and Synthesis	Ernst Schering Research Foundation, Workshop 42, 2003 , H. Waldmann, M. Koppitz (eds.)	2003		167-188
D. Huster, A. Vogel, C. Katzka, T. Gutberlet, S. Dante, H. Waldmann, K. Arnold	Membrane Binding of a Lipidated Ras Peptide Studied by Solid-State NMR, and Neutron Diffraction.	Bioph. Journal	2003	84	131A
R. P. Breinbauer, M. A. Koch, H. Waldmann	Naturstoffgetriebene kombinatorische Chemie	BIOspektrum, 2003 , <i>Sonderausgabe 9.</i> <i>Jahrgang</i>	2003		478-479
R. Reents, D. A. Jeyaraj, H. Waldmann	Biocatalyzed reactions on polymeric supports: Enzyme-labile linker groups	in: Polymeric Materials in Organic Synthesis and Catalysis, M. R. Buchmeiser (Ed.), Wiley-VCH	2003		445-466
C. Peters, A. Wolf, M. Wagner, J. Kuhlmann,	The Cholesterol Membrane Anchor of the Hedgehog Protein Confers Stable Membrane Association to Lipid-Modified Proteins	Proc. Natl. Ac. Sci	2004	101	8531-8536



H. Waldmann

- M. A. Koch, L.-O. Wittenberg, S. Basu, D.A. Jeyaraj, E. Gourzoulidou, K. Reinecke, A. Odermatt, H. Waldmann **Compound library development guided by protein structure similarity clustering and natural product structure** Proc. Natl. Ac. Sci **2004** 101 16721-16726
- C. Rosenbaum, P. Baumhof, R. Mazitschek, O. Müller, A. Giannis, H. Waldmann Synthesis and Biological Evaluation of an Indomethacin Library Reveals a New Class of Angiogenesis-Related Kinase Inhibitors Angew. Chem. / Angew. Chem. Int. Ed. **2004** 116 / 43 226-230 / 224-228
- O. Müller, E. Gourzoulidou, M. Carpintero, I.-M. Karaguni, A. Langerak, C. Herrmann, T. Möröy, L. Klein-Hitpaß, H. Waldmann Identification of Potent Ras Signalling Inhibitors by Pathway-Selective Phenotype-Based Screening Angew. Chem. / Angew. Chem. Int. Ed. **2004** 116 / 43 456-460 / 450-454
- H. Waldmann, I.-M. Karaguni, M. Carpintero, E. Gourzoulidou, C. Herrmann, C. Brockmann, H. Oschkinat, O. Müller Sulindac-Derived Ras Pathway Inhibitors Target the Ras-Raf Interaction and Downstream Effectors in the Ras Pathway Angew. Chem. / Angew. Chem. Int. Ed. **2004** 116 / 43 460-464 / 454-458
- R. Reents, M. Wagner, J. Kuhlmann, H. Waldmann Synthesis and Application of Fluorescence-Labeled Ras-Proteins for Live-Cell Imaging Angew. Chem. / Angew. Chem. Int. Ed. **2004** 116 / 43 2765-2768 / 2711-2714
- O. Barun, S. Sommer, H. Waldmann Asymmetric Solid-Phase Synthesis of 6,6-Spiroketal Angew. Chem. / Angew. Chem. Int. Ed. **2004** 116 / 43 3258-3261 / 3195-3199
- G. Kragol, M. Lumbierres, J. M. Palomo, H. Waldmann Solid-Phase Synthesis of Lipidated Peptides Angew. Chem. / Angew. Chem. Int. Ed. **2004** 116 / 43 5963-5966 / 5839-5842
- S. Janosch, C. Nicolini, B. Ludolph, C. Peters, M. Völkert, Th. L. Hazlet, E. Gratton, H. Waldmann Partitioning of Dual-lipidated Peptides into Membrane Microdomains – Lipid Sorting vs. Peptide Aggregation J. Am. Chem. Soc. **2004** 126 7496-7503



R. Winter

T. Durek, K. Alexandrov, R. S. Goody, A. Hildebrand, I. Heinemann, H. Waldmann	Synthesis of Fluorescently Labeled Mono- and Doubly Prenylated Rab7 GTPase	J. Am. Chem. Soc.	2004	126	16368- 16378
A. Fürstner, J. Ruiz-Caro, H. Prinz, H. Waldmann	Structure Assignment, Total Synthesis and Evaluation of the Phosphatase Modulating Activity of Glucolipin A	J. Org. Chem.	2004	2	459-467
L. Bialy, H. Waldmann	Total Synthesis and Biological Evaluation of the Protein Phosphatase 2A Inhibitor Cytostatin and Analogues	Chem. Eur. J.	2004	10	2759-2780
I.-M. Karaguni, E. Gourzoulidou, M. Carpintero, A. Langerak, L. Klein- Hitpaß, T. Möröy, G. Winde, H. Waldmann, O. Müller	SMAF-1 Inhibits the APC/ β -Catenin Pathway and Shows Properties Similar to Those of the Tumour Suppressor Protein APC	ChemBioChem	2004	5	1267-1270
L. Soulière, C. Aldrich, O. Daumke, R. Gail, L. Kissau, A. Wittinghofer, H. Waldmann	Synthesis of GTP-Derived Ras-Ligands	ChemBioChem	2004	10	1448-1453
A. Fürstner, K. Reinecke, H. Prinz, H. Waldmann	The Core Structures of Roseophilin and the Prodigiosin Alkaloids Define a New Class of Protein Tyrosine Phosphatase Inhibitors	ChemBioChem	2004	11	1575-1579
A. Fürstner, F. Feyen, H. Prinz, H. Waldmann	Synthesis and Evaluation of the Antitumor Agent TMC-69-6H and a focused library of Analogs	Tetrahedron	2004	60	9543-9558
R. Breinbauer, H. Waldmann	Solid and Solution Phase Combinatorial Chemistry – Preface	Tetrahedron	2004	60	8589
I. Heinemann, M. Völkert, H. Waldmann	Synthesis of Lipidated Peptides	Methods in Molecular Biology, Vol. 283: Bioconjugation Protocols: Strategies and Methods, C. Niemeyer (ed)., Humana Press	2004		221-232
O. Rocks, A. Peyker, M. Kahms, P. J. Verveer, C. Koerner, M. Lumbierres, J.	An Acylation Cycle Regulates Localization and Activity of Palmitoylated Ras Isoforms	Science	2005	307	1746-1752



Kuhlmann, H.
Waldmann, A.
Wittinghofer, P. I. H.
Bastiaens

M. A. Koch, A. Schuffenhauer, M. Scheck, M. Casaulta, A. Odermatt, P. Ertl, H. Waldmann	Charting Biologically Relevant Chemical Space: A Structural Classification of Natural Products (SCONP)	Proc. Natl. Ac. Sci.,	2005	102	17272-17277
L. Bialy, H. Waldmann	Inhibitors of Protein Tyrosine Phosphatase: Next Generation Drugs?"	Angewandte Chem./Angew. Chem. Int. Ed.	2005	117/44	2-28/3815-3839
P. Deck, D. Pendzialek, M. Biel, M. Wagner, B. Popkirova, B. Ludolph, G. Kragol, J., Kuhlmann, A. Giannis, H. Waldmann	Development and Biological Evaluation of Acyl Protein Thioesterase 1 (APT1) Inhibitors	Angew. Chem./Angew. Chem. Int. Ed.	2005	117/44	5055-5060/4975-4980
A. Vogel, C. Katzka, H. Waldmann, K. Arnold, M. F. Brown, D. Huster	Lipid Modifications of a Ras Peptide Exhibit Altered Packing and Mobility versus Host Membrane as Detected by ² H Solid-State NMR	J. Am. Chem. Soc	2005	127	12263-12272
E. Bayer, M. Thutewohl, C. Christner, T. Tradler, F. Osterkamp, H. Waldmann, P. Bayer	Identification of hPin1 inhibitors that induce apoptosis in a mammalian Ras transformed cell line	Chem. Comm.	2005	4	516-518
S. Sommer, H. Waldmann	Solid Phase Synthesis of a Spiro [5.5]ketal Library	Chem.Comm.	2005	45	5684-5686
L. Brunsveld, A. Watzke, T. Durek, K. Alexandrov, R. S. Goody, H. Waldmann	Synthesis of Functionalized RabGTPases by Combination of Solution and Solid Phase Lipopeptide Synthesis with Expressed Protein Ligation	Chem. Eur. J.	2005	11	2756-2772
M. Lumbierres, J. M. Palomo, G. Kragol, S. Roehrs, O. Müller, H. Waldmann	Solid-Phase Synthesis of Lipidated Peptides	Chem. Eur. J.	2005	11	7405-7415
R. Reents, M. Wagner, St. Schlummer, J. Kuhlmann, H. Waldmann	Synthesis and Application of Fluorescent Ras-Proteins for Live Cell Imaging	ChemBioChem	2005	6	86-94
E. Gourzoulidou, M. Carpintero,	Inhibition of Angiogenesis-Relevant Receptor Tyrosine Kinases by Sulindac	ChemBioChem	2005	6	527-531



P. Baumhof, A. Giannis, H. Waldmann	Analogues				
M. Manger, M. Scheck, H. Prinz, J.-P. von Kries, T. Langer, K. Saxena, H. Schwalbe, A. Fürstner, J. Rademann, H. Waldmann	Discovery of Mycobacterium Tuberculosis Protein Tyrosine Phosphatase A (MtpA) Inhibitors Based on Natural Products and a Fragment- Based Approach	ChemBioChem	2005	6	1749-1753
A. Watzke, L. Brunsveld, T. Durek, K. Alexandrov, A. Rak, R.S. Goody H. Waldmann	Chemical Biology of Protein Lipidation: Semi-Synthesis and Structure Elucidation of Prenylated RabGTPases	Org. Biomol. Chem.	2005	3	11-57-1164
O. Barun, K. Kumar, S. Sommer, A. Langerak, T. U. Mayer, O. Müller. H. Waldmann	Natural Product Guided Synthesis of a Spiroacetal Collection Reveals Modulators of Tubulin Cytoskeleton Integrity	Eur. J. Org. Chem.	2005	22	4773-4788
C. Rosenbaum, S. Röhrs, O. Müller, H. Waldmann M. A. Koch, H. Waldmann	Modulation of MRP-1-Mediated Multidrug Resistance by Indomethacin Analogues	J. Med. Chem.	2005	48	1179-1187
	Protein structure similarity clustering and natural product structure as guiding principles in drug discovery	Drug Disc. Today	2005	10	471-483
B. E. Dursina, R. Reents, A. Niculae, A. Veligodsky, R. Breitling, K. Pyatkov, H. Waldmann, R. S. Goody, K. Alexandrov	A genetically encodable microtag for chemo-enzymatic derivatization and purification of recombinant proteins	Protein Expression and Purification	2005	39	71-81
R. Balamurugan, F. J. Dekker, H. Waldmann	Design of compound libraries based on natural product scaffolds and protein structure similarity clustering (PSSC)	Molecular BioSystems	2005	1	36-45
F. J. Dekker, M. A. Koch, H. Waldmann	Protein structure similarity clustering (PSSC) and natural product structure as inspiration sources for drug development and chemical genomics	Curr. Op. in Chem. Biol.	2005	9	232-239
R. S. Goody, T. Durek, H. Waldmann, L. Brunsveld, K. Alexandrov	Application of Protein Semi-synthesis for the Construction of Functionalised Posttranslationally Modified RabGTPases	Methods in Enzymology	2005	403	29-42
A. Rak, O. Pylypenko, T. Durek, A. Watzke, H. Waldmann, R.S.	Protein semi-synthesis and its application to studies of structure and function of GTPases	Biopolymers	2005	80	490



Goody, K. Alexandrov

A. Vogel, C. Katzka, H. Waldmann, K. Arnold, M. F. Brown, D. Huster	Lipid Modifications of a Ras Peptide lead to alteration of bilayer packing and softness as detected by ^2H Solid-State NMR	Bioph. Journal	2005	88	26A
S.E. Feller, A. Vogel, H. Waldmann, Y. Arnold, D. Huster	Molecular dynamics simulation of a lipidated Ras peptide in membranes	Bioph. Journal	2005	88	422A
J. Hoffmann, Y. Feng, F. v. Hagen, A. Hillenbrand, J. Lin, R. Erber, P. Vajkoczy, E. Gourzoulidou, H Waldmann, A. Giannis, H. Wolburg, M. Shani, V. Jaeger, H.A. Weich, K.T. Preissner, S. Hoffmann, U. Deutsch, H.-P. Hammes	Endothelial survival factors and spatial completion, but not pericyte coverage of retinal capillaries determine vessel plasticity	FASEB Journal	2005	19 (14)	2035-2036
A. Nören-Müller, I. Reis Corrêa Jr., H. Prinz, C. Rosenbaum, K. Saxena, H. Schwalbe D. Vestweber, G. Cagna, S. Schunk, O. Schwarz, H. Schiewe, H. Waldmann	Discovery of New Protein Phosphatase Inhibitor Classes by Biology Oriented Synthesis (BIOS)	Proc. Natl. Acad. Sci.	2006	103	10606-10611
M. H. Gelb, L. Brunsveld, C. A. Hrycyna, S. Michaelis, F. Tamanoi, W. C. Van Voorhis, H. Waldmann	Therapeutic Intervention based on protein prenylation and associated modifications	Nature Chem. Biol.	2006	2	518-528
O. Pylypenko, A. Rak, T. Durek, S. Kushnir, B. E. Dursina, N. H. Thomae, A. T. Constantinescu, L. Brunsveld, A. Watzke, H. Waldmann, R. S. Goody, K. Alexandrov	Structure of doubly prenylated Ypt1: GDI complex and the mechanism of GDI mediated Rab recycling	Embo J.	2006	25	13-23
A. Dantas de Araújo, J. M. Palomo, J. Cramer, M. Köhn, H. Schröder, R. Wacker, C. M. Niemeyer, K.	Diels-Alder Ligation and Surface Immobilization of Proteins	Angew. Chem./Angew. Chem. Int. Ed.	2006	118/ 45	302-307/ 296-301



Alexandrov, H.
Waldmann

J. M. Palomo, M. Lumbierres, H. Waldmann	Efficient Solid-Phase Lipopeptide Synthesis Employing the Ellman Sulfonamide Linker	Angew. Chem./Angew. Chem. Int. Ed.	2006	118/ 45	491- 495/477-481
A. Watzke, M. Köhn, M. Gutierrez- Rodriguez, R. Wacker, H. Schröder, R. Breinbauer, J. Kuhlmann, K. Alexandrov, C. M. Niemeyer, R. S. Goody, H. Waldmann	Site-Selective Protein Immobilization by Staudinger Ligation	Angew. Chem. Int. Ed.	2006	45	1408-1412
L. Brunsveld, J. Kuhlmann, K. Alexandrov, A. Wittinghofer, R. S. Goody, H. Waldmann	Lipidated Ras and Rab Peptides and Proteins – Synthesis, Structure and Function (Review)	Angew. Chem. Int. Ed.	2006	45	6622-6646
G. Reuther, K.-T. Tan, J. Köhler, C. Nowak, A. Pampel, K. Arnold, J. Kuhlmann, H. Waldmann, D. Huster	Structural Model of the Membrane- Bound C-Terminus of Lipid-Modified Human N-Ras Protein	Angew. Chem. Int. Ed.	2006	45	5387-5390
B. D. Charette, R. G. MacDonald, S. Wetzel, D. B. Berkowitz, H. Waldmann	Protein Structure Similarity Clustering: Dynamic Treatment of PDB Structures Facilitates Clustering	Angew. Chem. Int. Ed.	2006	45	7766-7770
B. Dursina, R. Reents, M. Thutewohl, A. Veligodsky, A. Kalinin, V. Evstifeev, D. Ciobanu, S. E. Szedlacsek, H. Waldmann, R. S. Goody, K. Alexandrov	Identification and Specificity Profiling of Protein Prenyltransferase Inhibitors Using New Fluorescent Phosphoisoprenoids	J. Am. Chem. Soc.	2006	128	2822-2835
C. Nicolini, J. Baranski, S. Schlummer, J. Palomo, M. Lumbierres-Burgues, M. Kahms, J. Kuhlmann, S. Sanchez, E. Gratton, H. Waldmann, R. Winter	Visualizing Association of N-Ras in Lipid Microdomains: Influence of Domain Structure and interfacial Adsorption	J. Am. Chem. Soc.	2006	128	192-201
G. Reuther, K.-T. Tan,	The Lipidated Membrane Anchor of Full	J. Am. Chem. Soc.	2006	128	13840-



A. Vogel, C. Nowak, K. Arnold, J. Kuhlmann, H. Waldmann, D. Huster	Length N-Ras Protein Shows an Extensive Dynamics as Revealed by Solid-State NMR Spectroscopy				13846
A.-B. Garcia, T. Lessmann, J. Umarye, V. Mamane, S. Sommer, H. Waldmann	Stereocomplementary synthesis of a natural product-derived compound collection on the solid phase	ChemComm	2006		3868-3870
D. Gottlieb, C. Grunwald, C. Nowak, J. Kuhlmann, H. Waldmann	Intein-Mediated in vitro Synthesis of Lipidated Ras Proteins	ChemComm.	2006		260-262
T. Lessmann, H. Waldmann	Enantioselective Synthesis on the Solid Phase	ChemComm	2006		3380-3389
M. Biel, P. Deck, A. Giannis, H. Waldmann	Synthesis and Evaluation of Acyl-Protein Thioesterase 1 (APT1) Inhibitors	Chem. Eur. J.	2006	12	4121-4143
A. Dantas de Araújo, J. M. Palomo, J. Cramer, O. Seitz, K. Alexandrov, H. Waldmann	Diels-Alder Ligation of Peptides and Proteins	Chem. Eur. J.	2006	12	6095-6109
S. Basu, H. Waldmann	The Regioselectivity in the Formation of Small and Medium Sized Cyclic Ethers by Diene-Ene Ring Closing Metathesis	J. Org. Chem.	2006	71	3977-3979
M. A. Sanz, T. Voigt, H. Waldmann	Enantioselective Catalysis on the Solid Phase: Synthesis of Natural Product Derived Tetrahydropyrans Employing the Enantioselective Oxa-Diels-Alder Reaction [^]	Adv. Synth. Catal,	2006	348	1511-1515
S. Röttger, H. Waldmann	Solid Phase Synthesis of Decalin Scaffolds by Robinson Annulation with Immobilised Nazarov Reagents	Eur. J. Org. Chem.	2006		2093-2099
M. Lumbierres, J. M. Palomo, G. Kragol, H. Waldmann	Solid-Phase Synthesis of palmitoylated and farnesylated lipopeptides employing the fluoride-labile PTMSEL linker	Tetrahedron Lett.	2006	47	2671-2674
D. Müller, A. Krick, S. Kehraus, C. Mehner, M. Hart, F. C. Küpper, K. Saxena, H. Prinz, H. Schwalbe, P. Janning, H. Waldmann, G. M. König	Brunsvicamide A-C, Sponge-related Cyanobacterial Peptides with <i>Mycobacterium Tuberculosis</i> Protein Tyrosine Phosphatase Inhibitory Activity	J. Med. Chem	2006	49	4871-4878
S. Schlummer, R.	Influence of Serine-O-Glycosylation or	ChemBioChem	2006	7	88-97



Vetter, N. Kuder, A. Henkel, Y.-X. Chen, Y.-M. Li, J. Kuhlmann, H. Waldmann	O-Phosphorylation Close to the vJun Nuclear Localization Sequence on Nuclear Import				
Y.-W. Wu, H. Waldmann, R. Reents, F. H. Ebetino, R. S. Goody, K. Alexandrov	A Protein Fluorescence Amplifier: Continuous Fluorometric Assay for Rab Geranylgeranyltransferase	ChemBioChem	2006	7	1859-1861
T. Weide, L. Arve, H. Prinz, H. Waldmann, H. Kessler	3-Substituted Indolizine-1-carbonitrile Derivatives as Phosphatase Inhibitors	Bioorg. Med. Chem. Lett.	2006	16	59-63
A. Watzke, M. Gutierrez-Rodriguez, M. Köhn, R. Wacker, H. Schröder, R. Breinbauer, J. Kuhlmann, K. Alexandrov, C. M. Niemeyer, R. S. Goody, H. Waldmann	A Generic Building Block for C- and N- Terminal Protein-Labeling and Protein- Immobilization	Bioorg. Med. Chem.	2006	14	6288-6306
S. F. Seibert, E. Eguereva, A. Krick, S. Kehraus, E. Voloshina, G. Raabe, J. Fleischhauer, E. Leistner, M. Wiese, H. Prinz, K. Alexandrov, P. Janning, H. Waldmann, G. M. König	Polyketides from the marine-derived fungus <i>Ascochyta salicorniae</i> and their potential to inhibit protein phosphatases	Org. Biomol. Chem.	2006	4	2233-2240
A. Hemmerlin, R. Reents, J. Mutterer, J.-F. Feldtrauer, H. Waldmann, T. J. Bach	Monitoring farnesol-induced toxicity in tobacco BY-2 cells with a fluorescent analog	Arch. Biochem. and Biophys.	2006	448	93-103
A. Meister, C. Nicolini, H. Waldmann, J. Kuhlmann, A. Kerth, R. Winter, A. Blume	Insertion of lipidated Ras Proteins Into lipid monolayers studied by infrared reflection absorption spectroscopy (IRRAS)	Biophys. J.	2006	91	1388-1401
L. Arve. T. Voigt, H. Waldmann	Charting Biological and Chemical Space: PSSC and SCONP as Guiding Principles for the Development of Natural Product Scaffold Based Compound Collections	QSAR Comb. Sci.	2006	25	449-456
M. A. Koch, H. Waldmann	Protein Domain Fold Similarity and Natural Product Structure as Guiding Principles for Compound Library Design	Ernst Schering Research Foundation, Workshop 58, S. Jaroch, H. Weinmann (eds.)	2006		89-109



L. Brunsveld, J. Kuhlmann, H. Waldmann	Synthesis of Palmitoylated Ras-Peptides and -Proteins	Methods	2006	40	151-165
H. Waldmann, R. Breinbauer, S. Arndt	Das Chemical Genomics Centre	Laborwelt	2006	7	42-43
H.-D. Arndt, C.M. Niemeyer, H. Waldmann	Chemical Biology Education in Dortmund: A Joint Endeavour with a Max Planck Institute	ACS Chem. Biol.	2006	1	407-410
F. Dekker, S. Wetzel, Hr. Waldmann	Natural Product Scaffolds and Protein Structure Similarity Clustering (PSSC) as Inspiration Sources for Compound Library Design in Chemogenomicx and Drug Development	Chemogenomics: An Emergine Strategy for Rapid Target and Drug Discovery	2006		59-84
Y.-W. Wu, K.-T. Tan, H. Waldmann, R. S. Goody, K. Alexandrov	Interaction analysis of prenylated Rab GTPase with Rab escort protein and GDP dissociation inhibitor explains the need for both regulators	PNAS	2007	104	12294-12299
D. Rauh, H. Waldmann	Linking Chemistry and Biology for the Study of Protein Function	Angew. Chem. Int. Ed.	2007	46	826-829
H. Schroeder, B. Ellinger, C. F. W. Becker, H. Waldmann, C. M. Niemeyer	Generation of Life-Cell Microarrays by Means of DNA-Directed Immobilization of Specific Cell- Surface Ligands	Angew. Chem. Int. Ed.	2007	46	4180-4183
M. Köhn, P. Jonkheijm, M. Gutierrez-Rodriguez, S. Wetzel, R. Wacker, H. Schröder, H. Prinz, C. M. Niemeyer, R. Breinbauer, S. E. Szedlacsek, H. Waldmann	A Microarray Strategy for Mapping of Protein Tyrosine Phosphatase Substrate Specificity	Angew. Chem. Int. Ed.	2007	46	7700-7703
B. Bulic, M. Pickhardt, I. Khlistunova, J. Biernat, E.-M. Mandelkow, E. Mandelkow, H. Waldmann	Rhodanine-Based Tau Aggregation Inhibitors in Cell Models of Tauopathy	Angew. Chem. Int. Ed.	2007	46	9215-9219
T.-S. Hu, R. Tannert, H.-D. Arndt, H. Waldmann	Solid-phase based synthesis of Jaspilakinolide analogs by intramolecular azide/alkyne cycloadditions	Chem. Comm.	2007	38	3942-3944
T. Lessmann, M. G. Leuenberger, S. Menninger, M. Lopez-Canet, O. Müller, S.	Natural Product-Derived Modulators of Cell Cycle Progression and Viral Entry by Enantioselective Oxa-Diels-Alder Reactions on the Solid Phase	Chem. & Biol.	2007	14	443-451



Hümmer, J. Bormann,
K. Korn, E. Fava, M.
Zerial, T. U. Mayer, H.
Waldmann

J. D. Umarye, T. Lessmann, A. B. Garcia, V. Mamane, S. Sommer, H. Waldmann	Biology-Oriented Synthesis of Stereochemically Diverse Natural- Product Derived Compound Collections by Iterative Allylations on the Solid Support	Chem. Eur. J.	2007	13	3305-3319
I. R. Corrêa Jr., A. Nören-Müller, H.-D. Ambrosi, S. Jakupovic, K. Saxena, H. Schwalbe, H. Prinz, M. Kaiser, H. Waldmann	Identification of Inhibitors for Mycobacterial Protein Tyrosine Phosphatase B (MptpB) by Biology- Oriented Synthesis (BIOS)	Chem. Asian J.	2007	2	1109-1126
U. T. T. Nguyen, J. Cramer, J. Gomis, R. Reents, M. Gutierrez- Rodriguez, R. S. Goody, K. Alexandrov, H. Waldmann	Exploiting the Substrate Tolerance of Farnesyltransferase for Site-Selective Protein Derivatization	ChemBioChem	2007	8	408-423
A. Schuffenhauer, P. Ertl, S. Roggo, S. Wetzel, M. A. Koch, H. Waldmann	The Scaffold Tree – Visualization of the Scaffold Universe by Hierarchical Scaffold Classification	J. Chem. Inf. Model.	2007	47	47-58
V. Mamane, A. B. García, J. D. Umarye, T. Lessmann, S. Sommer, H. Waldmann	Stereoselective allylation of aldehydes on solid support and its application in biology-oriented synthesis (BIOS)	Tetrahedron	2007	63	5754-5767
K. Hübel, A. Krämer, H. Waldmann	Effektive Arzneimittelentwicklung mit „natürlichen“ Substanz-bibliotheken	Laborwelt	2007	3	33-35
F. Bringezu, M. Majerowicz, S. Wen, G. Reuther, K.-T. Tan, J. Kuhlmann, H. Waldmann, D. Huster	Membrane Binding of a Lipidated N-Ras Protein Studied in Lipid Monolayers	Eur Biophys J	2007	36	491-498
S. Wetzel, A. Schuffenhauer, S. Roggo, P. Ertl, H. Waldmann	Cheminformatic Analysis of Natural Products and their Chemical Space	Chimia	2007	61	355-360
K. Bierhals, A. C. Sondersorg, C.-T. Lin, C. Rosenbaum, H. Waldmann, F. Wehner	The ϵ -Isoform of PKC Mediates the Hypertonic Activation of Cation Channels in Confluent Monolayers of Rat Hepatocytes	Cell Physiol Biochem	2007	20	397-404



A. Ulaczyk-Lesanko, E. Pelletier, M. Lee, H. Prinz, H. Waldmann, D.G. Hall	Optimization of Three- and Four- Component Reactions for Polysubstituted Piperidines. Application to the Synthesis and Preliminary Biological Screening of a Prototype Library	J. Comb. Chem.	2007	9	695-703
S. Broussy, H. Waldmann	Solid Phase Synthesis of Highly Substituted Tetrahydropyrans by Tandem ene- Reaction/Intramolecular Sakurai cyclisation	J. Comb. Chem.	2007	9	1138-1143
A. Vogel, K.-T Tan, H. Waldmann, S. E. Feller, M. F. Brown, D. Huster	Flexibility of Ras Lipid Modifications Studied by ² H Solid-State NMR and Molecular Dynamics Simulations	Biophys. J.	2007	93	2697-2712
R. Tannert, M. Schürmann, H. Preut, H.-D. Arndt, H. Waldmann	(S)-3-[(R)-2,4-Dimethylpent-4-enoyl]-4- isopropyl-5,5-diphenyl-1,3-oxazolidin-2- one	Acta Crystallographica Section E	2007	E63	04381
A. Pyriochou, S. Tsigkos, T. Vassilakopoulos, T. Cottin, Z. Zhou, E. Gourzoulidou, C. Roussos, H. Waldmann, A. Giannis, A. Papapetropoulos	Anti-angiogenic properties of a sulindac analogue	Br J Pharmacol	2007	152	1207-1214
Z. Guo, Y.-W. Wu, K.- T. Tan, R. S. Bon, E. Guiu-Rozas, C. Delon, U. T. Nguyen, S. Wetzel, S. Arndt, R. S. Goody, W. Blankenfeldt, K. Alexandrov, H. Waldmann	Development of Selective RabGGTase Inhibitors and Crystal Structure of a RabGGTase-Inhibitor Complex	Angew. Chem. Angew. Chem. Int. Ed.	2008	120 47	3807-3810 3747-3750
P. Jonkheijm, D. Weinrich, M. Köhn, H. Engelkamp, P.C.M. Christianen, J. Kuhlmann, J.C. Maan ³ , D. Nüsse, H. Schroeder, R. Wacker, R. Breinbauer, C.M. Niemeyer, H. Waldmann	Photochemical Surface Patterning by Thiol-Ene Reaction	Angew. Chem. Angew. Chem. Int. Ed.	2008	120 47	4493-4496 4421-4424
R. Mishra, B. Bulic, D. Sellin, S. Jha, H.	Small-Molecule Inhibitors of Islet Amyloid Polypeptide Fibril Formation	Angew. Chem. Int. Ed.,	2008	47	4679-4682



Waldmann, and R.
Winter

A. Nören-Müller, W. Wilk, K. Saxena, H. Schwalbe, M. Kaiser, H. Waldmann	Discovery of a New Class of <i>Mycobacterium tuberculosis</i> Protein Tyrosine Phosphatase B by Biology Oriented Synthesis	Angew. Chem. Angew. Chem. Int. Ed.	2008	120 47	6061-6066 5973-5977
H. Waldmann, T.-S. Hu, S. Renner, S. Menninger R. Tannert, T. Oda, H.-D. Arndt	Total Synthesis of Condramide C and its Binding Mode to F-Actin	Angew. Chem. Int. Ed.	2008	47	6473-6477
H. Waldmann, V. Khedkar, H. Dücker, M. Schürmann, I. M. Oppel, K. Kumar	Asymmetric Synthesis of Natural Product Inspired Tricyclic Benzopyrones by an Organocatalyzed Annulation Reaction	Angew. Chem. Int. Ed	2008	47	6869-6872
P. Jonkheijm, D. Weinrich, H. Schroeder, C. M. Niemeyer, H. Waldmann	Chemical Strategies for Generating Protein Biochips	Angew. Chem. Int. Ed., Angew. Chem.	2008	47/ 120	9618-9647/ 9762-9792 (dt.)
E. Wong, V. Okhonin, M. V. Berezovski, T. Nozaki, H. Waldmann, K. Alexandrov, S. N. Krylov	"Inject-Mix-React-Separate-and-Quantitate" (IMReSQ) Method for Screening Enzyme Inhibitors	J. Am Chem. Soc.	2008	36	11862-11863
Z. Wang, C. Gu, T. Colby, T. Shindo, R. Balamurugan, H. Waldmann, M. Kaiser, R. A. L. v. d. Hoorn	β -Lactone probes identify a papain-like peptide ligase in <i>Arabidopsis thaliana</i>	Nat. Chem. Biol.	2008	4	557-563
Z. Guo, Y.-W. Wu, D. Das, C. Delon, J. Cramer, S. Yu, S. Thuns, N. Lupilova, H. Waldmann, L. Brunsveld, R. S. Goody, K. Alexandrov, W. Blankenfeldt	Structures of RabGGTase-substrate/product complexes provide insights into the evolution of protein prenylation	EMBO J.	2008	27	2444-2456
H. Waldmann, M. Kühn, W. Liu, K. Kumar	Reagent controlled domino synthesis of skeletally diverse compound collections	ChemComm	2008	10	1211-1213
T. Govindaraju, P. Jonkheijm, L. Gogolin, H. Schroeder, C. F. W. Becker, C. M. Niemeyer, H. Waldmann	Surface immobilization of biomolecules by click sulfonamide reaction	ChemComm	2008	10	3723-3725



H. Waldmann, Y. He, H. Tan, L. Arve, H.-D. Andt	Flexible total synthesis of biphenomycin B	ChemCommun.	2008		5562-5564
N. Bisek, S. Wetzel, H.-D. Arndt, H. Waldmann	Synthesis and Conformational Analysis of Stevastelin C3 Analogs and Their Activity Against the Dual- Specific Vaccinia H1-Related Phosphatase	Chem. Eur J.	2008	14	8847-8860
F. Wehner, A. Nören- Müller, O. Müller, I. Reis-Correa jr, A. Giannis, H. Waldmann	Indoloquinolizidine Derivatives as Novel and Potent Apoptosis Inducers and Cell- Cycle Blockers	ChemBioChem	2008	9	401-405
S. Sommer, M. Kühn, H. Waldmann	Solid-Phase Synthesis of [5.5]- Spiroketals	Adv. Synth. Catal.	2008	350	1736-1750
T. Walther, H.-D. Arndt, H. Waldmann	Solid-Support Based Total Synthesis and Stereochemical Correction of Brunsvicamide A	Org. Lett.	2008	10	3199-3202
H. Waldmann, G. V. Karanukar, K. Kumar	Gold (III)-Mediated Aldol Condensations Provide Efficient Access to Nitrogen Heterocycles	Org. Lett.	2008	10	2159-2162
G. Triola, L. Brunsveld, H. Waldmann	Racemization-free synthesis of S- alkylated cysteines <i>via</i> thiol-ene reaction	J. Org. Chem.	2008	73	3646-3649
M. Scheck, M. A. Koch, H. Waldmann	Synthesis of a dysidiolide-inspired compound library and discovery of acetylcholinesterase inhibitors based on protein structure similarity clustering (PSSC)	Tetrahedron	2008	64	4792-4802
J. Güldenaupt, Y. Adigüzel, J. Kuhlmann, H. Waldmann, C. Kötting, K. Gerwert	Secondary structure of lipidated Ras bound to a lipid bilayer	FEBS Journal	2008	23	5910-5918
K. Hübel, T. Leßmann, H. Waldmann	Chemical biology - identification of small molecule modulators of cellular activity by natural product inspired synthesis	Chem. Soc. Rev.	2008	37	1361-1374
M. Kaiser, S. Wetzel, K. Kumar, H. Waldmann	Biology-Inspired synthesis of compound libraries	Cell.Mol. Life Sci:	2008	65	1186-1201
T. Lessmann, H. Waldmann	Asymmetric Synthesis on the Solid Phase	<i>Asymmetric Synthesis – The Essentials</i> (Eds.: M. Christmann, S. Bräse), Wiley-VCH, Weinheim	2008	2 nd ed.	299-304



K. Kumar, S. Wetzel, H. Waldmann	Biology Oriented Synthesis (BIOS) and Diversity Oriented Synthesis (DOS) in Compound Collection Development	The Practice of Medicinal Chemistry, 3 rd Edition, Ed. Camille Georges Wermuth	2008	3 rd Ed.	187-209
U. T. T. Nguyen, Z. Guo, C. Delon, Y. Wu, C. Deraeve, B. Fränzel, R. S. Bon, W. Blankenfeldt, R. S. Goody, H. Waldmann, D. Wolters, K. Alexandrov	Analysis of the eukaryotic prenylome by isoprenoid affinity-tagging"	Nat. Chem. Biol.	2009	5	227-235
S. Wetzel, K. Klein, S. Renner, D. Rauh, T. I. Oprea, P. Mutzel, H. Waldmann	Interactive exploration of chemical space with Scaffold Hunter	Nat. Chem. Biol.	2009	5	581-583
S. Renner, W. A. L.. van Otterlo, M. Dominguez Seoane, S. Möcklinghoff, B. Hofmann, S. Wetzel, A. Schuffenhauer, P. Ertl, T. I. Oprea, D. Steinhilber, L. Brunsveld, D. Rauh, H. Waldmann	Bioactivity-guided mapping and navigation of chemical space	Nat. Chem. Biol.	2009	5	585-592
H. Waldmann	Killing 84 birds with one stone	Nat. Chem. Biol. (News and Views)	2009	5	76-77
T. I. Oprea, C. G. Bologa, S. Boyer, R. F. Curpan, R. C. Glen, A. L. Hopkins, C. A. Lipinski, G. R. Marshall, Y. c. Martin, L. Ostopovici-Halip, G. Rishton, O. Ursu, R. J. Vaz, C. Waller, H. Waldmann, L. A. Sklar	A crowdsourcing evaluation of the NIH chemical probes	Nat. Chem. Biol.	2009	5	441-447
G. Siegel, G. Obernosterer, R. Fiore, S. Bicker, M. Christensen, S. Khudayberdiev, P. F. Leuschner, C. J. L. Busch, C. Kane, K. Hübel, F. Dekker, C. Hedberg, B. Rengarajan, C. Drepper, H.	A functional screen implicates microRNA-138-dependent regulation of the depalmitoylation enzyme APT1 in dendritic spine morphogenesis	Nature Cell Biology	2009	11	705-730



Waldmann, S.
Kauppinen, M. E.
Greenberg, A.
Draguhn, M.
Rehmsmeier, J.
Martinez, G. M.
Schratt

B. Bulic, M. Pickhardt, B. Schmidt, E.-M. Mandelkow, H. Waldmann, E. Mandelkow	Development of Tau Aggregation Inhibitors for Alzheimer's Disease	Angew. Chem. Int. Ed./ Angew. Chem.	2009	48/ 121	1740-1752/ 1772-1785
K. Kumar, H. Waldmann	Synthesis of Natural Product Inspired Compound Collections	Angew. Chem. Int. Ed./	2009	48/ 121	3224-3242/ 3272-3290
D. Weinrich, P. Jonkheijm, C. M. Niemeyer, H. Waldmann	Applications of Protein Biochips in Biomedical and Biotechnical Research	Angew. Chem. Angew. Chem. Int. Ed./ Angew. Chem.	2009	48/ 121	7744-7751/ 7880-7888
T. Knoth, K. Warburg, C. Katzka, A. Rai, A. Wolf, A. Brockmeyer, P. Janning, T. F. Reubold, S. Eschenburg, D. J. Manstein, K. Hübel, M. Kaiser, H. Waldmann	The Ras Pathway Modulator Meloplin A Targets Dynamins	Angew. Chem. Int. Ed / Angew. Chem.	2009	48 / 121	7240-7245 / 7376-7381
U. Westerlind, H. Schröder, A. Hobel, N. Gaidzik, A. Kaiser, C. M. Niemeyer, E. Schmitt, H. Waldmann, H. Kunz	Tumor-Associated MUC1 Tandem- Repeat Glycopeptide Microarrays to Evaluate Serum- and Monoclonal- Antibody Specificity	Angew. Chem. Int. Ed. / Angew. Chem.	2009	48 / 121	8263-8267 / 8413-8417
A. Vogel, G. Reuther, K. Weise, G. Triola, J. Nikolaus, K.-T. Tan, C. Nowak, A. Herrmann, H. Waldmann, R. Winter, D. Huster	The Lipid Modifications of Ras that Sense Membrane Environments and Induce Local Enrichment	Angew. Chem. Int. Ed./ Angew. Chem.	2009	48 / 121	8784-8787 / 8942-8945
K. Weise, G. Triola, L. Brunsveld, H. Waldmann, R. Winter	Influence of the Lipidation motif on the Partitioning and Association of N-Ras in Model Membrane Subdomains	J. Am. Chem. Soc	2009	131 (4)	1557-1564
R. Tannert, T.-S. Hu, H.-D. Arndt, H. Waldmann	Solid-phase based total synthesis of Jasplakinolide by means of ring-closing metathesis methodology	ChemComm.	2009	12	1493-1495
M. Yoshida, C.	Traceless solid phase synthesis of	ChemComm.	2009	20	2926-2928



Hedberg, M. Kaiser, H. Waldmann	natural product inspired <i>cis</i> -1,2- dehydrodecalins				
W. Wilk, A. Nören- Müller, M. Kaiser, H. Waldmann	Biology-Oriented Combined Solid- and Solution-Phase Synthesis of a Macroline-Like Compound Collection	Chem. a Eur. J.	2009	15	11976- 11984
K.-T. Tan, E. Guiu- Rozas, R. S. Bon, Z. Guo, C. Delon, S. Wetzel, S. Arndt, K. Alexandrov, H. Waldmann, R.S. Goody, Y.-W.Wu, W. Blankenfeldt	Design, Synthesis and Characterization of Peptide-Based Rab Geranylgeranyl Transferase Inhibitors	J. Med. Chem.	2009	52	8025-8037
M. Alexander, M. Gerauer, M. Pechlivanis, B. Popkirova, R. Dvorsky, L. Brunsveld, H. Waldmann, J. Kuhlmann+	Mapping the Isoprenoid-binding pocket of PDE δ by a Semisynthetic, Photoactivatable N-Ras ILpoprotein	ChemBioChem	2009	1	98-108
T. Walther, S. Renner, H. Waldmann, H.-D. Arndt	Synthesis and Structure-Activity Correlation of a Brunsvicamide-Inspired Cyclopeptide Collection	ChemBioChem	2009	10	1153-1162
G. Triola, S. Wetzel, B. Ellinger, M. A. Koch, K. Hübel, D. Rauh, H. Waldmann	ATP competitive lihibitors of D-Alanine- D-Alanine Ligase Based on Protein Kinase Inhibitor Scaffolds	Bioorg. Med. Chem.	2009	17	1079-1087
W. Wilk, H. Waldmann, M. Kaiser	γ -Pyrone natural products – A privileged compound class provided by nature	Biorg. Med. Chem.	2009	17	2304-2309
B. Ludolph, H. Waldmann	Solid-pPhase Synthesis of benzodiazepinediones mimicking the C- terminus of the H-Ras protein	Tetrahedron Letters	2009	50	3148-3150
V. V. Vintonyak, A. P. Antonchick, D. Rauh, H. Waldmann	The therapeutic potential of phosphatase inhibitors	Current Opinion in Chemical Biology	2009	13	272-283
L. Brunsveld, H. Waldmann, D .Huster	Membrane Binding of lipidated Ras peptides and proteins – the structural point of view	Biochimica et Biochimica Acta – Biomembranes	2009	1788	273-288
M. Gerauer, S. Koch, H. Waldmann	Lipidated peptide Synthesis	In: Wiley Encyclopedia of Chemical Biology, Publisher: John Wiley&Sons, Ed.: T. P. Begley	2009		520-530
H. Waldmann, P. Janning (eds.)	Chemical Biology – Learning through Case Studies	Wiley-VCH, Weinheim	2009		



F. J. Dekker, O. Rocks, N. Vartak, S. Menninger, C. Hedberg, R. Balamurugan, S. Wetzels, S. Renner, M. Gerauer, B. Schölermann, M. Rusch, J. W. Kramer, D. Rauh, G. W. Coates, L. Brunsveld, PIH Bastiaens, H. Waldmann	Small-molecule inhibition of APT1 affects Ras localization and signaling	Nat. Chem. Biol.	2010	6	449-456
Y.-Wu, L. K. Oesterlin, K.-T. Tan, H. Waldmann, K. Alexandrov, R. S. Goody	Membrane targeting mechanism of Rab GTPases elucidated by semisynthetic protein probes	Nat. Chem. Biol.	2010	6	534-540
A. P. Antonchick, C. Gerding-Reimers, M. Catarinella, M. Schürmann, H. Preut, S. Ziegler, D. Rauh, H. Waldmann	Highly enantioselective synthesis and cellular evaluation of spirooxindoles inspired by natural products	Nat. Chem.	2010	2	735-740
O. Rocks, M. Gerauer, N. Vartak, S. Koch, Z.-P. Huang, M. Pechlivanis, J. Kuhlmann L. Brunsveld, A. Chandra, B. Ellinger, H. Waldmann, P. I. H. Bastiaens	The Palmitoylation Machinery is a Spatially Organizing System for Peripheral Membrane Proteins	Cell	2010	141	458-471
D. Weinrich, P.-C. Lin, P. Jonkheijm, U. T.T. Nguyen, H. Schröder, C. M. Niemeyer, K. Alexandrov, R. Goody, H. Waldmann	Oriented Immobilization of Farnesylated Proteins by the Thiol-Ene Reaction	Angew. Chem./ Angew. Chem. Int. Ed.,	2010	122/ 49	1274-1279/ 1252-1257
S. Wetzels, W. Wilk, S. Chammaa, B. Sperl, A. G. Roth, A. Yektaoglu, S. Renner, T. Berg, C. Arenz, A. Giannis, T. I. Oprea, D. Rauh, M. Kaiser, H. Waldmann	A Scaffold-Tree-Merging Strategy for Prospective Bioactivity Annotation of γ -Pyrones	Angew.Chem./ Angew. Chem Int. Ed.	2010	122/ 49	3748-3752 3666-3670
R. Rose, S. Erdmann, S. Bovens, A. Wolf, M.	Identification and Structure of Small-Molecule Stabilizers of 14-3-3 Protein-	Angew.Chem. Int. Ed.	2010	24	4129-4132



Rose, S. Hennig, H. Waldmann, C. Ottmann	Protein Interactions				
Y.-C. Chen, S. Koch K. Uhlenbrock, K. Weise, D. Das, L. Gremer, L. Brunsveld, A. Wittinghofer, R. Winter, G. Triola, H. Waldmann	Synthesis of the Rheb and K-Ras4B GTPases	Angew. Chem./ Angew. Chem. Int. Ed. –	2010	122/ 49	6226-6231 6090-6095
V.V. Vintonyak, K. Warburg, H. Kruse, S. Grimme, K. Hübel, D. Rauh, H. Waldmann	Identification of Thiazolidinones Spiro- Fused to Indolin-2-ones as Potent and Selective Inhibitors of <i>Mycobacterium tuberculosis</i> Protein Tyrosine Phosphatase B	Angew. Chem. Int. Ed.	2010	49	5902-5905
L. Yi, H. Sun, Y.-W. Wu, G. Triola, H. Waldmann, R. S. Goody	A Highly Efficient Strategy for Modification of Proteins at the C- Terminus	Angew. Chem. / Angew. Chem. Int. Ed.	2010	122/ 49	9607-9611 9417-9421
R. Tannert, L.-G. Milroy, B. Ellinger, T.- S. Hu, H.-D. Arndt, H. Waldmann	Synthesis and Structure-Activity Correlation of Natural-Product Inspired Cyclodepsipeptides Stabilizing F-Actin	J. Am. Chem. Soc.	2010	9	3063-3077
H. Waldmann, L. Eberhardt, K. Wittstein, K. Kumar	Silver catalyzed cascade synthesis of alkaloid ring systems: concise total synthesis of fascaplysin, homofascaplysin C and analogues	ChemComm, invited communication for thematic issue: "Enzymes and Proteins"	2010	46	4622-4624
G. Triola, M. Gerauer, K. Görmer, L. Brunsveld, H. Waldmann	Solid-Phase Synthesis of Lipidated Ras Peptides Employing the Ellman Sulfonamide Linker	Chem. – Eur. J. (Conference issue)	2010	16	9585-9591
D. Weinrich, M. Köhn, P. Jonkheijm, U. Westerlind, L. Dehmelt, H. Engelkamp, P. C. M. Christianen, J. Kuhlmann, J. C. Maan, D. Nüsse, H. Schröder, R. Wacker, E. Voges, R. Breinbauer, H. Kunz, C. M. Niemeyer, H. Waldmann	Preparation of Biomolecule Microstructures and Microarrays by Thiol-ene-Photoimmobilization	ChemBioChem	2010	11	235-247
A. Robubi, H. Waldmann, D. Rauh	RAF Kinase Inhibitors in Cancer Treatment: Like a Bull in a China Shop?	ChemBioChem	2010	11	1645-1648
R. Bon, H. Waldmann	Bioactivity-Guided Navigation of Chemical Space	Accounts of Chemical Research	2010	43	1103-1114



Po-Chiao Lin, D. Weinrich, H. Waldmann	Protein Biochips: Oriented Surface Immobilization of Proteins	Macromol. Chem. Phys., Special Issue dedicated to the 80th birthday of Prof. H. Ringsdorf	2010	2	136-144
H. Lachance, S. Wetzel, H. Waldmann	Role of Natural Products in Drug Discovery	Book Chapter in: "Lead Generation Approaches in Drug Discovery, eds. Zoran Rakovic, Richard Morphy, Wiley & Sons	2010		187-229
K. Görmer, L. Brunsveld, H. Waldmann	Lipidation of Peptides and Proteins	In: Comprehensive Natural Products II Chemistry and Biology; Mander, L., Lui, H.-W., Eds, Elsevier, Oxford	2010		531-586
K. Görmer, H. Waldmann, G. Triola	Efficient Microwave-Assisted Synthesis of Unsymmetrical Disulfides	J. Org. Chem.	2010	75	1811-1813
A. Gohlke, G. Triola, H. Waldmann, R. Winter	Influence of the Lipid Anchor Motif of N-Ras on the Interaction with Lipid Membranes – a Surface Plasmon Resonance Study	Bioph. J.	2010	98	2226-2235
W. Wilk, T. J. Zimmermann, M. Kaiser, H. Waldmann	Principles, implementation and application of biology-oriented synthesis (BIOS)	Biol. Chem.	2010	391	491-497
K. Weise, G. Triola, S. Janosch, H. Waldmann, R. Winter	Visualizing association of lipidated signaling proteins in heterogeneous membranes – Partitioning into subdomains, lipid sorting, interfacial adsorption, and protein association	Biochimica et Biophysica Acta - Biomembranes	2010	1798	1409-1417
A. Vogel, G. Reuther, M. B. Roark, K.-T. Tan, H. Waldmann, D. Huster	Backbone conformational flexibility of the lipid modified membrane anchor of the human N-Ras protein investigated by solid-state NMR and molecular dynamics simulation	Biochimica et Biophysica Acta – Biomembranes	2010	1798	275-285
D. Weinrich, H. Waldmann	Methods for Site-Selective Chemical Protein Immobilization	Revisita de la Societat Catalana de Química	2010	9	47-50
A. Hakkim, T.A. Fuchs, N. E. Martinez, S. Hess, H. Prinz, A. Zychlinsky, H. Waldmann	Activation of the Raf-MEK-ERK-pathway is required for neutrophil extracellular trap formation	Nat. Chem. Biol	2011	7	75-77



S. A. Ismail, Y.-X. Chen, A. Rusinova, A. Chandra, M. Bierbaum, L. Gremer, G. Triola, H. Waldmann, P. I. H. Bastiaens, A. Wittinghofer	Arl2-GTP and Arl3-GTP regulate a GDI-like transport system for farnesylated cargo	Nature Chem. Biol.	2011	7	942-949
S. Basu, B. Ellinger, S. Rizzo, C. Deraeve, M. Schürmann, H. Preut, H.-D. Arndt, H. Waldmann	Biology-oriented synthesis of a natural-product inspired oxepane collection yields a small-molecule activator of the Wnt-pathway	Proc. Natl. Acad. Sci.	2011	108	6805-6810
L. Radtke, M. Willot, H. Sun, S. Ziegler, S. Sauerland, C. Strohmam, R. Fröhlich, P. Habenberger, H. Waldmann, M. Christmann	Total Synthesis and Biological Evaluation of (-)-Englerin A and B: Synthesis of Analogues with Improved Activity Profile	Angew. Chem. Int. Ed.	2011	17	3998-4002
R. S. Bon, Z. Guo, E. A. Stigter, S. Wetzel, S. Menninger, A. Wolf, A. Choidas, K. Alexandrov, W. Blankenfeldt, R. S. Goody, H. Waldmann	Structure-Guided Development of Selective RabGGTase Inhibitors	Angew. Chem. Int. Ed. Angew. Chem.	2011	50 / 123	4957-4961 / 5059-5063
Yi, L., Sun, H., Itzen, A., Triola, G., Waldmann, H., Goody, R.S., Wu, Y.-W.	One-Pot Dual-Labeling of a Protein via Two Chemoselective Reactions	Angew. Chem. Int. Ed.	2011	50	8287-8290
J. Stöckigt, A. P. Antonchick, F. Wu, H. Waldmann	Review: The Pictet-Spengler Reaction in Nature and in Organic Chemistry	Angew. Chem. Int. Ed./ Angew. Chem.	2011	50 / 123	8538-8564/ 8692-8719
K. Wittstein, K. Kumar, H. Waldmann	Gold(I) Catalyzed Synthesis of Benzoxocines via an 8-endo-dig Cyclization	Angew. Chem. Int. Ed. Angew. Chem.	2011	50 123	9076-9079 9242-9246
C. Hedberg, F. J. Dekker, M. Rusch, S. Renner, S. Wetzel, N. Vartak, C. Gerding-Reimers, R. S. Bon, P. I. Bastiaens, H. Waldmann	Development of Highly Potent Inhibitors of the Ras-Targeting Human Acyl Protein Thioesterases Based on Substrate Similarity Design	Angew. Chem. Int. Ed.	2011	50	9832-9837
M. Rusch, T. J. Zimmermann, M. Bürger, F. J. Dekker,	Identification of Acyl Protein Thioesterases 1 and 2 as the Cellular Targets of the Ras-Signalling Modulators	Angew. Chem. Int. Ed	2011	50	9838-9842



K. Görmer, G. Triola, A. Brockmeyer, P. Janning, T. Böttcher, S. A. Sieber, I. R. Vetter, C. Hedberg, H. Waldmann	Palmostatin B and M				
S. Wetzel, R. Bon, K. Kumar, H. Waldmann	Review: Biology Oriented Synthesis	Angew. Chem. Int. Ed., Angew. Chem.	2011	50 / 123	10800-10826 / 10990-11018
K. Weise, S. Kapoor, C. Denter, J. Nikolaus, N. Opitz, S. Koch, G. Triola, A. Herrmann, H. Waldmann, R. Winter	Membrane-Mediated Induction and Sorting of K-Ras Microdomain Signaling Platforms	J. Am. Chem. Soc.	2011	133	880-887
H. Schuster, R. Martinez, H. Bruss, A. P. Antonchick, M. Kaiser, M. Schürmann, H. Waldmann	Synthesis of the B- <i>seco</i> Limonoid Scaffold	Chem. Commun.	2011	47 (23)	6545-6547
H. Dückert, V. Khedkar, H. Waldmann, K. Kumar	Lewis Base Catalyzed [4 + 2]-Annulation of Electron-Deficient Chromone-Derived Heterodienes and Acetylenes	Chem. – Eur. J.	2011	17	5130-5137
Y.-P. He, Ha. Tan, L. Arve, S. Baumann, H. Waldmann, H.-D. Arndt	Biphenomycin B and Derivatives: Total Synthesis and Translation Inhibition	Chem. – Asian J.	2011	6	1546-1556
S. Möcklinghoff, W.A.L. van Otterlo, R. Rose, S. Fuchs, T. Zimmermann, M. Dominguez Seoane, H. Waldmann, C. Ottmann, L. Brunsveld	Design and Evaluation of Fragment-Like Estrogen Receptor Tetrahydroisoquinoline Ligands from a Scaffold-Detection Approach	J. Med. Chem.	2011	54	2005-2011
V. V. Vintonyak, H. Waldmann, D. Rauh	Using small molecules to target protein phosphatases	Bioorg. Med. Chem.	2011	19	2145-2155
B. K. Wagner, T. J. Gilbert, J.-i. Hanai, S. Imamura, N. E. Bodycombe, R. S. Bon, H. Waldmann, P. A. Clemons, V. P. Sukhatme, V. S. Mootha	A small-molecule screening strategy to identify suppressors of statin myopathy	ACS Chem. Biol.	2011	6	900-904
A. Richter, C. Hedberg, H. Waldmann	Enantioselective Synthesis of the C ₁₀ -to C ₂₀ -Fragment of Fusicoccin A	J. Org. Chem.	2011	76	6694-6702



V. Vintonyak, K. Warburg, B. Over, K. Hübel, D. Rauh, H. Waldmann	Identification and further development of thiazolidinones spiro-fused to indolin-2-ones as potent and selective inhibitors of <i>Mycobacterium tuberculosis</i> protein tyrosine phosphatase B	Tetrahedron	2011	67	6713-6729
A. P. Antonchick, H. Schuster, H. Bruss, M. Schürmann, H. Preut, D. Rauh, H. Waldmann	Enantioselective synthesis of the dpirotryprostatin A scaffold	Tetrahedron	2011	67	10195-10202
H. Waldmann, H. Bruss, H. Dückert, K. Kumar	Synthesis of novel electron-deficient chromone-fused dienes <i>via</i> phosphine catalyzed [4+2] annulation	Tet. Lett	2011	52	2265-2267
S. Eisenberg, A. J. Beckett, I. A. Prior, F. J. Dekker, C. Hedberg, H. Waldmann, M. Ehrlich, Y. I. Henis	Raft Protein Clustering Alters N-Ras Membrane Interactions and Activation Pattern	Mol. Cell. Biol.	2011	31	3938-3952
T. Hakki, K. Hübel, H. Waldmann, R. Bernhardt	The development of a whole-cell based medium throughput screening system for the discovery of human aldosterone synthase (CYP11B2) inhibitors: Old drugs disclose new applications for the therapy of congestive heart failure, myocardial fibrosis and hypertension	J. Steroid Biochem. Mol Biol	2011	125	120-128
Y.-X. Chen, G. Triola, H. Waldmann	Bioorthogonal Chemistry for Site-Specific Labeling and Surface Immobilization of Proteins	Acc. Chem. Res.	2011	44	762-773
L. Eberhardt, K. Kumar, H. Waldmann	Exploring and Exploiting Biologically Relevant Chemical Space	Current Drug Targets	2011	12	1531-1546
A. Stigter, G. Triola, Roger S. Goody, H. Waldmann	Inhibition of Rab prenylation	The Enzymes Volume 30 Part B: Protein Prenylation, eds. C. A. Hrycyna, M. O. Berge, F. Tamanoi	2011		179-204
H. Dückert, V. Pries, V. Khedkar, S. Menninger, H. Bruss, A. W. Bird, Z. Maliga, A. Brockmeyer, P. Janning, A. Hyman, S. Grimme, M. Schürmann, H. Preut, K. Hübel, S. Ziegler, K. Kumar, H. Waldmann	Natural product-inspired cascade synthesis yields modulators of centrosome integrity	Nat. Chem. Biol.	2012	8	179-184



M. Potowski, M. Schürmann, H. Preut, A. P. Antonchick, H. Waldmann	Programmable enantioselective one-pot synthesis of molecules with eight stereocenters	Nature Chem. Biol.	2012	8	428-430
H. Waldmann	Drug Discovery ... The Third in the Band! (Angewandte Editorial)	Angew. Chem. Int. Ed.	2012	51	6284-6285
M. Potowski, J. O. Bauer, C. Strohmann, A. P. Antonchick, H. Waldmann	Highly Enantioselective Catalytic [6+3] Cycloadditions of Azomethine Ylides	Angew. Chem. Int. Ed.	2012	51	9512-9516
S. Kapoor, G. Triola, I. R. Vetter, M. Ernkamp, H. Waldmann, R. Winter	Revealing conformational substates of lipidated N-Ras protein by pressure modulation	Proc. Natl. Acad. Sci.	2012	109	460-465
J. D. Stewart, R. Marchan, M. S. Lesjak, J. Lambert, R. Hergenroeder, J. K. Ellis, C.-H. Lau, H. C. Keun, G. Schmitz, J. Schiller, M. Eibisch, C. Hedberg, H. Waldmann, E. Lausch, B. Tanner, J. Sehouli, J. Sagemueller, H. Staude, E. Steiner, J. G. Hengstler	Choline-releasing glycerophosphodiesterase EDI3 drives tumor cell migration and metastasis"	Proc. Natl. Acad. Sci.	2012	109	8155-8160
M. L. Sos, F. Dietlein, M. Peifer, J. Schöttle, H. Balke-Want, C. Müller, M. Koker, A. Richters, S. Heynck, F. Malchers, J. M. Heuckmann, D. Seidel, P. A. Eyers, R. T. Ullrich, A. P. Antonchick, V. V. Vintonyak, P. M. Schneider, T. Ninomiya, H. Waldmann, R. Büttner, D. Rauh, L. C. Heukamp, R. K. Thomas	A framework for identification of actionable cancer genome dependencies in small cell lung cancer	Proc. Natl. Acad. Sci.	2012	109	17034-17039
L.-G. Milroy, S. Rizzo, A. Calderon, B. Ellinger, S. Erdmann, J. Mondry, P. Vermeer,	Selective Chemical Imaging of Static Actin in Live Cells	J. Am. Chem. Soc.	2012	134	8480-8486



P. Bastiaens, H.
Waldmann, L.
Dehmelt, H.-D. Arndt

C. Deraeve, Z. Guo, R. S. Bon, W. Blankenfeldt, R. DiLucrezia, A. Wolf, S. Menninger, E. A. Stigter, S. Wetzel, A. Choidas, K. Alexandrov, H. Waldmann, R. S. Goody, Y.-W. Wu	Psoromic Acid is a Selective and Covalent Rab-Prenylation Inhibitor Targeting Autoinhibited RabGGTase	J. Am. Chem. Soc.	2012	134	7384-7391
K. Weise, S. Kapoor, A. Werkmüller, S. Möbitz, G. Zimmermann, G. Triola, Hr. Waldmann, R. Winter	Dissociation of the K Ras4B/PDE5 Complex upon Contact with Lipid Membranes: Membrane Delivery Instead of Extraction	J. Am. Chem. Soc.	2012	134	11503- 11510
L Yi, Y.-X. Chen, P-C. Lin, H. Schröder, C. M. Niemeyer, Y.-W Wu, R. S. Goody, G. Triola, H. Waldmann	Direct immobilization of oxyamine- modified proteins from cell lysates	ChemComm	2012	48	10829- 10831
A. Richter, R. Rose, C. Hedberg, H. Waldmann, C. Ottmann	An Optimised Small-molecule Stabilizer of the 14-3-3-PMA2 Protein-Protein Interaction	Chem – Eur. J.	2012	18	6520-6527
J. Xu, C. Hedberg, F. J. Dekker, Q. Li, K. M. Haigis, E. Hwang, H. Waldmann, K. Shannon	Inhibiting the pamitoylation/depamitoylation cycle selectively reduces the growth of hematopoietic cells expressing oncogenic <i>Nras</i>	Blood	2012	119	1032-1035
D. Willmann, SY Lim, S. Wetzel, E. Metzger, A. Jandausch, W. Wilk, M. Jung, I. Forne, A. Imhof, A. Janzer, J. Kirfel, H. Waldmann, R. Schüle, R. Buettner	Impairment of prostate cancer cell growth by a selective and reversible lysine-specific demethylase 1 inhibitor	Internat. J. of Cancer	2012	131	2704-2709
E. A. Stigter, Z. Guo, R. S. Bon, Y.-W. Wu, A. Choidas, A. Wolf, S. Menninger, H. Waldmann, W. Blankenfeldt, R. S. Goody	Development of Selective, Potent RabGGTase Inhibitors	J. Med.Chem.	2012	55	8330-8340



H. Lachance, S. Wetzel, K. Kumar, H. Waldmann	Charting, Navigating and Populating Natural Product Chemical Space for Drug Discovery	J. Med. Chem.	2012	55	5989-6001
K. Görmer, M. Bürger, J. A. W. Kruijtzter, I. Vetter, N. Vartak, L. Brunsveld, P. I. H. Bastiaens, R. Liskamp, G. Triola, H. Waldmann	Chemical-Biological Exploration of the Limits of the Ras De- and Repalmitoylating Machinery	ChemBioChem	2012	13	1017-1023
T. R. Samatov, A. Wolf, P. Odenwälder, S. Bessonov, C. Deraeve, R. S. Bon, H. Waldmann, R. Lührmann	Psoromic Acid Derivatives: A New Family of Small-molecule pre-mRNA splicing Inhibitors Discovered by a Stage-Specific High-Throughput in Vitro Splicing Assay	ChemBioChem	2012	13	640-644
D. Das, Z. Tnimov, U.T.T. Nguyen, G. Thimmaiah, H. Lo, D. Abankwa, Y. Wu, R. S. Goody, H. Waldmann, K. Alexandrov	Flexible and General Synthesis of Functionalized Phosphoisoprenoids for the Study of Prenylation in vivo and in vitro	ChemBioChem	2012	13	674-683
K. Matcha, A. V. R. Madduri, S. Roy, S. Ziegler, H. Waldmann, A. K. H. Hirsch, A. J. Minnaard	Total Synthesis of (-)-Doliculide, Structure-Activity Relationship Studies and its Binding to F-Actin	ChemBioChem	2012	13	2537-2548
G. Triola, H. Waldmann, C. Hedberg	Chemical Biology of Lipidated Proteins	ACS Chemical Biology	2012	8	87-99
Z. Tnimov, Z. Guo, Y. Gambin, U. T.T. Nguyen, Y.-W. Wu, D. Abankwa, A. Stigter, B. M. Collins, H. Waldmann, R. S. Goody, K. Alexandrov	Quantitative Analysis of Prenylated RhoA Interaction with Its Chaperone, RhoGDI	J. Biol. Chem.	2012	287	26549-26562
M. Bürger, T. J. Zimmermann, Y. Kondoh, P. Stege, N. Watanabe, H. Osada, H. Waldmann, I. R. Vetter	Crystal structure of the predicted phospholipase LYPLAL1 reveals unexpected functional plasticity despite close relationship to acyl protein thioesterases	J. of Lipid Research	2012	53	43-50
R. Toth, C. Gerding-Reimers, M. Deeks, S. Menninger, R.	Prieurianin/ endosidin 1 is an actin-stabilizing small molecule identified from a chemical genetic screen for circadian	The Plant Journal	2012	71	338-352



Martinez Gallegos, I. A. N. Tonaco, K. Hübel, P. J. Hussey, H. Waldmann, G. Coupland	clock effectors in <i>Arabidopsis thaliana</i>				
S. Kapoor, K. Weise, M. Erkkamp, G. Triola, H. Waldmann, R. Winter	The role of G-domain orientation and nucleotide state on the Ras isoform-specific membrane interaction	Eur. Biophys. J.	2012	41	801-813
S. Ziegler, H. Waldmann	Hunting the Targets of Natural Product-Inspired Compounds	M. Shibasaki et al (eds.), Chembiochemical Science: At the Frontier of Chemistry and Biology, Springer Verlag	2012		229-238
K. Alexandrov, Y.-W. Wu, W. Blankenfeldt, H. Waldmann, R. S. Goody	Organization and function of the Rab prenylation and recycling machinery	The Enzymes Volume 29: Protein Prenylation Part A Edited by Fuyuhiko Tamanoi, Christine A. Hrycyna and Martin O. Bergo	2012		147-162
J. Güldenaupt, T. Rudack, P. Bachler, D. Mann, G. Triola, H. Waldmann, C. Kötting, K. Gerwert	N-Ras Forms Dimers at POPC Membranes	Biophysical Journal	2012	103	1585-1593
G. Zimmermann, B. Papke, S. Ismail, N. Vartak, A. Chandra, M. Hoffmann, S. A. Hahn, G. Triola, A. Wittinghofer, P. I. H. Bastiaens, H. Waldmann	Small molecule inhibition of the Kras-PDE δ interaction impairs oncogenic KRAS signalling	Nature	2013	497	638-642
B. Over, S. Wetzel, C. Grütter, Y. Nakai, S. Renner, D. Rauh, H. Waldmann	Natural-product-derived fragments for fragment-based ligand discovery	Nat. Chem.	2013	5	21-28
T. Voigt, C. Gerding-Reimers, T. T. N. Tran, S. Bergmann, H. Lachance, B. Schölermann, A. Brockmeyer, P. Janning, S. Ziegler, H. Waldmann	A Natural Product Inspired Tetrahydropyran Collection Yields Mitosis Modulators that Synergistically Target CSE1L and Tubulin	Angew. Chem. Int. Ed.	2013	52	410-414
M. Gersch, F. Gut, V.	The Mechanism of Caseinolytic	Angew. Chem. Int.	2013	52	3009-3014



S. Korotkov, J. Lehmann, T. Böttcher, M. Rusch, C. Hedberg, H. Waldmann, G. Klebe, S. A. Sieber	Protease (ClpP) Inhibition	Ed.			
S. Ziegler, V. Pries, C. Hedberg, H. Waldmann	Review: Target Identification for Small Bioactive Molecules: Finding the Needle in the Haystack:	Angew. Chem. Int. Ed.	2013	52	2744-2792
P.-Y. Dakas, J. A. Parga, S. Höing, H. R. Schöler, J. Sterneckert, K. Kumar, H. Waldmann	Discovery of Neuritogenic Compound Classes Inspired by Natural Products	Angew. Chem. Angew. Chem. Int. Ed.	2013	36 52	9755-9760 9576-9581
H. Takayama, Z.-J. Jia, Lea Kremer, J. O. Bauer, C. Strohmann, S. Ziegler, A. P. Antonchick, H. Waldmann	Discovery of Inhibitors of the Wnt and Hedgehog Signaling Pathways through the Catalytic Enantioselective Synthesis of an Iridoid-Inspired Compound Collection	Angew. Chem. Int. Ed.	2013	52	12404- 12408
R. Narayan, J. O. Bauer, C. Strohmann, A. P. Antonchick, H. Waldmann	Catalytic Enantioselective Synthesis of Functionalized Tropanes Reveals Novel Inhibitors of Hedgehog Signaling	Angew. Chem. Int. Ed.	2013	52	12892- 12896
S. Kapoor, A. Werkmüller, R. S. Goody, H. Waldmann, R. Winter	Pressure modulation of Ras-membrane interactions and intervesicle transfer	J. Am. Chem. Soc.	2013		6149-6156
M. Potowski, A. P. Antonchick, H. Waldmann	Catalytic asymmetric exo-selective [6+3] cycloaddition of iminoesters with fulvenes	Chem Comm.	2013	49	7800-7802
A. P. Antonchick, S. López-Tosco, J. Parga, S. Sievers, M. Schürmann, H. Preut, S. Höing, H. R. Schöler, J. Sterneckert, D. Rauh, H. Waldmann	Highly Enantioselective Catalytic Synthesis of Neurite Growth-Promoting Secoyohimbanes	Chemistry & Biology	2013	20	500-509
V. Eschenbrenner- Lux, H. Dücker, V. Khedkar, H. Bruss, H. Waldmann, K. Kumar	Cascade Syntheses Routes to the Centrocountins	Chem. – Eur. J.	2013	19	2294-2304
T. J. Zimmermann, M. Bürger, E. Tashiro, Y. Kondoh, N. E. Martinez, K. Görmer,	Boron-Based Inhibitors of Acyl Protein Thioesterases 1 and 2	ChemBioChem	2013	14	115-122



S. Rosin-Steiner, T.
Shimizu, S. Ozaki, N.
Watanabe, D. Hall, I.
R. Vetter, H. Osada,
C. Hedberg, H.
Waldmann

T. J. Zimmermann, S.
Roy, N. E. Martinez,
S. Ziegler, C.
Hedberg, H.
Waldmann

K. Weise, D. Huster,
S. Kapoor, G. Triola,
H. Waldmann, R.
Winter

A. Werkmüller, G.
Triola, H. Waldmann,
R. Winter

M. Köhnke, C. Delon,
M. L. Hastie, U. T.T.
Nguyen; Y.-W. Wu, H.
Waldmann, R. S.
Goody, J. J. Gorman,
K. Alexandrov

J. Spiegel, P. M.
Cromm, G.
Zimmermann, T. N.
Grossmann, H.
Waldmann

J. Spiegel, P. M.
Cromm, A. Itzen, R. S.
Goody, T. N.
Großmann, H.
Waldmann

V. Eschenbrenner-
Lux, P. Küchler, S.
Ziegler, K. Kumar, H.
Waldmann

F. Kolundžić, A.
Murali, P. Pérez-
Galan, J. O. Bauer, C.
Strohmann, K. Kumar,
H. Waldmann

V. Eschenbrenner-
Lux, K. Kumar, H.
Waldmann

Biology-Oriented Synthesis of a
Tetrahydroisoquinoline-Based
Compound Collection Targeting
Microtubule Polymerization

Gibbs energy determinants of lipoprotein
insertion into lipid membranes: the case
study of Ras proteins

Rotational and Translational Dynamics of
Ras Proteins upon Binding to Model
Membrane Systems

Rab GTPase Prenylation Hierarchy and
Its Potential Role in Choroideremia
Disease. PLoS ONE 8(12): e81758.
doi:10.1371/journal.pone.0081758

Small molecule modulation of Ras
signaling

Direct Targeting of Rab-GTPase-Effector
Interactions

An Enantioselective Inverse-Electron-
Demand Imino-Diels-Alder Reaction

A Cyclization-Rearrangement Cascade
for the Synthesis of Structurally Complex
Chiral Gold(I)-Aminocarbene Complexes

The Asymmetric Hetero- Diels-Alder
Reaction in the Syntheses of Biologically
Relevant Compounds

ChemBioChem

Faraday Discuss.

ChemPhysChem

PLoS One

Nat. Chem. Biol.

Angew. Chem. Int.
Ed.

Angew. Chem Int. Ed.

Angew. Chem.

Angew. Chem. Int.
Ed.

Angew. Chem. Int.
Ed./
Angew. Chemie

2013

2013

2013

2013

2014

2014

2014

2014

2014

14

161

14

12

10

53

53

53

53/
126

295-300

549-561

3698-3705

doi:10.1371/
journal.pone
.0081758

613-622

2498-2503

2134-2137

2166-2169

8122-8126

11146-
11157
11326-
11337



G. Lukinavičius, L. Reymond, E. D'Este, A. Masharina, F. Göttfert, H. Ta, A. Güther, M. Fournier, S. Rizzo, H. Waldmann, C. Blaukopf, C. Sommer, D. W. Gerlich, H.-D. Arndt, S. W. Hell, K. Johnsson	Fluorogenic probes for live-cell imaging of the cytoskeleton	Nature Methods	2014	11	731-737
H.van Hattum, H. Waldmann	Biology-Oriented Synthesis: Harnessing the Power of Evolution	J. Am. Chem. Soc.	2014	136	11853-11859
G. Zimmermann, C. Schultz-Fademrecht, P. Kuchler, S. Murarka, S. Ismail, G. Triola, P. Nussbaumer, A. Wittinghofer, H. Waldmann	Structure Guided Design and Kinetic Analysis of Highly Potent Benzimidazole Inhibitors Targeting the PDE δ Prenyl Binding Site	J. Med. Chem.	2014	57	5435-5448
R. Narayan, M. Potowski, Z.-J. Jia, A. P. Antonchick, H. Waldmann	Catalytic Enantioselective 1,3-Dipolar Cycloadditions of Azomethine Ylides for Biology Oriented Synthesis	Acc. Chem. Res.	2014	47	1296-1310
S. Rizzo, H. Waldmann	Development of a Natural Product-Derived Chemical Toolbox for Modulation of Protein Function	Chem. Rev.	2014	114	4621-4639
N. Vartak, B. Papke, H. E. Grecco, L. Rossmannek, H. Waldmann, C. Hedberg, P. I. H. Bastiaens	The Autodepalmitoylating Activity of APT Maintains the Spatial Organization of Palmitoylated Membrane Proteins	Biophys. J.	2014	106	93-105
H.van Hattum, H. Waldmann	Chemical Biology Tools for Regulating RAS Signaling Complexity in Space and Time	Chemistry & Biology	2014	21	1185-1195
T. T.N. Tran, C. Gerding-Reimers, B. Schölermann, B. Stanitzki, T. Henkel, H. Waldmann, S. Ziegler	Podoverine A – a novel microtubule destabilizing natural product from the <i>Podophyllum</i> species	Bioorg. Med. Chem.	2014	22	5110-5116
S. Basu, H. Waldmann	Polymer supported synthesis of a natural product-inspired oxepane library	Bioorg. Med. Chem.	2014	22	4430-4444
H. Bruss, H. Schuster, R. Martinez, M. Kaiser, A. P.	Synthesis of the B-seco limonoid core scaffold	Beilstein J. Org. Chem.	2014	10	194-208



Antonchick, H.
Waldmann

V. Berg, M. Rusch, N. Vartak, A. Schauss, H. Waldmann, C. Hedberg, M. Hallek, C. P. Pallasch, P.I.H. Bastiaens, C.-M. Wendtner, L.P. Frenzel	miRs-138/ -424 control CD95-mediated cell death by targeting acyl protein thioesterases 1 and 2	Oncology Res. and Treatm.	2014	37	237
C. Klaus, U. Schneider, C. Hedberg, A.K. Schütz, J. Bernhagen, H. Waldmann, N. Gassler, E. Kaemmerer	Modulating effects of Acyl-CoA synthetase 5-derived mitochondrial Wnt2B palmitoylation on intestinal Wnt activity	World Journal of Gastroenterology	2014	20	14855-14864
S. Rizzo, V. Wakchaure, H. Waldmann	Natural Product-Derived and Natural Product-Inspired Compound Collections	Book chapter, S. Hanessian, ed. "Natural Products in Medicinal Chemistry, Wiley VCH	2014	60	43-80
Eschenbrenner-Lux, V., H. Waldmann, K. Kumar	Domino Reactions in Library Synthesis	Book Chapter, "Domino Reactions", Concepts for Efficient Organic Synthesis, ed. by Lutz Tietze, Wiley-VCH, 2014	2014		497-521
A. Vogel, J. Nikolaus, K. Weise, G. Triola, H. Waldmann, R. Winter, A. Herrmann, D. Huster	Interaction of the human N-Ras protein with lipid raft model membranes of varying degrees of complexity	Biol. Chem.	2014	395	779-789
S. Rao Vidadala, C. Golz, C. Strohmann, C.-G. Daniliuc, H. Waldmann	Highly Enantioselective Intramolecular 1,3-Dipolar Cycloaddition: A Route to Piperidino-Pyrrolizidines	Angew. Chem. Int. Ed.	2015	54	651-655
Y. Akbulut, H. J. Gaunt, K. Muraki, M. J. Ludlow, M. S. Amer, A. Bruns, N. S. Vasudev, L. Radtke, M. Willot, S. Hahn, T. Seitz, S. Ziegler, M. Christmann, D. J. Beech, H. Waldmann	(-)-Englerin is a Potent and Selective Activator of TRPC4 and TRPC5 Calcium Channels	Angew. Chem. Int. Ed.	2015	54	3787-3791
J. Švenda, M. Sheremet, L. Kremer,	Biology-Oriented Synthesis of a Withanolide-Inspired Compound	Angew. Chem. Int. Ed.	2015	54	5596-5602



L. Maier, J. O. Bauer, C. Strohmann, S. Ziegler, K. Kumar, H. Waldmann	Collection Reveals Novel Modulators of Hedgehog Signaling				
S. Murarka, Z.-J. Jia, C. Merten, C.-G. Daniliuc, A. P. Antonchick, H. Waldmann	Rhodium(II)-Catalyzed Enantioselective Synthesis of Troponoids	Angew. Chem. Int. Ed.	2015	54	7653-7656
P. Schröder, F. Förster, S. Kleine, C. Becker, A. Richters, S. Ziegler, D. Rauh, K. Kumar, H. Waldmann	Neuritogenic Militarone-Inspired 4- Hydroxypyridones Target the Stress Pathway Kinase Map4K4	Angew. Chem. Int. Ed.	2015	54	12398- 12403
P. M. Cromm, J. Spiegel, T. N. Großmann, H. Waldmann	Review: Direct Modulation of Small GTPase Activity and Function	Angew. Chem. Int. Ed.	2015	54/ 127	13516- 13537/ 13718- 13741
A. Pahl, M. Lakemeyer, M.-T. Vielberg, M. W. Hackl, J. Vomacka, V. S. Korotkov, M. L. Stein, C. Fetzner, K. Lorenz-Baath, K. Richter, H. Waldmann, M. Groll, S. A. Sieber	Reversible Inhibitors Arrest ClpP in a Defined Conformational State that Can Be Revoked by ClpX Association	Angew. Chem. Int. Ed.	2015	54	15892- 15896
M. W. Hackl, M. Lakemeyer, M. Dahmen, M. Glaser, A. Pahl, K. Lorenz- Baath, T. Menzel, S. Sievers, T. Boettcher, I. Antes, H. Waldmann, S. A. Sieber	Phenyl Esters are Potent Inhibitors of Caseinolytic Protease P and Reveal a Stereogenic Switch for Deoligomerization	J. Am. Chem. Soc.	2015	137	8475-8483
A. Danda, K. Kumar, H. Waldmann	A general catalytic reaction sequence to access alkaloid-inspired indole polycycles	ChemComm	2015	51	7536-7539
Z.-J. Jia, C. G. Daniliuc, A. P. Antonchick, H. Waldmann	Phosphine-catalyzed dearomatizing [3+2] annulations of isoquinolinium methylides with allenes	ChemComm	2015	51	1054-1057
M. Potowski, C. Merten, A. P. Antonchick, H. Waldmann	Catalytic Aerobic Oxidation and Tandem Enantioselective Cycloaddition in Cascade Multicomponent Synthesis	ChemEurJ	2015	21	4913-4917



H.-D. Arndt, S. Rizzo, C. Nöcker, V. N. Wakchaure, L.-G. Milroy, V. Bieker, A. Calderon, T. T. N. Tran, S. Brand, L. Dehmelt, H. Waldmann	Divergent solid-phase synthesis of natural product-inspired bipartite cyclodepsipeptides - Total synthesis of Seragamide A	Chem. – Eur. J.	2015	21	5311-5316
R. Meiß, K. Kumar, H. Waldmann	Divergent Gold()-Catalyzed Skeletal Rearrangements of 1,7-Enynes	Chem. – Eur. J.	2015	21	13526- 13530
C. R. Bertozzi, B. R. Stockwell, S. Kubicek, B. C. Dickinson, C. J. Chang, C. Schultz, P. A. Silver, J. E. Gestwicki, G. Chiosis, A. Chattopadhyay, R. A. Butcher, S. B. Park, B. K. Shoichet, S. L. Flitsch, J. Zhang, D. R. Liu, Y. Onishi, E. Weerapana, A. H. Williams, C. He, A. Moroni, G. Thiel, Y.-T. Chang, H. Waldmann, M. Bogyo, L. B. Oddershede, A. Christopoulos, B. Imperiali, H. Ehrlich, X. Chen, J. A. Prescher	Voices of chemical biology	Nat. Chem. Biol.	2015	11	378-379
V. Berg, M. Rusch, N. Vartak, C. Jüngst, A. Schauss, H. Waldmann, C. Hedberg, C. P. Pallasch, P. I. H. Bastiaens, M. Hallek, C.-M. Wendtner, L. P. Frenzel	miRs-138 and -424 control palmitoylation-dependent CD95- mediated cell death by targeting acyl protein thioesterases 1 and 2 in CLL	Blood	2015	125	2948-2957
G. Vendrell-Navarro, F. Rúa, J. Bujons, A. Brockmeyer, P. Janning, S. Ziegler, A. Messeguer, H. Waldmann	Positional Scanning Synthesis of a Peptoid Library Yields New Inducers of Apoptosis that Target Karyopherins and Tubulin	ChemBioChem	2015	16	1580-1587
T. Mejuch, H. v. Hattum, G. Triola, M. Jaiswal, H. Waldmann	Specificity of Lipoprotein Chaperones for the Characteristic Lipidated Structural Motifs of their Cognate Lipoproteins	ChemBioChem	2015	16	2460-2465
A. Ursu, H. Waldmann	Hide and seek: identification and	Bioorg. Med. Chem.	2015	25	3079-3086



	confirmation of small molecule protein targets	Letts.			
M. Potowski, C. Golz, C. Strohmann, A. P. Antonchick, H. Waldmann	Biology-oriented synthesis of benzopyrano[3,4-c]pyrrolidines	Bioorg. & Med. Chem.	2015	23	2895-2903
M. G. Sankar, L. Mantilli, J. Bull, F. Giordanetto, J. O. Bauer, C. Strohmann, H. Waldmann, K. Kumar	Stereoselective synthesis of a natural product inspired tetrahydroindolo[2,3-a]quinolizine compound library	Bioorg. & Med. Chem.	2015	23	2614-2620
S. Rao, H. Waldmann	One-pot synthesis of a natural product inspired pyrrolocoumarine compound collection by means of an intramolecular 1,3-dipolar cycloaddition as key step	Tetrah.Lett.	2015	56	3358-3360
G. Vendrell-Navarro, A. Brockmeyer, H. Waldmann, P. Janning, S. Ziegler	Identification of the targets of biologically active small molecules using quantitative proteomics	Chemical Biology: Methods and Protocols (ed. by J. E. Hempel, C. H. Williams, C. C. Hong)	2015		263-286
J. Gilleron, R. Paramasivam, A. Zeigerer, W. Querbes, G. Marsico, C. Andree, S. Seifert, P. Amaya, M. Stöter, V. Koteliansky, H. Waldmann, K. Fitzgerald, Y. Kalaidzidis, A. Akinc, M. A. Maier, M. Manoharan, M. Bickle, M. Zerial	Identification of siRNA delivery enhancers by a chemical library screen	Nucleic Acids Res.	2015	43	7984-8001
H. Xu, C. Golz, C. Strohmann, A. P. Antonchick, H. Waldmann	Enantiodivergent Combination of Natural Product Scaffolds Enabled by Catalytic Enantioselective Cycloaddition	Angew. Chem. Int. Ed.	2016	55	7761-7765
Y. Qian, M. Schürmann, P. Janning, C. Hedberg, H. Waldmann	Activity-Based Proteome Profiling Probes on Woodward's Reagent K with Distinct Target Selectivity	Angew. Chem. Int. Ed.	2016	55	7766-7771
L. Kötzner, M. Leutzsch, S. Sievers, S. Patil, H. Waldmann, Y. Zheng, W. Thiel, B. List	The Organocatalytic Approach to Enantiopure 2 <i>H</i> - and 3 <i>H</i> -Pyrroles: Inhibitors of the Hedgehog Signaling Pathway	Angew. Chem. Int. Ed.	2016	55	7693-7697
B.Papke, S. Murarka,	Identification of Pyrazolopyridazinones	Nature	2016		7:11360



H. A. Vogel, P. Martin-Gago, M. Kovacevic, D. C. Truxius, E. K. Fansa, S. Ismail, G. Zimmermann, K. Heinelt, C. Schultz-Fademrecht, A. Al Saabi, M. Baumann, P. Nussbaumer, A. Wittinghofer, H. Waldmann, P. I. H. Bastiaens	as PDE δ Inhibitors	Communications			
P. M. Cromm, S. Schaubach, J. Spiegel, A. Fürstner, T. N. Großmann, H. Waldmann	Orthogonal ring-closing alkyne and olefin metathesis for the synthesis of small GTPase-targeting bicyclic peptides	Nature Communications	2016	7	11300
A. Ursu, D. J. Illich, Y. Takemoto, A. T. Porfetye, M. Zhang, A. Brockmeyer, P. Janning, N. Watanabe, H. Osada, I. R. Vetter, S. Ziegler, H. R. Schöler, H. Waldmann	Epiblastin A Induces Reprogramming of Epiblast Stem Cells Into Embryonic Stem Cells by Inhibition of Casein Kinase 1	Cell Chemical Biology	2016	23	494-507
D. J. Illich, M. Zhang, A. Ursu, R. Osorno, K.-P. Kim, J. Yoon, M. J. Arauzo-Bravo, G. Wu, D. Esch, D. Sabour, D. Colby, K. S. Grassme, J. Chen, B. Greber, S. Höing, W. Herzog, S. Ziegler, I. Chambers, S. Gao, H. Waldmann, H.R. Schöler	Distinct Signaling Requirements for the Establishment of ESC Pluripotency in Late-Stage EpiSCs	Cell Reports	2016	15	787-800
M. Schürmann, P. Janning, S. Ziegler, H. Waldmann	Small-Molecule Target Engagement in Cells	Cell Chemical Biology	2016	23	435-441
K. Tschapalda, Y.-Q. Zhang, L. Liu, K. Golovnina, T. Schlemper, T. O. Eichmann, M. Lal-Nag, U. Sreenivasan, J. McLenithan, S. Ziegler, C. Sztalryd, A. Lass, D. Auld, B. Oliver, H. Waldmann,	A Class of Diacylglycerol Acyltransferase 1 Inhibitors Identified by a Combination of Phenotypic High-throughput Screening, Genomics, and Genetics	EBioMedicine	2016	8	49-59



Z. Li, M. Shen, M. B.
Boxer, M. Beller

M. Sellstedt, M. Schwalfenberg, S. Ziegler, A. P. Antonchick, H. Waldmann	Trienamine catalyzed asymmetric synthesis and biological investigation of a cytochalasin B-inspired compound collection	Org. & Biomol. Chem.	2016	14	50-54
P. Pérez-Gálan, H. Waldmann, K. Kumar	Building polycyclic indole scaffolds via gold(I)-catalyzed intra- and inter-molecular cyclization reactions of 1,6-enynes	Tetrahedron special issue Dr. Garg	2016	72	3647-3652
B. Sperlich, S. Kapoor, H. Waldmann, R. Winter, K. Weise	Regulation of K Ras4B Membrane Binding by Calmodulin	Biophysical Journal	2016	111	113-122
N. Erwin, B. Sperlich, G. Garivet, H. Waldmann, K. Weise, R. Winter	Lipoprotein Insertion into Membranes of Various Complexity: Lipid Sorting, Interfacial Adsorption and Protein Clustering	Phys. Chem. Chem. Phys.	2016	18	8954-8962
T. Mejuch, H. Waldmann	Synthesis of Lipidated Proteins (Review)	Bioconj. Chem.	2016	27	1771-1783
P. Schröder, J. O. Bauer, C. Strohmann, K. Kumar, H. Waldmann	Synthesis of an Iridoid-Inspired Compound Collection and Discovery of Autophagy Inhibitors	JOC	2016	81	10242-10255
M. Jaiswal, E. K. Fansa, S. K. Kösling, T. Mejuch, H. Waldmann, A. Wittinghofer	Novel Biochemical and Structural Insights into the Interaction of Myristoylated Cargo with Unc119 Protein and Their Release by Arl2/3	J. of Biol. Chem.	2016	291	20766-20778
P. M. Cromm, J. Spiegel, P. Kuchler, L. Dietrich, J. Kriegesmann, M. Wendt, R. Goody, H. Waldmann, T.N. Grossmann	Protease-resistant and cell-permeable double-stapled peptides targeting the Rab8a GTPase	ACS Chemical Biology	2016	11	2375-2382
P.-Y. Dakas, H. Waldmann, K. Kumar	Natural Product Inspired Enantioselective Synthesis of Hexahydro-Aza-Pentalenones	Heterocycles	2016	Vol. 93	465-473
E. Valeur, S. M. Guéret, H. Adihou, R. Gopalakrishnan, M. Lemurell, H. Waldmann, Tom N. Großmann, A. T. Plowright	Review: New Modalities for Challenging Targets in Drug Discovery	Angew. Chem. Int. Ed/ Angew. Chem.	2017	56/12 9	10294-10323/ 10428-10459



L. Laraia, K. Ohsawa, G. Konstantinidis, L. Robke, Y. Wu, K. Kumar, H. Waldmann	Discovery of Novel Cinchona-Alkaloid-Inspired Oxazatwistane Autophagy Inhibitors	Angew. Chem. Int. Ed.	2017	56	2145-2150
P. Martín-Gago, E.K. Fansa, C. H. Klein, S. Murarka, P. Janning, M. Schürmann, M. Metz, S. Ismail, C. Schultz-Fademrecht, M. Baumann, P.I.H. Bastiaens, A. Wittinghofer, H. Waldmann	A PDE6 δ -KRas Inhibitor Chemotype with up to Seven H-Bonds and Picomolar Affinity that Prevents Efficient Inhibitor Release by Arl2	Angew. Chem. Int. Ed.	2017	56	2423-2428
Z.-J. Jia, C. Merten, R. Gontla, C. G. Daniliuc, A. P. Antonchick, H. Waldmann	General Enantioselective C–H Activation Through Efficiently Tunable Cyclopentadienyl Ligands	Angew. Chem. Int. Ed.	2017	56	2429-2434
T. Mejuch, G. Garivet, W. Hofer, N. Kaiser, E. K. Fansa, C. Ehrt, O. Koch, M. Baumann, S. Ziegler, A. Wittinghofer, H. Waldmann	Small Molecule Inhibition of the UNC119-Cargo Interaction	Angew. Chem. Int. Ed.	2017	56	6181-6186
L. Robke, L. Laraia, M. A. Carnero Corrales, G. Konstantinidis, M. Muroi, A. Richters, M. Winzker, T. Engbring, S. Tomassi, N. Watanabe, H. Osada, D. Rauh, H. Waldmann, Y.-W. Wu, J. Engel	Phenotypic Identification of a Novel Autophagy Inhibitor Chemotype Targeting Lipid Kinase VPS34	Angew. Chem. Int. Ed.	2017	56	8153-8157
M. Dwivedi; T. Mejuch; H. Waldmann, R. Winter	Lateral Organization of Host Heterogenous Raft-like Membranes Altered by the Myristoyl Modification of Tyrosine Kinase c-Src	Angew. Chem. Int. Ed.	2017	56	10511-10515
H. Xu, L. Laraia, L. Schneider, K. Louven, C. Strohmam, A. P. Antonchick, H. Waldmann	Highly Enantioselective Catalytic Vinylogous Propargylation of Coumarins Yields a Novel Autophagy Inhibitor Class	Angew. Chem. Int. Ed. Angew. Chem.	2017	56 129	11232-11236 11384-11388
L. Kremer, C. Schultz-Fademrecht, M. Baumann, P.	Discovery of a Novel Hedgehog Signaling Pathway through Cell-based Compound Discovery and Target	Angew. Chem. Int. Ed.	2017	56	13021-13025



Habenberger, A. Choidas, B. Klebl, S. Kordes, H. R. Schöler, J. Sternecker, S. Ziegler, G. Schneider, H. Waldmann	Prediction				
Y.-C. Lee, S. Patil, C. Golz, C. Strohmann, S. Ziegler, K. Kumar, H. Waldmann	A ligand-directed divergent catalytic approach to establish structural and functional scaffold diversity	Nature Commun.	2017	8	DOI:10.1038/ncomms14043
A. Ursu, H. Schöler, H. Waldmann	Small-molecule phenotypic screening with stem cells	Nat. Chem. Biol.	2017	13	560-563
P. Martín-Gago, E.K. Fansa, M. Winzker, S. Murarka, P. Janning, C. Schuttz- Fademrecht, M.. Baumann, A. Wittinghofer, H. Waldmann	Covalent Protein Labeling at Glutamic Acids	Cell Chem. Biol.	2017	24/5	589-597e5
A. T. Plowright, C. Ottmann, M. Arkin, Y. P. Auberson, H. Timmerman, H. Waldmann	Joining Forces: The Chemical Biology-Medicinal Chemistry Continuum	Cell Chem. Biol.	2017	24	1058-1065
A. Sidarovich, C. L. Will, M. A. Anokhina, J. Ceballos, S. Sievers, D. E. Agafonov, T. Samatov, P. Bao, B. Kastner, H. Urlaub, H. Waldmann, R. Lührmann	Identification of a small molecule inhibitor that stalls splicing at an early step of spliceosome activation	eLife	2017	6	E23533
S. Mosalaganti, J. Keller, A. Altenfeld, M. Winzker, P. Rombaut, M. Saur, A. Petrovic, A. Wehenkel, S. Wohlgemuth, F. Müller, S. Maffini, T. Bange, F. Herzog, H. Waldmann, S. Raunser, A. Musacchio	Structure of the RZZ complex and molecular basis of its interaction with Spindly	J. Cell Biol.	2017	216	961-981
M. Dwivedi, T. Mejuch, H. Waldmann, R. Winter	The Myristoyl Modification of C-SRC Alters the Lateral Organization of Host Heterogenous Raft-Like Membranes	Bioph. J.	2017	112 (3)	326a



H. N. Rubaiy, M. J. Ludlow, M. Henrot, H. J. Gaunt, K. Miteva, S. Y. Cheung, Y. Tanahashi, N. Hamzah, K. E. Musialowski, N. M. Blythe, H. L. Appleby, M. A. Bailey, L. McKeown, R. Taylor, R. Foster, H. Waldmann, P. Nussbaumer, M. Christmann, R. S. Bon, K. Muraki, D. J. Beech	Picomolar, selective and subtype-specific small-molecule inhibition of TRPC1/4/5 channels	J. of Biol. Chem.	2017	292 (20)	8158-8173
S. Murarka, P. Martín-Gago, C. Schultz-Fademrecht, A. Al Saabi, M. Baumann, E. K. Fansa, S. Ismail, P. Nussbaumer, A. Wittinghofer, H. Waldmann	Development of Pyridazinone Chemotypes Targeting the PDE δ Prenyl Binding Site	Chem. Eur. J.	2017	23/25	6083-6093
N. E. Martinez, T.J. Zimmermann, C. Goosmann, T. Alexander, C. Hedberg, S. Ziegler, A. Zychlinsky, H. Waldmann	Tetrahydroisoquinolines: New Inhibitors of Neutrophil Extracellular Trap (NET) Formation	ChemBioChem	2017	18 (10)	888-893
T. Förster, S. López-Tosco, S. Ziegler, A. P. Antonchick, H. Waldmann	Enantioselective Organocatalytic Synthesis of a Secoyohimbane-Inspired Compound Collection with Neuritogenic Activity	ChemBioChem	2017	18	1098-1108
M. Sheremet, S. Kapoor, P. Schröder, K. Kumar, S. Ziegler, H. Waldmann	Small Molecules Inspired by the Natural Product Withanolides as Potent Inhibitors of Wnt Signaling	ChemBioChem	2017	18	1797-1806
L. Laraia, H. Waldmann	Natural Product Inspired Compound Collections: Evolutionary Principle, Chemical Synthesis, Phenotypic Screening, and Target Identification	DDT: Technologies	2017	23	75-82
P. Martín-Gago, E. K. Fansa, A. Wittinghofer, H. Waldmann	Structure-based development of PDE δ inhibitors	Biol. Chem.	2017	398 (5-6)	535-545
A. Pahl, H. Waldmann, K. Kumar	Exploring natural product fragments for drug and probe discovery	Chimia Int. J. Chem.	2017	71	653-660



K. Muraki, K. Ohnishi, A. Takezawa, H. Suzuki, N. Hatano, Y. Muraki, N. Hamzah, R. Foster, H. Waldmann, P. Nussbaumer, M. Christmann, R. S. Bon, D. J. Beech	Na ⁺ entry through heteromeric TRPC4/C1 channels mediates (-) Englerin A-induced cytotoxicity in synovial sarcoma cells	Nature – Scientific Reports	2017	7	DOI:10.1038/s41598-017-17303-3
Z.-J. Jia, C. Merten, S. Murarka, L. Knauer, C. Strohmann, H. Waldmann	Biology-Oriented Synthesis of Decahydro-4,8-epoxyazulene Scaffolds	Synlett (special issue, 80 th birthday V. Snieckus)	2017	28	2918-2922
S. Murarka, C. Golz, C. Strohmann, A. P. Antonchick, H. Waldmann	Biology-Oriented Synthesis of 3,3-Spiro (2-tetrahydrofuran)-oxindoles	Synthesis	2017	49	87-95
G. Karageorgis, E. S. Reckzeh, J. Ceballos, M. Schwalfenberg, S. Sievers, C. Ostermann, A. Pahl, S. Ziegler, H. Waldmann	Chromopyrones are pseudo natural product glucose uptake inhibitors targeting glucose transporters GLUT-1 and -3	Nat. Chem.	2018	10	1103-1111
Y.-C. Lee, K. Kumar, H. Waldmann	Minireview: Ligand-Directed Divergent Synthesis of Carbo- and Heterocyclic Ring Systems	Angew. Chem. Int. Ed.	2018	57/130	5212-5226/5308-5322
G. Shan, J. Flegel, H. Li, C. Merten, S. Ziegler, A. P. Antonchick, H. Waldmann	C-H Bond Activation for the Synthesis of Heterocyclic Atropisomers Yields Hedgehog Pathway Inhibitors	Angew. Chem. Int. Ed.	2018	57	14250-14254
Z.-J. Jia, G. Shan, C. G. Daniliuc, A. P. Antonchick, H. Waldmann	Enantioselective Synthesis of the Spirotranyloxindole Scaffold through Bimetallic Relay Catalysis	Angew. Chem. Int. Ed.	2018	57	14493-14497
L. Robke, Y. Futamura, G. Konstantinidis, J. Wilke, H. Aono, Z. Mahmoud, N. Watanabe, Y.-W. Wu, H. Osada, L. Laraia, H. Waldmann	Discovery of the novel autophagy inhibitor Aumitin that targets mitochondrial complex I	Chem. Sci.	2018	9	3014 - 3022
V. Pries, C. Nöcker, D. Khan, P. Johnen, Z. Hong, A. Tripathi,	Target Identification and Mechanism of Action of Picolinamide and Benzamide Chemotypes with Antifungal Properties	Cell Chem. Biol.	2018	25	279-290



A.-L. Keller, M. Fitz, F.
Perruccio, I. Filipuzzi,
S. Thavam, T. Aust,
R. Riedl, S. Ziegler, F.
Bono, G. Schaaf, V.
A. Bankaitis, H.
Waldmann, D.
Hoepfner

S. Höing, T.-Y. Yeh, M. Baumann, N.E. Martinez, P. Habenberger, L. Kremer, H. C.A. Drexler, P. Kuchler, P. Reinhardt, A. Choidas, M.-L. Zischinsky, G. Zischinsky, S. Nandini, A. P. Ledray, S. A. Ketcham, L. Reinhardt, M. Abo-Rady, M. Glatza, S. J. King, P. Nussbaumer, S. Ziegler, B. Klebl, T. A. Schroer, H. R. Schöler, H. Waldmann, J. Sternecker	Dynarrestin, a Novel Inhibitor of Cytoplasmic Dynein	Cell Chem. Biol.	2018	25	357-369
S. Brand, S. Roy, P. Schröder, B. Rathmer, J. Roos, S. Kapoor, S. Patil, C. Pommerenke, T. Maier, P. Janning, S. Eberth, D. Steinhilber, D. Schade, G. Schneider, K. Kumar, S. Ziegler, H. Waldmann	Combined Proteomic and <i>In Silico</i> Target Identification Reveal a Role for 5-Lipoxygenase in Developmental Signaling Pathways	Cell Chem. Biol	2018	25	1095-1106
L. Laraia, L. Robke, H. Waldmann	Review: Bioactive compound collections: from design to target identification	Chem	2018	4	705-730
N. Erwin, M. Dwivedi, T. Mejuch, H. Waldmann, R. Winter	UNC119A Decreases the Membrane Binding of Myristoylated c-Src	ChemBioChem	2018	19	1482-1487
Z.-J. Jia, H. Takayama, Y. Futamura, H. Aono, J. O. Bauer, C. Strohmann, A. P. Antonchick, H. Osada, H. Waldmann	Catalytic Enantioselective Synthesis of a Pyrrolizidine-Alkaloid-Inspired Compound Collection with Antiplasmodial Activity	J. Org. Chem.	2018	83	7033-7041



S. Y. Cheung, M. Henrot, M. A-Saad, M. Baumann, H. Muller, A. Unger, H. N. Rubaiy, I. Mathar, K. Dinkel, P. Nussbaumer, B. Klebl, M. Freichel, B. Rode, S. Trainor, S. J. Clapcote, M. Christmann, H. Waldmann, S. K. Abbas, D. J. Beech, N.S. Vasudev	TRPC4/TRPC5 channels mediate adverse reaction to the cancer cell cytotoxic agent (-)-Englerin A	Oncotarget	2018	9	29634- 29643
Y.-C. Lee, J.L. Knauer, K. Louven, C. Golz, C. Strohmann, H. Waldmann, K. Kumar	Gold(I)-Catalyzed and Nucleophile- Guided Ligand-directed Divergent Synthesis	Eur. J. Org. Chem..	2018		5688-5699
L. Robke, T. Rodriguez, P. Schröder, D. J. Foley, G. J. L. Bernardes, L. Laraia, H. Waldmann	Discovery of 2,4-dimethoxypyridines as novel autophagy inhibitors	Tetrahedron	2018	74	4531-4537
J. Wilke, T. Kawamura, N. Watanabe, H. Osada, S. Ziegler, H. Waldmann	Identification of cytotoxic, glutathione- reactive moieties inducing accumulation of reactive oxygen species via glutathione depletion	Bioorg. Med. Chem.	2018	26	1453-1461
P. Kuchler, G. Zimmermann, M. Winzker, J. Janning, H. Waldmann, S. Ziegler	Identification of novel PDE δ interacting proteins	Bioorg. Med. Chem	2018	26	1426-1434
K. Kumar, H. Waldmann	Nature Inspired Small Molecules for Chemical Biology	Israel Journal of Chemistry Doi:10.1002/ijch.201800 105	2018		
H. N. Rubaiy, T. Seitz, S. Hahn, A. Choidas, P. Habenberger, B. Klebl, K. Dinkel, P. Nussbaumer, H. Waldmann, M. Christmann	Identification of an (-)-englerin A analogue, which antagonizes (-)-englerin A at TRPC1/4/5 channels	B. J. Pharm.	2018	175	830-839
G. Karageorgis, H. Waldmann	Biology-Oriented Synthesis	Book Chapter in: "Chemical and Biological Synthesis:	2018		45-73



Enabling Approaches
for Understanding
Biology, ed. by Nick J.
Westwood and Adam
Nelson

L. Laraia, H. Waldmann	Synthesis and target identification of natural product inspired compound collections	Book chapter "Chemical Biology of Natural Products"	2018		
E. S. Reckzeh, A. Brockmeyer, M. Metz, H. Waldmann, P. Janning	Target Engagement of Small Molecules: Thermal Profiling Approaches on Different Levels	In: Systems Chemical Biology, Methods and Protocols (Ziegler, Waldmann eds)	2018		73-92
S. Ziegler, H. Waldmann (editors)	Systems Chemical Biology, Methods and Protocols (Methods in Molecular Biology, Series Editor: John M. Walker)	Humana Press	2018		
H. Li, R. Gontla, J. Flegel, C. Merten, S. Ziegler, A. P. Antonchick, H. Waldmann	Enantioselective Formal C(sp ³)-H Bond Activation in the Synthesis of Bioactive Spiropyrazolone Derivatives	Angew. Chem. Int. Ed.	2019	58	307-311
V. Nemeč, M. Hylsová, L. Maier, J. Flegel, S. Sievers, S. Ziegler, M. Schröder, B.-T. Berger, A. Chaikuad, B. Valčíková, S. Uldrijan, S. Drápela, K. Souček, H. Waldmann, S. Knapp, K. Paruch	Furo[3,2-b]pyridine – A Privileged Scaffold for Highly Selective Kinase Inhibitors and Effective Modulators of the Hedgehog Pathway	Angew. Chem. Int. Ed	2019	58	1062-1066
A. Friese, S. Kapoor, T. Schneidewind, S. Rao Vidadala, J. Sardana, A. Brause, T. Förster, M. Bischoff, J. Wagner, P. Janning, S. Ziegler, H. Waldmann	Chemical Genetics Reveals a Role of dCTP Pyrophosphatase 1 in Wnt Signaling	Angew. Chem. Int. Ed. https://doi.org/10.1002/anie.201905977	2019	58	13009- 13013
A. Christoforow, J. Wilke, A. Binici, A. Pahl, C. Ostermann, S. Sievers, H. Waldmann	Design, Synthesis and Phenotypic Profiling of Pyrano-Furo-Pyridone Pseudo Natural Products	Angew. Chem. Int. Ed. 10.1002/anie.201907 853 Angew. Chem. 10.1002/ange.201907 853	2019	58	14715- 14723
L. Kremer, E. Hennes, A. Brause, A. Ursu, L.	Discovery of the Hedgehog Pathway Inhibitor Pipinib	Angew. Chem. Int. Ed.	2019	58	16617- 16628



Robke, H. T. Matsubayashi, Y. Nihongaki, J. Flegel, I. Mejdrová, J. Eickhoff, M. Baumann, R. Nencka, P. Janning, S. Kordes, H. R. Schöler, J. Sterneckert, T. Inoue, S. Ziegler, H. Waldmann	that Targets PI4KIII β				
J. Ceballos, M. Schwalfenberg, G. Karageorgis, E. S. Reckzeh, S. Sievers, C. Ostermann, A. Pahl, M. Sellstedt, J. Nowacki, M. A. Carnero Corrales, J. Wilke, L. Laraia, K. Tschapalda, M. Metz, D. A. Sehr, S. Brand, K. Winklhofer, P. Janning, S. Ziegler, H. Waldmann	Synthesis of Indomorphan Pseudo Natural Product Inhibitors of Glucose Transporters GLUT-1 and -3	Angew. Chem. Int. Ed.	2019	58	17016- 17035
L. Laraia, A. Friese, D. P. Corkery, G. Konstantinidis, N. Erwin, W. Hofer, H. Karatas, L. Klewer, A. Brockmeyer, M. Metz, B. Schölermann, M. Dwivedi, L. Li, P. Rios-Munoz, M. Köhn, R. Winter, I. R. Vetter, S. Ziegler, P. Janning, Y.-W. Wu, H. Waldmann	The cholesterol transfer protein GRAMD1A regulates autophagosome biogenesis	Nat. Chem. Biol.	2019	15	710-720
T. Schneidewind, S. Kapoor, G. Garivet, G. Karageorgis, R. Narayan, G. Vendrell- Navarro, A. P. Antonchick, S. Ziegler, H. Waldmann	The Pseudo Natural Product Myokinasib is a Myosin Light Chain Kinase 1 Inhibitor with Unprecedented Chemotype	Cell Chem. Biol.	2019	26	512-523
E. S. Reckzeh, G. Karageorgis, M. Schwalfenberg, J. Ceballos, J. Nowacki, M.C.M. Stroet, A. Binici, L. Knauer, S. Brand, A. Choidas, C.	Inhibition of Glucose Transporters and Glutaminase Synergistically Impairs Tumor Cell Growth	Cell Chem. Biol. https://doi.org/10.1016/j.chembiol.2019.06.005	2019	26	1214-1228



Strohmann, S. Ziegler,
H. Waldmann

G. Garivet, W. Hofer, A. Konitsiotis, C. Klein, N. Kaiser, T. Mejuch, E. Fansa, A. Wittinghofer, P. I.H. Bastiaens, H. Waldmann	Small-Molecule Inhibition of the UNC-Src Interaction Impairs Dynamic Src Localization in Cells	Cell Chem. Biol.	2019	26	842-851.e7
A. Friese, A. Ursu, A. Hochheimer, H. R. Schöler, H. Waldmann, J. M. Bruder	The Convergence of Stem Cell Technologies and Phenotypic Drug Discovery	Cell Chem. Biol.	2019	26	1050-1066
N. Kaiser, T. Mejuch, R. Fedoryshchak, P. Janning, E. W. Tate, H. Waldmann	Photoactivatable Myristic Acid Probes for UNC119-Cargo Interactions	ChemBioChem	2019	20	134-139
K. Kumar, H. Waldmann	Nature Inspired Small Molecules for Chemical Biology	Israel Journal of Chemistry Doi:10.1002/ijch.201800 105	2019	59	41-51
T. Förster, E. Shang, K. Shimizu, E. Sanada, B. Schölermann, M. Hybecker, G. Hahne, M. Pascual Lopez- Alberca, P. Janning, N. Watanabe, S. Sievers, F. Giordanetto, T. Shimizu, S. Ziegler, H. Osada, H. Waldmann	2-Sulfonylpyrimidines target the kinesin HSET via cysteine alkylation	EJOC https://doi.org/10.1002/ejoc.201900586	2019		5486-5496
E. S.Reckzeh, A. Brockmeyer, M. Metz, H. Waldmann, P. Janning	Target Engagement of Small Molecules: Thermal Profiling Approaches on Different Levels	Methods Mol. Biol.	2019	1888	73-98
N. Kaiser, D. Corkery, Y. Wu, L. Laraia, H. Waldmann	Modulation of autophagy by the novel mitochondrial complex I inhibitor Authipyrin	Bioorg. Med. Chem.	2019	27	2444-2448
Y.-W. Wu, H. Waldmann	Toward the role of cholesterol and cholesterol transfer protein in autophagosome biogenesis	Autophagy DOI: 10.1080/15548627.20 19.1666595	2019	15:12	2167-2168
P. Cromm, H. Adihou, S. Kapoor, M.	Lipidated Stapled Peptides Targeting the Acyl Binding Protein UNC 119	ChemBioChem	2019	20	2987-2990



Vazquez-Chantada,
P. Davey, D.
Longmire, E. Hennes,
W. Hofer, P. Kuchler,
E. Chiarparin, H.
Waldmann, T. N.
Grossmann

T. Furuta, Y. Mizukami, L. Asano, K. Kotake, S. Ziegler, H. Yoshida, M. Watanabe, S. Sato, H. Waldmann, M. Nishikawa, M. Uesugi	Nutrient-Based Chemical Library as a Source of Energy Metabolism Modulators	ACS Chem. Biol.	2019	14	1860-1865
A.V. Pobbati, T. Mejuch, S. Charkaborty, H. Karatas, S. R. Bharath, S. M. Gueret, P.-A. Goy, G. Hahne, A. Pahl, S. Sievers, E. Guccione, H. Song, H. Waldmann, W. Hong	Identification of Quinolinols as Activators of TEAD-Dependent Transcription	ACS Chem. Biol.	2019	14	2909-2921
E. S. Reckzeh, H. Waldmann	Development of Glucose Transporter (GLUT) Inhibitors	EurJoc	2019	16	2321-2329
P. t'Hart, J. Openy, A. Krzyzanowski, H. Adihou, H. Waldmann	Hot-spot guided design of macrocyclic inhibitors of the LSD1-CoREST1 interaction	Tetrahedron	2019	75	130685
L. Laraia, G. Garivet, D. J. Foley, N. Kaiser, S. Müller, S. Zinken, T. Pinkert, J. Wilke, D. Corkery, A. Pahl, S. Sievers, P. Janning, C. Arenz, Y.-W. Wu, R. Rodriguez, H. Waldmann	Image-Based Morphological Profiling Identifies a Lysosomotropic, Iron- Sequestering Autophagy Inhibitor	Angew. Chem. Int. Ed. doi: 10.1002/anie.201913 712.	2020	59	5721-5729
M. Winzker, A. Friese, U. Koch, P. Janning, S. Ziegler, H. Waldmann	Development of a PDE δ Targeting PROTACs that Impair Lipid Metabolism	Angew. Chem. Int. Ed.	2020	59	5595-5601
D. J. Foley, S. Zinken, D. Corkery, L. Laraia, A. Pahl, Y. Wu, H. Waldmann	Phenotyping Reveals the Targets of a Pseudo-Natural Product Autophagy Inhibitor	Angew. Chem. Int. Ed. / Angew. Chem.	2020	59 / 132	12470- 12476 / 12570- 12576
S. M. Guéret, S. Thavam, R. J. Carbajo, M.	Macrocyclic Modalities Combining Peptide Epitopes and Natural Product Fragments	J. Am. Chem. Soc. http://dx.doi.org/10.1021/jacs.0c00269 .	2020	142	4904-4915



Potowski, N. Larsson,
G. Dahl, A. Dellsén, T.
N. Grossmann, A. T.
Plowright, E. Valeur,
M. Lemurell, H.
Waldmann

G. Karageorgis, D. J. Foley, L. Laraia, H. Waldmann Principle and design of pseudo-natural products: Nat. Chem. **2020** 12 227-235

H. Adihou, R. Gopalakrishnan, T. Förster, S. M. Guéret, R. Gasper, S. Geschwindner, C. C. García, H. Karatas, A. V. Pobbati, M. Vazquez-Chantada, P. Davey, C. M. Wassvik, J. Kah Sheng Pang, B. Seng Soh, W. Hong, E. Chiarparin, D. Schade, A. T. Plowright, E. Valeur, M. Lemurell, T. N. Grossmann, H. Waldmann A protein tertiary structure mimetic modulator of the Hippo signalling pathway Nat. Commun. **2020** 11 5425

E. S. Reckzeh, H. Waldmann Small-Molecule Inhibition of Glucose Transporters GLUT-1-4 ChemBioChem **2020** 21 45-52

H. Adihou, R. Gopalakrishnan, T. Förster, S. M. Guéret, R. Gasper, S. Geschwindner, C. Carillo Garcia, H. Karatas, A. V. Pobbati, M. Vazquez-Chantada, P. Davey, C. M. Wassvik, J. K. Sheng Pang, B. Seng Soh, W. Hong, E. Chiarparin, D. Schade, A. T. Plowright, E. Valeur, M. Lemurell, T. N. Grossmann, H. Waldmann A protein tertiary structure mimetic modulator of the Hippo signalling pathway Nature Communications **2020** 11 1-10
<https://doi.org/10.1038/s41467-020-19224-8>

M. Grigalunas, A. Burhop, A. Christoforow, H. Waldmann Pseudo-natural products and natural product-inspired methods in chemical biology and drug discovery Curr. Op.Chem. Biol. **2020** 56 111-118



Waldmann

S. Shaaban, C. Davies, C. Merten, J. Flegel, F. Otte, C. Strohmann, H. Waldmann	RhIII-catalyzed C-H Activation of Aryl-Hydroxamates for the Synthesis of Isoindolinones	Chem. – A Eur. J.	2020	26	10729-10734
S. Shaaban, C. Davies, H. Waldmann	Applications of Chiral Cyclopentadienyl (Cp*) Metal Complexes in Asymmetric Catalysis (Minireview)	Eur. J. Org. Chem.	2020		6512-6524
T. Kawamura, Y. Futumura, E. Shang, M. Muroi, P. Janning, M. Ueno, J. Wilke, S. Takeda, Y. Kondoh, S. Ziegler, N. Watanabe, H. Waldmann, H. Osada	Discovery of small-molecule modulator of heterotrimeric Gi-protein by integrated phenotypic profiling and chemical proteomics	Bioscience, Biotechnology and Biochemistry	2020		DOI: 10.1080/09168451.2020.1812375
H. Karatas, M. Akbarzadeh, H. Adihou, G. Hahne, A. V. Pobbati, E. Yihui Ng, S. M. Guéret, S. Sievers, A. Pahl, M. Metz, S. Zinken, L. Dötsch, C. Nowak, S. Thavam, A. Friese, C. Kang, W. Hong, H. Waldmann	Discovery of Covalent Inhibitors Targeting the Transcriptional Associate Domain Central Pocket	J. Med.Chem	2020	63	11972-11989
T. Schneidewind, A. Brause, A. Pahl, A. Burhop, T. Mejuch, S. Sievers, H. Waldmann, S. Ziegler	Morphological Profiling Identifies a Common Mode of Action for Small Molecules with Different Targets	ChemBioChem	2020	21	3197-3207
G. S. Cremosnik, J. Liu, H. Waldmann	Guided by Evolution: from Biology Oriented Synthesis to Pseudo Natural Products	Natural Product Reports	2020	37	1497-1510
S. Shaaban, H. Li, F. Otte, C. Strohmann, A. Antonchick, H. Waldmann	Enantioselective Synthesis of 5-Membered-Ring Atropisomers with Chiral Rh(III) Complexes	Org. Lett.	2020	22 (23)	9199-9202
P. t-Hart, P. Hommen, A. Noisier, A. Krzyzanowski, D. Schüler, A. T. Porfetye, M. Akbarzadeh, I. R. Vetter, H. Adihou, H. Waldmann	Structure based design of bicyclic peptide inhibitors of RbAp48	Angew. Chem. Int. Ed.	2021	60	1813-1820



J.Liu, G. S. Cremosnik, F. Otte, A. Pahl, S. Sievers, C. Strohmann, H. Waldmann	Design, Synthesis and Biological Evaluation of Chemically and Biologically Diverse Pyrroquinoline Pseudo Natural Products	Angew. Chem. Int. Ed.	2021	60	4648-4656
E. Hennes, P. Lampe, L. Dötsch, N. Bruning, L.-M. Pulvermacher, S. Sievers, S. Ziegler, H. Waldmann	Cell-Based Identification of New IDO1 Modulator Chemotypes	Angew. Chem. Int. Ed.	2021	60	9869-9874
G. Karageorgis, D. J. Foley, L. Laraia, S. Brakmann, H. Waldmann	Minireview: Pseudo Natural Products – Chemical Evolution of Natural Product Structure	Angew. Chem. Int. Ed.	2021	60	15705- 15723
O. Yildirim, M. Grigalunas, L. Brieger, C. Strohmann, A. P. Antonchick, H. Waldmann	Dynamic Catalytic Highly Enantioselective 1,3-Dipolar Cycloadditions	Angew. Chem. Int. Ed.	2021	60	20012- 20020
J. Liu, J. Flegel, F. Otte, A. Pahl, S. Sievers, C. Strohmann, H. Waldmann	Combination of Pseudo-Natural Product Design and Formal Natural Product Ring Distortion Yields Stereochemically and Biologically Diverse Pseudo Sesquiterpenoid Alkaloids	Angew. Chem. Int. Ed.	2021	60	21384- 21395
M. Grigalunas, A. Burhop, S. Zinken, A. Pahl, N. Wild, Y. Mantel, S. Sievers, D. J. Foley, R. Scheel, A. P. Antonchick, H. Waldmann	Natural Product Fragment Combination to Performance-Diverse Pseudo-Natural Products	Nat. Commun.	2021	12	1883 https://doi.org/10.1038/s41467-021-22174-4
A. Burhop, S. Bag, M. Grigalunas, S. Woitalla, P. Bodenbinder, L. Brieger, C. Strohmann, A. Pahl, S. Sievers H. Waldmann	Synthesis of Indofulvin Pseudo-Natural Products Yields a New Autophagy Inhibitor Chemotype	Advanced Science, DOI: 10.1002/ adv.202102042	2021	8	20102042
S. Ziegler, S. Sievers, H. Waldmann	Morphological Profiling of Small Molecules (Review)	Cell Chem Biol	2021	28	300-319
T. Schneidewind, A. Brause, S. Sievers, A. Pahl, M.G. Sankar, M. Winzker, P.Janning, K. Kumar, S. Ziegler, H. Waldmann	Combined morphological and proteome profiling reveals target-independent impairment of cholesterol homeostasis	Cell Chem. Biol. https://doi.org/10.1016/j.chembiol.2021.06.003	2021	28	1780-1794



J. Wilke, T. Kawamura, H. Xu, A. Brause, A. Friese, M. Metz, D. Schepmann, B. Wünsch, A. Artacho-Cordón, F. R. Nieto, N. Watanabe, H. Osada, S. Ziegler, H. Waldmann	Discovery of a novel σ 1 receptor antagonist by combination of unbiased Cell Painting and thermal proteome profiling	Cell Chem. Biol. https://doi.org/10.1016/j.chembiol.2021.01.009	2021	28	848-854
M.A. Carnero Corrales, S. Zinken, G. Konstantinidis, M. Rafehi, A. Abdelrahman, Y.-W. Wu, P. Janning, C. E. Müller, L. Laraia, H. Waldmann	Thermal Proteome Profiling Identifies the Membrane-Bound Purinergic Receptor P2X4 as a Target of the Autophagy Inhibitor Indophagolin	Cell Chem. Biol.	2021	28	1750-1754
A. Lopéz-Pérez, S. Freischem, I. Grimm, O. Weiergräber, A. Dingley, M. Pascual López-Alberca, H. Waldmann, W. Vollmer, K. Kumar, C. Vuong	Discovery of pyrrolidine-2,3-diones as novel inhibitors of <i>P. aeruginosa</i> PBP3	Antibiotics https://doi.org/10.3390/antibiotics10050529	2021	10	529
A. Krzyzanowski, R. Gasper, H. Adihou, P. Hart, H. Waldmann	Biochemical Investigation of the Interaction of pICln, RioK1 and COPR5 with the PRMT5–MEP50 Complex	ChemBioChem.	2021	22	1908-1944
S. Shaaban, H. Li, C. Merten, A. P. Antonchick, H. Waldmann	Rhodium(III)-catalyzed Enantioselective Benzamidation of Cyclopropenes	Synthesis	2021	53	2192-2200
C. Nöcker, N. Kaiser, D. Foley, S. Sievers, P. Janning, H. Waldmann, L. Laraia	Thermal proteome profiling efficiently identifies ribosome destabilizing oxazolidinones	Tetrahedron https://doi.org/10.1016/j.tet.2021.132118	2021	87	132118
J.M. Gally, A. Pahl, H. Waldmann	Identifying Bioactivity of Pseudo-Natural Products using the Cell Painting Assay	ARKIVOC	2021		Part iv, 89-104
M. Akbarzadeh, J. Flegel, S. Patil, E. Shang, R. Narayan, M. Buchholzer, N. S. Kazemein Jasemi, M. Grigalunas, A. Krzyzanowski, D. Abegg, A. Shuster, M. Potowski, H. Karatas, G. Karageorgis, N.	The Pseudo-Natural Product Rhonin Targets RHOGDI1	Angew. Chem. Int.Ed.	2022	61	e202115193



Mosaddeghzadeh, M.-
L. Zischinsky, C.
Merten, C. Golz, L.
Brieger, C.
Strohmann, A. P.
Antonchick, P.
Janning, A. Adibekian,
R. S. Goody, M. R.
Ahmadian, S. Ziegler,
H. Waldmann

G. Niggemeyer, A. Knyazeva, R. Gasper, D. Corkery, P. Bodenbinder, J. Holstein, S. Sievers, Y..W. Wu, H. Waldmann	Synthesis of 20-Membered Macrocyclic Pseudo-Natural Products Yields Inducers of LC3-Lipidation	Angew. Chem. Int. Ed.	2022	61	E202114328
Davies, C., Dötsch, L., Ciulla, M.G., Hennes, E., Yoshida, K., Gasper, R., Scheel, R., Sievers, S., Strohmann, C., Kumar, K., Ziegler, S., Waldmann, H.	Identification of a Novel Pseudo-Natural Product Type IV IDO1 Inhibitor Chemotype	Angew. Chem. Int. Ed.	2022	61	E202209374
Young, R.Y., Flitsch, S.L., Grigalunas, M., Leeson, P.D., Quinn, R.J., Turner, N.J., Waldmann, H.	The Time and Place for Nature in Drug Discovery	J. Am. Chem. Soc. Au	2022	2	2400-2416
M. Grigalunas, S. Brakmann, H. Waldmann	Perspective: Chemical Evolution of Natural Product Structure	J. Am. Chem. Soc.	2022	144	3314-3329
M. Akbarzadeh, I. Deipenwisch, B. Schölermann, A. Pahl, S. Siever, S. Ziegler, H. Waldmann	Morphological Profiling by Means of Cell Painting Assay Enables Identification of Tubulin-Targeting Compounds	Cell Chem. Biol.	2022	29	1053- 1064.e3
Wesseler, F., Riege, D., Puthanveedu, M., Halver, J., Müller, E., Bertrand, J., Antonchick, A., Sievers, S., Waldmann, H., Schade, D.	Probing Embryonic Development Enables the Discovery of Unique Small-Molecule Bone Morphogenetic Protein Potentiators	J. Med. Chem.	2022	65	3978-3990
Wesseler, F., Lohmann, S., Riege,	Phenotypic Discovery of Triazolo[1,5-c]quinazolines as a First-In-	J. Med. Chem.	2022	65	15263- 15281



D., Halver, J., Roth, A., Pichlo, C., Weber, S., Takamiya, M., Müller, E., Ketzel, J., Flegel, J., Gihring, A., Rastegar, S., Bertrand, J., Baumann, U., Knippschild, U., Peifer, C., Sievers, S., Waldmann, H., Schade, D.	Class Bone Morphogenetic Protein Amplifier Chemotype				
Krzyzanowski, A., Esser, L.M., Willaume, A., Prudent, R., Peter, C., 't Hart, P., Waldmann, H	Development of Macrocyclic PRMT5-Adaptor Protein Interaction Inhibitors	J. Med. Chem.	2022	65	15300-15311
Flegel, J., Shaaban, S., Jia, Z., Schulte, B., Lian, Y., Krzyzanowski, A., Metz, M., Schneidewind, T., Wessler, F., Flegel, A., Reich, A., Brause, A., Xue, G., Zhang, M., Dötsch, L., Stender, I., Hoffmann, J.-E., Scheel, R., Janning, P., Rastinejad, F., Schade, D., Strohmman, C., Antonchick, A., Sievers, S., Moura-Alves, P., Ziegler, S., Waldmann, H.	The Highly Potent AhR Agonist Picoberin Modulates Hh-Dependent Osteoblast Differentiation	J. Med. Chem.	2022	65	16268-16289
S. Shaaban, C. Merten, H. Waldmann	Catalytic Atroposelective C7-Functionalisation of Indolines and Indoles	Chem. Eur. J.	2022	28	e202103365
Grigalunas, M., Patil, S., Krzyzanowski, A., Pahl, A., Flegel, J., Schölermann, B., Xie, J., Sievers, S., Ziegler, S., Waldmann, H.	Unprecedented Combination of Polyketide Natural Product Fragments Identifies the New Hedgehog Signaling Pathway Inhibitor Grisonone	Chem. Eur. J.	2022	28	e202202164
Yoshioka, H., Kawamura, T., Muroi, M., Kondoh, Y., Honda, K., Kawatani,	Identification of a Small Molecule That Enhances Ferroptosis via Inhibition of Ferroptosis Suppressor Protein 1 (FSP1)	ACS Chem. Biol.	2022	17	483-491



M., Aono, H.,
Waldmann, H.,
Watanabe, H., Osada,
H.

Picard, L.K., Littwitz-
Salomon, E.,
Waldmann, H., Watzl,
C. Inhibition of glucose uptake blocks
proliferation but not cytotoxic activity of
NK cells Cells **2022** 11 3489-3506

Foley, D.J.,
Waldmann, H. Ketones as strategic building blocks for
the synthesis of natural product-inspired
compounds Chem. Soc. Rev. **2022** 51 4094-4120

C. Davies, S.
Shaaban, H.
Waldmann Asymmetric Catalysis with Chiral
Cyclopentadienyl Complexes to Access
Privileged Scaffolds Trends in Chemistry **2022** 4 318-330

Jones, P.S.,
Boucharens, S.,
McElroy, S.P.,
Morrison, A.,
Honarnejad, S., van
Boeckel, S., van den
Hurk, H., Basting, D.,
Huser, J., Jaroch, S.,
et al. IMI European Lead Factory —
democratizing access to highthroughput
screening Nat. Rev. Drug
Discov. **2022** 21 245-246

Pahl, A.,
Schölermann, B.,
Rusch, M., Dow, M.,
Hedberg, C., Nelson,
A., Sievers, S.,
Waldmann, H.,
Ziegler, S. Morphological Subprofile Analysis for
Bioactivity Annotation of Small Molecules bioRxiv **2023**
<https://doi.org/10.1101/2022.08.15.503944>

Bag, S., Grigalunas,
M., Liu, J., Patil,
S.D., Bonowski, J.,
Schölermann, B.,
Zhang, R., Wang, L.,
Pahl, A., Sievers, S.,
Brieger, L,
Strohmann, C.,
Ziegler, S.,
Waldmann, H. A divergent intermediate strategy yields
biologically diverse pseudo-natural
products Nature Chemistry subm
itted

Liu, J., Mallick, S.,
Xie, Y., Lucas, B.,
Schölermann, B.,
Scheel, R.,
Strohmann, C.,
Protzel, C., Berg, T., Morphological profiling identifies the
motor protein Eg5 as cellular target of
spirooxindoles Angew. Chem. subm
itted

Prof. Dr. Dr. h.c. Herbert
Waldmann

MAX PLANCK INSTITUTE
OF MOLECULAR PHYSIOLOGY



tu technische universität
dortmund

Ziegler, S.,
Waldmann, H.