

Jörg Rademann

List of Publications by Year in descending order

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152
papers

4,883
citations

108046

37
h-index

145109

60
g-index

184
all docs

184
docs citations

184
times ranked

6717
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Peptideâ€“Bismuth Bicycles: In Situ Access to Stable Constrained Peptides with Superior Bioactivity. <i>Angewandte Chemie - International Edition</i> , 2022, 61, . | 7.2 | 11 |
| 2 | Peptideâ€“Bismuth Bicycles: In Situ Access to Stable Constrained Peptides with Superior Bioactivity. <i>Angewandte Chemie</i> , 2022, 134, . | 1.6 | 3 |
| 3 | Pentafluorophosphatoâ€“Phenylalanines:Â“Amphiphilic Phosphotyrosine Mimetics Displaying Fluorineâ€“Specific Protein Interactions. <i>Angewandte Chemie - International Edition</i> , 2022, , . | 7.2 | 3 |
| 4 | A Formylglycineâ€“Peptide for the Siteâ€“Directed Identification of Phosphotyrosineâ€“Mimetic Fragments**. <i>Chemistry - A European Journal</i> , 2022, 28, . | 1.7 | 4 |
| 5 | 2-Cyanoisonicotinamide Conjugation: A Facile Approach to Generate Potent Peptide Inhibitors of the Zika Virus Protease. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 732-737. | 1.3 | 21 |
| 6 | Chemische Evolution antiviraler Wirkstoffe gegen Enterovirus D68 durch Proteintemplatâ€“gesteuerte Knoevenagelreaktionen. <i>Angewandte Chemie</i> , 2021, 133, 13405-13413. | 1.6 | 1 |
| 7 | Chemical Evolution of Antivirals Against Enterovirus D68 through Proteinâ€“Templated Knoevenagel Reactions. <i>Angewandte Chemie - International Edition</i> , 2021, 60, 13294-13301. | 7.2 | 9 |
| 8 | Small-molecule inhibitors of the PDZ domain of Dishevelled proteins interrupt Wnt signalling. <i>Magnetic Resonance</i> , 2021, 2, 355-374. | 0.8 | 5 |
| 9 | Tuning the network charge of biohybrid hydrogel matrices to modulate the release of SDF-1. <i>Biological Chemistry</i> , 2021, 402, 1453-1464. | 1.2 | 4 |
| 10 | Structural insights into the modulation of PDGF/PDGFR-Î² complexation by hyaluronan derivatives. <i>Biological Chemistry</i> , 2021, 402, 1441-1452. | 1.2 | 9 |
| 11 | Insights into structure, affinity, specificity, and function of GAG-protein interactions through the chemoenzymatic preparation of defined sulfated oligohyaluronans. <i>Biological Chemistry</i> , 2021, 402, 1375-1384. | 1.2 | 3 |
| 12 | Identification of intracellular glycosaminoglycan-interacting proteins by affinity purification mass spectrometry. <i>Biological Chemistry</i> , 2021, 402, 1427-1440. | 1.2 | 5 |
| 13 | Characterization of defined sulfated heparinâ€“like oligosaccharides by electrospray ionization ion trap mass spectrometry. <i>Journal of Mass Spectrometry</i> , 2021, 56, e4692. | 0.7 | 10 |
| 14 | Nanoparticulate Inhibitors of Flavivirus Proteases from Zika, West Nile and Dengue Virus Are Cell-Permeable Antivirals. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 1955-1961. | 1.3 | 3 |
| 15 | Sulfation pattern dependent Iron (III) mediated interleukinâ€“8 glycan binding. <i>ChemBioChem</i> , 2021, , . | 1.3 | 4 |
| 16 | IR action spectroscopy of glycosaminoglycan oligosaccharides. <i>Analytical and Bioanalytical Chemistry</i> , 2020, 412, 533-537. | 1.9 | 24 |
| 17 | Peptideâ€“mediated surface coatings for the release of woundâ€“healing cytokines. <i>Journal of Tissue Engineering and Regenerative Medicine</i> , 2020, 14, 1738-1748. | 1.3 | 9 |
| 18 | Catching a Moving Target: Comparative Modeling of Flaviviral NS2B-NS3 Reveals Small Molecule Zika Protease Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 514-520. | 1.3 | 10 |

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|----|---|-----|-----------|
| 19 | Side-Chain Modification of Peptides Using a Phosphoranylidene Amino Acid. <i>Organic Letters</i> , 2020, 22, 2976-2980. | 2.4 | 8 |
| 20 | Identification and validation of a novel dual small-molecule TLR2/8 antagonist. <i>Biochemical Pharmacology</i> , 2020, 177, 113957. | 2.0 | 5 |
| 21 | Biological Characterization, Mechanistic Investigation and Structure-Activity Relationships of Chemically Stable TLR2 Antagonists. <i>ChemMedChem</i> , 2020, 15, 1364-1371. | 1.6 | 8 |
| 22 | Modulation of Human CXCL12 Binding Properties to Glycosaminoglycans To Enhance Chemotactic Gradients. <i>ACS Biomaterials Science and Engineering</i> , 2019, 5, 5128-5138. | 2.6 | 10 |
| 23 | Syntheses of defined sulfated oligohyaluronans reveal structural effects, diversity and thermodynamics of GAG-protein binding. <i>Chemical Science</i> , 2019, 10, 866-878. | 3.7 | 30 |
| 24 | Sulfation Patterns of Saccharides and Heavy Metal Ion Binding. <i>Chemistry - A European Journal</i> , 2019, 25, 12083-12090. | 1.7 | 16 |
| 25 | An Intrinsic Hydrophobicity Scale for Amino Acids and Its Application to Fluorinated Compounds. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 8216-8220. | 7.2 | 30 |
| 26 | Eine intrinsische Hydrophobieskala für Aminosäuren und ihre Anwendung auf fluorierte Verbindungen. <i>Angewandte Chemie</i> , 2019, 131, 8300-8304. | 1.6 | 2 |
| 27 | How Proteins Catalyze Ligand Formation: Protein-Templated Fragment Ligation Employed in the Validation of Cancer Targets. <i>Proceedings (mdpi)</i> , 2019, 22, 110. | 0.2 | 0 |
| 28 | Dual Action of Sulfated Hyaluronan on Angiogenic Processes in Relation to Vascular Endothelial Growth Factor-A. <i>Scientific Reports</i> , 2019, 9, 18143. | 1.6 | 28 |
| 29 | Hyaluronan/collagen hydrogels containing sulfated hyaluronan improve wound healing by sustained release of heparin-binding EGF-like growth factor. <i>Acta Biomaterialia</i> , 2019, 86, 135-147. | 4.1 | 113 |
| 30 | The transcription factor STAT5 catalyzes Mannich ligation reactions yielding inhibitors of leukemic cell proliferation. <i>Nature Communications</i> , 2019, 10, 66. | 5.8 | 25 |
| 31 | Phenylthiomethyl Ketone-Based Fragments Show Selective and Irreversible Inhibition of Enteroviral 3C Proteases. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 1218-1230. | 2.9 | 20 |
| 32 | Identification of a pyrogallol derivative as a potent and selective human TLR2 antagonist by structure-based virtual screening. <i>Biochemical Pharmacology</i> , 2018, 154, 148-160. | 2.0 | 20 |
| 33 | Structural Basis for Binding of Fluorescent CMP-Neu5Ac Mimetics to Enzymes of the Human ST8Sia Family. <i>ACS Chemical Biology</i> , 2018, 13, 2320-2328. | 1.6 | 6 |
| 34 | Protein-Templated Fragment Ligations—From Molecular Recognition to Drug Discovery. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 7358-7378. | 7.2 | 60 |
| 35 | C-type lectin receptor DCIR modulates immunity to tuberculosis by sustaining type I interferon signaling in dendritic cells. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, E540-E549. | 3.3 | 67 |
| 36 | Proteintemplat-gesteuerte Fragmentligationen – von der molekularen Erkennung zur Wirkstofffindung. <i>Angewandte Chemie</i> , 2017, 129, 7464-7485. | 1.6 | 19 |

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|----|--|-----|-----------|
| 37 | Sulfated Hyaluronan Alters Endothelial Cell Activation in Vitro by Controlling the Biological Activity of the Angiogenic Factors Vascular Endothelial Growth Factor-A and Tissue Inhibitor of Metalloproteinase-3. <i>ACS Applied Materials & Interfaces</i> , 2017, 9, 9539-9550. | 4.0 | 23 |
| 38 | Protein-Templated Formation of an Inhibitor of the Blood Coagulation Factor Xa through a Background-Free Amidation Reaction. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 3718-3722. | 7.2 | 28 |
| 39 | Sulfated Hyaluronan Derivatives Modulate TGF- β 1:Receptor Complex Formation: Possible Consequences for TGF- β 21 Signaling. <i>Scientific Reports</i> , 2017, 7, 1210. | 1.6 | 30 |
| 40 | Proteintemplat-gesteuerte Bildung eines Inhibitors des Koagulationsfaktors Xa durch eine Amidierung ohne Hintergrundreaktion. <i>Angewandte Chemie</i> , 2017, 129, 3772-3776. | 1.6 | 7 |
| 41 | Biotransformation of 2,4-toluenediamine in human skin and reconstructed tissues. <i>Archives of Toxicology</i> , 2017, 91, 3307-3316. | 1.9 | 4 |
| 42 | Rhamnolipids form drug-loaded nanoparticles for dermal drug delivery. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2017, 116, 31-37. | 2.0 | 36 |
| 43 | Benzyl Mono- <i>P</i> -Fluorophosphonate and Benzyl Penta- <i>P</i> -Fluorophosphate Anions Are Physiologically Stable Phosphotyrosine Mimetics and Inhibitors of Protein Tyrosine Phosphatases. <i>Chemistry - A European Journal</i> , 2017, 23, 15387-15395. | 1.7 | 17 |
| 44 | Pharmacological inhibition of focal segmental glomerulosclerosis-related, gain of function mutants of TRPC6 channels by semi-synthetic derivatives of larixol. <i>British Journal of Pharmacology</i> , 2017, 174, 4099-4122. | 2.7 | 16 |
| 45 | Mapping Protein-Protein Interactions of the Resistance-Related Bacterial Zeta Toxin-Epsilon Antitoxin Complex (μ 21 τ 2) with High Affinity Peptide Ligands Using Fluorescence Polarization. <i>Toxins</i> , 2016, 8, 222. | 1.5 | 11 |
| 46 | Incorporation of Amino Acids with Long-Chain Terminal Olefins into Proteins. <i>Molecules</i> , 2016, 21, 287. | 1.7 | 10 |
| 47 | Chemoenzymatic Synthesis of Nonasulfated Tetrahyaluronan with a Paramagnetic Tag for Studying Its Complex with Interleukin-10. <i>Chemistry - A European Journal</i> , 2016, 22, 5563-5574. | 1.7 | 35 |
| 48 | Eintopfsynthese ungeschützter anomerer Glykosylthiole in Wasser für Glykanligationen mit hochfunktionalisierten Zuckern. <i>Angewandte Chemie</i> , 2016, 128, 15736-15740. | 1.6 | 7 |
| 49 | Structural and functional insights into the interaction of sulfated glycosaminoglycans with tissue inhibitor of metalloproteinase-3 - A possible regulatory role on extracellular matrix homeostasis. <i>Acta Biomaterialia</i> , 2016, 45, 143-154. | 4.1 | 31 |
| 50 | Sulfated Hyaluronan Alters the Interaction Profile of TIMP-3 with the Endocytic Receptor LRP-1 Clusters II and IV and Increases the Extracellular TIMP-3 Level of Human Bone Marrow Stromal Cells. <i>Biomacromolecules</i> , 2016, 17, 3252-3261. | 2.6 | 20 |
| 51 | One-Pot Synthesis of Unprotected Anomeric Glycosyl Thiols in Water for Glycan Ligation Reactions with Highly Functionalized Sugars. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 15510-15514. | 7.2 | 40 |
| 52 | Irreversible inhibitors of the 3C protease of Coxsackie virus through templated assembly of protein-binding fragments. <i>Nature Communications</i> , 2016, 7, 12761. | 5.8 | 30 |
| 53 | The structural investigation of glycosaminoglycan binding to CXCL12 displays distinct interaction sites. <i>Glycobiology</i> , 2016, 26, 1209-1221. | 1.3 | 27 |
| 54 | Identification of the Glycosaminoglycan Binding Site of Interleukin-10 by NMR Spectroscopy. <i>Journal of Biological Chemistry</i> , 2016, 291, 3100-3113. | 1.6 | 32 |

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|----|---|-----|-----------|
| 55 | Cholesteryl esters stabilize human CD1c conformations for recognition by self-reactive T cells. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, E1266-75. | 3.3 | 41 |
| 56 | Identification and Validation of Larixyl Acetate as a Potent TRPC6 Inhibitor. <i>Molecular Pharmacology</i> , 2016, 89, 197-213. | 1.0 | 67 |
| 57 | Peptide-polymer ligands for a tandem WW-domain, an adaptive multivalent protein-protein interaction: lessons on the thermodynamic fitness of flexible ligands. <i>Beilstein Journal of Organic Chemistry</i> , 2015, 11, 837-847. | 1.3 | 11 |
| 58 | Light-switched inhibitors of protein tyrosine phosphatase PTP1B based on phosphonocarbonyl phenylalanine as photoactive phosphotyrosine mimetic. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 2839-2847. | 1.4 | 15 |
| 59 | Shp2 signaling suppresses senescence in <i>PyMT</i> -induced mammary gland cancer in mice. <i>EMBO Journal</i> , 2015, 34, 1493-1508. | 3.5 | 31 |
| 60 | Selective Inhibitors of the Protein Tyrosine Phosphatase SHP2 Block Cellular Motility and Growth of Cancer Cells <i>in vitro</i> and <i>in vivo</i> . <i>ChemMedChem</i> , 2015, 10, 815-826. | 1.6 | 65 |
| 61 | Nonlinear topology optimization of centrifugally loaded aero-engine part with newly developed optimality-criteria based algorithm. <i>Aerospace Science and Technology</i> , 2014, 39, 705-711. | 2.5 | 6 |
| 62 | Fluorescent Mimetics of CMP-Neu5Ac Are Highly Potent, Cell-Permeable Polarization Probes of Eukaryotic and Bacterial Sialyltransferases and Inhibit Cellular Sialylation. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 5700-5705. | 7.2 | 18 |
| 63 | Catalytic activation of pre-substrates via dynamic fragment assembly on protein templates. <i>Nature Communications</i> , 2014, 5, 5170. | 5.8 | 18 |
| 64 | Chemoselective Wittig and Michael Ligations of Unprotected Peptidyl Phosphoranes in Water Furnish Potent Inhibitors of Caspase-3. <i>Organic Letters</i> , 2014, 16, 4428-4431. | 2.4 | 3 |
| 65 | Multivalent presentation of the cell-penetrating peptide nona-arginine on a linear scaffold strongly increases its membrane-perturbing capacity. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 2014, 1838, 3097-3106. | 1.4 | 17 |
| 66 | Flexible, polymer-supported synthesis of sphingosine derivatives provides ceramides with enhanced biological activity. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 5506-5512. | 1.4 | 6 |
| 67 | Fluoreszente Mimetika von CMP-Neu5Ac sind hochaffine, zellgängige Polarisationssonden eukaryotischer und bakterieller Sialyltransferasen und inhibieren die zelluläre Sialylierung. <i>Angewandte Chemie</i> , 2014, 126, 5808-5813. | 1.6 | 4 |
| 68 | Alzheimer's Disease: Identification and Development of β -Secretase (BACE1) Binding Fragments and Inhibitors by Dynamic Ligand Screening (DLS). <i>ChemMedChem</i> , 2013, 8, 1041-1056. | 1.6 | 14 |
| 69 | Effects of Charge and Charge Distribution on the Cellular Uptake of Multivalent Arginine-Containing Peptide-Polymer Conjugates. <i>ChemBioChem</i> , 2013, 14, 1982-1990. | 1.3 | 15 |
| 70 | Highly Functionalized Terpyridines as Competitive Inhibitors of AKAP-PKA Interactions. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 12187-12191. | 7.2 | 46 |
| 71 | Propargyl Amides as Irreversible Inhibitors of Cysteine Proteases-A Lesson on the Biological Reactivity of Alkynes. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 8210-8212. | 7.2 | 23 |
| 72 | Hoch funktionalisierte Terpyridine als kompetitive Inhibitoren von AKAP-PKA-Wechselwirkungen. <i>Angewandte Chemie</i> , 2013, 125, 12409-12413. | 1.6 | 6 |

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|----|--|-----|-----------|
| 73 | Fmoc-Based Synthesis of Peptide Thioacids for Azide Ligations via 2-Cyanoethyl Thioesters. <i>Organic Letters</i> , 2012, 14, 5038-5041. | 2.4 | 22 |
| 74 | Extra- and Intracellular Imaging of Human Matrix Metalloprotease 11 (hMMP-11) with a Cell-penetrating FRET Substrate. <i>Journal of Biological Chemistry</i> , 2012, 287, 37857-37867. | 1.6 | 19 |
| 75 | Multivalent Design of Apoptosis-Inducing BH3 Peptide-Oligosaccharides Boosts the Intracellular Activity at Identical Overall Peptide Concentrations. <i>Chemistry - A European Journal</i> , 2012, 18, 16708-16715. | 1.7 | 29 |
| 76 | Soluble Peptidyl Phosphoranes for Metal-Free, Stereoselective Ligations in Organic and Aqueous Solution. <i>Organic Letters</i> , 2012, 14, 14-17. | 2.4 | 13 |
| 77 | New Tacrine-4-Oxo-4-chromene Hybrids as Multifunctional Agents for the Treatment of Alzheimer's Disease, with Cholinergic, Antioxidant, and β -Amyloid-Reducing Properties. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 1303-1317. | 2.9 | 244 |
| 78 | Benzoylphosphonate-Based Photoactive Phosphopeptide Mimetics for Modulation of Protein Tyrosine Phosphatases and Highly Specific Labeling of SH2 Domains. <i>Angewandte Chemie - International Edition</i> , 2012, 51, 9441-9447. | 7.2 | 20 |
| 79 | High yield expression of catalytically active USP18 (UBP43) using a Trigger Factor fusion system. <i>BMC Biotechnology</i> , 2012, 12, 56. | 1.7 | 14 |
| 80 | Fmoc-Based Synthesis of Peptide Thioesters for Native Chemical Ligation Employing a tert-Butyl Thiol Linker. <i>Organic Letters</i> , 2011, 13, 1606-1609. | 2.4 | 37 |
| 81 | Acyated cholesteryl galactosides are ubiquitous glycolipid antigens among <i>Borrelia burgdorferi</i> sensu lato: Figure 1. <i>FEMS Immunology and Medical Microbiology</i> , 2011, 63, 140-143. | 2.7 | 14 |
| 82 | Peptide aldehyde inhibitors challenge the substrate specificity of the SARS-coronavirus main protease. <i>Antiviral Research</i> , 2011, 92, 204-212. | 1.9 | 112 |
| 83 | Peptide-Heterocycle Chimera: New Classes of More Drug-Like Peptidomimetics by Ligations of Peptide-Bis(electrophiles) with Various Bis(nucleophiles). <i>European Journal of Organic Chemistry</i> , 2011, 2011, 730-739. | 1.2 | 10 |
| 84 | Discovery, Structure-Activity Relationship Studies, and Crystal Structure of Nonpeptide Inhibitors Bound to the Shank3 PDZ Domain. <i>ChemMedChem</i> , 2011, 6, 1411-1422. | 1.6 | 34 |
| 85 | Dynamic Substrate Enhancement for the Identification of Specific, Second-Site Binding Fragments Targeting a Set of Protein Tyrosine Phosphatases. <i>ChemBioChem</i> , 2011, 12, 2640-2646. | 1.3 | 25 |
| 86 | Coupling to Polymeric Scaffolds Stabilizes Biofunctional Peptides for Intracellular Applications. <i>Molecular Pharmacology</i> , 2011, 79, 692-700. | 1.0 | 16 |
| 87 | Design of chemical libraries with potentially bioactive molecules applying a maximum common substructure concept. <i>Molecular Diversity</i> , 2010, 14, 401-408. | 2.1 | 69 |
| 88 | Chemoenzymatic Synthesis of a Glycolipid Library and Elucidation of the Antigenic Epitope for Construction of a Vaccine Against Lyme Disease. <i>Chemistry - A European Journal</i> , 2010, 16, 3536-3544. | 1.7 | 20 |
| 89 | Cyclative Cleavage through Dipolar Cycloaddition: Polymer-Bound Azidopeptidylphosphoranes Deliver Locked cis-Triazolylcyclopeptides as Privileged Protein Binders. <i>Angewandte Chemie - International Edition</i> , 2010, 49, 5378-5382. | 7.2 | 45 |
| 90 | Efficient access to peptidyl ketones and peptidyl diketones via C-alkylations and C-acylations of polymer-supported phosphorus ylides followed by hydrolytic and/or oxidative cleavage. <i>Biopolymers</i> , 2010, 94, 220-228. | 1.2 | 13 |

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|-----|---|-----|-----------|
| 91 | Lysosomal Pathology and Osteopetrosis upon Loss of H ⁺ -Driven Lysosomal Cl ⁻ Accumulation. <i>Science</i> , 2010, 328, 1401-1403. | 6.0 | 210 |
| 92 | Phosphoinositol 3-kinase- β mediates antineutrophil cytoplasmic autoantibody-induced glomerulonephritis. <i>Kidney International</i> , 2010, 77, 118-128. | 2.6 | 64 |
| 93 | Dynamic template-assisted strategies in fragment-based drug discovery. <i>Trends in Biotechnology</i> , 2009, 27, 512-521. | 4.9 | 45 |
| 94 | Metal-Free, Regioselective Triazole Ligations that Deliver Locked <i>cis</i> Peptide Mimetics. <i>Angewandte Chemie - International Edition</i> , 2009, 48, 5042-5045. | 7.2 | 64 |
| 95 | Selective Identification of Cooperatively Binding Fragments in a High-Throughput Ligation Assay Enables Development of a Picomolar Caspase-3 Inhibitor. <i>Angewandte Chemie - International Edition</i> , 2009, 48, 6346-6349. | 7.2 | 45 |
| 96 | Resin-Bound Aminofluorescein for C-Terminal Labeling of Peptides: High-Affinity Polarization Probes Binding to Polyproline-Specific GYF Domains. <i>ChemBioChem</i> , 2008, 9, 2452-2462. | 1.3 | 13 |
| 97 | Sensitized Detection of Inhibitory Fragments and Iterative Development of Non-Peptidic Protease Inhibitors by Dynamic Ligation Screening. <i>Angewandte Chemie - International Edition</i> , 2008, 47, 3275-3278. | 7.2 | 64 |
| 98 | HPMA as a Scaffold for the Modular Assembly of Functional Peptide Polymers by Native Chemical Ligation. <i>Bioconjugate Chemistry</i> , 2008, 19, 2081-2087. | 1.8 | 19 |
| 99 | Specific inhibitors of the protein tyrosine phosphatase Shp2 identified by high-throughput docking. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2008, 105, 7275-7280. | 3.3 | 199 |
| 100 | Small-Molecule Scaffolds for CYP51 Inhibitors Identified by High-Throughput Screening and Defined by X-Ray Crystallography. <i>Antimicrobial Agents and Chemotherapy</i> , 2007, 51, 3915-3923. | 1.4 | 70 |
| 101 | C-Acylation of Polymeric Phosphoranylidene Acetates for C-Terminal Variation of Peptide Carboxylic Acids. <i>Organic Letters</i> , 2007, 9, 949-952. | 2.4 | 23 |
| 102 | Biophysical characterization of synthetic rhamnolipids. <i>FEBS Journal</i> , 2006, 273, 5101-5112. | 2.2 | 36 |
| 103 | Chemical Synthesis of a Glycolipid Library by a Solid-Phase Strategy Allows Elucidation of the Structural Specificity of Immunostimulation by Rhamnolipids. <i>Chemistry - A European Journal</i> , 2006, 12, 7116-7124. | 1.7 | 55 |
| 104 | An Efficient Method for the Synthesis of Peptide Aldehyde Libraries Employed in the Discovery of Reversible SARS Coronavirus Main Protease (SARS-CoV M ^{pro}) Inhibitors. <i>ChemBioChem</i> , 2006, 7, 1048-1055. | 1.3 | 50 |
| 105 | The Potential of P1 Site Alterations in Peptidomimetic Protease Inhibitors as Suggested by Virtual Screening and Explored by the Use of C α -C-Coupling Reagents. <i>ChemMedChem</i> , 2006, 1, 445-457. | 1.6 | 33 |
| 106 | Endotoxin-like properties of a rhamnolipid exotoxin from Burkholderia (<i>Pseudomonas</i>) plantarii: immune cell stimulation and biophysical characterization. <i>Biological Chemistry</i> , 2006, 387, 301-10. | 1.2 | 77 |
| 107 | Reversible Cross-Linking of Hyperbranched Polymers: A Strategy for the Combinatorial Decoration of Multivalent Scaffolds. <i>Angewandte Chemie - International Edition</i> , 2005, 44, 1560-1563. | 7.2 | 21 |
| 108 | Stereospecific Synthesis of Chiral 2,3-Dihydro-1,4-benzodithiine and Methyl-2,3-dihydro-1,4-benzodithiine Derivatives and their Toxic Effects on <i>Trypanosoma brucei</i> . <i>ChemBioChem</i> , 2005, 6, 1438-1441. | 1.3 | 1 |

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|-----|---|-----|-----------|
| 109 | Discovery of Mycobacterium Tuberculosis Protein Tyrosine Phosphatase A (MtpA) Inhibitors Based on Natural Products and a Fragment-Based Approach. ChemBioChem, 2005, 6, 1749-1753. | 1.3 | 60 |
| 110 | Polymer-Supported Synthetic Methodsâ€”Solid-Phase Synthesis (SPS) and Polymer-Assisted Solution-Phase (PASP) Synthesis. ChemInform, 2005, 36, no. | 0.1 | 0 |
| 111 | Novel Polymer and Linker Reagents for the Preparation of Protease Inhibitor Libraries. ChemInform, 2005, 36, no. | 0.1 | 0 |
| 112 | Progress in the preparation of peptide aldehydes via polymer supported IBX oxidation and scavenging by threonyl resin. Journal of Peptide Science, 2005, 11, 142-152. | 0.8 | 19 |
| 113 | Understanding Supported Reactions in Spherical Compartments: A General Algorithm To Model and Determine Rate Constants, Diffusion Coefficients, and Spatial Product Distributions. ACS Combinatorial Science, 2005, 7, 929-941. | 3.3 | 6 |
| 114 | Hydrophobically Assisted Switching Phase Synthesis: The Flexible Combination of Solid-Phase and Solution-Phase Reactions Employed for Oligosaccharide Preparation. Journal of the American Chemical Society, 2005, 127, 7296-7297. | 6.6 | 59 |
| 115 | High loading polymer reagents based on polycationic Ultraresins. Polymer-supported reductions and oxidations with increased efficiency. Tetrahedron, 2004, 60, 8703-8709. | 1.0 | 23 |
| 116 | Polymer-bound alkyltriazenes for mild racemization-free esterification of amino acid and peptide derivatives. Journal of Peptide Science, 2004, 10, 603-611. | 0.8 | 4 |
| 117 | Organic Protein Chemistry: Drug Discovery through the Chemical Modification of Proteins. Angewandte Chemie - International Edition, 2004, 43, 4554-4556. | 7.2 | 32 |
| 118 | Tailoring Ultraresins Based on the Cross-Linking of Polyethylene Imines. Comparative Investigation of the Chemical Composition, the Swelling, the Mobility, the Chemical Accessibility, and the Performance in Solid-Phase Synthesis of Very High Loaded Resins. ACS Combinatorial Science, 2004, 6, 340-349. | 3.3 | 13 |
| 119 | Addition to Carbon-Hetero Multiple Bonds. , 2004, , 322-345. | | 1 |
| 120 | Ein Phosphoran als polymergebundenes AcylanionenÄquivalent: Linkerreagentien für schonende und vielseitige C-C-Kupplungen. Angewandte Chemie, 2003, 115, 2595-2598. | 1.6 | 26 |
| 121 | A Phosphorane as Supported Acylanion Equivalent: Linker Reagents for Smooth and Versatile C-C Coupling Reactions. Angewandte Chemie - International Edition, 2003, 42, 2491-2494. | 7.2 | 37 |
| 122 | Trimellitic anhydride linker (TAL)â€”highly orthogonal conversions of primary amines employed in the parallel synthesis of labeled carbohydrate derivatives. Tetrahedron Letters, 2003, 44, 5019-5023. | 0.7 | 19 |
| 123 | Advanced Polymer Reagents Based on Activated Reactants and Reactive Intermediates: Powerful Novel Tools in Diversity-Oriented Synthesis. Methods in Enzymology, 2003, 369, 366-390. | 0.4 | 1 |
| 124 | SPOCC-194, a New High Functional Group Density PEG-Based Resin for Solid-Phase Organic Synthesis. ACS Combinatorial Science, 2002, 4, 523-529. | 3.3 | 33 |
| 125 | ULTRA-hochbeladene Harze durch Quervernetzung von linearem Polyethylenimin. Steigerung der Atomökonomie in der Polymer-unterstützten Chemie Wir danken der Firma Personal Chemistry GmbH, Konstanz, für die zeitweilige Überlassung eines Mikrowellen-Syntheseautomaten. J.R. bedankt sich für Unterstützung bei Prof. G. Jung und Prof. M. E. Maier, Universität Tübingen, dem | | |

| # | ARTICLE | IF | CITATIONS |
|-----|---|-----|-----------|
| 127 | ULTRA Loaded Resins Based on the Cross-Linking of Linear Poly(ethylene imine). Improving the Atom Economy of Polymer-Supported Chemistry. <i>Angewandte Chemie - International Edition</i> , 2002, 41, 3313-3313. | 7.2 | 0 |
| 128 | Epitope-mapping of transglutaminase with parallel label-free optical detection. <i>Biosensors and Bioelectronics</i> , 2002, 17, 937-944. | 5.3 | 26 |
| 129 | Investigation of enzyme activity and inhibition in the interior of novel solid supports. , 2002, , 14-20. | | 0 |
| 130 | SPOCC resins: Polar and chemically inert resins for organic synthesis and library enzyme assays. , 2002, , 176-178. | | 0 |
| 131 | Î±-Ketocarbonyl Peptides: A General Approach to Reactive Resin-Bound Intermediates in the Synthesis of Peptide Isosteres for Protease Inhibitor Screening on Solid Support. <i>Journal of the American Chemical Society</i> , 2001, 123, 2176-2181. | 6.6 | 65 |
| 132 | Spatially Resolved Single Bead Analysis: Homogeneity, Diffusion, and Adsorption in Cross-Linked Polystyrene. <i>Chemistry - A European Journal</i> , 2001, 7, 3884-3889. | 1.7 | 42 |
| 133 | Alkylating Polymers: Resin-Released Carbenium Ions as Versatile Reactive Intermediates in Polymer-Assisted Solution-Phase Synthesis. <i>Angewandte Chemie - International Edition</i> , 2001, 40, 381-385. | 7.2 | 52 |
| 134 | Oxoammonium Resins as Metal-Free, Highly Reactive, Versatile Polymeric Oxidation Reagents. <i>Angewandte Chemie - International Edition</i> , 2001, 40, 1436-1439. | 7.2 | 83 |
| 135 | Combinatorial Chemistry in the Commercial Jungle. <i>Angewandte Chemie - International Edition</i> , 2001, 40, 2721-2721. | 7.2 | 1 |
| 136 | Oxidizing Polymers: A Polymer-Supported, Recyclable Hypervalent Iodine(V) Reagent for the Efficient Conversion of Alcohols, Carbonyl Compounds, and Unsaturated Carbamates in Solution J.R. gratefully acknowledges generous support from Prof. M. E. Maier, TÄ¼bingen, the Strukturfonds of the University of TÄ¼bingen, the Fonds der Chemischen Industrie, and the DFG. We thank Graeme Nicholson, Dietmar Schmid, and Daniel Bischoff for analytical support.. <i>Angewandte Chemie - International Edition</i> , 2001, 40, 4395. | 7.2 | 135 |
| 137 | Solid-Supported Triazenes as Alkylating Polymers Employed for the Versatile Derivatization of Peptides. , 2001, , 267-268. | | 0 |
| 138 | Alkylating Polymers: Resin-Released Carbenium Ions as Versatile Reactive Intermediates in Polymer-Assisted Solution-Phase Synthesis J.R. gratefully acknowledges support from Prof. M. E. Maier, TÄ¼bingen, the Strukturfonds of the University of TÄ¼bingen, and Merck KGaA, Darmstadt, Germany. J.S. received a grant from the DFG graduate college "Analytical Chemistry".. <i>Angewandte Chemie - International Edition</i> , 2001, 40, 381-385. | 7.2 | 0 |
| 139 | Oxoammonium Resins as Metal-Free, Highly Reactive, Versatile Polymeric Oxidation Reagents J.R. gratefully acknowledges generous support from Prof. M. E. Maier, TÄ¼bingen, the Strukturfonds of the UniversitÄ¼t TÄ¼bingen, and Merck KGaA, Darmstadt, Germany.. <i>Angewandte Chemie - International Edition</i> , 2001, 40, 1436-1439. | 7.2 | 3 |
| 140 | DRUG DISCOVERY: Integrating Combinatorial Synthesis and Bioassays. <i>Science</i> , 2000, 287, 1947-1948. | 6.0 | 48 |
| 141 | Physical Properties of Poly(ethylene glycol) (PEG)-Based Resins for Combinatorial Solid Phase Organic Chemistry: A Comparison of PEG-Cross-Linked and PEG-Grafted Resins. <i>ACS Combinatorial Science</i> , 2000, 2, 108-119. | 3.3 | 86 |
| 142 | Solid-Phase Synthesis of Peptide Isosteres by Nucleophilic Reactions with N-Terminal Peptide Aldehydes on a Polar Support Tailored for Solid-Phase Organic Chemistry. <i>Chemistry - A European Journal</i> , 1999, 5, 1218-1225. | 1.7 | 30 |
| 143 | SPOCC: A Resin for Solid-Phase Organic Chemistry and Enzymatic Reactions on Solid Phase. <i>Journal of the American Chemical Society</i> , 1999, 121, 5459-5466. | 6.6 | 142 |
| 144 | Solid-Phase Supported Synthesis of the Branched Pentasaccharide Moiety That Occurs in Most Complex Type N-Glycan Chains. <i>Angewandte Chemie - International Edition</i> , 1998, 37, 1241-1245. | 7.2 | 56 |

| # | ARTICLE | IF | CITATIONS |
|-----|---|-----|-----------|
| 145 | Oligosaccharide Synthesis on Controlled-Pore Glass as Solid Phase Material. Synlett, 1998, 1998, 171-173. | 1.0 | 34 |
| 146 | Repetitive Solid Phase Glycosylation on an Alkyl Thiol Polymer Leading to Sugar Oligomers Containing 1,2-trans- and 1,2-cis-Glycosidic Linkages. Journal of Organic Chemistry, 1997, 62, 3650-3653. | 1.7 | 103 |
| 147 | An N-acetylglucosamine containing glycopeptide synthesis and structure assignment. Carbohydrate Research, 1997, 304, 21-28. | 1.1 | 12 |
| 148 | A new method for the solid phase synthesis of oligosaccharides. Tetrahedron Letters, 1996, 37, 3989-3990. | 0.7 | 97 |
| 149 | Solid-phase synthesis of a glycosylated hexapeptide of human sialophorin, using the trichloroacetimidate method. Carbohydrate Research, 1995, 269, 217-225. | 1.1 | 44 |
| 150 | Oxidizing and Reducing Agents. , 0, , 81-99. | | 0 |
| 151 | Integration of C-Acylation in the Solid-Phase Synthesis of Peptides and Peptidomimetics employing Meldrum's Acid, Phosphorus and Sulfur Ylides. Synthesis, 0, 0, . | 1.2 | 1 |
| 152 | Pentafluorophosphato-Phenylalanines: Amphiphilic Phosphotyrosine Mimetics Displaying Fluorine-Specific Protein Interactions. Angewandte Chemie, 0, , . | 1.6 | 0 |