List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1867699/publications.pdf Version: 2024-02-01

		22153	24982
316	15,310	59	109
papers	citations	h-index	g-index
337	337	337	13194
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Synthesis of a tricyclic hexapeptide –via two consecutive ruthenium-catalyzed macrocyclization steps– with a constrained topology to mimic vancomycin's binding properties toward D-Ala-D-Ala dipeptide. Bioorganic and Medicinal Chemistry Letters, 2022, 73, 128887.	2.2	4
2	Design and Synthesis of HCV-E2 Glycoprotein Epitope Mimics in Molecular Construction of Potential Synthetic Vaccines. Viruses, 2021, 13, 326.	3.3	2
3	TMTHSI, a superior 7-membered ring alkyne containing reagent for strain-promoted azide–alkyne cycloaddition reactions. Chemical Science, 2020, 11, 9011-9016.	7.4	19
4	Improving the aqueous solubility of HCV‣2 glycoprotein epitope mimics by cyclization using POLAR hinges. Journal of Peptide Science, 2020, 26, e3222.	1.4	2
5	Molecular Construction of Sulfonamide Antisense Oligonucleotides. Journal of Organic Chemistry, 2019, 84, 10635-10648.	3.2	4
6	Synthesis of tris-tertiary amine CycloTriVeratrilene (TACTV) derivatives as water soluble pre-organized three aromatic ring containing molecular scaffolds for the construction of protein mimics. Tetrahedron Letters, 2019, 60, 151245.	1.4	2
7	A new perspective on fungal metabolites: identification of bioactive compounds from fungi using zebrafish embryogenesis as read-out. Scientific Reports, 2019, 9, 17546.	3.3	26
8	Design, Synthesis, and Evaluation of a Diazirine Photoaffinity Probe for Ligand-Based Receptor Capture Targeting G Protein–Coupled Receptors. Molecular Pharmacology, 2019, 95, 196-209.	2.3	15
9	Immobilization by Surface Conjugation of Cyclic Peptides for Effective Mimicry of the HCV-Envelope E2 Protein as a Strategy toward Synthetic Vaccines. Bioconjugate Chemistry, 2018, 29, 1091-1101.	3.6	12
10	Tailoring Polyethers for Post-polymerization Functionalization by Cross Metathesis. Organic Letters, 2018, 20, 2253-2256.	4.6	7
11	Targeted Covalent Inhibition of Prolyl Oligopeptidase (POP): Discovery of Sulfonylfluoride Peptidomimetics. Cell Chemical Biology, 2018, 25, 1031-1037.e4.	5.2	36
12	Potent and Highly Selective Inhibitors of the Proteasome Trypsin-like Site by Incorporation of Basic Side Chain Containing Amino Acid Derived Sulfonyl Fluorides. Journal of Medicinal Chemistry, 2018, 61, 5395-5411.	6.4	48
13	Synthesis and cellular penetration properties of new phosphonium based cationic amphiphilic peptides. MedChemComm, 2018, 9, 982-987.	3.4	6
14	Synthetic antibody protein mimics of infliximab by molecular scaffolding on novel CycloTriVeratrilene (CTV) derivatives. Organic and Biomolecular Chemistry, 2018, 16, 5254-5274.	2.8	10
15	An orthogonally protected CycloTriVeratrylene (CTV) as a highly pre-organized molecular scaffold for subsequent ligation of different cyclic peptides towards protein mimics. Bioorganic and Medicinal Chemistry, 2017, 25, 5008-5015.	3.0	8
16	Polar Hinges as Functionalized Conformational Constraints in (Bi)cyclic Peptides. ChemBioChem, 2017, 18, 387-395.	2.6	18
17	Synthesis of bicyclic tripeptides inspired by the ABC-ring system of vancomycin through ruthenium-based cyclization chemistries. Tetrahedron Letters, 2017, 58, 4542-4546.	1.4	12
18	Potential peptidic proteasome inhibitors by incorporation of an electrophilic trap based on amino acid derived α-substituted sulfonyl fluorides. Bioorganic and Medicinal Chemistry, 2017, 25, 5055-5063.	3.0	16

#	Article	IF	CITATIONS
19	Triple-targeting Gram-negative selective antimicrobial peptides capable of disrupting the cell membrane and lipid A biosynthesis. RSC Advances, 2016, 6, 65418-65421.	3.6	8
20	Proteasome inhibition by new dual warhead containing peptido vinyl sulfonyl fluorides. Bioorganic and Medicinal Chemistry, 2016, 24, 3429-3435.	3.0	39
21	Peptide Microarrays for Real-Time Kinetic Profiling of Tyrosine Phosphatase Activity of Recombinant Phosphatases and Phosphatases in Lysates of Cells or Tissue Samples. Methods in Molecular Biology, 2016, 1447, 67-78.	0.9	2
22	Highly potent antimicrobial peptide derivatives of bovine cateslytin. RSC Advances, 2016, 6, 94840-94844.	3.6	5
23	Molecular construction of HIV-gp120 discontinuous epitope mimics by assembly of cyclic peptides on an orthogonal alkyne functionalized TAC-scaffold. Organic and Biomolecular Chemistry, 2016, 14, 701-710.	2.8	16
24	The presence of C/EBPα and its degradation are both required for TRIB2-mediated leukaemia. Oncogene, 2016, 35, 5272-5281.	5.9	25
25	Synthesis of pyrazole containing α-amino acids via a highly regioselective condensation/aza-Michael reaction of β-aryl α,β-unsaturated ketones. Organic and Biomolecular Chemistry, 2015, 13, 4514-4523.	2.8	28
26	Synthesis of nisin AB dicarba analogs using ring-closing metathesis: influence of sp <sup>3</sup> versus sp <sup>2</sup> hybridization of the α-carbon atom of residues dehydrobutyrine-2 and dehydroalanine-5 on the lipid II binding affinity. Organic and Biomolecular Chemistry, 2015, 13, 5997-6009.	2.8	13
27	A Proteinaceous Fraction of Wheat Bran May Interfere in the Attachment of Enterotoxigenic E. Coli K88 (F4+) to Porcine Epithelial Cells. PLoS ONE, 2014, 9, e104258.	2.5	10
28	Convenient Stereoselective Synthesis of Substituted Ureido Glycosides Using Stable 4-Chlorophenylcarbamates without the Requirement of Lewis Acids. Synlett, 2014, 25, 205-208.	1.8	0
29	pH-controlled aggregation polymorphism of amyloidogenic Aβ(16–22): Insights for obtaining peptide tapes and peptide nanotubes, as function of the N-terminal capping moiety. European Journal of Medicinal Chemistry, 2014, 88, 55-65.	5.5	8
30	A versatile spectrophotometric protein tyrosine phosphatase assay based on 3-nitrophosphotyrosine containing substrates. Analytical Biochemistry, 2014, 448, 9-13.	2.4	9
31	New properties of wheat bran: antiâ€biofilm activity and interference with bacteria quorumâ€sensing systems. Environmental Microbiology, 2014, 16, 1346-1353.	3.8	24
32	Versatile convergent synthesis of a three peptide loop containing protein mimic of whooping cough pertactin by successive Cu(I)â€catalyzed azide alkyne cycloaddition on an orthogonal alkyne functionalized TACâ€scaffold. Journal of Peptide Science, 2014, 20, 235-239.	1.4	26
33	Synthesis of the TACO Scaffold as a New Selectively Deprotectable Conformationally Restricted Triazacyclophane Based Scaffold. Organic Letters, 2014, 16, 3106-3109.	4.6	2
34	Efficient Synthesis of Protein Mimics by Sequential Native Chemical Ligation. Organic Letters, 2014, 16, 2138-2141.	4.6	9
35	Scaffolded multiple cyclic peptide libraries for protein mimics by native chemical ligation. Organic and Biomolecular Chemistry, 2014, 12, 4471-4478.	2.8	16
36	Characterization and Activity of an Immobilized Antimicrobial Peptide Containing Bactericidal PEG-Hydrogel. Biomacromolecules, 2014, 15, 3390-3395.	5.4	57

#	Article	IF	CITATIONS
37	Semi-synthesis of biologically active nisin hybrids composed of the native lanthionine ABC-fragment and a cross-stapled synthetic DE-fragment. Bioorganic and Medicinal Chemistry, 2014, 22, 5345-5353.	3.0	17
38	Convenient Preparation of Bactericidal Hydrogels by Covalent Attachment of Stabilized Antimicrobial Peptides Using Thiol–ene Click Chemistry. ACS Macro Letters, 2014, 3, 477-480.	4.8	64
39	Bicycling into cells. Nature Chemistry, 2014, 6, 855-857.	13.6	12
40	Selective Inhibition of the Immunoproteasome by Ligandâ€Induced Crosslinking of the Active Site. Angewandte Chemie - International Edition, 2014, 53, 11969-11973.	13.8	71
41	Synthesis and evaluation of linear CuAAC-oligomerized antifreeze neo-glycopeptides. MedChemComm, 2014, 5, 1159-1165.	3.4	14
42	A conformationally constrained fused tricyclic nisin AB-ring system mimic toward an improved pyrophosphate binder of lipid II. Tetrahedron, 2014, 70, 7691-7699.	1.9	5
43	Azide–alkyne cycloaddition affording enzymatically tunable bisubstrate based inhibitors of histone acetyltransferase PCAF. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 113-116.	2.2	0
44	Expedient synthesis of a novel asymmetric selectively deprotectable derivative of the ATAC scaffold. Tetrahedron, 2014, 70, 4002-4007.	1.9	6
45	Improving the biological activity of the antimicrobial peptide anoplin by membrane anchoring through a lipophilic amino acid derivative. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 3749-3752.	2.2	27
46	Unusual binding of Grb2 protein to a bivalent polyproline-ligand immobilized on a SPR sensor: Intermolecular bivalent binding. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2013, 1834, 524-535.	2.3	1
47	Synthesis, Antimicrobial Activity, and Membrane Permeabilizing Properties of C-Terminally Modified Nisin Conjugates Accessed by CuAAC. Bioconjugate Chemistry, 2013, 24, 2058-2066.	3.6	25
48	Synthesis and structural characterization of the individual diastereoisomers of a cross-stapled alkene-bridged nisin DE-ring mimic. Organic and Biomolecular Chemistry, 2013, 11, 7486.	2.8	13
49	Scalable purification of the lantibiotic nisin and isolation of chemical/enzymatic cleavage fragments suitable for semiâ€synthesis. Journal of Peptide Science, 2013, 19, 692-699.	1.4	30
50	Synthesis of 1,5-triazole bridged vancomycin CDE-ring bicyclic mimics using RuAAC macrocyclization. Chemical Communications, 2013, 49, 4498.	4.1	35
51	Scaffold optimization in discontinuous epitope containing protein mimics of gp120 using smart libraries. Organic and Biomolecular Chemistry, 2013, 11, 2676.	2.8	16
52	Cell-Penetrating Bisubstrate-Based Protein Kinase C Inhibitors. ACS Chemical Biology, 2013, 8, 1479-1487.	3.4	15
53	Enzymatic Fragment Condensation of Side Chainâ€Protected Peptides using Subtilisin A in Anhydrous Organic Solvents: A General Strategy for Industrial Peptide Synthesis. Advanced Synthesis and Catalysis, 2013, 355, 287-293.	4.3	9
54	Realâ€Time Monitoring of the Dephosphorylating Activity of Protein Tyrosine Phosphatases Using Microarrays with 3â€Nitrophosphotyrosine Substrates. ChemPlusChem, 2013, 78, 1349-1357.	2.8	6

#	Article	IF	CITATIONS
55	Broad-Spectrum Antimalarial Activity of Peptido Sulfonyl Fluorides, a New Class of Proteasome Inhibitors. Antimicrobial Agents and Chemotherapy, 2013, 57, 3576-3584.	3.2	24
56	Activity-based probes for rhomboid proteases discovered in a mass spectrometry-based assay. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 2472-2477.	7.1	60
57	CHAPTER 10. Chemical Approaches for Localization, Characterization and Mimicry of Peptide Epitopes. RSC Drug Discovery Series, 2013, , 263-284.	0.3	2
58	Synthesis of Cyclic Peptides Containing a Thioester Handle for Native Chemical Ligation. Journal of Organic Chemistry, 2012, 77, 10058-10064.	3.2	17
59	Triazacyclophane (TAC)-scaffolded histidine and aspartic acid residues as mimics of non-heme metalloenzyme active sites. Organic and Biomolecular Chemistry, 2012, 10, 1088-1092.	2.8	1
60	Mutual influence of backbone proline substitution and lipophilic tail character on the biological activity of simplified analogues of caspofungin. Organic and Biomolecular Chemistry, 2012, 10, 7491.	2.8	10
61	Peptido Sulfonyl Fluorides as New Powerful Proteasome Inhibitors. Journal of Medicinal Chemistry, 2012, 55, 10995-11003.	6.4	67
62	Directed Modulation of Protein Kinaseâ€C Isozyme Selectivity with Bisubstrateâ€Based Inhibitors. ChemMedChem, 2012, 7, 2113-2121.	3.2	11
63	A combinatorial approach toward smart libraries of discontinuous epitopes of HIV gp120 on a TAC synthetic scaffold. Chemical Communications, 2012, 48, 10007.	4.1	22
64	Thermosensitive Peptide-Hybrid ABC Block Copolymers Obtained by ATRP: Synthesis, Self-Assembly, and Enzymatic Degradation. Macromolecules, 2012, 45, 842-851.	4.8	32
65	Trivalent Ultrashort Lipopeptides are Potent pH Dependent Antifungal Agents. Journal of Medicinal Chemistry, 2012, 55, 1296-1302.	6.4	29
66	Imprinted Polymers Displaying High Affinity for Sulfated Protein Fragments. Angewandte Chemie - International Edition, 2012, 51, 8326-8329.	13.8	59
67	Glucocorticoidâ€Loaded Core rossâ€Linked Polymeric Micelles with Tailorable Release Kinetics for Targeted Therapy of Rheumatoid Arthritis. Angewandte Chemie - International Edition, 2012, 51, 7254-7258.	13.8	102
68	Chemicalâ€Biological Exploration of the Limits of the Ras De―and Repalmitoylating Machinery. ChemBioChem, 2012, 13, 1017-1023.	2.6	22
69	Enzymatic C-terminal amidation of amino acids and peptides. Tetrahedron Letters, 2012, 53, 3777-3779.	1.4	20
70	Probing the Lipid-Protein Interface Using Model Transmembrane Peptides with a Covalently Linked Acyl Chain. Biophysical Journal, 2011, 101, 1959-1967.	0.5	9
71	Cu(I)- and Ru(II)-Mediated "Click―Cyclization of Tripeptides Toward Vancomycin-Inspired Mimics. Organic Letters, 2011, 13, 3438-3441.	4.6	40
72	Cell permeable ITAM constructs for the modulation of mediator release in mast cells. Organic and Biomolecular Chemistry, 2011, 9, 820-833.	2.8	3

#	Article	IF	CITATIONS
73	A convenient [2+2] cycloaddition–cycloreversion reaction for the synthesis of 1,1-dicyanobuta-1,3-diene-scaffolded peptides as new imaging chromophores. Tetrahedron Letters, 2011, 52, 6963-6967.	1.4	6
74	Enzymatic synthesis of activated esters and their subsequent use in enzyme-based peptide synthesis. Journal of Molecular Catalysis B: Enzymatic, 2011, 71, 79-84.	1.8	27
75	Synthesis and evaluation of novel macrocyclic antifungal peptides. Bioorganic and Medicinal Chemistry, 2011, 19, 6505-6517.	3.0	15
76	A peptide mimic of the chemotaxis inhibitory protein of Staphylococcus aureus: towards the development of novel anti-inflammatory compounds. Amino Acids, 2011, 40, 731-740.	2.7	8
77	Synthesis and biological evaluation of novel irreversible serine protease inhibitors using amino acid based sulfonyl fluorides as an electrophilic trap. Bioorganic and Medicinal Chemistry, 2011, 19, 2397-2406.	3.0	43
78	Fully Enzymatic <i>N</i> → <i>C</i> â€Directed Peptide Synthesis Using <i>C</i> â€Terminal Peptide αâ€Carboxamide to Ester Interconversion. Advanced Synthesis and Catalysis, 2011, 353, 1039-1044.	4.3	15
79	Enantioselective Cu <sup>II</sup> atalyzed Diels–Alder and Michael Addition Reactions in Water Using Bioâ€Inspired Triazacyclophaneâ€Based Ligands. European Journal of Organic Chemistry, 2011, 2011, 1714-1720.	2.4	16
80	Spacer Effects on in vivo Properties of DOTAâ€Conjugated Dimeric [Tyr3]Octreotate Peptides Synthesized by a "Cu <sup>I</sup> â€Click―and "Sulfoâ€Click―Ligation Method. ChemBioChem, 2011 750-760.	, 1 <b>2,</b> 6	17
81	Peptides and Proteins as a Continuing Exciting Source of Inspiration for Peptidomimetics. ChemBioChem, 2011, 12, 1626-1653.	2.6	144
82	A Simple Large-Scale Synthesis of Cbz-Protected Taurylsulfonyl Azide. Synlett, 2011, 2011, 2228-2230.	1.8	0
83	The role of the disulfide bond in the interaction of islet amyloid polypeptide with membranes. European Biophysics Journal, 2010, 39, 1359-1364.	2.2	18
84	Towards the synthesis of sulfonamide-based RNA mimetics. Tetrahedron: Asymmetry, 2010, 21, 469-475.	1.8	3
85	Direct Structural Comparison of a Rigid Cyclic Peptidic Scaffold Using Crystallography and NMR in Strained PH Polymer Gels. European Journal of Organic Chemistry, 2010, 2010, 4501-4507.	2.4	5
86	Fully Enzymatic Peptide Synthesis using <i>C</i> â€Terminal <i>tert</i> â€Butyl Ester Interconversion. Advanced Synthesis and Catalysis, 2010, 352, 2399-2404.	4.3	25
87	Rapid Screening of Lectins for Multivalency Effects with a Glycodendrimer Microarray. ChemBioChem, 2010, 11, 1896-1904.	2.6	65
88	Synthesis and characterization of tailorable biodegradable thermoresponsive methacryloylamide polymers based on l-serine and l-threonine alkyl esters. Polymer, 2010, 51, 2479-2485.	3.8	11
89	Nanomolar affinity, iminosugar-based chemical probes for specific labeling of lysosomal glucocerebrosidase. Bioorganic and Medicinal Chemistry, 2010, 18, 267-273.	3.0	24
90	CHIPS binds to the phosphorylated N-terminus of the C5a-receptor. Bioorganic and Medicinal Chemistry Letters, 2010, 20, 3338-3340.	2.2	7

#	Article	IF	CITATIONS
91	â€~Sulfoâ€click' for ligation as well as for siteâ€specific conjugation with peptides, fluorophores, and metal chelators. Journal of Peptide Science, 2010, 16, 1-5.	1.4	48
92	Thermodynamics of phosphotyrosine peptide–peptoid hybrids binding to the p56 <sup>lck</sup> SH2 domain. Journal of Peptide Science, 2010, 16, 322-328.	1.4	1
93	Synthesis of DOTA-Conjugated Multimeric [Tyr <sup>3</sup> ]Octreotide Peptides via a Combination of Cu(I)-Catalyzed "Click―Cycloaddition and Thio Acid/Sulfonyl Azide "Sulfo-Click―Amidation and Their in Vivo Evaluation. Journal of Medicinal Chemistry, 2010, 53, 3944-3953.	6.4	77
94	Synthesis and Characterization of Enzymatically Biodegradable PEG and Peptide-Based Hydrogels Prepared by Click Chemistry. Biomacromolecules, 2010, 11, 1608-1614.	5.4	112
95	The N-terminal fragment of human islet amyloid polypeptide is non-fibrillogenic in the presence of membranes and does not cause leakage of bilayers of physiologically relevant lipid composition. Biochimica Et Biophysica Acta - Biomembranes, 2010, 1798, 1805-1811.	2.6	26
96	Detection of pathogenic Streptococcus suis bacteria using magnetic glycoparticles. Organic and Biomolecular Chemistry, 2010, 8, 2425.	2.8	46
97	Preparation of novel alkylated arginine derivatives suitable for click-cycloaddition chemistry and their incorporation into pseudosubstrate- and bisubstrate-based kinase inhibitors. Organic and Biomolecular Chemistry, 2010, 8, 1629.	2.8	12
98	Nanostructure Determines Antifungal Activity of De Novo Designed pH Dependent Histidine Containing Ultra-Short Lipopeptides. Biophysical Journal, 2010, 98, 278a-279a.	0.5	1
99	Structure of the Tyrosine-sulfated C5a Receptor N Terminus in Complex with Chemotaxis Inhibitory Protein of Staphylococcus aureus. Journal of Biological Chemistry, 2009, 284, 12363-12372.	3.4	40
100	Versatile Selective α-Carboxylic Acid Esterification of N-Protected Amino Acids and Peptides by Alcalase. Synthesis, 2009, 2009, 809-814.	2.3	5
101	The Filament-specific Rep1-1 Repellent of the Phytopathogen Ustilago maydis Forms Functional Surface-active Amyloid-like Fibrils. Journal of Biological Chemistry, 2009, 284, 9153-9159.	3.4	38
102	The State of the Art of Chemical Biology. ChemBioChem, 2009, 10, 16-29.	2.6	41
103	Synthesis and Evaluation of New Thiodigalactosideâ€Based Chemical Probes to Label Galectinâ€3. ChemBioChem, 2009, 10, 1724-1733.	2.6	36
104	Development of Selective Bisubstrateâ€Based Inhibitors Against Protein Kinase C (PKC) Isozymes By Using Dynamic Peptide Microarrays. ChemBioChem, 2009, 10, 2042-2051.	2.6	33
105	Synthesis of β-aminoethanesulfonyl fluorides or 2-substituted taurine sulfonyl fluorides as potential protease inhibitors. Tetrahedron Letters, 2009, 50, 3391-3393.	1.4	41
106	Switching between low and high affinity for the Syk tandem SH2 domain by irradiation of azobenzene containing ITAM peptidomimetics. Journal of Peptide Science, 2009, 15, 685-691.	1.4	20
107	A versatile and selective chemo-enzymatic synthesis of β-protected aspartic and γ-protected glutamic acid derivatives. Tetrahedron Letters, 2009, 50, 2719-2721.	1.4	15
108	Photocrosslinking and Click Chemistry Enable the Specific Detection of Proteins Interacting with Phospholipids at the Membrane Interface. Chemistry and Biology, 2009, 16, 3-14.	6.0	83

#	Article	IF	CITATIONS
109	Potential scorpionate antibiotics: Targeted hydrolysis of lipid II containing model membranes by vancomycin–TACzyme conjugates and modulation of their antibacterial activity by Zn-ions. Bioorganic and Medicinal Chemistry Letters, 2009, 19, 3721-3724.	2.2	4
110	A general sequence independent solid phase method for the site specific synthesis of multiple sulfated-tyrosine containing peptides. Chemical Communications, 2009, , 2999.	4.1	23
111	Enzymatic Synthesis of <i>C</i> -Terminal Arylamides of Amino Acids and Peptides. Journal of Organic Chemistry, 2009, 74, 5145-5150.	3.2	35
112	Microwave-assisted click polymerization for the synthesis of Aβ(16–22) cyclic oligomers and their self-assembly into polymorphous aggregates. Organic and Biomolecular Chemistry, 2009, 7, 4517.	2.8	18
113	Synthesis and Applications of Biomedical and Pharmaceutical Polymers via Click Chemistry Methodologies. Bioconjugate Chemistry, 2009, 20, 2001-2016.	3.6	266
114	Versatile Conjugation of Octreotide to Dendrimers by Cycloaddition ("Clickâ€ <del>)</del> Chemistry to Yield High-Affinity Multivalent Cyclic Peptide Dendrimers. Bioconjugate Chemistry, 2009, 20, 1323-1331.	3.6	34
115	A general approach for the non-stop solid phase synthesis of TAC-scaffolded loops towards protein mimics containing discontinuous epitopes. Chemical Communications, 2009, , 821-823.	4.1	22
116	ITAM-derived phosphopeptide-containing dendrimers as multivalent ligands for Syk tandem SH2 domain. Organic and Biomolecular Chemistry, 2009, 7, 4088.	2.8	13
117	pH Controlled Aggregation Morphology of Aβ(16–22): Formation of Peptide Nanotubes, Helical Tapes and Amyloid Fibrils. Advances in Experimental Medicine and Biology, 2009, 611, 239-240.	1.6	4
118	Alkene/Alkane-Bridged Mimics of the Lantibiotic Nisin: Toward Novel Peptide-Based Antibiotics. Advances in Experimental Medicine and Biology, 2009, 611, 533-534.	1.6	1
119	Peptidomimetic Ligands for the Tandem SH2 Domain of the Syk Protein Involved in Signal Transduction. Advances in Experimental Medicine and Biology, 2009, 611, 81-82.	1.6	Ο
120	Multivalent Carbohydrate Recognition on a Glycodendrimerâ€Functionalized Flowâ€Through Chip. ChemBioChem, 2008, 9, 1836-1844.	2.6	83
121	A photoswitchable ITAM peptidomimetic: Synthesis and real time surface plasmon resonance (SPR) analysis of the effects of cis–trans isomerization on binding. Bioorganic and Medicinal Chemistry, 2008, 16, 1393-1399.	3.0	19
122	Delayed fibril formation of amylin(20–29) by incorporation of alkene dipeptidosulfonamide isosteres obtained by solid phase olefin cross metathesis. Bioorganic and Medicinal Chemistry Letters, 2008, 18, 78-84.	2.2	22
123	Mirror image supramolecular helical tapes formed by the enantiomeric-depsipeptide derivatives of the amyloidogenic peptide amylin(20–29). Tetrahedron Letters, 2008, 49, 987-991.	1.4	5
124	Structural Insight into the Recognition of the H3K4me3 Mark by the TFIID Subunit TAF3. Structure, 2008, 16, 1245-1256.	3.3	123
125	Influence of Trifluoroethanol on Membrane Interfacial Anchoring Interactions of Transmembrane α-Helical Peptides. Biophysical Journal, 2008, 94, 1315-1325.	0.5	22
126	The Vancomycinâ~'Nisin(1â~'12) Hybrid Restores Activity against Vancomycin Resistant Enterococci. Biochemistry, 2008, 47, 12661-12663.	2.5	82

#	Article	IF	CITATIONS
127	Synthesis of multivalent Streptococcus suis adhesion inhibitors by enzymatic cleavage of polygalacturonic acid and †click' conjugation. Organic and Biomolecular Chemistry, 2008, 6, 1425.	2.8	33
128	TAC-Scaffolded Tripeptides as Artificial Hydrolytic Receptors: A Combinatorial Approach Toward Esterase Mimics. ACS Combinatorial Science, 2008, 10, 814-824.	3.3	26
129	Preparation of <i>N</i> <sup>G</sup> -Substituted <scp>l</scp> -Arginine Analogues Suitable for Solid Phase Peptide Synthesis. Journal of Organic Chemistry, 2008, 73, 7849-7851.	3.2	30
130	Synthesis and Characterization of Biodegradable Peptide-Based Polymers Prepared by Microwave-Assisted Click Chemistry. Biomacromolecules, 2008, 9, 2834-2843.	5.4	69
131	Multivalent Presentation Strategies in Novel Inhibitors of Bacterial (Toxin) Adhesion and Synthetic Vaccines. Anti-Infective Agents in Medicinal Chemistry, 2008, 7, 193-200.	0.6	7
132	A novel strategy to mimic discontinuous protective epitopes using a synthetic scaffold. Vaccine, 2007, 25, 6807-6817.	3.8	25
133	Effects of linker variation on the in vitro and in vivo characteristics of an 1111n-labeled RGD peptide. Nuclear Medicine and Biology, 2007, 34, 29-35.	0.6	76
134	β-Sheet Structured β-Amyloid(1-40) Perturbs Phosphatidylcholine Model Membranes. Journal of Molecular Biology, 2007, 368, 982-997.	4.2	75
135	Resin-Bound Sulfonyl Azides:  Efficient Loading and Activation Strategy for the Preparation of the N-Acyl Sulfonamide Linker. Journal of Organic Chemistry, 2007, 72, 4574-4577.	3.2	41
136	Step-wise and pre-organization induced synthesis of a crossed alkene-bridged nisin Z DE-ring mimic by ring-closing metathesis. Organic and Biomolecular Chemistry, 2007, 5, 924.	2.8	31
137	Synthesis of DOTA-conjugated multivalent cyclic-RGD peptide dendrimers via 1,3-dipolar cycloaddition and their biological evaluation: implications for tumor targeting and tumor imaging purposes. Organic and Biomolecular Chemistry, 2007, 5, 935.	2.8	180
138	Strong inhibition of cholera toxin binding by galactose dendrimers. Chemical Communications, 2007, , 5043.	4.1	75
139	Scaffolded amino acids as a close structural mimic of type-3 copper binding sites. Chemical Communications, 2007, , 4895.	4.1	18
140	Synthesis of Peptide-Based Polymers by Microwave-Assisted Cycloaddition Backbone Polymerization. Biomacromolecules, 2007, 8, 327-330.	5.4	48
141	Enhanced Membrane Pore Formation by Multimeric/Oligomeric Antimicrobial Peptides. Biochemistry, 2007, 46, 13437-13442.	2.5	74
142	Synthesis and Evaluation of TACâ€Based Inhibitors of Papain as Mimics of Cystatin B. ChemBioChem, 2007, 8, 1950-1956.	2.6	10
143	Synthesis of Bicyclic Alkene…Alkaneâ€Bridged Nisin Mimics by Ring losing Metathesis and their Biochemical Evaluation as Lipid II Binders: toward the Design of Potential Novel Antibiotics. ChemBioChem, 2007, 8, 1540-1554.	2.6	48
144	Strong Inhibition of Cholera Toxin by Multivalent GM1 Derivatives. ChemBioChem, 2007, 8, 1500-1503.	2.6	101

#	Article	IF	CITATIONS
145	αvβ3 Integrin-targeting of intraperitoneally growing tumors with a radiolabeled RGD peptide. International Journal of Cancer, 2007, 120, 605-610.	5.1	61
146	Solid-phase carbohydrate synthesis via on-bead protecting group chemistry. Tetrahedron, 2007, 63, 4290-4296.	1.9	11
147	Synthesis and evaluation of chloromethyl sulfoxides as a new class of selective irreversible cysteine protease inhibitors. Bioorganic and Medicinal Chemistry, 2007, 15, 6985-6993.	3.0	3
148	Transformation of the amyloidogenic peptide amylin(20–29) into its corresponding peptoid and retropeptoid: Access to both an amyloid inhibitor and template for self-assembled supramolecular tapes. Bioorganic and Medicinal Chemistry Letters, 2007, 17, 1837-1842.	2.2	33
149	Application of the 1,3â€Dipolar Cycloaddition Reaction in Chemical Biology: Approaches Toward Multivalent Carbohydrates and Peptides and Peptideâ€Based Polymers. QSAR and Combinatorial Science, 2007, 26, 1181-1190.	1.4	65
150	Backboneâ€modified amylin derivatives: implications for amyloid inhibitor design and as template for selfâ€assembling bionanomaterials. Journal of Peptide Science, 2007, 13, 709-716.	1.4	4
151	Improved targeting of the αvβ3 integrin by multimerisation of RGD peptides. European Journal of Nuclear Medicine and Molecular Imaging, 2007, 34, 267-273.	6.4	195
152	Detection of galectin-3 by novel peptidic photoprobes. Bioorganic and Medicinal Chemistry Letters, 2007, 17, 376-378.	2.2	19
153	Synthesis of Alkyne-Bridged Cyclic Tripeptides toward Constrained Mimics of Vancomycin. Journal of Organic Chemistry, 2006, 71, 1817-1824.	3.2	30
154	Synthesis and biological evaluation of potent αvβ3-integrin receptor antagonists. Nuclear Medicine and Biology, 2006, 33, 953-961.	0.6	45
155	A new chemical probe for the detection of the cancer-linked galectin-3. Organic and Biomolecular Chemistry, 2006, 4, 4387.	2.8	52
156	Synthesis and structural investigations of N-alkylated β-peptidosulfonamide–peptide hybrids of the amyloidogenic amylin(20–29) sequence: implications of supramolecular folding for the design of peptide-based bionanomaterials. Organic and Biomolecular Chemistry, 2006, 4, 3587-3597.	2.8	27
157	Investigating the Dynamic Nature of the Interactions between Nuclear Proteins and Histones upon DNA Damage Using an Immobilized Peptide Chemical Proteomics Approach. Journal of Proteome Research, 2006, 5, 2380-2388.	3.7	6
158	Islet Amyloid Polypeptide Inserts into Phospholipid Monolayers as Monomer. Journal of Molecular Biology, 2006, 356, 783-789.	4.2	170
159	Interference with Protein-Protein Interactions Involved in Protease Inhibitor Complex Formation. , 2006, , 212-213.		0
160	Peptide-Derived (Sulfonyl)Azides as Versatile Synthons in Chemoselective Bioconjugations. , 2006, , 198-199.		0
161	Synthesis of Cyclic Peptides via Transition Metal Catalyzed C-C Bond Formation. , 2006, , 50-52.		0
162	Selective enrichment of Ser-/Thr-phosphorylated peptides in the presence of Ser-/Thr-glycosylated peptides. Proteomics, 2006, 6, 6394-6399.	2.2	19

#	Article	IF	CITATIONS
163	Alkene- and alkyne-bridged mimics of nisin as potential peptide-based antibiotics. Journal of Molecular Catalysis A, 2006, 254, 68-77.	4.8	31
164	Tannic acid mimicking dendrimers as small intestine submucosa stabilizing nanomordants. Biomaterials, 2006, 27, 745-751.	11.4	12
165	Self-Assembly of Amylin(20–29) Amide-Bond Derivatives into Helical Ribbons and Peptide Nanotubes rather than Fibrils. Chemistry - A European Journal, 2006, 12, 3714-3725.	3.3	41
166	Synthesis and Applications of $\hat{l}^2$ -Aminoethanesulfonyl Azides. Synthesis, 2006, 2006, 455-460.	2.3	8
167	A New Efficient Post-Assembly Strategy for the Synthesis of Sulfated Peptides. , 2006, , 158-159.		0
168	Binding of a Diphosphorylated-ITAM peptide to spleen tyrosine kinase (Syk) induces distal conformational changes: A hydrogen exchange mass spectrometry study. Journal of the American Society for Mass Spectrometry, 2005, 16, 1039-1051.	2.8	19
169	Synthesis of a novel potent cyclic peptide MC4-ligand by ring-closing metathesis. Bioorganic and Medicinal Chemistry, 2005, 13, 4221-4227.	3.0	20
170	Synthesis of cyclic peptidosulfonamides as scaffolds for MC4 pharmacophoric groups. Bioorganic and Medicinal Chemistry Letters, 2005, 15, 287-290.	2.2	14
171	Novel multivalent mannose compounds and their inhibition of the adhesion of type 1 fimbriated uropathogenic E. coli. Tetrahedron: Asymmetry, 2005, 16, 361-372.	1.8	62
172	The synthesis of amides and dipeptides from unprotected amino acids by a simultaneous protection–activation strategy using boron trifluoride diethyl etherate. Tetrahedron Letters, 2005, 46, 653-656.	1.4	20
173	A convenient solid phase synthesis of S-palmitoyl transmembrane peptides. Tetrahedron Letters, 2005, 46, 3341-3345.	1.4	22
174	Brilliant lipids. Nature Methods, 2005, 2, 14-15.	19.0	19
175	Synthesis of Novel Dendrimeric Systems Containing NLO Ligands. European Journal of Organic Chemistry, 2005, 2005, 487-495.	2.4	18
176	High-Yielding Microwave-Assisted Synthesis of Triazole-Linked Glycodendrimers by Copper-Catalyzed [3+2] Cycloaddition. European Journal of Organic Chemistry, 2005, 2005, 3182-3185.	2.4	99
177	A New Chemical Probe for Proteomics of Carbohydrate-Binding Proteins. ChemBioChem, 2005, 6, 291-295.	2.6	63
178	Protein Flexibility and Ligand Rigidity: A Thermodynamic and Kinetic Study of ITAM-Based Ligand Binding to Syk Tandem SH2. ChemBioChem, 2005, 6, 2261-2270.	2.6	32
179	Development of a Novel Chemical Probe for the Selective Enrichment of Phosphorylated Serine- and Threonine-Containing Peptides. ChemBioChem, 2005, 6, 2271-2280.	2.6	64
180	Peptide Transformation Leading to Peptide-Peptidosulfonamide Hybrids and Oligo Peptidosulfonamides. ChemInform, 2005, 36, no.	0.0	0

#	Article	IF	CITATIONS
181	A facile synthesis of the GalNAcβ1→4Gal target sequence of respiratory pathogens. Carbohydrate Research, 2005, 340, 2436-2442.	2.3	10
182	Peptoidâ^'Peptide Hybrids as Potent Novel Melanocortin Receptor Ligands. Journal of Medicinal Chemistry, 2005, 48, 4224-4230.	6.4	30
183	Pre-organization induced synthesis of a crossed alkene-bridged nisin Z DE-ring mimic by ring-closing metathesis. Chemical Communications, 2005, , 192.	4.1	33
184	Highly Efficient Coupling of β-Substituted Aminoethane Sulfonyl Azides with Thio Acids, toward a New Chemical Ligation Reaction. Organic Letters, 2005, 7, 1125-1128.	4.6	91
185	Synthesis of Leukotriene B4 Antagonists Labeled with In-111 or Tc-99m to Image Infectious and Inflammatory Foci. Journal of Medicinal Chemistry, 2005, 48, 6442-6453.	6.4	8
186	Ring-Closing Alkyne Metathesis Approach toward the Synthesis of Alkyne Mimics of Thioether A-, B-, C-, and DE-Ring Systems of the Lantibiotic Nisin Z. Organic Letters, 2005, 7, 2961-2964.	4.6	62
187	Influence of Flanking Residues on Tilt and Rotation Angles of Transmembrane Peptides in Lipid Bilayers. A Solid-State2H NMR Study. Biochemistry, 2005, 44, 1004-1012.	2.5	95
188	The Structure of the C5a Receptor-blocking Domain of Chemotaxis Inhibitory Protein of Staphylococcus aureus is Related to a Group of Immune Evasive Molecules. Journal of Molecular Biology, 2005, 353, 859-872.	4.2	57
189	Possibilities and limitations in the rational design of modified peptides for T cell mediated immunotherapy. Molecular Immunology, 2005, 42, 365-373.	2.2	18
190	Limited plasticity in T cell recognition of modified T cell receptor contact residues in MHC class II bound peptides. Molecular Immunology, 2005, 42, 355-364.	2.2	17
191	Efficient microwave-assisted synthesis of multivalent dendrimeric peptides using cycloaddition reaction (click) chemistry. Chemical Communications, 2005, , 4581.	4.1	120
192	Surface Plasmon Resonance Thermodynamic and Kinetic Analysis as a Strategic Tool in Drug Design. Distinct Ways for Phosphopeptides to Plug into Src- and Grb2 SH2 Domains. Journal of Medicinal Chemistry, 2005, 48, 753-763.	6.4	42
193	N-Terminal Residues of the Chemotaxis Inhibitory Protein of <i>Staphylococcus aureus</i> Are Essential for Blocking Formylated Peptide Receptor but Not C5a Receptor. Journal of Immunology, 2004, 173, 5704-5711.	0.8	76
194	Accelerating sensory recovery after sciatic nerve crush: non-selective versus melanocortin MC4 receptor-selective peptides. European Journal of Pharmacology, 2004, 495, 145-152.	3.5	2
195	Peptide transformation leading to peptide-peptidosulfonamide hybrids and oligo peptidosulfonamides. Molecular Diversity, 2004, 8, 79-87.	3.9	30
196	Combinatorial solid-phase synthesis and screening of a diverse tripodal triazacyclophane (TAC)-based synthetic receptor library showing a remarkable selectivity towards a d-Ala-d-Ala containing ligand. Tetrahedron, 2004, 60, 8691-8697.	1.9	25
197	Selection of synthetic receptors capable of sensing the difference in binding of d-Ala-d-Ala or d-Ala-d-Lac ligands by screening of a combinatorial CTV-based library. Tetrahedron, 2004, 60, 11145-11157.	1.9	20
198	Synthesis, Screening and Evaluation of a Combined Library of Tweezer- and Tripodal Synthetic Receptors. QSAR and Combinatorial Science, 2004, 23, 546-559.	1.4	16

#	Article	IF	CITATIONS
199	TheαM1 transmembrane segment of the nicotinic acetylcholine receptor interacts strongly with model membranes. Magnetic Resonance in Chemistry, 2004, 42, 148-154.	1.9	19
200	Structure-Activity Studies on the Corticotropin Releasing Factor Antagonist Astressin, leading to a Minimal Sequence necessary for Antagonistic Activity. ChemBioChem, 2004, 5, 340-348.	2.6	18
201	Synthesis of Cyclic Peptidosulfonamides by Ring-Closing Metathesis ChemInform, 2004, 35, no.	0.0	0
202	Synthesis and biological activity of N-terminal lipidated and/or fluorescently labeled conjugates of astressin as corticotropin releasing factor antagonists. Bioorganic and Medicinal Chemistry, 2004, 12, 5099-5106.	3.0	2
203	Microwave-assisted, tin-mediated, regioselective 3-O-alkylation of galactosides. Tetrahedron Letters, 2004, 45, 6685-6687.	1.4	27
204	Synthesis and cholera toxin binding properties of multivalent GM1 mimicsElectronic supplementary information (ESI) available: characterization of the polyvalent compounds ? imide by-products. See http://www.rsc.org/suppdata/ob/b4/b405344c/. Organic and Biomolecular Chemistry, 2004, 2, 2113.	2.8	77
205	Ring-closing metathesis for the synthesis of side chain knotted pentapeptides inspired by vancomycin. Organic and Biomolecular Chemistry, 2004, 2, 2658.	2.8	18
206	Inhibition ofStreptococcussuisAdhesion by Dendritic Galabiose Compounds at Low Nanomolar Concentration. Journal of Medicinal Chemistry, 2004, 47, 6499-6508.	6.4	85
207	Photo-Crosslinking Analysis of Preferential Interactions between a Transmembrane Peptide and Matching Lipids. Biochemistry, 2004, 43, 4482-4489.	2.5	31
208	Synthesis of Cyclic Peptidosulfonamides by Ring-Closing Metathesis. Journal of Organic Chemistry, 2004, 69, 3662-3668.	3.2	41
209	Tilt Angles of Transmembrane Model Peptides in Oriented and Non-Oriented Lipid Bilayers as Determined by 2H Solid-State NMR. Biophysical Journal, 2004, 86, 3709-3721.	0.5	172
210	Probing the Self-Assembly and the Accompanying Structural Changes of Hydrophobin SC3 on a Hydrophobic Surface by Mass Spectrometry. Biophysical Journal, 2004, 87, 1919-1928.	0.5	34
211	The αM1 segment of the nicotinic acetylcholine receptor exhibits conformational flexibility in a membrane environment. Biochimica Et Biophysica Acta - Biomembranes, 2004, 1665, 40-47.	2.6	19
212	Stabilization of peptide guinea pig myelin basic protein 72?85 by N-terminal acetylation?implications for immunological studies. Molecular Immunology, 2004, 40, 943-948.	2.2	14
213	Adhesion Inhibition of F1C-Fimbriated Escherichia coli and Pseudomonas aeruginosa PAK and PAO by Multivalent Carbohydrate Ligands. ChemBioChem, 2003, 4, 1317-1325.	2.6	57
214	Chemoselective coupling of peptide fragments using the Staudinger ligation. Tetrahedron Letters, 2003, 44, 4515-4518.	1.4	53
215	Amino propynyl benzoic acid building block in rigid spacers of divalent ligands binding to the syk SH2 domains with equally high affinity as the natural ligand. Bioorganic and Medicinal Chemistry Letters, 2003, 13, 1241-1244.	2.2	21
216	Structural analysis of high affinity divalent phosphopeptide hybrids of spleen tyrosine kinase. International Journal of Mass Spectrometry, 2003, 228, 879-890.	1.5	5

#	Article	IF	CITATIONS
217	Role of solution conformation and flexibility of short peptide ligands that bind to the p56lck SH2 domain. Bioorganic and Medicinal Chemistry, 2003, 11, 941-949.	3.0	13
218	Approaches to the Solid Phase of a Cyclotriveratrylene Scaffold-Based Tripodal Library as Potential Artificial Receptors. ACS Combinatorial Science, 2003, 5, 794-801.	3.3	16
219	Discovery and in vivo evaluation of new melanocortin-4 receptor-selective peptides. Peptides, 2003, 24, 271-280.	2.4	20
220	Synthesis of novel trivalent amino acid glycoconjugates based on the cyclotriveratrylene (â€~CTV') scaffold. Organic and Biomolecular Chemistry, 2003, 1, 2661-2669.	2.8	32
221	Cyclic phosphopeptides for interference with Grb2 SH2 domain signal transduction prepared by ring-closing metathesis and phosphorylation. Organic and Biomolecular Chemistry, 2003, 1, 3297.	2.8	35
222	Rigidified multivalent lactose molecules and their interactions with mammalian galectins: a route to selective inhibitors. Organic and Biomolecular Chemistry, 2003, 1, 803-810.	2.8	111
223	Synthesis and Binding Studies of Aminothiazoline-Carbohydrate Conjugates. , 2003, , 94.		0
224	Synthesis and Cholera Toxin Binding Properties of a Lactose-2-aminothiazoline Conjugate. Organic Letters, 2002, 4, 1807-1808.	4.6	18
225	Synthesis and Screening of Libraries of Synthetic Tripodal Receptor Molecules with Three Different Amino Acid or Peptide Arms:  Identification of Iron Binders. ACS Combinatorial Science, 2002, 4, 275-284.	3.3	27
226	Lipid Dependence of Membrane Anchoring Properties and Snorkeling Behavior of Aromatic and Charged Residues in Transmembrane Peptidesâ€. Biochemistry, 2002, 41, 7190-7198.	2.5	106
227	The Effects of Hydrophobic Mismatch between Phosphatidylcholine Bilayers and Transmembrane α-Helical Peptides Depend on the Nature of Interfacially Exposed Aromatic and Charged Residuesâ€. Biochemistry, 2002, 41, 8396-8404.	2.5	94
228	Synthesis of Cyclic (α2β)-Tripeptides as Potential Peptide Turn Mimetics. Organic Letters, 2002, 4, 2173-2176.	4.6	19
229	A convenient preparation of several N-linked glycoamino acid building blocks for efficient solid-phase synthesis of glycopeptides. Journal of the Chemical Society, Perkin Transactions 1, 2002, , 1042-1049.	1.3	27
230	Inhibition of Amyloid Fibril Formation of Human Amylin by N-Alkylated Amino Acid and -Hydroxy Acid Residue Containing Peptides. Chemistry - A European Journal, 2002, 8, 4285-4291.	3.3	68
231	Cyclotriveratrylene (CTV) as a New Chiral Triacid Scaffold Capable of Inducing Triple Helix Formation of Collagen Peptides Containing either a Native Sequence or Pro-Hyp-Gly Repeats. Chemistry - A European Journal, 2002, 8, 4613-4621.	3.3	39
232	Replacement of the Intervening Amino Acid Sequence of a Syk-Binding Diphosphopeptide by a Nonpeptide Spacer with Preservation of High Affinity. ChemBioChem, 2002, 3, 238-242.	2.6	33
233	Characterization of a phosphorylated peptide and peptoid and peptoid-peptide hybrids by mass spectrometry. Journal of Mass Spectrometry, 2002, 37, 47-55.	1.6	24
234	An optimized solid phase synthesis strategy? including on-resin lactamization?of astressin, its retro-, inverso-, and retro-inverso isomers as corticotropin releasing factor antagonists. Biopolymers, 2002, 63, 141-149.	2.4	7

#	Article	IF	CITATIONS
235	Synthesis and biological activity of polygalloyl-dendrimers as stable tannic acid mimics. Bioorganic and Medicinal Chemistry Letters, 2002, 12, 1567-1570.	2.2	24
236	Major histocompatibility complex class II binding characteristics of peptoid–peptide hybrids. Bioorganic and Medicinal Chemistry, 2002, 10, 1939-1945.	3.0	29
237	A convenient synthesis of azido peptides by post-assembly diazo transfer on the solid phase applicable to large peptides. Tetrahedron Letters, 2002, 43, 3657-3660.	1.4	30
238	Synthesis of amides from unprotected amino acids by a simultaneous protection–activation strategy using dichlorodialkyl silanes. Tetrahedron Letters, 2002, 43, 9203-9207.	1.4	48
239	Catalytic conversions of diazosugars. Tetrahedron Letters, 2002, 43, 9601-9603.	1.4	16
240	Synthesis and Biological Activity of Polygalloylâ€Dendrimers as Stable Tannic Acid Mimics ChemInform, 2002, 33, 66-66.	0.0	0
241	Solid-Phase Synthesis of Oligourea Peptidomimetics Employing the Fmoc Protection Strategy. Journal of Organic Chemistry, 2001, 66, 8454-8462.	3.2	72
242	A Selectively Deprotectable Triazacyclophane Scaffold for the Construction of Artificial Receptors. Organic Letters, 2001, 3, 3499-3502.	4.6	39
243	Bio-inspired synthetic receptor molecules towards mimicry of vancomycin. Bioorganic and Medicinal Chemistry Letters, 2001, 11, 1521-1525.	2.2	24
244	Synthesis of Lactose Dendrimers and Multivalency Effects in Binding to the Cholera Toxin B Subunit. European Journal of Organic Chemistry, 2001, 2001, 4685.	2.4	70
245	Peptoid-Peptide Hybrids That Bind Syk SH2 Domains Involved in Signal Transduction. ChemBioChem, 2001, 2, 171-179.	2.6	47
246	Wedgelike Glycodendrimers as Inhibitors of Binding of Mammalian Galectins to Glycoproteins, Lactose Maxiclusters, and Cell Surface Glycoconjugates. ChemBioChem, 2001, 2, 822.	2.6	206
247	Sensitivity of Single Membrane-Spanning α-Helical Peptides to Hydrophobic Mismatch with a Lipid Bilayer:  Effects on Backbone Structure, Orientation, and Extent of Membrane Incorporation. Biochemistry, 2001, 40, 5000-5010.	2.5	171
248	Peptoid-Peptide Hybrids: Design, Synthesis and MHC Binding. , 2001, , 1045-1046.		0
249	Peptide Transformation and Synthesis of Oligopeptidomimetics. , 2001, , 232-233.		0
250	Covalent Control of Shape and Folding in Peptides by Ring-Closing Metathesis. , 2001, , 127-129.		0
251	Peptoid–Peptide Hybrids That Bind Syk SH2 Domains Involved in Signal Transduction. ChemBioChem, 2001, 2, 171-179.	2.6	0
252	A practical solid phase synthesis of oligopeptidosulfonamide foldamers. Tetrahedron Letters, 2000, 41, 7991-7995.	1.4	44

#	Article	IF	CITATIONS
253	Peptidomimetic building blocks for the synthesis of sulfonamide peptoids. Tetrahedron Letters, 2000, 41, 1103-1106.	1.4	20
254	An Efficient Synthesis of N-Protected Î <sup>2</sup> -Aminoethanesulfonyl Chlorides: Versatile Building Blocks for the Synthesis of Oligopeptidosulfonamides. Synthesis, 2000, 2000, 1579-1584.	2.3	35
255	Combinatorial Chemistry for Ligand Development in Catalysis:  Synthesis and Catalysis Screening of Peptidosulfonamide Tweezers on the Solid Phase. Journal of Organic Chemistry, 2000, 65, 1750-1757.	3.2	64
256	Synthesis of Cyclic Peptides by Ring-Closing Metathesis. Journal of Organic Chemistry, 2000, 65, 6187-6195.	3.2	93
257	Different Membrane Anchoring Positions of Tryptophan and Lysine in Synthetic Transmembrane α-Helical Peptides. Journal of Biological Chemistry, 1999, 274, 20839-20846.	3.4	298
258	Solution phase combinatorial chemistry using cyclotriveratrylene-based tripodal scaffolds. Tetrahedron Letters, 1999, 40, 9347-9351.	1.4	14
259	Increased stability of peptidesulfonamide peptidomimetics towards protease catalyzed degradation. Bioorganic and Medicinal Chemistry, 1999, 7, 1043-1047.	3.0	51
260	Synthesis and catalytic application of amino acid based dendritic macromolecules. Tetrahedron Letters, 1999, 40, 1413-1416.	1.4	55
261	Solid-Phase Synthesis of Oligourea Peptidomimetics. European Journal of Organic Chemistry, 1999, 1999, 2127-2135.	2.4	15
262	Rolling Loop Scan: An Approach Featuring Ring-Closing Metathesis for Generating Libraries of Peptides with Molecular Shapes Mimicking Bioactive Conformations or Local Folding of Peptides and Proteins. Angewandte Chemie - International Edition, 1999, 38, 3684-3687.	13.8	68
263	A Versatile Method for the Conjugation of Proteins and Peptides to Poly[2-(dimethylamino)ethyl methacrylate]. Bioconjugate Chemistry, 1999, 10, 687-692.	3.6	45
264	Site-specific N-alkylation of peptides on the solid phase. Tetrahedron Letters, 1998, 39, 1243-1246.	1.4	87
265	Solid-Phase Syntheses of Peptoids using Fmoc-ProtectedN-Substituted Glycines: The Synthesis of (Retro)Peptoids of Leu-Enkephalin and Substance P. Chemistry - A European Journal, 1998, 4, 1570-1580.	3.3	102
266	Tweezers with Different Bite: Increasing the Affinity of Synthetic Receptors by Varying the Hinge Part. Angewandte Chemie - International Edition, 1998, 37, 1846-1850.	13.8	38
267	Sequencing of peptoid peptidomimetics by Edman degradation. Tetrahedron Letters, 1998, 39, 3589-3592.	1.4	26
268	Combinatorial chemistry of hydantoins. Bioorganic and Medicinal Chemistry Letters, 1998, 8, 2375-2380.	2.2	59
269	Sizing of amino acid based dendrimers in Langmuir monolayers. Journal of the Chemical Society Perkin Transactions II, 1998, , 1535-1538.	0.9	12
270	Solid-Phase Syntheses of Peptoids using Fmoc-Protected N-Substituted Glycines: The Synthesis of (Retro)Peptoids of Leu-Enkephalin and Substance P. , 1998, 4, 1570.		1

#	Article	IF	CITATIONS
271	Synthesis of a novel amino acid based dendrimer. Tetrahedron Letters, 1997, 38, 631-634.	1.4	51
272	Molecular Diversity of Novel Amino Acid Based Dendrimers. Tetrahedron Letters, 1997, 38, 3085-3088.	1.4	46
273	Approaches to the synthesis of ureapeptoid peptidomimetics. Tetrahedron Letters, 1997, 38, 5335-5338.	1.4	58
274	The rational design of TAP inhibitors using peptide substrate modifications and peptidomimetics. European Journal of Immunology, 1997, 27, 898-904.	2.9	50
275	Comparing Mass Spectrometric Characteristics of Peptides and Peptoids—2â€. Journal of Mass Spectrometry, 1997, 32, 697-704.	1.6	24
276	Reaction of N-trityl amino acids with BOP: Efficient synthesis of t-butyl esters as well as N-trityl serine- and threonine-β-lactones. Tetrahedron Letters, 1996, 37, 4237-4240.	1.4	34
277	Synthetic receptors based on peptidosulfonamide peptidomimetics. Tetrahedron Letters, 1996, 37, 8253-8256.	1.4	32
278	Molecular diversity of peptidomimetics: Approaches to the solid-phase synthesis of peptidosulfonamides. Bioorganic and Medicinal Chemistry, 1996, 4, 667-672.	3.0	48
279	Solid-phase synthesis of peptidosulfonamide containing peptides derived from Leu-enkephalin. Bioorganic and Medicinal Chemistry Letters, 1996, 6, 3035-3040.	2.2	40
280	Synthesis in Solution of Peptoids using Fmoc-protected N-substituted Glycines. Tetrahedron Letters, 1995, 36, 6969-6972.	1.4	27
281	Synthesis of Peptidosulfinamides and Peptidosulfonamides: Peptidomimetics Containing the Sulfinamide or Sulfonamide Transition-State Isostere. Journal of Organic Chemistry, 1995, 60, 5157-5169.	3.2	136
282	Synthesis in solution of peptoids using fmoc-protected n-substituted glycines. Tetrahedron Letters, 1995, 36, 6969-6972.	1.4	24
283	A New Application of Modified Peptides and Peptidomimetics: Potential Anticancer Agents. Angewandte Chemie International Edition in English, 1994, 33, 305-307.	4.4	27
284	Opportunities for New Chemical Libraries: Unnatural Biopolymers and Diversomers. Angewandte Chemie International Edition in English, 1994, 33, 633-636.	4.4	81
285	Modifizierte Peptide und Peptidmimetica als potentielle Tumorhemmer. Angewandte Chemie, 1994, 106, 313-315.	2.0	5
286	Syntheses of Amino Acid Based Phosphodiester Linkage-Containing Cryptands as well as Diphosphorylated Macrocycles. Journal of Organic Chemistry, 1994, 59, 2399-2408.	3.2	11
287	Conformationally restricted amino acids and dipeptides, (non)peptidomimetics and secondary structure mimetics. Recueil Des Travaux Chimiques Des Pays-Bas, 1994, 113, 1-19.	0.0	167
288	Synthesis of peptides containing a sulfinamide or a sulfonamide transition-state isostere. Tetrahedron, 1993, 49, 1133-1150.	1.9	82

#	Article	IF	CITATIONS
289	Peptides containing the novel methylphosphinamide transition-state isostere. Tetrahedron, 1993, 49, 11055-11064.	1.9	14
290	Synthesis and structure of cyclic phosphopeptides containing a phosphodiester linkage. Journal of Organic Chemistry, 1993, 58, 3722-3730.	3.2	6
291	Solid-phase synthesis of O-phosphorothioylserine- and -threonine-containing peptides as well as of O-phosphoserine- and -threonine-containing peptides. Journal of Organic Chemistry, 1993, 58, 1309-1317.	3.2	37
292	Computer-aided molecular modeling and design of DNA-inserting molecules. Journal of Computer-Aided Molecular Design, 1992, 6, 33-46.	2.9	4
293	Synthesis of alkene dipeptide isosteres employing the Wittig-Still rearrangement. Tetrahedron, 1992, 48, 6425-6438.	1.9	42
294	Synthesis of peptides containing the β-substituted aminoethane sulfinamide or sulfonamide transition-state isostere derived from amino acids. Tetrahedron Letters, 1992, 33, 6389-6392.	1.4	46
295	Peptides containing a sulfinamide or a sulfonamide moiety: New transition-state analogues. Tetrahedron Letters, 1991, 32, 409-412.	1.4	77
296	A novel N-myristylated synthetic octapeptide inhibits protein kinase C activity and partially reverses murine fibrosarcoma cell resistance to Adriamycin. Investigational New Drugs, 1991, 9, 169-179.	2.6	33
297	Inhibition of IL-2 receptor induction and IL-2 production in the human leukemic cell line Jurkat by a novel peptide inhibitor of protein kinase C. Cellular Immunology, 1990, 131, 242-252.	3.0	23
298	Inhibition of protein kinase C and calmodulin by the geometric isomerscis- andtrans-tamoxifen. Biopolymers, 1990, 29, 97-104.	2.4	32
299	Macromodel?an integrated software system for modeling organic and bioorganic molecules using molecular mechanics. Journal of Computational Chemistry, 1990, 11, 440-467.	3.3	3,727
300	N-myristyl-Lys-Arg-Thr-Leu-Arg: A novel protein kinase C inhibitor. Biochemical Pharmacology, 1990, 39, 49-57.	4.4	32
301	Specificity and Function of the Individual Amino Acids of an Important Determinant of Human Immunodeficiency Virus Type 1 that Induces Neutralizing Activity. Journal of General Virology, 1989, 70, 1505-1512.	2.9	96
302	Synthesis of a cyclic phosphopeptide containing a phosphodiester linkage. Journal of the American Chemical Society, 1989, 111, 9103-9105.	13.7	16
303	Structure-activity relationships of sparsomycin and its analogs. Inhibition of peptide bond formation in cell-free systems and of L1210 and bacterial cell growth. Journal of Medicinal Chemistry, 1987, 30, 325-333.	6.4	34
304	Total synthesis and absolute configuration of the natural dipeptide .gammaglutamylmarasmine. Journal of Organic Chemistry, 1987, 52, 1511-1517.	3.2	23
305	Solid-phase synthesis of a naturally occurring β-(1→5)-linked d-galactofuranosyl heptamer containing the artificial linkage arm L-homoserine. Tetrahedron Letters, 1987, 28, 6695-6698.	1.4	64
306	Computer-assisted molecular modeling of tumor promoters: rationale for the activity of phorbol esters, teleocidin B, and aplysiatoxin Proceedings of the National Academy of Sciences of the United States of America, 1986, 83, 241-245.	7.1	117

11

#	Article	IF	CITATIONS
307	Cellular uptake and localization of fluorescent derivatives of phorbol ester tumor promoters. Biochemical and Biophysical Research Communications, 1985, 131, 920-927.	2.1	40
308	Structure-activity relationships of sparsomycin and its analogs: octylsparsomycin: The first analog more active than sparsomycin. Journal of Medicinal Chemistry, 1984, 27, 301-306.	6.4	27
309	Sulfoxide configuration in sparsomycin determines time-dependent and competitive inhibition of peptidyl transferase. Biochemical and Biophysical Research Communications, 1984, 125, 784-789.	2.1	6
310	Determination of sparsomycin in plasma and urine of the dog by means of reversed-phase high-performance liquid chromatography and first pharmacokinetic results. Biomedical Applications, 1983, 275, 145-153.	1.7	5
311	Conformational analysis of functionalized sultines by nuclear magnetic resonance and x-ray crystallography. Application of a generalized Karplus equation. Journal of the American Chemical Society, 1983, 105, 5406-5414.	13.7	14
312	Flash vacuum thermolysis of functionalized .gammasultines. Journal of Organic Chemistry, 1983, 48, 2733-2736.	3.2	14
313	Absolute configuration of sparsomycin. A chiroptical study of sulfoxides. Journal of the American Chemical Society, 1981, 103, 1720-1723.	13.7	31
314	Synthesis and ring-opening reactions of functionalized sultines. New approach to sparsomycin. Journal of Organic Chemistry, 1981, 46, 5408-5413.	3.2	48
315	Total synthesis of the antibiotic sparsomycin, a modified uracil amino acid monoxodithioacetal. Journal of Organic Chemistry, 1981, 46, 3273-3283.	3.2	66

Bioactive Macrocyclic Peptides and Peptide Mimics. , 0, , 1-27.