

Fernando Albericio

List of Publications by Year in descending order

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835
papers

30,367
citations

6233

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12910

131
g-index

942
all docs

942
docs citations

942
times ranked

24476
citing authors

#	ARTICLE	IF	CITATIONS
1	The Pharmaceutical Industry in 2021. An Analysis of FDA Drug Approvals from the Perspective of Molecules. <i>Molecules</i> , 2022, 27, 1075.	1.7	60
2	Linkers: An Assurance for Controlled Delivery of Antibody-Drug Conjugate. <i>Pharmaceutics</i> , 2022, 14, 396.	2.0	48
3	Understanding OxymaPure as a Peptide Coupling Additive: A Guide to New Oxyma Derivatives. <i>ACS Omega</i> , 2022, 7, 6007-6023.	1.6	6
4	2021 FDA TIDES (Peptides and Oligonucleotides) Harvest. <i>Pharmaceutics</i> , 2022, 15, 222.	1.7	48
5	Amino-Li-Resinâ€”A Fiber Polyacrylamide Resin for Solid-Phase Peptide Synthesis. <i>Polymers</i> , 2022, 14, 928.	2.0	4
6	Chemoselective Disulfide Formation by Thiol-Disulfide Interchange in SIT-Protected Cysteiny Peptides. <i>Journal of Organic Chemistry</i> , 2022, 87, 708-712.	1.7	7
7	<i>In situ</i> Fmoc removal â€” a sustainable solid-phase peptide synthesis approach. <i>Green Chemistry</i> , 2022, 24, 4887-4896.	4.6	6
8	Morphological behavior of fullereneâ€”steroid hybrid derivatives. <i>Surface and Interface Analysis</i> , 2022, 54, 1041-1051.	0.8	2
9	Synthesis and Antiproliferative Activity of a New Series of Mono- and Bis(dimethylpyrazolyl)- <i>s</i> -triazine Derivatives Targeting EGFR/PI3K/AKT/mTOR Signaling Cascades. <i>ACS Omega</i> , 2022, 7, 24858-24870.	1.6	14
10	Liquid-Phase Peptide Synthesis (LPPS): A Third Wave for the Preparation of Peptides. <i>Chemical Reviews</i> , 2022, 122, 13516-13546.	23.0	35
11	NOXA upregulation by the prohibitinâ€”binding compound fluorizoline is transcriptionally regulated by integrated stress responseâ€”induced ATF3 and ATF4. <i>FEBS Journal</i> , 2021, 288, 1271-1285.	2.2	18
12	Lactoferrin-dual drug nanoconjugate: Synergistic anti-tumor efficacy of docetaxel and the NF-Î²B inhibitor celastrol. <i>Materials Science and Engineering C</i> , 2021, 118, 111422.	3.8	27
13	1,3,5-Triazine as core for the preparation of dendrons. <i>Arkivoc</i> , 2021, 2020, 64-73.	0.3	2
14	The Pharmaceutical Industry in 2020. An Analysis of FDA Drug Approvals from the Perspective of Molecules. <i>Molecules</i> , 2021, 26, 627.	1.7	87
15	Harnessing polarity and viscosity to identify green binary solvent mixtures as viable alternatives to DMF in solid-phase peptide synthesis. <i>Green Chemistry</i> , 2021, 23, 3295-3311.	4.6	32
16	Green solvents in the biotechnology-based pharmaceutical industry. , 2021, , 87-104.		2
17	Nature-inspired dimerization as a strategy to modulate neuropeptide pharmacology exemplified with vasopressin and oxytocin. <i>Chemical Science</i> , 2021, 12, 4057-4062.	3.7	12
18	Replacing DMF in solid-phase peptide synthesis: varying the composition of green binary solvent mixtures as a tool to mitigate common side-reactions. <i>Green Chemistry</i> , 2021, 23, 3312-3321.	4.6	24

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19	Structure-Acid Lability Relationship of N-alkylated β,β -dialkylglycine Obtained via a Ugi Multicomponent Reaction. <i>Molecules</i> , 2021, 26, 197.	1.7	2
20	A native mass spectrometry platform identifies HOP inhibitors that modulate the HSP90 α -HOP protein α -protein interaction. <i>Chemical Communications</i> , 2021, 57, 10919-10922.	2.2	3
21	2020 FDA TIDES (Peptides and Oligonucleotides) Harvest. <i>Pharmaceuticals</i> , 2021, 14, 145.	1.7	51
22	Propylphosphonic Anhydride (T3P $\hat{\circ}$) as Coupling Reagent for Solid α -Phase Peptide Synthesis. <i>ChemistrySelect</i> , 2021, 6, 2649-2657.	0.7	9
23	Liquid Phase Peptide Synthesis via One α -Pot Nanostar Sieving (PEPSTAR). <i>Angewandte Chemie - International Edition</i> , 2021, 60, 7786-7795.	7.2	20
24	An Androsterone α -H α ₂ @C α ₆₀ hybrid: Synthesis, Properties and Molecular Docking Simulations with SARS α -Cov α 2. <i>ChemPlusChem</i> , 2021, 86, 972-981.	1.3	9
25	The Antiproliferative and Apoptotic Effect of a Novel Synthesized S-Triazine Dipeptide Series, and Toxicity Screening in Zebrafish Embryos. <i>Molecules</i> , 2021, 26, 1170.	1.7	7
26	Liquid Phase Peptide Synthesis via One α -Pot Nanostar Sieving (PEPSTAR). <i>Angewandte Chemie</i> , 2021, 133, 7865-7874.	1.6	4
27	s-Triazine: A Privileged Structure for Drug Discovery and Bioconjugation. <i>Molecules</i> , 2021, 26, 864.	1.7	31
28	Refractive Index: The Ultimate Tool for Real-Time Monitoring of Solid-Phase Peptide Synthesis. Greening the Process. <i>Organic Process Research and Development</i> , 2021, 25, 1047-1053.	1.3	9
29	Improving the Gastrointestinal Stability of Linaclotide. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 8384-8390.	2.9	14
30	Targeting Energy Expenditure α -Drugs for Obesity Treatment. <i>Pharmaceuticals</i> , 2021, 14, 435.	1.7	16
31	An Androsterone α -H α 2 @C 60 hybrid: Synthesis, Properties and Molecular Docking Simulations with SARS α -Cov α 2. <i>ChemPlusChem</i> , 2021, 86, 970-971.	1.3	2
32	Scope and Limitations of Barbituric and Thiobarbituric Amino Acid Derivatives as Protecting Groups for Solid α -Phase Peptide Synthesis: Towards a Green Protecting Group. <i>ChemistrySelect</i> , 2021, 6, 6626-6630.	0.7	3
33	Super-Cationic Peptide Dendrimers α -Synthesis and Evaluation as Antimicrobial Agents. <i>Antibiotics</i> , 2021, 10, 695.	1.5	5
34	Latest Advances on Synthesis, Purification, and Characterization of Peptides and Their Applications. <i>Applied Sciences (Switzerland)</i> , 2021, 11, 5593.	1.3	3
35	A novel α -smart α ™ PNIPAM-based copolymer for breast cancer targeted therapy: Synthesis, and characterization of dual pH/temperature-responsive lactoferrin-targeted PNIPAM-co-AA. <i>Colloids and Surfaces B: Biointerfaces</i> , 2021, 202, 111694.	2.5	38
36	Effects of elderflower extract enriched with polyphenols on antioxidant defense of salmon leukocytes. <i>Electronic Journal of Biotechnology</i> , 2021, 52, 13-20.	1.2	5

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37	Fully Automated Screening of a Combinatorial Library to Avoid False Positives: Application to Tetanus Toxoid Ligand Identification. <i>ACS Omega</i> , 2021, 6, 18756-18762.	1.6	1
38	Rhodiasolv PolarClean " a greener alternative in solid-phase peptide synthesis. <i>Green Chemistry Letters and Reviews</i> , 2021, 14, 545-550.	2.1	11
39	Synthesis of New Peptide-Based Ligands with 1,2-HOPO Pendant Chelators and Thermodynamic Evaluation of Their Iron(III) Complexes**. <i>ChemistrySelect</i> , 2021, 6, 7674-7681.	0.7	1
40	Tea Bags for Fmoc Solid-Phase Peptide Synthesis: An Example of Circular Economy. <i>Molecules</i> , 2021, 26, 5035.	1.7	19
41	Amide Formation: Choosing the Safer Carbodiimide in Combination with OxymaPure to Avoid HCN Release. <i>Organic Letters</i> , 2021, 23, 6900-6904.	2.4	14
42	Novel Biomimetic Human TLR2-Derived Peptides for Potential Targeting of Lipoteichoic Acid: An In Silico Assessment. <i>Biomedicines</i> , 2021, 9, 1063.	1.4	1
43	Evaluation of the tert-butyl group as a probe for NMR studies of macromolecular complexes. <i>Journal of Biomolecular NMR</i> , 2021, 75, 347-363.	1.6	4
44	<i>In vivo</i> micro computed tomography detection and decrease in amyloid load by using multifunctionalized gold nanorods: a neurotheranostic platform for Alzheimer's disease. <i>Biomaterials Science</i> , 2021, 9, 4178-4190.	2.6	14
45	Minimizing side reactions during amide formation using DIC and oxymapure in solid-phase peptide synthesis. <i>Tetrahedron Letters</i> , 2021, 85, 153462.	0.7	8
46	NIR and glutathione trigger the surface release of methotrexate linked by Diels-Alder adducts to anisotropic gold nanoparticles. <i>Materials Science and Engineering C</i> , 2021, 131, 112512.	3.8	10
47	A short peptide fragment of the vascular endothelial growth factor as a novel ligand for bevacizumab purification. <i>Protein Expression and Purification</i> , 2020, 165, 105500.	0.6	4
48	Di- and tri-substituted s-triazine derivatives: Synthesis, characterization, anticancer activity in human breast-cancer cell lines, and developmental toxicity in zebrafish embryos. <i>Bioorganic Chemistry</i> , 2020, 94, 103397.	2.0	17
49	Synthesis and Antimicrobial Activity of a New Series of Thiazolidine-2,4-diones Carboxamide and Amino Acid Derivatives. <i>Molecules</i> , 2020, 25, 105.	1.7	16
50	Protocol for bevacizumab purification using Ac-PHQGQHIGVSK-agarose. <i>MethodsX</i> , 2020, 7, 100769.	0.7	2
51	Hydroxamate siderophores: Natural occurrence, chemical synthesis, iron binding affinity and use as Trojan horses against pathogens. <i>European Journal of Medicinal Chemistry</i> , 2020, 208, 112791.	2.6	50
52	Novel formulation of antimicrobial peptides enhances antimicrobial activity against methicillin-resistant <i>Staphylococcus aureus</i> (MRSA). <i>Amino Acids</i> , 2020, 52, 1439-1457.	1.2	20
53	Exploiting azido-dichloro-triazine as a linker for regioselective incorporation of peptides through their N, O, S functional groups. <i>Bioorganic Chemistry</i> , 2020, 104, 104334.	2.0	3
54	Disulfide-Based Protecting Groups for the Cysteine Side Chain. <i>Organic Letters</i> , 2020, 22, 9644-9647.	2.4	10

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55	Solid-phase synthesis of peptides containing 1-Hydroxypyridine-2-one (1,2-HOPO). <i>Tetrahedron Letters</i> , 2020, 61, 152299.	0.7	2
56	Butylpyrrolidinone for Solid-Phase Peptide Synthesis is Environmentally Friendlier and Synthetically Better than DMF. <i>ChemSusChem</i> , 2020, 13, 5288-5294.	3.6	29
57	Radical Dendrimers Based on Biocompatible Oligoethylene Glycol Dendrimers as Contrast Agents for MRI. <i>Pharmaceutics</i> , 2020, 12, 772.	2.0	18
58	Novel 4,6-Disubstituted s-Triazin-2-yl Amino Acid Derivatives as Promising Antifungal Agents. <i>Journal of Fungi (Basel, Switzerland)</i> , 2020, 6, 237.	1.5	8
59	Editorial: Chemical Design and Biomedical Applications of Disulfide-rich Peptides: Challenges and Opportunities. <i>Frontiers in Chemistry</i> , 2020, 8, 586377.	1.8	4
60	The tea-bag protocol for comparison of Fmoc removal reagents in solid-phase peptide synthesis. <i>Amino Acids</i> , 2020, 52, 1201-1205.	1.2	18
61	Protocol for efficient solid-phase synthesis of peptides containing 1-hydroxypyridine-2-one (1,2-HOPO). <i>MethodsX</i> , 2020, 7, 101082.	0.7	2
62	Insights into the chemistry of the amphibactin-metal (M ³⁺) interaction and its role in antibiotic resistance. <i>Scientific Reports</i> , 2020, 10, 21049.	1.6	3
63	Identification of New Ocellatin Antimicrobial Peptides by cDNA Precursor Cloning in the Frame of This Family of Intriguing Peptides. <i>Antibiotics</i> , 2020, 9, 751.	1.5	3
64	Peptide Therapeutics 2.0. <i>Molecules</i> , 2020, 25, 2293.	1.7	98
65	Synthetic peptides to produce antivenoms against the Cys-rich toxins of arachnids. <i>Toxicon: X</i> , 2020, 6, 100038.	1.2	4
66	Enamine Barbiturates and Thiobarbiturates as a New Class of Bacterial Urease Inhibitors. <i>Applied Sciences (Switzerland)</i> , 2020, 10, 3523.	1.3	5
67	Protocol for synthesis of di- and tri-substituted s-triazine derivatives. <i>MethodsX</i> , 2020, 7, 100825.	0.7	2
68	2019 FDA TIDES (Peptides and Oligonucleotides) Harvest. <i>Pharmaceutics</i> , 2020, 13, 40.	1.7	54
69	Synthesis and characterisation of thiobarbituric acid enamine derivatives, and evaluation of their α -glucosidase inhibitory and anti-glycation activity. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020, 35, 692-701.	2.5	17
70	Synthesis of Stable Cholesteryl-Polyethylene Glycol-Peptide Conjugates with Non-Disperse Polyethylene Glycol Lengths. <i>ACS Omega</i> , 2020, 5, 5508-5519.	1.6	3
71	Breaking a Couple: Disulfide Reducing Agents. <i>ChemBioChem</i> , 2020, 21, 1947-1954.	1.3	39
72	Use of a phosphopeptide as a ligand to purify phospholipase A2 from the venom of <i>Crotalus durissus terrificus</i> by affinity chromatography. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2020, 1146, 122070.	1.2	6

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73	From Ugi Multicomponent Reaction to Linkers for Bioconjugation. ACS Omega, 2020, 5, 7424-7431.	1.6	10
74	Somuncurins: Bioactive Peptides from the Skin of the Endangered Endemic Patagonian Frog <i>Pleurodema somuncurense</i> . Journal of Natural Products, 2020, 83, 972-984.	1.5	8
75	Crystal Structure and Theoretical Investigation of Thiobarbituric Acid Derivatives as Nonlinear Optical (NLO) Materials. Crystals, 2020, 10, 442.	1.0	2
76	Revisiting NO ₂ as Protecting Group of Arginine in Solid-Phase Peptide Synthesis. International Journal of Molecular Sciences, 2020, 21, 4464.	1.8	7
77	Barbiturate- and Thiobarbiturate-Based <i>s</i> -Triazine Hydrazone Derivatives with Promising Antiproliferative Activities. ACS Omega, 2020, 5, 15805-15811.	1.6	21
78	The Pharmaceutical Industry in 2019. An Analysis of FDA Drug Approvals from the Perspective of Molecules. Molecules, 2020, 25, 745.	1.7	121
79	Gold nanoparticle based double-labeling of melanoma extracellular vesicles to determine the specificity of uptake by cells and preferential accumulation in small metastatic lung tumors. Journal of Nanobiotechnology, 2020, 18, 20.	4.2	68
80	Greening Fmoc- <i>t</i> -Bu solid-phase peptide synthesis. Green Chemistry, 2020, 22, 996-1018.	4.6	85
81	Phenol as a Modulator in the Chemical Reactivity of 2,4,6-Trichloro-1,3,5-triazine: Rules of the Game II. Australian Journal of Chemistry, 2020, 73, 352.	0.5	5
82	Carpino's protecting groups, beyond the Boc and the Fmoc. Peptide Science, 2020, 112, e24164.	1.0	7
83	Cleaving protected peptides from 2-chlorotrityl chloride resin. Moving away from dichloromethane. Green Chemistry, 2020, 22, 2840-2845.	4.6	11
84	Solid-Phase Synthesis of Head to Side-Chain Tyr-Cyclodepsipeptides Through a Cyclative Cleavage From Fmoc-MeDbz/MeNbz-resins. Frontiers in Chemistry, 2020, 8, 298.	1.8	7
85	Naturally Occurring Oxazole-Containing Peptides. Marine Drugs, 2020, 18, 203.	2.2	34
86	Gold Nanoparticles Mediate Improved Detection of β -amyloid Aggregates by Fluorescence. Nanomaterials, 2020, 10, 690.	1.9	28
87	Successful development of a method for the incorporation of Fmoc-Arg(Pbf)-OH in solid-phase peptide synthesis using <i>N</i> -butylpyrrolidinone (NBP) as solvent. Green Chemistry, 2020, 22, 3162-3169.	4.6	22
88	OxymaPure Coupling Reagents: Beyond Solid-Phase Peptide Synthesis. Synthesis, 2020, 52, 3189-3210.	1.2	6
89	<i>s</i> -Triazine: A Multidisciplinary and International Journey. Chemistry Proceedings, 2020, 3, .	0.1	0
90	β -Valerolactone (GVL): An eco-friendly anchoring solvent for solid-phase peptide synthesis. Tetrahedron Letters, 2019, 60, 151058.	0.7	19

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91	Identification of Antimicrobial Peptides from the Microalgae <i>Tetraselmis suecica</i> (Kylin) Butcher and Bactericidal Activity Improvement. <i>Marine Drugs</i> , 2019, 17, 453.	2.2	74
92	Calculating Resin Functionalization in Solid-Phase Peptide Synthesis Using a Standardized Method based on Fmoc Determination. <i>ACS Combinatorial Science</i> , 2019, 21, 717-721.	3.8	7
93	Chemical Modification of Microcin J25 Reveals New Insights on the Stereospecific Requirements for Antimicrobial Activity. <i>International Journal of Molecular Sciences</i> , 2019, 20, 5152.	1.8	11
94	Scope and Limitations of $\hat{\text{I}}^3$ -Valerolactone (GVL) as a Green Solvent to be Used with Base for Fmoc Removal in Solid Phase Peptide Synthesis. <i>Molecules</i> , 2019, 24, 4004.	1.7	20
95	Antibiotic Resistance: From the Bench to Patients. <i>Antibiotics</i> , 2019, 8, 129.	1.5	91
96	Investigating Triorthogonal Chemoselectivity. Effect of Azide Substitution on the Triazine Core. <i>Organic Letters</i> , 2019, 21, 7888-7892.	2.4	9
97	Green Transformation of Solid-Phase Peptide Synthesis. <i>ACS Sustainable Chemistry and Engineering</i> , 2019, 7, 3671-3683.	3.2	67
98	Large-Area Biomolecule Nanopatterns on Diblock Copolymer Surfaces for Cell Adhesion Studies. <i>Nanomaterials</i> , 2019, 9, 579.	1.9	6
99	Bypassing Osmotic Shock Dilemma in a Polystyrene Resin Using the Green Solvent Cyclopentyl methyl Ether (CPME): A Morphological Perspective. <i>Polymers</i> , 2019, 11, 874.	2.0	8
100	Optimized Stepwise Synthesis of the API Liraglutide Using BAL Resin and Pseudoprolines. <i>ACS Omega</i> , 2019, 4, 8674-8680.	1.6	8
101	2018 FDA Tides Harvest. <i>Pharmaceuticals</i> , 2019, 12, 52.	1.7	39
102	Carbosilane Dendronâ€“Peptide Nanoconjugates as Antimicrobial Agents. <i>Molecular Pharmaceutics</i> , 2019, 16, 2661-2674.	2.3	27
103	Pseudo-Wang Handle for the Preparation of Fully Protected Peptides. Synthesis of Liraglutide by Fragment Condensation. <i>Organic Letters</i> , 2019, 21, 2459-2463.	2.4	11
104	Design and synthesis of mono-and di-pyrazolyl-s-triazine derivatives, their anticancer profile in human cancer cell lines, and in vivo toxicity in zebrafish embryos. <i>Bioorganic Chemistry</i> , 2019, 87, 457-464.	2.0	37
105	Troubleshooting When Using $\hat{\text{I}}^3$ -Valerolactone (GVL) in Green Solid-Phase Peptide Synthesis. <i>Organic Process Research and Development</i> , 2019, 23, 1096-1100.	1.3	29
106	The Pharmaceutical Industry in 2018. An Analysis of FDA Drug Approvals from the Perspective of Molecules. <i>Molecules</i> , 2019, 24, 809.	1.7	95
107	2-(Dibenzylamino)butane-1,4-dithiol (DABDT), a Friendly Disulfide-Reducing Reagent Compatible with a Broad Range of Solvents. <i>Organic Letters</i> , 2019, 21, 10111-10114.	2.4	7
108	Report of <i>in vitro</i> antileishmanial properties of Iberian macroalgae. <i>Natural Product Research</i> , 2019, 33, 1778-1782.	1.0	5

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109	OctaGel Resin - A New PEG-PS-based Solid Support for Solid-Phase Peptide Synthesis. <i>Letters in Organic Chemistry</i> , 2019, 16, 935-940.	0.2	4
110	Efficient Route for Synthesis of Enamines from 1,3-Alkyl-2-Thioxodihydropyrimidine-4,6(1H,5H)-dione Enols. <i>Letters in Organic Chemistry</i> , 2019, 16, 538-540.	0.2	0
111	Toward the Synthesis of Phormidolides. <i>ACS Omega</i> , 2018, 3, 2351-2362.	1.6	5
112	Bioconjugation through Mesitylene Thiol Alkylation. <i>Bioconjugate Chemistry</i> , 2018, 29, 1199-1208.	1.8	5
113	Solid-Phase Synthesis of Pyrrole Derivatives through a Multicomponent Reaction Involving Lys-Containing Peptides. <i>ACS Combinatorial Science</i> , 2018, 20, 187-191.	3.8	14
114	1,3,5-Triazino Peptide Derivatives: Synthesis, Characterization, and Preliminary Antileishmanial Activity. <i>ChemMedChem</i> , 2018, 13, 725-735.	1.6	23
115	Application of Decafluorobiphenyl (DFBP) Moiety as a Linker in Bioconjugation. <i>Bioconjugate Chemistry</i> , 2018, 29, 225-233.	1.8	7
116	Identification of New Activators of Mitochondrial Fusion Reveals a Link between Mitochondrial Morphology and Pyrimidine Metabolism. <i>Cell Chemical Biology</i> , 2018, 25, 268-278.e4.	2.5	84
117	3D Electrophoresis-Assisted Lithography (3DEAL): 3D Molecular Printing to Create Functional Patterns and Anisotropic Hydrogels. <i>Advanced Functional Materials</i> , 2018, 28, 1703014.	7.8	13
118	Single step recombinant human growth hormone (rhGH) purification from milk by peptide affinity chromatography. <i>Biotechnology Progress</i> , 2018, 34, 999-1005.	1.3	3
119	Microwave-Assisted Green Solid-Phase Peptide Synthesis Using γ -Valerolactone (GVL) as Solvent. <i>ACS Sustainable Chemistry and Engineering</i> , 2018, 6, 8034-8039.	3.2	65
120	Solid-phase synthesis of homodetic cyclic peptides from Fmoc-MeDbz-resin. <i>Tetrahedron Letters</i> , 2018, 59, 1779-1782.	0.7	14
121	Exploring the influence of Diels-Alder linker length on photothermal molecule release from gold nanorods. <i>Colloids and Surfaces B: Biointerfaces</i> , 2018, 166, 323-329.	2.5	11
122	N -methylation in amino acids and peptides: Scope and limitations. <i>Biopolymers</i> , 2018, 109, e23110.	1.2	41
123	Teixobactin as a scaffold for unlimited new antimicrobial peptides: SAR study. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 2788-2796.	1.4	40
124	Improved pharmacokinetic profile of lipophilic anti-cancer drugs using ^{125}I -targeted polyurethane-polyurea nanoparticles. <i>Nanomedicine: Nanotechnology, Biology, and Medicine</i> , 2018, 14, 257-267.	1.7	13
125	Crystal structure, spectroscopic studies and theoretical studies of thiobarbituric acid derivatives: understanding the hydrogen-bonding patterns. <i>Acta Crystallographica Section C, Structural Chemistry</i> , 2018, 74, 1703-1714.	0.2	4
126	Greening the Solid-Phase Peptide Synthesis Process. 2-MeTHF for the Incorporation of the First Amino Acid and Precipitation of Peptides after Global Deprotection. <i>Organic Process Research and Development</i> , 2018, 22, 1809-1816.	1.3	33

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127	Leptodactylus latrans Amphibian Skin Secretions as a Novel Source for the Isolation of Antibacterial Peptides. <i>Molecules</i> , 2018, 23, 2943.	1.7	7
128	Perfluorophenyl Derivatives as Unsymmetrical Linkers for Solid Phase Conjugation. <i>Frontiers in Chemistry</i> , 2018, 6, 589.	1.8	5
129	Welcome to the New Journal Methods and Protocols. <i>Methods and Protocols</i> , 2018, 1, 1.	0.9	5
130	Hydroxylamine Derivatives as a New Paradigm in the Search of Antibacterial Agents. <i>ACS Omega</i> , 2018, 3, 17057-17069.	1.6	10
131	A Lasso-Inspired Bicyclic Peptide: Synthesis, Structure and Properties. <i>Chemistry - A European Journal</i> , 2018, 24, 19250-19257.	1.7	7
132	Synthesis, carbonic anhydrase inhibitory activity and antioxidant activity of some 1,3-oxazine derivatives. <i>Drug Development Research</i> , 2018, 79, 352-361.	1.4	10
133	Peptide Ligations by Using Aryloxycarbonyl-methylaminoanilides: Chemical Synthesis of Palmitoylated Sonic Hedgehog. <i>Angewandte Chemie - International Edition</i> , 2018, 57, 16120-16125.	7.2	22
134	Bacteria Hunt Bacteria through an Intriguing Cyclic Peptide. <i>ChemMedChem</i> , 2018, 14, 24-51.	1.6	7
135	Peptide Ligations by Using Aryloxycarbonyl-methylaminoanilides: Chemical Synthesis of Palmitoylated Sonic Hedgehog. <i>Angewandte Chemie</i> , 2018, 130, 16352-16357.	1.6	8
136	Exploring the Orthogonal Chemoselectivity of 2,4,6-Trichloro-1,3,5-Triazine (TCT) as a Trifunctional Linker With Different Nucleophiles: Rules of the Game. <i>Frontiers in Chemistry</i> , 2018, 6, 516.	1.8	30
137	2017 FDA Peptide Harvest. <i>Pharmaceuticals</i> , 2018, 11, 42.	1.7	44
138	Investigation of the Biosynthesis of the Lasso Peptide Chaxapeptin Using an <i>E. coli</i> -Based Production System. <i>Journal of Natural Products</i> , 2018, 81, 2050-2056.	1.5	25
139	One-Pot Peptide Ligation-Oxidative Cyclization Protocol for the Preparation of Short-/Medium-Size Disulfide Cyclopeptides. <i>Organic Letters</i> , 2018, 20, 4306-4309.	2.4	9
140	Investigating green ethers for the precipitation of peptides after global deprotection in solid-phase peptide synthesis. <i>Current Opinion in Green and Sustainable Chemistry</i> , 2018, 11, 99-103.	3.2	21
141	In Vitro Antibacterial Activity of Teixobactin Derivatives on Clinically Relevant Bacterial Isolates. <i>Frontiers in Microbiology</i> , 2018, 9, 1535.	1.5	25
142	Further applications of classical amide coupling reagents: Microwave-assisted esterification on solid phase. <i>Journal of Peptide Science</i> , 2018, 24, e3111.	0.8	9
143	Identification of Peptides in Flowers of <i>Sambucus nigra</i> with Antimicrobial Activity against Aquaculture Pathogens. <i>Molecules</i> , 2018, 23, 1033.	1.7	21
144	Formation of N-terminal 2-dialkyl amino oxazoles from guanidinated derivatives under mild conditions. <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 5661-5666.	1.5	3

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145	Exploiting the Thiobarbituric Acid Scaffold for Antibacterial Activity. <i>ChemMedChem</i> , 2018, 13, 1923-1930.	1.6	12
146	The Pharmaceutical Industry in 2017. An Analysis of FDA Drug Approvals from the Perspective of Molecules. <i>Molecules</i> , 2018, 23, 533.	1.7	94
147	Chemical Synthesis and Functional Analysis of VarvA Cyclotide. <i>Molecules</i> , 2018, 23, 952.	1.7	8
148	On the Importance of Polyurethane and Polyurea Nanosystems for Future Drug Delivery. <i>Current Drug Delivery</i> , 2018, 15, 37-43.	0.8	31
149	Choosing the Right Coupling Reagent for Peptides: A Twenty-Five-Year Journey. <i>Organic Process Research and Development</i> , 2018, 22, 760-772.	1.3	108
150	Natural Snake Venom Inhibitors and their Pharmaceutical Uses: Challenges and Possibilities. <i>Current Pharmaceutical Design</i> , 2018, 24, 1737-1747.	0.9	7
151	Diethylphosphoryl-OxymaB (DEPO-B) as a Solid Coupling Reagent for Amide Bond Formation. <i>Letters in Organic Chemistry</i> , 2018, 16, 30-33.	0.2	2
152	Green Solid-Phase Peptide Synthesis (GSPPS) 3. Green Solvents for Fmoc Removal in Peptide Chemistry. <i>Organic Process Research and Development</i> , 2017, 21, 365-369.	1.3	52
153	The synthesis of an EDTA-like chelating peptidomimetic building block suitable for solid-phase peptide synthesis. <i>Chemical Communications</i> , 2017, 53, 2634-2636.	2.2	1
154	Microwave-Assisted Synthesis of Antimicrobial Peptides. <i>Methods in Molecular Biology</i> , 2017, 1548, 51-59.	0.4	6
155	Novel Globular Polymeric Supports for Membrane-Enhanced Peptide Synthesis. <i>Macromolecules</i> , 2017, 50, 1626-1634.	2.2	19
156	Tetrahydropyranyl: A Nonaromatic, Mildly Acid-Labile Group for Hydroxyl Protection in Solid-Phase Peptide Synthesis. <i>ChemistryOpen</i> , 2017, 6, 206-210.	0.9	4
157	Understanding Tetrahydropyranyl as a Protecting Group in Peptide Chemistry. <i>ChemistryOpen</i> , 2017, 6, 168-177.	0.9	15
158	Oxidative couplings on tryptophan-based diketopiperazines leading to fused and bridged chemotypes. <i>Chemical Communications</i> , 2017, 53, 2740-2743.	2.2	6
159	Stapled Peptides by Late-Stage C(sp ³) ³ -H Activation. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 314-318.	7.2	132
160	Stapled Peptides by Late-Stage C(sp ³) ³ -H Activation. <i>Angewandte Chemie</i> , 2017, 129, 320-324.	1.6	50
161	Facile solid-phase synthesis of head-side chain cyclotidepeptides through a cyclative cleavage from MeDbz-resin. <i>Tetrahedron Letters</i> , 2017, 58, 2788-2791.	0.7	16
162	Role of the Nozaki-Hiyama-Takai-Kishi Reaction in the Synthesis of Natural Products. <i>Chemical Reviews</i> , 2017, 117, 8420-8446.	23.0	136

#	ARTICLE	IF	CITATIONS
163	Novel pyrazolyl-s-triazine derivatives, molecular structure and antimicrobial activity. <i>Journal of Molecular Structure</i> , 2017, 1145, 244-253.	1.8	45
164	The prohibitin-binding compound fluorizoline induces apoptosis in chronic lymphocytic leukemia cells through the upregulation of NOXA and synergizes with ibrutinib, 5-aminoimidazole-4-carboxamide riboside or venetoclax. <i>Haematologica</i> , 2017, 102, 1587-1593.	1.7	19
165	Effect of TLR ligands co-encapsulated with multiepitopic antigen in nanoliposomes targeted to human DCs via Fc receptor for cancer vaccines. <i>Immunobiology</i> , 2017, 222, 989-997.	0.8	34
166	Synthesis, in vitro evaluation, and ⁶⁸ Ga radiolabeling of CDP1 toward PET/CT imaging of bacterial infection. <i>Chemical Biology and Drug Design</i> , 2017, 90, 572-579.	1.5	10
167	Intercalative DNA binding of the marine anticancer drug variolin B. <i>Scientific Reports</i> , 2017, 7, 39680.	1.6	19
168	A Trp-BODIPY cyclic peptide for fluorescence labelling of apoptotic bodies. <i>Chemical Communications</i> , 2017, 53, 945-948.	2.2	67
169	Improving gold nanorod delivery to the central nervous system by conjugation to the shuttle Angiopep-2. <i>Nanomedicine</i> , 2017, 12, 2503-2517.	1.7	41
170	Re-evaluating the stability of COMU in different solvents. <i>Journal of Peptide Science</i> , 2017, 23, 763-768.	0.8	18
171	Converting Teixobactin into a Cationic Antimicrobial Peptide (AMP). <i>Journal of Medicinal Chemistry</i> , 2017, 60, 7476-7482.	2.9	42
172	Fmoc-Amox, A Suitable Reagent for the Introduction of Fmoc. <i>Organic Process Research and Development</i> , 2017, 21, 1533-1541.	1.3	3
173	Green solid-phase peptide synthesis 4. ¹³ C-Valerolactone and N-formylmorpholine as green solvents for solid phase peptide synthesis. <i>Tetrahedron Letters</i> , 2017, 58, 2986-2988.	0.7	61
174	Gold nanoparticles as an efficient drug delivery system for GLP-1 peptides. <i>Colloids and Surfaces B: Biointerfaces</i> , 2017, 158, 25-32.	2.5	37
175	Preparation of a Trp-BODIPY fluorogenic amino acid to label peptides for enhanced live-cell fluorescence imaging. <i>Nature Protocols</i> , 2017, 12, 1588-1619.	5.5	58
176	Sudemycin K: A Synthetic Antitumor Splicing Inhibitor Variant with Improved Activity and Versatile Chemistry. <i>ACS Chemical Biology</i> , 2017, 12, 163-173.	1.6	23
177	Functionalization of CoCr surfaces with cell adhesive peptides to promote HUVECs adhesion and proliferation. <i>Applied Surface Science</i> , 2017, 393, 82-92.	3.1	42
178	Investigation of the N-Terminus Amino Function of Arg10-Teixobactin. <i>Molecules</i> , 2017, 22, 1632.	1.7	20
179	Phakellistatins: An Underwater Unsolved Puzzle. <i>Marine Drugs</i> , 2017, 15, 78.	2.2	23
180	The Pharmaceutical Industry in 2016. An Analysis of FDA Drug Approvals from a Perspective of the Molecule Type. <i>Molecules</i> , 2017, 22, 368.	1.7	28

#	ARTICLE	IF	CITATIONS
181	Sulfonamide-Linked Ciprofloxacin, Sulfadiazine and Amantadine Derivatives as a Novel Class of Inhibitors of Jack Bean Urease; Synthesis, Kinetic Mechanism and Molecular Docking. <i>Molecules</i> , 2017, 22, 1352.	1.7	42
182	Structure-Activity Relationship of Arg10-Teixobactin: A Recently Discovered Antimicrobial Peptide. <i>Proceedings (mdpi)</i> , 2017, 1, .	0.2	0
183	Synthesis, Characterization, and Tautomerism of 1,3-Dimethyl Pyrimidine-2,4,6-Trione s-Triazinyl Hydrazine/Hydrazone Derivatives. <i>Journal of Chemistry</i> , 2017, 2017, 1-10.	0.9	7
184	Dual Inhibition of AChE and BChE with the C-5 Substituted Derivative of Meldrum's Acid: Synthesis, Structure Elucidation, and Molecular Docking Studies. <i>Crystals</i> , 2017, 7, 211.	1.0	18
185	Synthesis, Crystal Structure and DFT Studies of 1,3-Dimethyl-5-propionylpyrimidine-2,4,6(1H,3H,5H)-trione. <i>Crystals</i> , 2017, 7, 31.	1.0	6
186	Synthesis, Crystal Structure, DFT Study of m-Methoxy-N ² -(3-Methoxybenzoyl)-N-Phenylbenzohydrazide. <i>Crystals</i> , 2017, 7, 19.	1.0	5
187	Deprotection Reagents in Fmoc Solid Phase Peptide Synthesis: Moving Away from Piperidine?. <i>Molecules</i> , 2016, 21, 1542.	1.7	69
188	Can macroalgae provide promising anti-tumoral compounds? A closer look at <i>Cystoseira tamariscifolia</i> as a source for antioxidant and anti-hepatocarcinoma compounds. <i>PeerJ</i> , 2016, 4, e1704.	0.9	33
189	Galactosidase Loaded Nanoliposomes with Enhanced Enzymatic Activity and Intracellular Penetration. <i>Advanced Healthcare Materials</i> , 2016, 5, 829-840.	3.9	40
190	Enhanced antimicrobial activity of a peptide derived from human lysozyme by arylation of its tryptophan residues. <i>Journal of Peptide Science</i> , 2016, 22, 123-128.	0.8	18
191	Ultrasonic promoted synthesis of novel s-triazine-Schiff base derivatives; molecular structure, spectroscopic studies and their preliminary anti-proliferative activities. <i>Journal of Molecular Structure</i> , 2016, 1125, 121-135.	1.8	41
192	Lysine Scanning of Arg ₁₀ -Teixobactin: Deciphering the Role of Hydrophobic and Hydrophilic Residues. <i>ACS Omega</i> , 2016, 1, 1262-1265.	1.6	51
193	A Facile Synthesis of NODASA-Functionalized Peptide. <i>Synlett</i> , 2016, 27, 1685-1688.	1.0	7
194	Combinatorial Library Screening Coupled to Mass Spectrometry to Identify Valuable Cyclic Peptides. <i>Current Protocols in Chemical Biology</i> , 2016, 8, 109-130.	1.7	6
195	Green Solid-Phase Peptide Synthesis 2. 2-Methyltetrahydrofuran and Ethyl Acetate for Solid-Phase Peptide Synthesis under Green Conditions. <i>ACS Sustainable Chemistry and Engineering</i> , 2016, 4, 6809-6814.	3.2	85
196	Synthesis of complex head-to-side-chain cyclodepsipeptides. <i>Nature Protocols</i> , 2016, 11, 1924-1947.	5.5	22
197	Constrained Cyclopeptides: Biaryl Formation through Pd-Catalyzed C-H Activation in Peptides: Structural Control of the Cyclization vs. Cyclodimerization Outcome. <i>Chemistry - A European Journal</i> , 2016, 22, 13114-13119.	1.7	63
198	Nanoencapsulated budesonide in self-stratified polyurethane-polyurea nanoparticles is highly effective in inducing human tolerogenic dendritic cells. <i>International Journal of Pharmaceutics</i> , 2016, 511, 785-793.	2.6	14

#	ARTICLE	IF	CITATIONS
199	Re-evaluation of the N-terminal substitution and the D-residues of teixobactin. RSC Advances, 2016, 6, 73827-73829.	1.7	34
200	Enantioselective Synthesis of the Polyhydroxylated Chain of Oscillariolide and Phormidolides Aâ€“C. Organic Letters, 2016, 18, 4485-4487.	2.4	9
201	Proximate biochemical composition and mineral content of edible species from the genus Cystoseira in Portugal. Botanica Marina, 2016, .	0.6	10
202	Short AntiMicrobial Peptides (SAMPs) as a class of extraordinary promising therapeutic agents. Journal of Peptide Science, 2016, 22, 438-451.	0.8	64
203	Peptides conjugated to silver nanoparticles in biomedicine â€“ a â€œvalue-addedâ€“ phenomenon. Biomaterials Science, 2016, 4, 1713-1725.	2.6	34
204	Spacer-free BODIPY fluorogens in antimicrobial peptides for direct imaging of fungal infection in human tissue. Nature Communications, 2016, 7, 10940.	5.8	112
205	Synthesis of (<i>E</i>)-4-bromo-3-methoxybut-3-en-2-one, the Key Fragment in the Polyhydroxylated Chain Common to Oscillariolide and Phormidolides Aâ€“C. Chemistry - A European Journal, 2016, 22, 7033-7035.	1.7	8
206	Oxyma-T, expanding the arsenal of coupling reagents. Tetrahedron Letters, 2016, 57, 3523-3525.	0.7	5
207	Synthesis of (E)-4-bromo-3-methoxybut-3-en-2-one, the Key Fragment in the Polyhydroxylated Chain Common to Oscillariolide and Phormidolides A-C. Chemistry - A European Journal, 2016, 22, 6993-6993.	1.7	0
208	One pot synthesis, molecular structure and spectroscopic studies (X-ray, IR, NMR, UVâ€“Vis) of novel 2-(4,6-dimethoxy-1,3,5-triazin-2-yl) amino acid ester derivatives. Spectrochimica Acta - Part A: Molecular and Biomolecular Spectroscopy, 2016, 159, 184-198.	2.0	13
209	Comparative proteomic analysis of growth hormone secretagogue A233 treatment of murine macrophage cells J774A.2 indicates it has a role in antiviral innate response. Biochemistry and Biophysics Reports, 2016, 5, 379-387.	0.7	7
210	CuAAC: An Efficient Click Chemistry Reaction on Solid Phase. ACS Combinatorial Science, 2016, 18, 1-14.	3.8	178
211	An improved and efficient strategy for the total synthesis of a colistin-like peptide. Tetrahedron Letters, 2016, 57, 1885-1888.	0.7	15
212	Inhibitory effect of short cationic homopeptides against Gram-negative bacteria. Amino Acids, 2016, 48, 1445-1456.	1.2	8
213	BbrzSP-32, the first serine protease isolated from Bothrops brazili venom: Purification and characterization. Comparative Biochemistry and Physiology Part A, Molecular & Integrative Physiology, 2016, 195, 15-25.	0.8	20
214	Isololiolide, a carotenoid metabolite isolated from the brown alga Cystoseira tamariscifolia, is cytotoxic and able to induce apoptosis in hepatocarcinoma cells through caspase-3 activation, decreased Bcl-2 levels, increased p53 expression and PARP cleavage. Phytomedicine, 2016, 23, 550-557.	2.3	55
215	Highly chemoselective ligation of thiol- and amino-peptides on a bromomaleimide core. Chemical Communications, 2016, 52, 2334-2337.	2.2	9
216	The road to the synthesis of â€œdifficult peptidesâ€“. Chemical Society Reviews, 2016, 45, 631-654.	18.7	171

#	ARTICLE	IF	CITATIONS
217	2-Methyltetrahydrofuran and cyclopentyl methyl ether for green solid-phase peptide synthesis. <i>Amino Acids</i> , 2016, 48, 419-426.	1.2	69
218	A comparative evaluation of biological activities and bioactive compounds of the seagrasses <i>Zostera marina</i> and <i>Zostera noltei</i> from southern Portugal. <i>Natural Product Research</i> , 2016, 30, 724-728.	1.0	14
219	Targeting prohibitins induces apoptosis in acute myeloid leukemia cells. <i>Oncotarget</i> , 2016, 7, 64987-65000.	0.8	19
220	Péptidos que atraviesan la membrana celular como potenciales transportadores de fármacos. <i>Revista Bionatura</i> , 2016, 1, .	0.1	2
221	Installing Multifunctionality on Titanium with RGD-Decorated Polyurethane-Polyurea Roxithromycin Loaded Nanoparticles: Toward New Osseointegrative Therapies. <i>Advanced Healthcare Materials</i> , 2015, 4, 1956-1960.	3.9	27
222	Injectable Hyaluronan Hydrogels with Peptide-Binding Dendrimers Modulate the Controlled Release of BMP-2 and TGF- β 1. <i>Macromolecular Bioscience</i> , 2015, 15, 1035-1044.	2.1	25
223	Review backbone N-modified peptides: How to meet the challenge of secondary amine acylation. <i>Biopolymers</i> , 2015, 104, 435-452.	1.2	26
224	Formylation of Electron-Rich Aromatic Rings Mediated by Dichloromethyl Methyl Ether and TiCl ₄ : Scope and Limitations. <i>Molecules</i> , 2015, 20, 5409-5422.	1.7	20
225	Synthesis and Preliminary Biological Evaluation of 1,3,5-Triazine Amino Acid Derivatives to Study Their MAO Inhibitors. <i>Molecules</i> , 2015, 20, 15976-15988.	1.7	24
226	Relaxivities of Dendrons Based on a OEG-DTPA Architecture: Effect of Gd ³⁺ -Placement and Dendron Functionalization. <i>Journal of Nanotechnology</i> , 2015, 2015, 1-8.	1.5	2
227	New peptide architectures through C-H activation stapling between tryptophan-phenylalanine/tyrosine residues. <i>Nature Communications</i> , 2015, 6, 7160.	5.8	235
228	Chemical Protein Synthesis Using a Second-Generation N-Acylurea Linker for the Preparation of Peptide-Thioester Precursors. <i>Journal of the American Chemical Society</i> , 2015, 137, 7197-7209.	6.6	179
229	The Larock Reaction in the Synthesis of Heterocyclic Compounds. <i>Advances in Heterocyclic Chemistry</i> , 2015, 116, 1-35.	0.9	24
230	Synthesis and Biological Evaluation of a Teixobactin Analogue. <i>Organic Letters</i> , 2015, 17, 6182-6185.	2.4	77
231	Stereoselective Allylstannane Addition for a Convergent Synthesis of a Complex Molecule. <i>Organic Letters</i> , 2015, 17, 6246-6249.	2.4	7
232	Fatty acid profile of different species of algae of the <i>Cystoseira</i> genus: a nutraceutical perspective. <i>Natural Product Research</i> , 2015, 29, 1264-1270.	1.0	30
233	In vitro antioxidant and inhibitory activity of water decoctions of carob tree (<i>Ceratonia</i>) Tj ETQq1 1 0.784314 rgBT /Overlock 1 2155-2159.	1.0	31
234	6-(Bromomaleimido)hexanoic Acid as a Connector for the Construction of Multiple Branched Peptide Platforms. <i>Organic Letters</i> , 2015, 17, 464-467.	2.4	6

#	ARTICLE	IF	CITATIONS
235	Semipermanent <i>C</i> -Terminal Carboxylic Acid Protecting Group: Application to Solubilizing Peptides and Fragment Condensation. <i>Organic Letters</i> , 2015, 17, 294-297.	2.4	22
236	Optimized Microwave Assisted Synthesis of LL37, a Cathelicidin Human Antimicrobial Peptide. <i>International Journal of Peptide Research and Therapeutics</i> , 2015, 21, 13-20.	0.9	7
237	Development of surface modified biodegradable polymeric nanoparticles to deliver GSE24.2 peptide to cells: A promising approach for the treatment of defective telomerase disorders. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2015, 91, 91-102.	2.0	25
238	A solid-phase combinatorial approach for indoloquinolizidine-peptides with high affinity at D1 and D2 dopamine receptors. <i>European Journal of Medicinal Chemistry</i> , 2015, 97, 173-180.	2.6	11
239	A new quinoxaline-containing peptide induces apoptosis in cancer cells by autophagy modulation. <i>Chemical Science</i> , 2015, 6, 4537-4549.	3.7	19
240	Methods, setup and safe handling for anhydrous hydrogen fluoride cleavage in Boc solid-phase peptide synthesis. <i>Nature Protocols</i> , 2015, 10, 1067-1083.	5.5	41
241	Tetrahydropyranyl, a Nonaromatic Acid-Labile Cys Protecting Group for Fmoc Peptide Chemistry. <i>Organic Letters</i> , 2015, 17, 1680-1683.	2.4	20
242	Chemical Platforms for Peptide Vaccine Constructs. <i>Advances in Protein Chemistry and Structural Biology</i> , 2015, 99, 99-130.	1.0	4
243	On the Mechanism of Phenolic Formylation Mediated by $TiCl_4$ Complexes: Existence of Diradical Intermediates Induced by Valence Tautomerism. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 2111-2118.	1.2	4
244	Hantzsch dihydropyridines: Privileged structures for the formation of well-defined gold nanostars. <i>Journal of Colloid and Interface Science</i> , 2015, 453, 260-269.	5.0	18
245	An efficient solid-phase strategy for total synthesis of naturally occurring amphiphilic marine siderophores: amphibactin-T and moanachelin ala-B. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 4760-4768.	1.5	10
246	EDC·HCl and Potassium Salts of Oxyma and Oxyma β as Superior Coupling Cocktails for Peptide Synthesis. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 3116-3120.	1.2	22
247	A synthetic peptide derived from the D1 domain of flagellin induced the expression of proinflammatory cytokines in fish macrophages. <i>Fish and Shellfish Immunology</i> , 2015, 47, 239-244.	1.6	12
248	The effect of N-methylation of amino acids (Ac-X-OMe) on solubility and conformation: a DFT study. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 9993-10006.	1.5	55
249	Multifunctionalized polyurethane β polyurea nanoparticles: hydrophobically driven self-stratification at the o/w interface modulates encapsulation stability. <i>Journal of Materials Chemistry B</i> , 2015, 3, 7604-7613.	2.9	15
250	BbMP-1, a new metalloproteinase isolated from Bothrops brazili snake venom with in vitro antiplasmodial properties. <i>Toxicon</i> , 2015, 106, 30-41.	0.8	18
251	Single-molecule kinetics and footprinting of DNA bis-intercalation: the paradigmatic case of Thio coraline. <i>Nucleic Acids Research</i> , 2015, 43, 2767-2779.	6.5	30
252	An immunochemical strategy based on peptidoglycan synthetic peptide epitopes to diagnose <i>Staphylococcus aureus</i> infections. <i>Analytica Chimica Acta</i> , 2015, 889, 203-211.	2.6	6

#	ARTICLE	IF	CITATIONS
253	Gated Mesoporous Silica Nanoparticles Using a Double-Role Circular Peptide for the Controlled and Target-Preferential Release of Doxorubicin in CXCR4-Expressing Lymphoma Cells. <i>Advanced Functional Materials</i> , 2015, 25, 687-695.	7.8	54
254	Peptide synthesis beyond DMF: THF and ACN as excellent and friendlier alternatives. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 2393-2398.	1.5	69
255	Addition of Vinylmetallic Reagents to Chiral α -Formyltetrahydrofuran. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 235-241.	1.2	7
256	Phenolic composition, antioxidant potential and in vitro inhibitory activity of leaves and acorns of <i>Quercus suber</i> on key enzymes relevant for hyperglycemia and Alzheimer's disease. <i>Industrial Crops and Products</i> , 2015, 64, 45-51.	2.5	80
257	A simple protocol for combinatorial cyclic depsipeptide libraries sequencing by matrix-assisted laser desorption/ionisation mass spectrometry. <i>Journal of Peptide Science</i> , 2015, 21, 40-45.	0.8	13
258	Phormidolides B and C, Cytotoxic Agents from the Sea: Enantioselective Synthesis of the Macrocyclic Core. <i>Chemistry - A European Journal</i> , 2015, 21, 150-156.	1.7	26
259	α -Ketoamino acid ester derivatives as promising MAO inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 70-74.	1.0	13
260	<i>Botryococcus braunii</i> and <i>Nannochloropsis oculata</i> extracts inhibit cholinesterases and protect human dopaminergic SH-SY5Y cells from H ₂ O ₂ -induced cytotoxicity. <i>Journal of Applied Phycology</i> , 2015, 27, 839-848.	1.5	31
261	Trends to Acid-Labile Cys Protecting Groups: Thp as an Efficient and Non-Aromatic Cys Protecting Group for Fmoc Chemistry. , 2015, , .		3
262	A novel prohibitin-binding compound induces the mitochondrial apoptotic pathway through NOXA and BIM upregulation. <i>Oncotarget</i> , 2015, 6, 41750-41765.	0.8	29
263	Palladium-catalyzed coupling reactions for the preparation of concatenated azoles. <i>Arkivoc</i> , 2015, 2015, 34-43.	0.3	0
264	Animal Toxins and Their Advantages in Biotechnology and Pharmacology. <i>BioMed Research International</i> , 2014, 2014, 1-2.	0.9	11
265	Airylation of Histidine Residues of Bothrops jararacussu Venom Proteins and Isolated Phospholipases $A \times 2$ A Biotechnological Tool to Improve the Production of Antibodies. <i>BioMed Research International</i> , 2014, 2014, 1-12.	0.9	10
266	Thiopeptide Antibiotics: Retrospective and Recent Advances. <i>Marine Drugs</i> , 2014, 12, 317-351.	2.2	151
267	A Novel Phospholipase A ₂ (D49) from the Venom of the <i>Crotalus oreganus abyssus</i> (North American) Tj ETQq1 1 0.784314 rgBT/Overl	0.9	10
268	α -Clicking-Porphyrins to Magnetic Nanoparticles for Photodynamic Therapy. <i>ChemPlusChem</i> , 2014, 79, 90-98.	1.3	25
269	Gold nanoparticles for photothermally controlled drug release. <i>Nanomedicine</i> , 2014, 9, 2023-2039.	1.7	45
270	Peptide Affinity Chromatography Based on Combinatorial Strategies for Protein Purification. <i>Methods in Molecular Biology</i> , 2014, 1129, 277-302.	0.4	15

#	ARTICLE	IF	CITATIONS
271	Mesopattern of immobilised bone morphogenetic protein-2 created by microcontact printing and dip-pen nanolithography influence C2C12 cell fate. <i>RSC Advances</i> , 2014, 4, 56809-56815.	1.7	10
272	Isolation and Biochemical Characterization of a New Thrombin-Like Serine Protease from <i>Bothrops pirajai</i> Snake Venom. <i>BioMed Research International</i> , 2014, 2014, 1-13.	0.9	18
273	Liposomes containing NY-ESO-1/tetanus toxoid and adjuvant peptides targeted to human dendritic cells via the Fc receptor for cancer vaccines. <i>Nanomedicine</i> , 2014, 9, 435-449.	1.7	32
274	Semi-synthesis of acylated triterpenes from olive-oil industry wastes for the development of anticancer and anti-HIV agents. <i>European Journal of Medicinal Chemistry</i> , 2014, 74, 278-301.	2.6	39
275	Covalent immobilization of hLf1-11 peptide on a titanium surface reduces bacterial adhesion and biofilm formation. <i>Acta Biomaterialia</i> , 2014, 10, 3522-3534.	4.1	125
276	Fatty acid composition and biological activities of <i>Isochrysis galbana</i> T-ISO, <i>Tetraselmis</i> sp. and <i>Scenedesmus</i> sp.: possible application in the pharmaceutical and functional food industries. <i>Journal of Applied Phycology</i> , 2014, 26, 151-161.	1.5	66
277	Structural glance into a novel anti-staphylococcal peptide. <i>Biopolymers</i> , 2014, 102, 49-57.	1.2	5
278	Immobilized N-Chlorosuccinimide as a Friendly Peptide Disulfide-Forming Reagent. <i>ACS Combinatorial Science</i> , 2014, 16, 160-163.	3.8	15
279	Stellatolides, a New Cyclodepsipeptide Family from the Sponge <i>Ecionemia acervus</i> : Isolation, Solid-Phase Total Synthesis, and Full Structural Assignment of Stellatolide A. <i>Journal of the American Chemical Society</i> , 2014, 136, 6754-6762.	6.6	37
280	Thiopeptide Engineering: A Multidisciplinary Effort towards Future Drugs. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 6602-6616.	7.2	80
281	BOP- <i>OXy</i> , BOP- <i>OBt</i> , and BOP- <i>OA</i> t: novel organophosphinic coupling reagents useful for solution and solid-phase peptide synthesis. <i>Journal of Peptide Science</i> , 2014, 20, 1-6.	0.8	8
282	Solid-phase peptide synthesis (SPPS), C-terminal vs. side-chain anchoring: a reality or a myth. <i>Amino Acids</i> , 2014, 46, 1827-1838.	1.2	13
283	Disulfide Formation Strategies in Peptide Synthesis. <i>European Journal of Organic Chemistry</i> , 2014, 2014, 3519-3530.	1.2	87
284	Synthesis of All the Diastereomers of 2-Amino-3-hydroxy-4,5-dimethylhexanoic Acid. <i>European Journal of Organic Chemistry</i> , 2014, 2014, 44-47.	1.2	4
285	The potential of N-alkoxymethyl groups as peptide backbone protectants. <i>Tetrahedron Letters</i> , 2014, 55, 184-188.	0.7	6
286	Multifaceted Roles of Disulfide Bonds. Peptides as Therapeutics. <i>Chemical Reviews</i> , 2014, 114, 901-926.	23.0	477
287	2-Methoxy-4-methylsulfinylbenzyl: A Backbone Amide Safety-Catch Protecting Group for the Synthesis and Purification of Difficult Peptide Sequences. <i>Chemistry - A European Journal</i> , 2014, 20, 15031-15039.	1.7	20
288	Triazene as a Powerful Tool for Solid-Phase Derivatization of Phenylalanine Containing Peptides: Zygosporamide Analogues as a Proof of Concept. <i>Journal of Organic Chemistry</i> , 2014, 79, 11409-11415.	1.7	13

#	ARTICLE	IF	CITATIONS
289	Immobilized Coupling Reagents: Synthesis of Amides/Peptides. ACS Combinatorial Science, 2014, 16, 579-601.	3.8	22
290	Proline N-oxides: modulators of the 3D conformation of linear peptides through α -NO-turns. Organic and Biomolecular Chemistry, 2014, 12, 4479.	1.5	14
291	Synthesis of cyclohexapeptides as antimalarial and anti-trypanosomal agents. MedChemComm, 2014, 5, 1309-1316.	3.5	12
292	Tackling Lipophilicity of Peptide Drugs: Replacement of the Backbone <i>N</i> -Methyl Group of Cilengitide by <i>N</i> -Oligoethylene Glycol (<i>N</i> -OEG) Chains. Bioconjugate Chemistry, 2014, 25, 11-17.	1.8	16
293	Polythiazole linkers as functional rigid connectors: a new RGD cyclopeptide with enhanced integrin selectivity. Chemical Science, 2014, 5, 3929.	3.7	10
294	Multivalent dendrimers presenting spatially controlled clusters of binding epitopes in thermoresponsive hyaluronan hydrogels. Acta Biomaterialia, 2014, 10, 4340-4350.	4.1	22
295	A Trifluorinated Thiazoline Scaffold Leading to Proapoptotic Agents Targeting Prohibitins. Angewandte Chemie, 2014, 126, 10314-10318.	1.6	1
296	Dissecting the Structure of Thiopeptides: Assessment of Thiazoline and Tail Moieties of Baringolin and Antibacterial Activity Optimization. Journal of Medicinal Chemistry, 2014, 57, 4185-4195.	2.9	23
297	Novel Peptide-Based Platform for the Dual Presentation of Biologically Active Peptide Motifs on Biomaterials. ACS Applied Materials & Interfaces, 2014, 6, 6525-6536.	4.0	73
298	A Trifluorinated Thiazoline Scaffold Leading to Proapoptotic Agents Targeting Prohibitins. Angewandte Chemie - International Edition, 2014, 53, 10150-10154.	7.2	35
299	Facile and Mild Synthesis of Linear and Cyclic Peptides via Thioesters. Organic Letters, 2014, 16, 3922-3925.	2.4	16
300	High Control, Fast Growth OEG-Based Dendron Synthesis via a Sequential Two-Step Process of Copper-Free Diazo Transfer and Click Chemistry. Macromolecules, 2014, 47, 2585-2591.	2.2	3
301	Thioester Bonds of Thiocoraline Can Be Replaced with NMe-Amide Bridges without Affecting Its DNA-Binding Properties. ACS Medicinal Chemistry Letters, 2014, 5, 45-50.	1.3	5
302	Selective Formation of a <i>Z</i> -Trisubstituted Double Bond Using a 1-(<i>tert</i> -Butyl)tetrazolyl Sulfone. Journal of Organic Chemistry, 2014, 79, 10648-10654.	1.7	9
303	Oxyma-B, an excellent racemization suppressor for peptide synthesis. Organic and Biomolecular Chemistry, 2014, 12, 8379-8385.	1.5	28
304	Microreactors for peptide synthesis: looking through the eyes of twenty first century !!! Amino Acids, 2014, 46, 2091-2104.	1.2	17
305	Linear versus branched poly-lysine/arginine as polarity enhancer tags. Organic and Biomolecular Chemistry, 2014, 12, 7194-7196.	1.5	8
306	Antimicrobial Peptides from Skin Secretions of <i>Hypsiboas pulchellus</i> (Anura: Hylidae). Journal of Natural Products, 2014, 77, 831-841.	1.5	27

#	ARTICLE	IF	CITATIONS
307	Controlling Multivalency and Multimodality: Up to Pentamodal Dendritic Platforms Based on Diethylenetriaminepentaacetic Acid Cores. <i>Organic Letters</i> , 2014, 16, 1318-1321.	2.4	13
308	Cysteine Pseudoprolines for Thiol Protection and Peptide Macrocyclization Enhancement in Fmoc-Based Solid-Phase Peptide Synthesis. <i>Organic Letters</i> , 2014, 16, 1772-1775.	2.4	22
309	Solid-Phase Library Synthesis of Bi-Functional Derivatives of Oleanolic and Maslinic Acids and Their Cytotoxicity on Three Cancer Cell Lines. <i>ACS Combinatorial Science</i> , 2014, 16, 428-447.	3.8	30
310	Amphiphilic Cationic Carbosilane-PEG Dendrimers: Synthesis and Applications in Gene Therapy. <i>European Journal of Medicinal Chemistry</i> , 2014, 76, 43-52.	2.6	35
311	Morphological characterization of fullerene-androsterone conjugates. <i>Beilstein Journal of Nanotechnology</i> , 2014, 5, 374-379.	1.5	9
312	TOMBU and COMBU as Novel Uronium-Type Peptide Coupling Reagents Derived from Oxyma-B. <i>Molecules</i> , 2014, 19, 18953-18965.	1.7	11
313	Structure-activity relationship of 1-desamino-8-D-arginine vasopressin as an antiproliferative agent on human vasopressin V2 receptor-expressing cancer cells. <i>Molecular Medicine Reports</i> , 2014, 9, 2568-2572.	1.1	13
314	Chiral Thiazoline and Thiazole Building Blocks for the Synthesis of Peptide-Derived Natural Products. <i>Current Topics in Medicinal Chemistry</i> , 2014, 14, 1244-1256.	1.0	14
315	Abstract 4601: Astrocytic elevated gene 1 (AEG1) a target for pharmacological anticancer intervention. , 2014, , .		1
316	Synthesis of C-2 Arylated Tryptophan Amino Acids and Related Compounds through Palladium-Catalyzed C-H Activation. <i>Journal of Organic Chemistry</i> , 2013, 78, 8129-8135.	1.7	107
317	Constella, (EU) Linzess, (USA): the last milestone in the long journey of the peptide linaclotide and its implications for the future of peptide drugs. <i>Future Medicinal Chemistry</i> , 2013, 5, 291-300.	1.1	10
318	Extracts from <i>Quercus</i> sp. acorns exhibit in vitro neuroprotective features through inhibition of cholinesterase and protection of the human dopaminergic cell line SH-SY5Y from hydrogen peroxide-induced cytotoxicity. <i>Industrial Crops and Products</i> , 2013, 45, 114-120.	2.5	32
319	N-chlorosuccinimide, an efficient peptide disulfide bond-forming reagent in aqueous solution. <i>RSC Advances</i> , 2013, 3, 14277.	1.7	16
320	Synthesis and NMR elucidation of pentacycloundecane-derived hydroxy acid peptides as potential anti-HIV-1 agents. <i>Structural Chemistry</i> , 2013, 24, 1461-1471.	1.0	3
321	Friendly Strategy to Prepare Encoded One Bead-One Compound Cyclic Peptide Library. <i>ACS Combinatorial Science</i> , 2013, 15, 525-529.	3.8	14
322	The first total synthesis of the cyclodepsipeptide pipecolidepsin A. <i>Nature Communications</i> , 2013, 4, 2352.	5.8	49
323	RADA16: A Tough Peptide - Strategies for Synthesis and Purification. <i>European Journal of Organic Chemistry</i> , 2013, 2013, 5871-5878.	1.2	9
324	The Backbone N-(4-Azidobutyl) Linker for the Preparation of Peptide Chimera. <i>Organic Letters</i> , 2013, 15, 4572-4575.	2.4	11

#	ARTICLE	IF	CITATIONS
325	Inhibitory effect of short cationic homopeptides against Gram-positive bacteria. <i>Journal of Peptide Science</i> , 2013, 19, 792-800.	0.8	26
326	Head-to-Side-Chain-Cyclodepsipeptides of Marine Origin. <i>Marine Drugs</i> , 2013, 11, 1693-1717.	2.2	50
327	Rescuing Biological Activity from Synthetic Phakellistatin 19. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 9780-9788.	2.9	25
328	Multifunctional Nanovesicle-Bioactive Conjugates Prepared by a One-Step Scalable Method Using CO ₂ -Expanded Solvents. <i>Nano Letters</i> , 2013, 13, 3766-3774.	4.5	40
329	Oxyma: a Strong Acylation-Promoting, CTC Resin-Friendly Coupling Additive. <i>European Journal of Organic Chemistry</i> , 2013, 2013, 6372-6378.	1.2	29
330	From 2,6-Dichloronicotinic Acid to Thiopeptide Cores. <i>European Journal of Organic Chemistry</i> , 2013, 2013, 6404-6419.	1.2	6
331	Efficient cysteine labelling of peptides with N-succinimidyl 4-[¹⁸ F]fluorobenzoate: stability study and in vivo biodistribution in rats by positron emission tomography (PET). <i>RSC Advances</i> , 2013, 3, 8028.	1.7	5
332	N-Triethylene glycol (N-TEG) as a surrogate for the N-methyl group: application to Sansalvamide A peptide analogs. <i>Chemical Communications</i> , 2013, 49, 6430.	2.2	17
333	Synthesis and biological evaluation of a post-synthetically modified Trp-based diketopiperazine. <i>MedChemComm</i> , 2013, 4, 1171.	3.5	16
334	N-Chlorosuccinimide, an Efficient Reagent for On-Resin Disulfide Formation in Solid-Phase Peptide Synthesis. <i>Organic Letters</i> , 2013, 15, 616-619.	2.4	45
335	Effective and Versatile Strategy for the Total Solid-Phase Synthesis of Alkanethiols for Biological Applications. <i>European Journal of Organic Chemistry</i> , 2013, 2013, 1233-1239.	1.2	2
336	Use of an Internal Reference for the Quantitative HPLC-UV Analysis of Solid-Phase Reactions: A Case Study of 2-Chlorotriyl Chloride Resin. <i>ACS Combinatorial Science</i> , 2013, 15, 229-234.	3.8	6
337	Tetrahydrofuran-Containing Macrolides: A Fascinating Gift from the Deep Sea. <i>Chemical Reviews</i> , 2013, 113, 4567-4610.	23.0	275
338	Stable Conjugates of Peptides with Gold Nanorods for Biomedical Applications with Reduced Effects on Cell Viability. <i>ACS Applied Materials & Interfaces</i> , 2013, 5, 4076-4085.	4.0	67
339	Handles for Fmoc Solid-Phase Synthesis of Protected Peptides. <i>ACS Combinatorial Science</i> , 2013, 15, 217-228.	3.8	66
340	Enzyme-Labile Protecting Groups for the Synthesis of Natural Products: Solid-Phase Synthesis of Thiocoraline. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 5726-5730.	7.2	18
341	Biocompatible, multifunctional, and well-defined OEG-based dendritic platforms for biomedical applications. <i>Organic and Biomolecular Chemistry</i> , 2013, 11, 4109.	1.5	14
342	COMU: scope and limitations of the latest innovation in peptide acyl transfer reagents. <i>Journal of Peptide Science</i> , 2013, 19, 408-414.	0.8	37

#	ARTICLE	IF	CITATIONS
343	Electrostatic Binding and Hydrophobic Collapse of Peptide–Nucleic Acid Aggregates Quantified Using Force Spectroscopy. <i>ACS Nano</i> , 2013, 7, 5102-5113.	7.3	26
344	Orthogonal Chemistry for the Synthesis of Thiocoraline–Triostin Hybrids. Exploring their Structure–Activity Relationship. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 5587-5600.	2.9	23
345	Wang Linker Free of Side Reactions. <i>Organic Letters</i> , 2013, 15, 246-249.	2.4	17
346	Imidazole-1-sulfonyl Azide-Based Diazo-Transfer Reaction for the Preparation of Azido Solid Supports for Solid-Phase Synthesis. <i>ACS Combinatorial Science</i> , 2013, 15, 331-334.	3.8	24
347	Polyproline–OEG Co–Oligomeric Dendrimers: A Family of Highly Branched Polyproline Macromolecules. <i>European Journal of Organic Chemistry</i> , 2013, 2013, 8279-8287.	1.2	0
348	Total Synthesis and Stereochemical Assignment of Baringolin. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 7818-7821.	7.2	37
349	OxymaPure/DIC: An Efficient Reagent for the Synthesis of a Novel Series of 4-[2-(2-Acetylamino-phenyl)-2-oxo-acetyl-amino] Benzoyl Amino Acid Ester Derivatives. <i>Molecules</i> , 2013, 18, 14747-14759.	1.7	20
350	Understanding Acid Lability of Cysteine Protecting Groups. <i>Molecules</i> , 2013, 18, 5155-5162.	1.7	14
351	Neurotoxicity of Prion Peptides Mimicking the Central Domain of the Cellular Prion Protein. <i>PLoS ONE</i> , 2013, 8, e70881.	1.1	20
352	Liquid phase organic synthesis of 3,5-disubstituted 1,3,5-thiadiazinane-2-thione derivatives on polyethylene glycol (PEG) support. <i>Arkivoc</i> , 2013, 2012, 326-338.	0.3	5
353	Low-epimerization Peptide Bond Formation with Oxyma Pure: Preparation of Z-L-Phg-Val-OMe. <i>Organic Syntheses</i> , 2013, 90, 306.	1.0	1
354	Synthesis and Thermal Properties of Novel Polyamides Containing $\hat{\pm}$ -Amino Acid Moieties: Structure-Property Relationship. <i>Journal of Macromolecular Science - Pure and Applied Chemistry</i> , 2012, 49, 41-54.	1.2	5
355	Enhancing immunogenicity and cross-reactivity of HIV-1 antigens by <i>in vivo</i> targeting to dendritic cells. <i>Nanomedicine</i> , 2012, 7, 1591-1610.	1.7	5
356	Drug Delivery: Surface-Adhered Composite Poly(Vinyl Alcohol) Physical Hydrogels: Polymer-Supported Delivery of Therapeutic Small Molecules (<i>Adv. Healthcare Mater.</i> 6/2012). <i>Advanced Healthcare Materials</i> , 2012, 1, 790-790.	3.9	2
357	Oxime-Based Carbonates as Useful Reagents for Both N-Protection and Peptide Coupling. <i>Molecules</i> , 2012, 17, 14361-14376.	1.7	1
358	Trimethoxyphenylthio as a Highly Labile Replacement for <i>tert</i> -Butylthio Cysteine Protection in Fmoc Solid Phase Synthesis. <i>Organic Letters</i> , 2012, 14, 5468-5471.	2.4	50
359	A universal strategy for preparing protected C-terminal peptides on the solid phase through an intramolecular click chemistry-based handle. <i>Chemical Communications</i> , 2012, 48, 2313.	2.2	9
360	Eco-Friendly Combination of the Immobilized PGA Enzyme and the <i>S</i> -Phacm Protecting Group for the Synthesis of Cys-Containing Peptides. <i>Chemistry - A European Journal</i> , 2012, 18, 16166-16176.	1.7	27

#	ARTICLE	IF	CITATIONS
361	H-bonding promotion of peptide solubility and cyclization by fluorinated alcohols. RSC Advances, 2012, 2, 2729.	1.7	13
362	Introducing an Asp-Pro Linker in the Synthesis of Random One-Bead-One-Compound Hexapeptide Libraries Compatible with ESI-MS Analysis. ACS Combinatorial Science, 2012, 14, 145-149.	3.8	7
363	Synthesis and In Vivo Evaluation of the Biodistribution of a ¹⁸ F-Labeled Conjugate Gold-Nanoparticle-Peptide with Potential Biomedical Application. Bioconjugate Chemistry, 2012, 23, 399-408.	1.8	100
364	The marine halophytes <i>Carpobrotus edulis</i> L. and <i>Arthrocnemum macrostachyum</i> L. are potential sources of nutritionally important PUFAs and metabolites with antioxidant, metal chelating and anticholinesterase inhibitory activities. Botanica Marina, 2012, 55, 281-288.	0.6	34
365	Efficient ¹³ C-amino-proline-derived cell penetrating peptide-superparamagnetic iron oxide nanoparticle conjugates via aniline-catalyzed oxime chemistry as bimodal imaging nanoagents. Chemical Communications, 2012, 48, 5322.	2.2	21
366	Surface-Adhered Composite Poly(Vinyl Alcohol) Physical Hydrogels: Polymersome-Aided Delivery of Therapeutic Small Molecules. Advanced Healthcare Materials, 2012, 1, 791-795.	3.9	36
367	Therapeutic peptides. Future Medicinal Chemistry, 2012, 4, 1527-1531.	1.1	261
368	Rapid and high-yielding cysteine labelling of peptides with N-succinimidyl 4-[¹⁸ F]fluorobenzoate. Chemical Communications, 2012, 48, 6118.	2.2	12
369	Acid-Labile Cys-Protecting Groups for the Fmoc-t-Bu Strategy: Filling the Gap. Organic Letters, 2012, 14, 5472-5475.	2.4	34
370	Vascular effects and electrolyte homeostasis of the natriuretic peptide isolated from <i>Crotalus oreganus abyssus</i> (North American Grand Canyon rattlesnake) venom. Peptides, 2012, 36, 206-212.	1.2	18
371	Targeting Nanoparticles to Dendritic Cells for Immunotherapy. Methods in Enzymology, 2012, 509, 143-163.	0.4	110
372	Thiadiazines, N,N-Heterocycles of Biological Relevance. Molecules, 2012, 17, 7612-7628.	1.7	23
373	Cancer Prognostics by Direct Detection of p53-Antibodies on Gold Surfaces by Impedance Measurements. Small, 2012, 8, 2106-2115.	5.2	20
374	Biosensors: Cancer Prognostics by Direct Detection of p53-Antibodies on Gold Surfaces by Impedance Measurements (Small 13/2012). Small, 2012, 8, 1962-1962.	5.2	0
375	Solid-Phase Synthesis of N-Me-IB-01212, a Highly N-Methylated Cyclic Peptide. Organic Letters, 2012, 14, 612-615.	2.4	35
376	Screening of N-Alkyl-Cyanoacetamido Oximes as Substitutes for N-Hydroxysuccinimide. ChemistryOpen, 2012, 1, 147-152.	0.9	14
377	Use of Oxyma as pH modulatory agent to be used in the prevention of base-driven side reactions and its effect on 2-chlorotriyl chloride resin. Biopolymers, 2012, 98, 89-97.	1.2	38
378	Microalgae of different phyla display antioxidant, metal chelating and acetylcholinesterase inhibitory activities. Food Chemistry, 2012, 131, 134-140.	4.2	91

#	ARTICLE	IF	CITATIONS
379	Cell adhesion and focal contact formation on linear RGD molecular gradients: study of non-linear concentration dependence effects. <i>Nanomedicine: Nanotechnology, Biology, and Medicine</i> , 2012, 8, 432-439.	1.7	39
380	Cell-penetrating $\hat{1}^3$ -peptide/antimicrobial undecapeptide conjugates with anticancer activity. <i>Tetrahedron</i> , 2012, 68, 4406-4412.	1.0	12
381	Cyanoacetamide-based oxime carbonates: an efficient, simple alternative for the introduction of Fmoc with minimal dipeptide formation. <i>Tetrahedron</i> , 2012, 68, 3056-3062.	1.0	10
382	Orthogonal Protecting Groups in the Synthesis of Tryptophanyl- $\hat{1}^6$ -Hexahydropyrroloindoles. <i>European Journal of Organic Chemistry</i> , 2012, 2012, 67-73.	1.2	10
383	Myoblast Cell Interaction with Polydopamine Coated Liposomes. <i>Biointerphases</i> , 2012, 7, 8.	0.6	43
384	Polymers and Drug Delivery Systems. <i>Current Drug Delivery</i> , 2012, 9, 367-394.	0.8	210
385	Progress on lamellarins. <i>MedChemComm</i> , 2011, 2, 689-697.	3.5	80
386	Study of Various Presentation Forms for a Peptide Mimetic of <i>Neisseria meningitidis</i> Serogroup B Capsular Polysaccharide. <i>Bioconjugate Chemistry</i> , 2011, 22, 33-41.	1.8	3
387	Trivalent PEGylated Platform for the Conjugation of Bioactive Compounds. <i>Bioconjugate Chemistry</i> , 2011, 22, 2172-2178.	1.8	1
388	Biotin Ergopeptide Probes for Dopamine Receptors. <i>Journal of Medicinal Chemistry</i> , 2011, 54, 1080-1090.	2.9	13
389	Affinity Chromatography Based on a Combinatorial Strategy for rErythropoietin Purification. <i>ACS Combinatorial Science</i> , 2011, 13, 251-258.	3.8	28
390	Antioxidant and Cytotoxic Activities of Carob Tree Fruit Pulps Are Strongly Influenced by Gender and Cultivar. <i>Journal of Agricultural and Food Chemistry</i> , 2011, 59, 7005-7012.	2.4	53
391	4-(4,6-Di[2,2,2-trifluoroethoxy]-1,3,5-triazin-2-yl)-4-methylmorpholinium Tetrafluoroborate. Triazine-Based Coupling Reagents Designed for Coupling Sterically Hindered Substrates. <i>Journal of Organic Chemistry</i> , 2011, 76, 4506-4513.	1.7	22
392	Targeting Nanosystems to Human DCs via Fc Receptor as an Effective Strategy to Deliver Antigen for Immunotherapy. <i>Molecular Pharmaceutics</i> , 2011, 8, 104-116.	2.3	85
393	Total Synthesis of Aeruginazole A. <i>Organic Letters</i> , 2011, 13, 4648-4651.	2.4	18
394	Acridine and quindoline oligomers linked through a 4-aminoproline backbone prefer G-quadruplex structures. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2011, 1810, 769-776.	1.1	14
395	Eco-Friendly Methodology to Prepare N-Heterocycles Related to Dihydropyridines: Microwave-Assisted Synthesis of Alkyl 4-Arylsubstituted-6-chloro-5-formyl-2-methyl-1,4-dihydropyridine-3-carboxylate and 4-Arylsubstituted-4,7-dihydrofuro[3,4-b]pyridine-2,5(1H,3H)-dione. <i>Molecules</i> , 2011, 16, 9620-9635.	1.7	8
396	A natural peptide and its variants derived from the processing of infectious pancreatic necrosis virus (IPNV) displaying enhanced antimicrobial activity: A novel alternative for the control of bacterial diseases. <i>Peptides</i> , 2011, 32, 852-858.	1.2	20

#	ARTICLE	IF	CITATIONS
397	Solid-Phase Synthesis of a Library of Amphipatic Hydantoins. Discovery of New Hits for TRPV1 Blockade. ACS Combinatorial Science, 2011, 13, 458-465.	3.8	10
398	Synthesis and Aminolysis of 2,4-Dinitrophenyl and 5-Nitropyridine <i>N</i> -Hydroxy Oxime Derivatives. Bulletin of the Chemical Society of Japan, 2011, 84, 633-639.	2.0	6
399	Aspartimide formation in peptide chemistry: occurrence, prevention strategies and the role of N-hydroxylamines. Tetrahedron, 2011, 67, 8595-8606.	1.0	76
400	Highly efficient, multigram and enantiopure synthesis of (S)-2-(2,4-bithiazol-2-yl)pyrrolidine. Tetrahedron Letters, 2011, 52, 5435-5437.	0.7	10
401	Leishmania mexicana: LACK (Leishmania homolog of receptors for activated C-kinase) is a plasminogen binding protein. Experimental Parasitology, 2011, 127, 752-761.	0.5	18
402	Peptide Coupling Reagents, More than a Letter Soup. Chemical Reviews, 2011, 111, 6557-6602.	23.0	922
403	Functionalization of gold surfaces: recent developments and applications. Journal of Materials Science, 2011, 46, 7643-7648.	1.7	23
404	Phytochemical Profile, Antioxidant and Cytotoxic Activities of the Carob Tree (Ceratonia siliqua L.) Germ Flour Extracts. Plant Foods for Human Nutrition, 2011, 66, 78-84.	1.4	64
405	Effect of a Pool of Peptides Isolated from Crotalus durissus terrificus (South American Rattlesnake) Venom on Glucose Levels of Mice Fed on a High-Fat Diet. International Journal of Peptide Research and Therapeutics, 2011, 17, 225-230.	0.9	3
406	Total synthesis of a depsidomycin analogue by convergent solid-phase peptide synthesis and macrolactonization strategy for antitubercular activity. Journal of Peptide Science, 2011, 17, 683-689.	0.8	26
407	Optimized Fmoc solid-phase synthesis of the cysteine-rich peptide linaclotide. Biopolymers, 2011, 96, 69-80.	1.2	50
408	Structure, Bioactivity and Synthesis of Natural Products with Hexahydropyrrolo[2,3-b]indole. Chemistry - A European Journal, 2011, 17, 1388-1408.	1.7	429
409	TRPV1 modulators: Structure-activity relationships using a rational combinatorial approach. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 3541-3545.	1.0	8
410	High-throughput preparation of alkyl 4-aryl substituted-2-methyl-6-thioxo-1,4,5,6-tetrahydropyridine-3-carboxylates under microwave irradiation. Arkivoc, 2011, 2011, 125-141.	0.3	4
411	Synthesis of orthogonally protected l-threo- β -ethoxyasparagine. Amino Acids, 2010, 39, 161-165.	1.2	8
412	Targeted PLGA nano- but not microparticles specifically deliver antigen to human dendritic cells via DC-SIGN in vitro. Journal of Controlled Release, 2010, 144, 118-126.	4.8	242
413	Solid-Phase Synthesis of Aza-Kahalalide F Analogues: (2 <i>R</i> ,3 <i>R</i>)-2-Amino-3-azidobutanoic Acid, as Precursor of the Aza-Threonine. European Journal of Organic Chemistry, 2010, 2010, 2536-2543.	1.2	13
414	Oxime Carbonates: Novel Reagents for the Introduction of Fmoc and Alloc Protecting Groups, Free of Side Reactions. European Journal of Organic Chemistry, 2010, 2010, 3275-3280.	1.2	16

#	ARTICLE	IF	CITATIONS
415	2,2,4,6,7â€Pentamethylâ€2,3â€dihydrobenzofuranâ€5â€methyl (Pbfm) as an Alternative to the Trityl Group for the Sideâ€Chain Protection of Cysteine and Asparagine/Glutamine. European Journal of Organic Chemistry, 2010, 2010, 3631-3640.	1.2	11
416	A Novel Family of Onium Salts Based Upon Isonitroso Meldrum's Acid Proves Useful as Peptide Coupling Reagents. European Journal of Organic Chemistry, 2010, 2010, 3641-3649.	1.2	32
417	Streamlined Access to Functionalized Chromenes and Quinolines using Domino Reactions of Salicylic Aldehydes and Methyl 4â€Chloroâ€2â€butynoate. European Journal of Organic Chemistry, 2010, 2010, 5373-5379.	1.2	16
418	Capsosomes with Multilayered Subcompartments: Assembly and Loading with Hydrophobic Cargo. Advanced Functional Materials, 2010, 20, 59-66.	7.8	111
419	Postsynthetic Modification of Peptides: Chemoselective Câ€Arylation of Tryptophan Residues. Chemistry - A European Journal, 2010, 16, 1124-1127.	1.7	159
420	Amideâ€toâ€Ester Substitution Allows Fineâ€Tuning of the Cyclopeptide Conformational Ensemble. Angewandte Chemie - International Edition, 2010, 49, 2732-2737.	7.2	16
421	Microwave assisted SPPS of amylin and its toxicity of the pure product to RINâ€5F cells. Biopolymers, 2010, 94, 323-330.	1.2	17
422	Sample preparation for sequencing hits from one-beadâ€one-peptide combinatorial libraries by matrix-assisted laser desorption/ionization time-of-flight mass spectrometry. Analytical Biochemistry, 2010, 400, 295-297.	1.1	23
423	COMU: A third generation of uroniumâ€type coupling reagents. Journal of Peptide Science, 2010, 16, 6-9.	0.8	97
424	A convenient microwaveâ€enhanced solidâ€phase synthesis of short chain <i>N</i>-methylâ€rich peptides. Journal of Peptide Science, 2010, 16, 136-140.	0.8	29
425	Improved antimicrobial activity of hâ€lysozyme (107â€115) by rational Ala substitution. Journal of Peptide Science, 2010, 16, 424-429.	0.8	12
426	ChemMatrix^{Â®} for complex peptides and combinatorial chemistry. Journal of Peptide Science, 2010, 16, 675-678.	0.8	53
427	Solid-Phase Synthesis of New Trp(Nps)-Containing Dipeptide Derivatives as TRPV1 Channel Blockers. Molecules, 2010, 15, 4924-4933.	1.7	1
428	Synthesis of 2-(4,6-Dimethoxy-1,3,5-triazin-2-yloxyimino) Derivatives: Application in Solution Peptide Synthesis. Molecules, 2010, 15, 9403-9417.	1.7	6
429	A Hybrid Indoloquinolizidine Peptide as Allosteric Modulator of Dopamine D1 Receptors. Journal of Pharmacology and Experimental Therapeutics, 2010, 332, 876-885.	1.3	13
430	The Sea as a Source of New Drugs. , 2010, , 237-249.		4
431	Isolation, Structural Assignment, and Total Synthesis of Barmumycin. Journal of Organic Chemistry, 2010, 75, 8508-8515.	1.7	33
432	Engineering Advanced Capsosomes: Maximizing the Number of Subcompartments, Cargo Retention, and Temperature-Triggered Reaction. ACS Nano, 2010, 4, 1351-1361.	7.3	139

#	ARTICLE	IF	CITATIONS
433	Total Regioselective Control of Tartaric Acid. <i>Journal of Organic Chemistry</i> , 2010, 75, 5746-5749.	1.7	4
434	The Antitumoral Depsipeptide IB-01212 Kills <i>Leishmania</i> through an Apoptosis-like Process Involving Intracellular Targets. <i>Molecular Pharmaceutics</i> , 2010, 7, 1608-1617.	2.3	23
435	Improving the brain delivery of gold nanoparticles by conjugation with an amphipathic peptide. <i>Nanomedicine</i> , 2010, 5, 897-913.	1.7	103
436	PyOxP and PyOxB: the Oxyma-based novel family of phosphonium salts. <i>Organic and Biomolecular Chemistry</i> , 2010, 8, 3665.	1.5	41
437	Amphiphilic peptides and their cross-disciplinary role as building blocks for nanoscience. <i>Chemical Society Reviews</i> , 2010, 39, 241-263.	18.7	236
438	Multifunctionalized Gold Nanoparticles with Peptides Targeted to Gastrin-Releasing Peptide Receptor of a Tumor Cell Line. <i>Bioconjugate Chemistry</i> , 2010, 21, 1070-1078.	1.8	70
439	A novel dipeptidomimetic containing a cyclic threonine. <i>Chemical Communications</i> , 2010, 46, 1266.	2.2	13
440	Synthesis and Aminolysis of N,N-Diethyl Carbamic Ester of HOBt Derivatives. <i>Bulletin of the Korean Chemical Society</i> , 2010, 31, 75-81.	1.0	10
441	Oxyma: An Efficient Additive for Peptide Synthesis to Replace the Benzotriazole-Based HOBt and HOAt with a Lower Risk of Explosion ^[1] . <i>Chemistry - A European Journal</i> , 2009, 15, 9394-9403.	1.7	326
442	COMU: A Safer and More Effective Replacement for Benzotriazole-Based Uronium Coupling Reagents. <i>Chemistry - A European Journal</i> , 2009, 15, 9404-9416.	1.7	260
443	Structure-Activity Relationships of SSAO/VAP-1 Arylalkylamine-Based Substrates. <i>ChemMedChem</i> , 2009, 4, 495-503.	1.6	16
444	Indoloquinolizidine-Peptide Hybrids as Multiple Agonists for D ₁ and D ₂ Dopamine Receptors. <i>ChemMedChem</i> , 2009, 4, 1514-1522.	1.6	16
445	Synthesis and Application of <i>N</i> -Hydroxylamine Derivatives as Potential Replacements for HOBt. <i>European Journal of Organic Chemistry</i> , 2009, 2009, 1499-1501.	1.2	27
446	Oxathiocoraline: Lessons to be Learned from the Synthesis of Complex <i>N</i> -Methylated Depsipeptides. <i>European Journal of Organic Chemistry</i> , 2009, 2009, 2957-2974.	1.2	14
447	Enhanced microwave-assisted method for on-bead disulfide bond formation: Synthesis of β -conotoxin MII. <i>Biopolymers</i> , 2009, 92, 23-34.	1.2	29
448	Design and facile solid-phase synthesis of peptide-based LPS inhibitors containing PEG-like functionalities. <i>Biopolymers</i> , 2009, 92, 508-517.	1.2	4
449	Optimized Fmoc solid-phase synthesis of Thymosin β 1 by side-chain anchoring onto a PEG resin. <i>Biopolymers</i> , 2009, 92, 565-572.	1.2	10
450	Solid-phase peptide synthesis using acetonitrile as a solvent in combination with PEG-based resins. <i>Journal of Peptide Science</i> , 2009, 15, 629-633.	0.8	47

#	ARTICLE	IF	CITATIONS
451	Optical Tweezers Study of Topoisomerase Inhibition. <i>Small</i> , 2009, 5, 1269-1272.	5.2	5
452	Microwave irradiation and COMU: a potent combination for solid-phase peptide synthesis. <i>Tetrahedron Letters</i> , 2009, 50, 6200-6202.	0.7	48
453	The ChEMBL project: building annotated molecular libraries for drug discovery. <i>New Biotechnology</i> , 2009, 25, S5.	2.4	0
454	Solution- and solid-phase synthesis and anti-HIV activity of maslinic acid derivatives containing amino acids and peptides. <i>Bioorganic and Medicinal Chemistry</i> , 2009, 17, 1139-1145.	1.4	63
455	Amino Acid-Protecting Groups. <i>Chemical Reviews</i> , 2009, 109, 2455-2504.	23.0	658
456	Siamese Depsipeptides: Constrained Bicyclic Architectures. <i>Angewandte Chemie - International Edition</i> , 2009, 48, 8564-8567.	7.2	5
457	Total Synthesis and Antiproliferative Activity Screening of (±)-Aplicyanins A, B and E and Related Analogues. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 6217-6223.	2.9	35
458	Lamellarin D Bioconjugates II: Synthesis and Cellular Internalization of Dendrimer and Nuclear Location Signal Derivatives. <i>Bioconjugate Chemistry</i> , 2009, 20, 1112-1121.	1.8	27
459	Solid-Phase Synthesis of Chiral Bicyclic Guanidinium Oligomers. <i>ACS Combinatorial Science</i> , 2009, 11, 410-421.	3.3	9
460	Lamellarin D Bioconjugates I: Synthesis and Cellular Internalization of PEG-Derivatives. <i>Bioconjugate Chemistry</i> , 2009, 20, 1100-1111.	1.8	23
461	Screening of One-Bead-One-Peptide Combinatorial Library Using Red Fluorescent Dyes. Presence of Positive and False Positive Beads. <i>ACS Combinatorial Science</i> , 2009, 11, 146-150.	3.3	44
462	Adenosine A _{2A} Receptor-Antagonist/Dopamine D ₂ Receptor-Agonist Bivalent Ligands as Pharmacological Tools to Detect A _{2A} -D ₂ Receptor Heteromers. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 5590-5602.	2.9	129
463	Conjugation of Kahalalide F with Gold Nanoparticles to Enhance in Vitro Antitumoral Activity. <i>Bioconjugate Chemistry</i> , 2009, 20, 138-146.	1.8	71
464	Antioxidant activity and <i>in vitro</i> inhibition of tumor cell growth by leaf extracts from the carob tree (<i>Ceratonia siliqua</i>). <i>Pharmaceutical Biology</i> , 2009, 47, 721-728.	1.3	27
465	<i>N</i> -Me Amide as a Synthetic Surrogate for the Thioester Moiety in Thiocoraline. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 834-839.	2.9	33
466	Manufacturing peptides as active pharmaceutical ingredients. <i>Future Medicinal Chemistry</i> , 2009, 1, 361-377.	1.1	151
467	Synthesis of the pyrrolo[2,3- <i>c</i>]carbazole core of the dictyodendrins. <i>Organic and Biomolecular Chemistry</i> , 2009, 7, 860.	1.5	38
468	Kahalalide F, an Antitumor Depsipeptide in Clinical Trials, and Its Analogues as Effective Antileishmanial Agents. <i>Molecular Pharmaceutics</i> , 2009, 6, 813-824.	2.3	39

#	ARTICLE	IF	CITATIONS
469	1,2-Dimethylindole-3-sulfonyl (MIS) as protecting group for the side chain of arginine. <i>Organic and Biomolecular Chemistry</i> , 2009, 7, 2565.	1.5	13
470	Solid-Phase Peptide Synthesis in Water Using Microwave-Assisted Heating. <i>Organic Letters</i> , 2009, 11, 4488-4491.	2.4	77
471	Phenyl-EDOTn derivatives as super acid labile carboxylic acid protecting groups for peptide synthesis. <i>Tetrahedron Letters</i> , 2008, 49, 3304-3307.	0.7	9
472	Solid Phase Preparation of 1,3-Disubstituted Indazole derivatives. <i>QSAR and Combinatorial Science</i> , 2008, 27, 1267-1273.	1.5	2
473	Protection by Conformationally Restricted Mobility: First Solid-Phase Synthesis of Triostin A. <i>Chemistry - A European Journal</i> , 2008, 14, 4475-4478.	1.7	14
474	Conformationally Restricted Hydantoin-Based Peptidomimetics as Inhibitors of Caspase-3 with Basic Groups Allowed at the S ₃ Enzyme Subsite. <i>ChemMedChem</i> , 2008, 3, 979-985.	1.6	11
475	Use of N-Methylpiperazine for the Preparation of Piperazine-Based Unsymmetrical Bis-Ureas as Anti-HIV Agents. <i>ChemMedChem</i> , 2008, 3, 1034-1037.	1.6	11
476	Nanostructure Formation Enhances the Activity of LPS-Neutralizing Peptides. <i>ChemMedChem</i> , 2008, 3, 1748-1755.	1.6	13
477	Synthesis of Natural Product Derivatives Containing 2,4-Concatenated Oxazoles. <i>European Journal of Organic Chemistry</i> , 2008, 2008, 3389-3396.	1.2	22
478	EDOTn and MIM, new peptide backbone protecting groups. <i>Biopolymers</i> , 2008, 90, 444-449.	1.2	23
479	Synthesis of Oligonucleotide Derivatives Using ChemMatrix Supports. <i>Chemistry and Biodiversity</i> , 2008, 5, 209-218.	1.0	9
480	Design, synthesis and antiproliferative properties of oligomers with chromophore units linked by amide backbones. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 2440-2444.	1.0	14
481	Solid-phase synthesis of oligomers carrying several chromophore units linked by phosphodiester backbones. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 2306-2310.	1.0	11
482	Stereomeric studies on the oxidation and alkylation of 4-thiazolidinones. <i>Tetrahedron Letters</i> , 2008, 49, 1569-1572.	0.7	15
483	[{Cu(pzPh)(Opo)} ₂ (μ -Cl) ₂]: A new dinuclear copper(II) complex with a chloride bridge and mixed blocking ligands. <i>Inorganica Chimica Acta</i> , 2008, 361, 2455-2461.	1.2	30
484	Design and Synthesis of FAJANU: a de Novo C ₂ -Symmetric Cyclopeptide Family. <i>Journal of Medicinal Chemistry</i> , 2008, 51, 3194-3202.	2.9	11
485	Trishomocubane Amino Acid as a Turn scaffold. <i>Chemical Biology and Drug Design</i> , 2008, 71, 125-130.	1.5	20
486	Tiratricol Neutralizes Bacterial Endotoxins and Reduces Lipopolysaccharide-Induced TNF α Production in the Cell. <i>Chemical Biology and Drug Design</i> , 2008, 72, 320-328.	1.5	3

#	ARTICLE	IF	CITATIONS
487	Solid-phase synthesis and characterization of N-methyl-rich peptides. <i>Chemical Biology and Drug Design</i> , 2008, 65, 153-166.	1.2	107
488	<i>N,N,N,N</i> -Tetramethylchloroformamidinium Hexafluorophosphate (TCFH), a Powerful Coupling Reagent for Bioconjugation. <i>Bioconjugate Chemistry</i> , 2008, 19, 1968-1971.	1.8	17
489	THAL, a Sterically Unhindered Linker for the Solid-Phase Synthesis of Acid-Sensitive Protected Peptide Acids. <i>Journal of Organic Chemistry</i> , 2008, 73, 7342-7344.	1.7	5
490	Morpholine-Based Immonium and Halogenoamidinium Salts as Coupling Reagents in Peptide Synthesis. <i>Journal of Organic Chemistry</i> , 2008, 73, 2731-2737.	1.7	61
491	Structure-Activity Relationship of Kahalalide F Synthetic Analogues. <i>Journal of Medicinal Chemistry</i> , 2008, 51, 4920-4931.	2.9	38
492	A Novel Protecting/Activating Strategy for β -Hydroxy Acids and Its Use in Convergent Peptide Synthesis. <i>Journal of Organic Chemistry</i> , 2008, 73, 2311-2314.	1.7	10
493	Synthesis and Antitumor Activity of Mechercharmycin A Analogues. <i>Journal of Medicinal Chemistry</i> , 2008, 51, 5722-5730.	2.9	23
494	Solid-Phase Combinatorial Synthesis of a Lysyl-tRNA Synthetase (LysRS) Inhibitory Library. <i>ACS Combinatorial Science</i> , 2008, 10, 391-400.	3.3	10
495	Cysteine-S-trityl a Key Derivative to Prepare <i>N</i> -Methyl Cysteines. <i>ACS Combinatorial Science</i> , 2008, 10, 69-78.	3.3	17
496	A New 6H-Pyran-3-one Scaffold Derived from a C-Glycoside (SUPPORTING INFORMATION). <i>Letters in Organic Chemistry</i> , 2008, 5, 374-378.	0.2	1
497	Crystal structure of hexakis(4-fluorophenylethylammonium)decavanadate(V) tetrahydrate, (C ₈ H ₁₁ FN) ₆ [V ₁₀ O ₂₈] · 4H ₂ O. <i>Zeitschrift Fur Kristallographie - New Crystal Structures</i> , 2008, 223, 45-47.	0.1	1
498	Asymmetric Synthesis of α -Unsubstituted β -Hydroxy Acids. <i>Current Organic Synthesis</i> , 2008, 5, 151-161.	0.7	11
499	Oral Insulin-Mimetic Compounds That Act Independently of Insulin. <i>Diabetes</i> , 2007, 56, 486-493.	0.3	60
500	Novel Synthesis of Arylethynyl Heterocycles. <i>Synthesis</i> , 2007, 2007, 1559-1565.	1.2	4
501	New developments in the synthesis of oligonucleotide-peptide conjugates. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2007, 26, 963-967.	0.4	4
502	Advances in Solid-Phase Cycloadditions for Heterocyclic Synthesis. <i>ACS Combinatorial Science</i> , 2007, 9, 521-565.	3.3	36
503	Peptides and metallic nanoparticles for biomedical applications. <i>Nanomedicine</i> , 2007, 2, 287-306.	1.7	129
504	Synthesis of IB-01211, a Cyclic Peptide Containing 2,4-Concatenated Thia- and Oxazoles, via Hantzsch Macrocyclization. <i>Organic Letters</i> , 2007, 9, 809-811.	2.4	42

#	ARTICLE	IF	CITATIONS
505	Solid-Phase Synthesis of Oxathiocoraline by a Key Intermolecular Disulfide Dimer. <i>Journal of the American Chemical Society</i> , 2007, 129, 5322-5323.	6.6	46
506	Novel Proton Acceptor Immonium-Type Coupling Reagents: Application in Solution and Solid-Phase Peptide Synthesis. <i>Organic Letters</i> , 2007, 9, 4475-4477.	2.4	39
507	A Nonacid Degradable Linker for Solid-Phase Synthesis. <i>Organic Letters</i> , 2007, 9, 4319-4322.	2.4	13
508	Smallest Peptoids with Antiproliferative Activity on Human Neoplastic Cells. <i>Journal of Medicinal Chemistry</i> , 2007, 50, 2443-2449.	2.9	19
509	p-Nitromandelic Acid as a Highly Acid-Stable Safety-Catch Linker for Solid-Phase Synthesis of Peptide and Depsipeptide Acids. <i>Organic Letters</i> , 2007, 9, 1429-1432.	2.4	14
510	Novel Ergopeptides as Dual Ligands for Adenosine and Dopamine Receptors. <i>Journal of Medicinal Chemistry</i> , 2007, 50, 3062-3069.	2.9	39
511	Design and Synthesis of Indole-Based Peptoids as Potent Noncompetitive Antagonists of Transient Receptor Potential Vanilloid 1. <i>Journal of Medicinal Chemistry</i> , 2007, 50, 6133-6143.	2.9	19
512	From the One-Bead-One-Compound Concept to One-Bead-One-Reactor. <i>ACS Combinatorial Science</i> , 2007, 9, 395-398.	3.3	9
513	Solid-Phase Synthesis of Sulfamate Peptidomimetics. <i>ACS Combinatorial Science</i> , 2007, 9, 501-506.	3.3	4
514	ds-Oligonucleotide-Peptide Conjugates Featuring Peptides from the Leucine-Zipper Region of Fos as Switchable Receptors for the Oncoprotein Jun. <i>ChemBioChem</i> , 2007, 8, 1110-1114.	1.3	24
515	Fmoc-2-mercaptobenzothiazole, for the introduction of the Fmoc moiety free of side-reactions. <i>Biopolymers</i> , 2007, 88, 733-737.	1.2	34
516	Simple machine-assisted protocol for solid-phase synthesis of depsipeptides. <i>Biopolymers</i> , 2007, 88, 823-828.	1.2	7
517	Amide-Ester Substitution in Coiled Coils: The Effect of Removing Hydrogen Bonds on Protein Structure. <i>Angewandte Chemie - International Edition</i> , 2007, 46, 7766-7769.	7.2	42
518	Synthesis of a 24-Membered Cyclic Peptide-Biphenyl Hybrid. <i>European Journal of Organic Chemistry</i> , 2007, 2007, 1301-1308.	1.2	17
519	Regioselective Monobromination of Free and Protected Phenols. <i>European Journal of Organic Chemistry</i> , 2007, 2007, 1921-1924.	1.2	19
520	Inhibition of VAP1: Quickly Gaining Ground as an Anti-Inflammatory Therapy. <i>ChemMedChem</i> , 2007, 2, 173-174.	1.6	14
521	Synthesis and Structure-Activity Relationship of Cytotoxic Marine Cyclodepsipeptide IB-01212 Analogues. <i>ChemMedChem</i> , 2007, 2, 1076-1084.	1.6	10
522	Formation of dihydrouracils via cyclization of N-substituted 3-thioureidopropanoic acids and facile desulfurization. <i>Tetrahedron</i> , 2007, 63, 8949-8953.	1.0	1

#	ARTICLE	IF	CITATIONS
523	Preparation of penta-azole containing cyclopeptides: challenges in macrocyclization. <i>Tetrahedron</i> , 2007, 63, 9862-9870.	1.0	24
524	Identification of protein-binding peptides by direct matrix-assisted laser desorption ionization time-of-flight mass spectrometry analysis of peptide beads selected from the screening of one bead- ϵ -one peptide combinatorial libraries. <i>Analytical Biochemistry</i> , 2007, 370, 215-222.	1.1	26
525	Chlorotriyl Chloride (CTC) Resin as a Reusable Carboxyl Protecting Group. <i>QSAR and Combinatorial Science</i> , 2007, 26, 1027-1035.	1.5	32
526	Understanding the Mechanism of Action of the Novel SSAO Substrate (C ₇ NH ₁₀) ₆ (V ₁₀ O ₂₈) \cdot 2H ₂ O, a Prodrug of Peroxovanadate Insulin Mimetics. <i>Chemical Biology and Drug Design</i> , 2007, 69, 423-428.	1.5	46
527	Chemical Synthesis of ¹⁹ F-labeled HIV-1 Protease using Fmoc-Chemistry and ChemMatrix Resin. <i>International Journal of Peptide Research and Therapeutics</i> , 2007, 13, 221-227.	0.9	20
528	Beyond Azathiocoraline: Synthesis of Analogues. <i>International Journal of Peptide Research and Therapeutics</i> , 2007, 13, 295-306.	0.9	3
529	Does the Solid-Phase Synthesis of a Tetrapeptide Represent a Challenge at the Onset of the XXI Century? The Case of Cyclo [(3R)-3-hydroxydecanoyl-L-seryl-(3R)-3-hydroxydecanoyl-L-seryl]. <i>International Journal of Peptide Research and Therapeutics</i> , 2007, 13, 313-327.	0.9	2
530	Partially Fluorinated Heterocycles from 4,4-Bis(trifluoromethyl)-hetero-1,3-dienes via C-F Bond Activation ϵ Synthesis of 2-Fluoro-3-(trifluoromethyl)furan. <i>Monatshefte für Chemie</i> , 2007, 138, 227-236.	0.9	9
531	Enolase as a plasminogen binding protein in <i>Leishmania mexicana</i> . <i>Parasitology Research</i> , 2007, 101, 1511-1516.	0.6	104
532	Cell-Penetrating Proline-Rich Peptidomimetics. <i>Methods in Molecular Biology</i> , 2007, 386, 241-267.	0.4	6
533	The synthesis of naturally occurring peptides and their analogs. <i>Current Opinion in Drug Discovery & Development</i> , 2007, 10, 768-83.	1.9	19
534	Convergent Approaches for the Synthesis of the Antitumoral Peptide, Kahalalide F. Study of Orthogonal Protecting Groups. <i>Journal of Organic Chemistry</i> , 2006, 71, 7196-7204.	1.7	27
535	IB-01212, a New Cytotoxic Cyclodepsipeptide Isolated from the Marine Fungus <i>Clonostachys</i> sp. ESNA-A009. <i>Journal of Organic Chemistry</i> , 2006, 71, 3335-3338.	1.7	49
536	Total Solid-Phase Synthesis of Marine Cyclodepsipeptide IB-01212. <i>Journal of Organic Chemistry</i> , 2006, 71, 3339-3344.	1.7	22
537	ChemMatrix, a Poly(ethylene glycol)-Based Support for the Solid-Phase Synthesis of Complex Peptides. <i>ACS Combinatorial Science</i> , 2006, 8, 213-220.	3.3	241
538	New Nomenclature for Complex Cyclopeptides. , 2006, , 142-143.		0
539	New Efficient Substrates for Semicarbazide-Sensitive Amine Oxidase/VAP-1 Enzyme: ϵ Analysis by SARs and Computational Docking. <i>Journal of Medicinal Chemistry</i> , 2006, 49, 6197-6208.	2.9	28
540	Hexafluoroacetone as Protecting and Activating Reagent: ϵ New Routes to Amino, Hydroxy, and Mercapto Acids and Their Application for Peptide and Glyco- and Depsipeptide Modification. <i>Chemical Reviews</i> , 2006, 106, 4728-4746.	23.0	40

#	ARTICLE	IF	CITATIONS
541	Solid-Phase Combinatorial Synthesis of Peptide-Biphenyl Hybrids as Calpain Inhibitors. <i>Organic Letters</i> , 2006, 8, 3621-3621.	2.4	0
542	Synthesis and Structure-Activity Relationship Study of Potent Cytotoxic Analogues of the Marine Alkaloid Lamellarin D. <i>Journal of Medicinal Chemistry</i> , 2006, 49, 3257-3268.	2.9	100
543	Enhancing Atom Economy of SPS: Recoverable and Reusable Building Blocks for Depsipeptide Synthesis. , 2006, , 108-109.		0
544	Total Solid Phase Synthesis of a Marine Cyclodepsipeptide IB-01212. , 2006, , 210-211.		0
545	Chlorotriyl Chloride (CTC) Resin as a Convenient Reusable Protecting Group. , 2006, , 220-221.		3
546	Solid-Phase Peptide Synthesis Using ChemMatrix [®] , a Polyethylenglycol (PEG)-based Solid. , 2006, , 114-115.		1
547	1-Hydroxy-6,7-dimethoxy-8-nitro-1,2,3,4-tetrahydroisoquinoline. <i>Acta Crystallographica Section E: Structure Reports Online</i> , 2006, 62, o2285-o2287.	0.2	0
548	Solid-Phase N-Electrophilic Amination of 2-Aminopyridines: Preparation of 2-Substituted-[1,2,4]triazolo[1,5-a]pyridine Derivatives. <i>QSAR and Combinatorial Science</i> , 2006, 25, 961-965.	1.5	7
549	Solid-Phase Synthesis and Structural Study of Substituted 1,4,5,6-Tetrahydro-6-oxopyridine-3-carboxylic Acids. <i>QSAR and Combinatorial Science</i> , 2006, 25, 921-927.	1.5	11
550	Design of a minimized cyclic tetrapeptide that neutralizes bacterial endotoxins. <i>Journal of Peptide Science</i> , 2006, 12, 491-496.	0.8	16
551	Facile solid-phase synthesis of biotinylated alkyl thiols. <i>Tetrahedron</i> , 2006, 62, 6876-6881.	1.0	18
552	Sulfoxidations in the solid phase. <i>Tetrahedron: Asymmetry</i> , 2006, 17, 3327-3331.	1.8	4
553	A convenient semicarbazide resin for the solid-phase synthesis of peptide ketones and aldehydes. <i>Tetrahedron Letters</i> , 2006, 47, 1657-1661.	0.7	18
554	Homologation of α -hydroxy acids to α -unsubstituted β -hydroxy carboxamides via Arndt-Eistert reaction. <i>Tetrahedron Letters</i> , 2006, 47, 4557-4560.	0.7	9
555	Microwave-assisted synthesis of 1,3-dihydro-[1,2,5]thiadiazolo[3,4-b]pyrazine-2,2-dioxides. <i>Tetrahedron Letters</i> , 2006, 47, 8603-8606.	0.7	11
556	Synthesis of α -trifluoromethyl α -amino acids with aromatic, heteroaromatic and ferrocenyl subunits in the side chain. <i>Amino Acids</i> , 2006, 31, 55-62.	1.2	9
557	Synthetic Approaches to Disulfide-free Circular Bovine Pancreatic Trypsin Inhibitor (c-BPTI) Analogues. <i>International Journal of Peptide Research and Therapeutics</i> , 2006, 12, 93-104.	0.9	2
558	Domino reactions with fluorinated five-membered heterocycles. α -Trifluoromethyl α -amino acids with unsaturated side-chains. <i>Amino Acids</i> , 2006, 31, 427-433.	1.2	4

#	ARTICLE	IF	CITATIONS
559	Domino Reactions with Fluorinated Five-Membered Heterocycles – Syntheses of Trifluoromethyl Substituted Butenolides and β -Ketoacids.. ChemInform, 2006, 37, no.	0.1	0
560	Total Solid-Phase Synthesis of the Azathiocoraline Class of Symmetric Bicyclic Peptides. Chemistry - A European Journal, 2006, 12, 9001-9009.	1.7	27
561	The synergy of ChemMatrix resin [®] and pseudoproline building blocks renders Rantes, a complex aggregated chemokine. Biopolymers, 2006, 84, 566-575.	1.2	59
562	Design and Synthesis of New Immonium-Type Coupling Reagents. European Journal of Organic Chemistry, 2006, 2006, 1563-1573.	1.2	23
563	p-Nitrobenzyloxycarbonyl (pNZ) as an Alternative to Fmoc for the Protection of Amines in Solid-Phase Peptide Synthesis. , 2006, , 116-117.		0
564	Synthesis of Partially Fluorinated Heterocycles from 4,4-Bis(trifluoromethyl) Substituted Hetero-1,3-dienes via C-F Bond Activation and Their Application as Trifluoromethyl Substituted Building Blocks. Heterocycles, 2006, 69, 569.	0.4	10
565	Role of the Acid Group in the Pictet-Spengler Reaction of β -Amino Acids. Synlett, 2006, 2006, 1903-1907.	1.0	10
566	Solid-Phase Chemistry in the Total Synthesis of Non-Peptidic Natural Products. Mini-Reviews in Medicinal Chemistry, 2006, 6, 11-25.	1.1	17
567	Suzuki coupling reaction for the solid-phase preparation of 5-substituted nicotinic acid derivatives. Tetrahedron Letters, 2005, 46, 581-585.	0.7	26
568	An efficient strategy for the preparation of one-bead-one-peptide libraries on a new biocompatible solid support. Tetrahedron Letters, 2005, 46, 1561-1564.	0.7	42
569	5,6-Dihydropyrrolo[2,1-b]isoquinolines as scaffolds for synthesis of lamellarin analogues. Tetrahedron Letters, 2005, 46, 2041-2044.	0.7	41
570	N-[Chloro(dimethylamino)methylene]-N-methylmethanaminium chloride (TMUCl Cl), the reagent of choice for the solid-phase synthesis of anilides. Tetrahedron Letters, 2005, 46, 5383-5386.	0.7	12
571	NO as temporary guanidino-protecting group provides efficient access to Pbf-protected argininic acid. Tetrahedron Letters, 2005, 46, 6733-6735.	0.7	9
572	Avoiding pyran ring opening during palladium acetate catalyzed C-glycosidation of peracetylated glycals. Tetrahedron Letters, 2005, 46, 7271-7274.	0.7	26
573	Semipermanent p-nitrobenzyloxycarbonyl (pNZ) protection of Orn and Lys side chains: prevention of undesired β -Fmoc removal and application to the synthesis of cyclic peptides. Tetrahedron Letters, 2005, 46, 7733-7736.	0.7	12
574	Use of p-nitrobenzyloxycarbonyl (pNZ) as a permanent protecting group in the synthesis of Kahalalide F analogs. Tetrahedron Letters, 2005, 46, 7737-7741.	0.7	18
575	A new approach to 3-hydroxyquinoline-2-carboxylic acid. Tetrahedron, 2005, 61, 1407-1411.	1.0	11
576	Combinatorial approaches towards the discovery of new tryptase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2005, 15, 1659-1664.	1.0	18

#	ARTICLE	IF	CITATIONS
577	Multicomponent Reactions with Dihydroazines: Efficient Synthesis of a Diverse Set of Pyrido-Fused Tetrahydroquinolines. <i>ACS Combinatorial Science</i> , 2005, 7, 33-41.	3.3	47
578	Abbreviated nomenclature for cyclic and branched homo- and hetero-detic peptides. <i>Chemical Biology and Drug Design</i> , 2005, 65, 550-555.	1.2	50
579	p-Nitrobenzyloxycarbonyl (pNZ) as a Temporary N-Protecting Group in Orthogonal Solid-Phase Peptide Synthesis - Avoiding Diketopiperazine and Aspartimide Formation. <i>European Journal of Organic Chemistry</i> , 2005, 2005, 3031-3039.	1.2	50
580	Synthesis and screening of a small library of proline-based biodendrimers for use as delivery agents. <i>Biopolymers</i> , 2005, 80, 800-814.	1.2	29
581	Developments in Peptide and Amide Synthesis. <i>ChemInform</i> , 2005, 36, no.	0.1	0
582	Qualitative Colorimetric Tests for Solid Phase Synthesis. <i>ChemInform</i> , 2005, 36, no.	0.1	0
583	Hexafluoroacetone as a Protecting and Activating Reagent. Regioselective Esterification of Aspartic, Malic, and Thiomalic Acid.. <i>ChemInform</i> , 2005, 36, no.	0.1	0
584	A New Approach to 3-Hydroxyquinoline-2-carboxylic Acid.. <i>ChemInform</i> , 2005, 36, no.	0.1	0
585	Multicomponent Reactions with Dihydroazines: Efficient Synthesis of a Diverse Set of Pyrido-Fused Tetrahydroquinolines.. <i>ChemInform</i> , 2005, 36, no.	0.1	0
586	Peptide and Amide Bond Containing Dendrimers. <i>ChemInform</i> , 2005, 36, no.	0.1	0
587	Directly Linked Polyazoles: Important Moieties in Natural Products. <i>ChemInform</i> , 2005, 36, no.	0.1	0
588	Hexafluoroacetone as a Protecting and Activating Reagent. N- and O-Glycosylation of Isoleucine and Isocysteine. <i>Monatshefte für Chemie</i> , 2005, 136, 577-595.	0.9	4
589	Orthogonally Protected, Carboxy-Activated L-Homoisoleucine, 2-Methyl-L-homoisoleucine, and Homoisocysteine Derivatives. New Building Blocks for Peptide and Dipeptide Modification. <i>Monatshefte für Chemie</i> , 2005, 136, 763-776.	0.9	4
590	Domino Reactions with Fluorinated Five-membered Heterocycles – Syntheses of Trifluoromethyl Substituted Butenolides and β -Ketoacids. <i>Monatshefte für Chemie</i> , 2005, 136, 1763-1779.	0.9	14
591	Solid-Phase Preparation of a Library Based on a Phenylalanine Scaffold. <i>QSAR and Combinatorial Science</i> , 2005, 24, 913-922.	1.5	9
592	Evolutionary combinatorial chemistry, a novel tool for SAR studies on peptide transport across the blood-brain barrier. Part 2. Design, synthesis and evaluation of a first generation of peptides. <i>Journal of Peptide Science</i> , 2005, 11, 789-804.	0.8	18
593	Chapter 1 Lamellarins: Isolation, activity and synthesis. <i>Progress in Heterocyclic Chemistry</i> , 2005, 16, 1-26.	0.5	21
594	Chloromethoxymethyl Polystyrene (CMM Resin), an Acid Labile Resin for Anchoring/Cleavage of N-Heterocycles and Oxygen Aromatic Compounds. <i>Letters in Organic Chemistry</i> , 2005, 2, 371-373.	0.2	3

#	ARTICLE	IF	CITATIONS
595	Backbone Amide Linker Strategies for the Solid-Phase Synthesis of C - Terminal Modified Peptides. , 2005, 298, 195-208.		7
596	Cell-Penetrating cis- β -Amino-L-Proline-Derived Peptides. Journal of the American Chemical Society, 2005, 127, 9459-9468.	6.6	102
597	A Straightforward Synthesis of 5'-Peptide Oligonucleotide Conjugates Using β -Fmoc-Protected Amino Acids. Organic Letters, 2005, 7, 4349-4352.	2.4	26
598	Preparation of de Novo Globular Proteins Based on Proline Dendrimers. Journal of Organic Chemistry, 2005, 70, 6274-6281.	1.7	23
599	A New Strategy for Solid-Phase Depsipeptide Synthesis Using Recoverable Building Blocks. Organic Letters, 2005, 7, 597-600.	2.4	23
600	Peptide and Amide Bond-Containing Dendrimers. Chemical Reviews, 2005, 105, 1663-1682.	23.0	321
601	Modular Total Synthesis of Lamellarin D. Journal of Organic Chemistry, 2005, 70, 8231-8234.	1.7	108
602	Evaluation of Solution and Solid-Phase Approaches to the Synthesis of Libraries of α,β -Disubstituted α -acylamino ketones. ACS Combinatorial Science, 2005, 7, 843-863.	3.3	11
603	Application of hexafluoroacetone as protecting and activating reagent in solid phase peptide and depsipeptide synthesis. Arkivoc, 2005, 2005, 191-199.	0.3	11
604	Incorporation of the β -Mercapto Acid Unit into Peptides. Synthesis, 2004, 2004, 1088-1092.	1.2	0
605	Re-Evaluation of a Solid-Phase Hydantoin Synthesis. Letters in Organic Chemistry, 2004, 1, 224-226.	0.2	20
606	From Production of Peptides in Milligram Amounts for Research to Multi-Tons Quantities for Drugs of the Future. Current Pharmaceutical Biotechnology, 2004, 5, 29-43.	0.9	174
607	Inhibition of beta-amyloid toxicity by short peptides containing N-methyl amino acids. Chemical Biology and Drug Design, 2004, 63, 324-328.	1.2	48
608	One-Pot Preparation of N-Carbamate Protected Amino Acids via the Azide. Organic Process Research and Development, 2004, 8, 920-924.	1.3	28
609	Synthesis of Polyheterocyclic Nitrogen-Containing Marine Natural Products. Monatshefte für Chemie, 2004, 135, 615-627.	0.9	41
610	Hexafluoroacetone as a Protecting and Activating Reagent. Regioselective Esterification of Aspartic, Malic, and Thiomalic Acid. Monatshefte für Chemie, 2004, 135, 1427-1443.	0.9	7
611	Gaining diversity in solid-phase synthesis by modulation of cleavage conditions from hydroxymethyl-based supports. Application to lamellarin synthesis. Tetrahedron, 2004, 60, 8669-8675.	1.0	24
612	A Combination of Different Spectroscopic Techniques to Monitor in situ Solid-phase Synthesis of Organic Molecules. QSAR and Combinatorial Science, 2004, 23, 61-68.	1.5	7

#	ARTICLE	IF	CITATIONS
613	A Re-evaluation of the Use of Rink, BAL, and PAL Resins and Linkers. <i>QSAR and Combinatorial Science</i> , 2004, 23, 145-152.	1.5	49
614	Solid-phase synthesis of second-generation polyproline dendrimers. <i>Biopolymers</i> , 2004, 76, 283-297.	1.2	26
615	Synthetic Circularized Analogues of Bovine Pancreatic Trypsin Inhibitor. <i>European Journal of Organic Chemistry</i> , 2004, 2004, 4541-4544.	1.2	5
616	Solid-Phase Synthesis of C-Terminal Modified Peptides. <i>ChemInform</i> , 2004, 35, no.	0.1	0
617	Industrial Application of Coupling Reagents in Peptides. <i>ChemInform</i> , 2004, 35, no.	0.1	1
618	Synthesis of Polyheterocyclic Nitrogen-Containing Marine Natural Products.. <i>ChemInform</i> , 2004, 35, no.	0.1	1
619	From Production of Peptides in Milligram Amounts for Research to Multi-tons Quantities for Drugs of the Future. <i>ChemInform</i> , 2004, 35, no.	0.1	0
620	Solid-phase synthesis of 4H-2-(3-hydroxy-4-methoxyphenyl)-naphtho[1,2-b]pyran-1-one. <i>Tetrahedron Letters</i> , 2004, 45, 7311-7314.	0.7	3
621	Solid-phase synthesis of lamellarins Q and O. <i>Tetrahedron</i> , 2004, 60, 8659-8668.	1.0	51
622	Developments in peptide and amide synthesis. <i>Current Opinion in Chemical Biology</i> , 2004, 8, 211-221.	2.8	234
623	Solid-Phase Combinatorial Synthesis of Peptide-Biphenyl Hybrids as Calpain Inhibitors. <i>Organic Letters</i> , 2004, 6, 4089-4092.	2.4	21
624	A Comparative Study of Different Presentation Strategies for an HIV Peptide Immunogen. <i>Bioconjugate Chemistry</i> , 2004, 15, 112-120.	1.8	46
625	A New Class of Foldamers Based on α -Amino- β -proline. <i>Journal of the American Chemical Society</i> , 2004, 126, 6048-6057.	6.6	97
626	Solid-Phase Syntheses of Furopyridine and Furoquinoline Systems. <i>Organic Letters</i> , 2004, 6, 1405-1408.	2.4	38
627	Monoclonal antibody purification by affinity chromatography with ligands derived from the screening of peptide combinatorial libraries. <i>Biotechnology Letters</i> , 2003, 25, 1545-1548.	1.1	33
628	Synthesis and antifungal activity of an acivicine-based dipeptide library. <i>International Journal of Peptide Research and Therapeutics</i> , 2003, 10, 645-653.	0.1	2
629	Synthesis and SAR of α -Acylaminoketone Ligands for Control of Gene Expression.. <i>ChemInform</i> , 2003, 34, no.	0.1	0
630	o-Formylation of Electron-Rich Phenols with Dichloromethyl Methyl Ether and TiCl ₄ .. <i>ChemInform</i> , 2003, 34, no.	0.1	0

#	ARTICLE	IF	CITATIONS
631	Solid-Phase Synthesis of the Cyclic Liponadepsipeptide [N-Mst(Ser1), D-Ser4, L-Thr6, L-Asp8, L-Thr9]Syringotoxin. <i>Chemistry - A European Journal</i> , 2003, 9, 1096-1103.	1.7	7
632	Solid-phase synthesis of C-terminal modified peptides. <i>Biopolymers</i> , 2003, 71, 454-477.	1.2	67
633	Saturated resins or stress of the resin. <i>Tetrahedron Letters</i> , 2003, 44, 1751-1754.	0.7	15
634	o-Formylation of electron-rich phenols with dichloromethyl methyl ether and TiCl ₄ . <i>Tetrahedron Letters</i> , 2003, 44, 4961-4963.	0.7	23
635	Solid-phase synthesis: a linker for side-chain anchoring of arginine. <i>Tetrahedron Letters</i> , 2003, 44, 5319-5321.	0.7	23
636	Synthesis of variolin B. <i>Tetrahedron Letters</i> , 2003, 44, 6191-6194.	0.7	20
637	BAL resin for the preparation of secondary amines. <i>Tetrahedron Letters</i> , 2003, 44, 6907-6910.	0.7	18
638	Synthesis and SAR of Î±-Acylaminoketone ligands for control of gene expression. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2003, 13, 475-478.	1.0	47
639	Development of a Genetic Algorithm to Design and Identify Peptides that can Cross the Blood-Brain Barrier. <i>QSAR and Combinatorial Science</i> , 2003, 22, 745-753.	1.5	19
640	Solid-Phase Total Synthesis of the Pentacyclic System Lamellarins U and L. <i>Organic Letters</i> , 2003, 5, 2959-2962.	2.4	74
641	Solution Structure of the Antitumor Candidate Trunkamide A by 2D NMR and Restrained Simulated Annealing Methods. <i>Journal of Organic Chemistry</i> , 2003, 68, 211-215.	1.7	34
642	Bicyclic Homodetic Peptide Libraries: Comparison of Synthetic Strategies for Their Solid-Phase Synthesis. <i>ACS Combinatorial Science</i> , 2003, 5, 760-768.	3.3	23
643	Synthesis and NMR Structure of P41icf, a Potent Inhibitor of Human Cathepsin L. <i>Journal of the American Chemical Society</i> , 2003, 125, 1508-1517.	6.6	24
644	Tentoxin as a Scaffold for Drug Discovery. Total Solid-Phase Synthesis of Tentoxin and a Library of Analogues. <i>Organic Letters</i> , 2003, 5, 2115-2118.	2.4	21
645	Qualitative Colorimetric Tests for Solid Phase Synthesis. <i>Methods in Enzymology</i> , 2003, 369, 21-35.	0.4	31
646	Total Syntheses of Variolin B and Deoxyvariolin B1. <i>Journal of Organic Chemistry</i> , 2003, 68, 10020-10029.	1.7	52
647	Fmoc Solid-Phase Synthesis of Peptide Thioesters by Masking as Trithioortho Esters. <i>Organic Letters</i> , 2003, 5, 2951-2953.	2.4	102
648	Small molecules targeting the vanilloid receptor complex as drugs for inflammatory pain. <i>Drugs of the Future</i> , 2003, 28, 787.	0.0	16

#	ARTICLE	IF	CITATIONS
649	Effects of l- and d-REKR amino acid-containing peptides on HIV and SIV envelope glycoprotein precursor maturation and HIV and SIV replication. <i>Biochemical Journal</i> , 2002, 366, 863-872.	1.7	4
650	Peptide Dendrimers Based on Polyproline Helices. <i>Journal of the American Chemical Society</i> , 2002, 124, 8876-8883.	6.6	111
651	Branched Poly(proline) Peptides: An Efficient New Approach to the Synthesis of Repetitive Branched Peptides. <i>European Journal of Organic Chemistry</i> , 2002, 2002, 1756-1762.	1.2	15
652	Monitoring the Chemical Assembly of a Transmembrane Bradykinin Receptor Fragment: Correlation Between Resin Solvation, Peptide Chain Mobility, and Rate of Coupling. <i>European Journal of Organic Chemistry</i> , 2002, 2002, 3686-3694.	1.2	26
653	Synthesis of peptides containing $\hat{1}\pm$, $\hat{1}^2$ -didehydroamino acids. Scope and limitations. <i>International Journal of Peptide Research and Therapeutics</i> , 2002, 9, 135-141.	0.1	3
654	Practical protocols for stepwise solid-phase synthesis of cysteine-containing peptides. <i>Chemical Biology and Drug Design</i> , 2002, 60, 292-299.	1.2	63
655	Structural, kinetic and cytotoxicity aspects of 12-28 γ -amyloid protein fragment: a reappraisal. <i>Journal of Peptide Science</i> , 2002, 8, 578-588.	0.8	22
656	Four-dimensional orthogonal solid-phase synthesis of new scaffolds based on cyclic tetra- $\hat{1}^2$ -peptides. <i>Tetrahedron Letters</i> , 2002, 43, 2029-2032.	0.7	21
657	Solid-phase syntheses of N-substituted carbamates. Reaction monitoring by gel-phase ^{13}C NMR using a ^{13}C enriched BAL-linker. <i>Tetrahedron Letters</i> , 2002, 43, 3543-3546.	0.7	17
658	Undesired removal of the Fmoc group by the free $\hat{1}\mu$ -amino function of a lysine residue. <i>Tetrahedron Letters</i> , 2002, 43, 7813-7815.	0.7	30
659	Solid phase synthesis of $\hat{1}\pm$ -acylamino- $\hat{1}\pm$, $\hat{1}\pm$ -disubstituted ketones. <i>Tetrahedron Letters</i> , 2002, 43, 7491-7494.	0.7	8
660	Combined solid phase and solution synthesis of a library of $\hat{1}\pm$, $\hat{1}\pm$ -disubstituted- $\hat{1}\pm$ -acylaminoketones. <i>Tetrahedron Letters</i> , 2002, 43, 7495-7498.	0.7	13
661	Synthesis of Fmoc-protected amino ketones bearing tert-butyl based side-chain protecting groups. <i>Tetrahedron Letters</i> , 2002, 43, 7499-7502.	0.7	19
662	Synthesis of peptides containing $\hat{1}\pm$, $\hat{1}^2$ -didehydroamino acids. Scope and limitations. <i>International Journal of Peptide Research and Therapeutics</i> , 2002, 9, 135-141.	0.1	9
663	Backbone amide linker (BAL) for solid-phase synthesis of 2,5-piperazinediones (DKP), useful scaffolds for combinatorial chemistry. , 2002, , 37-39.		0
664	Synthesis and applications of a bis-sulfonyl handle for solid-phase synthesis of peptides. , 2002, , 307-308.		0
665	Solid-phase peptide synthesis in the $\text{N}\hat{1}^+\text{C}$ direction. , 2002, , 78-79.		0
666	Backbone Amide Linker (BAL) methodology to accommodate C-terminal hindered, unreactive, and/or sensitive modifications. , 2002, , 102-103.		0

#	ARTICLE	IF	CITATIONS
667	Disulfide Bond Based Self-Assembly of Peptides Leading To Spheroidal Cyclic Trimers. , 2002, , 243-256.		0
668	AN IMPROVED SYNTHESIS OF N-[(9-HYDROXYMETHYL)-2-FLUORENYL]SUCCINAMIC ACID (HMFS), A VERSATILE HANDLE FOR THE SOLID-PHASE SYNTHESIS OF BIOMOLECULES. Synthetic Communications, 2001, 31, 225-232.	1.1	21
669	Solid-Phase Total Synthesis of Trunkamide A1. Journal of Organic Chemistry, 2001, 66, 7568-7574.	1.7	44
670	Synthesis and Structure Determination of Kahalalide F1,2. Journal of the American Chemical Society, 2001, 123, 11398-11401.	6.6	127
671	Synthesis of a Sulfahydantoin Library. ACS Combinatorial Science, 2001, 3, 290-300.	3.3	24
672	Backbone Amide Linker (BAL) Strategy for Solid-Phase Synthesis. , 2001, , 121-138.		3
673	Solid-phase peptide synthesis using N ^t -trityl-amino acids. International Journal of Peptide Research and Therapeutics, 2001, 8, 331-338.	0.1	2
674	A useful and sensitive color test to monitor aldehydes on solid-phase. Tetrahedron Letters, 2001, 42, 6691-6693.	0.7	25
675	Solid-phase syntheses of constrained RGD scaffolds and their binding to the $\alpha_5\beta_3$ integrin receptor. Tetrahedron Letters, 2001, 42, 7387-7391.	0.7	31
676	2-Mercaptopyridine-1-oxide-based peptide coupling reagents. Tetrahedron, 2001, 57, 9607-9613.	1.0	25
677	Solid-phase synthesis of 4-aminopiperidine analogues using the Alloc protecting group: an investigation of Alloc removal from secondary amines. Tetrahedron Letters, 2001, 42, 4471-4474.	0.7	28
678	Inhibition of HIV-2ROD replication in a lymphoblastoid cell line by the α_1 -antitrypsin Portland variant (α_1 -PDX) and the decRVKRcmk peptide: comparison with HIV-1LAI. Microbes and Infection, 2001, 3, 1073-1084.	1.0	3
679	Cu(OBt) ₂ and Cu(OAt) ₂ , copper(II)-based racemization suppressors ready for use in fully automated solid-phase peptide synthesis. Journal of Peptide Science, 2001, 7, 115-120.	0.8	20
680	Solid-Phase Synthesis of Peptides Containing α,β -Didehydroamino Acids. European Journal of Organic Chemistry, 2001, 2001, 45-48.	1.2	11
681	Solid-phase peptide synthesis using N ^t -trityl-amino acids. International Journal of Peptide Research and Therapeutics, 2001, 8, 331-338.	0.1	9
682	Proline: A Key Building Block in <i>de novo</i> -Designed Peptide Molecules. , 2001, , 432-434.		0
683	Design, synthesis, and conformational analysis of azacycloalkane amino acids as conformationally constrained probes for mimicry of peptide secondary structures. Biopolymers, 2000, 55, 101-122.	1.2	105
684	Orthogonal protecting groups for N ^t -amino and C-terminal carboxyl functions in solid-phase peptide synthesis. Biopolymers, 2000, 55, 123-139.	1.2	99

#	ARTICLE	IF	CITATIONS
685	Editorial: Current perspectives in peptide chemistry. II. Supports. <i>Biopolymers</i> , 2000, 55, 187-187.	1.2	0
686	An efficient solid-phase strategy for the construction of chemokines. <i>Journal of Peptide Science</i> , 2000, 6, 512-518.	0.8	15
687	Solid phase synthesis of sulfahydantoins. <i>Tetrahedron Letters</i> , 2000, 41, 3161-3163.	0.7	26
688	A modified backbone amide linker (BAL) solid-phase peptide synthesis strategy accommodating prolyl, N-alkylamino acyl, or histidyl derivatives at the C-terminus. <i>Tetrahedron Letters</i> , 2000, 41, 7277-7280.	0.7	33
689	Synthesis of aspartimide-free protected peptides on base-labile functionalized resins. <i>Tetrahedron Letters</i> , 2000, 41, 8093-8096.	0.7	12
690	Kahalalide B. Synthesis of a natural cyclodepsipeptide. <i>Tetrahedron Letters</i> , 2000, 41, 9765-9769.	0.7	25
691	Nsc and Fmoc N ⁺ -amino protection for solid-phase peptide synthesis: a parallel study. <i>Chemical Biology and Drug Design</i> , 2000, 56, 63-69.	1.2	23
692	Title is missing!. <i>International Journal of Peptide Research and Therapeutics</i> , 2000, 7, 187-194.	0.1	5
693	Exploring solid-phase approaches for the preparation of new β -lactams from amino acids. <i>Molecular Diversity</i> , 2000, 6, 75-84.	2.1	11
694	¹ H NMR spectroscopy with internal and external standards for the quantification of libraries. <i>Molecular Diversity</i> , 2000, 6, 165-168.	2.1	6
695	Allylic protection of thiols and cysteine. III. Use of Fmoc-Cys(Fsam)-OH for solid-phase peptide synthesis. <i>International Journal of Peptide Research and Therapeutics</i> , 2000, 7, 187-194.	0.1	0
696	Disulfide Bonded Cyclic Peptide Dimers and Trimers: An Easy Entry to High Symmetry Peptide Frameworks. <i>Synlett</i> , 2000, 2000, 172-181.	1.0	8
697	Solid-Phase Peptide Synthesis in the Reverse (N ⁺ →C) Direction. <i>Organic Letters</i> , 2000, 2, 1815-1817.	2.4	60
698	Substituted Guanidines: Introducing Diversity in Combinatorial Chemistry. <i>Organic Letters</i> , 2000, 2, 3539-3542.	2.4	34
699	Understanding the structure/reactivity of aminium/uronium salts as coupling reagents in peptide synthesis. <i>Tetrahedron Letters</i> , 1999, 40, 2641-2644.	0.7	13
700	Useful scaffolds and handles for creating diversity in the preparation of chemical libraries. <i>Reactive and Functional Polymers</i> , 1999, 41, 103-110.	2.0	10
701	Continuous-flow solid-phase peptide synthesis using polystyrene resins. <i>Chemical Biology and Drug Design</i> , 1999, 53, 682-683.	1.2	5
702	Title is missing!. <i>International Journal of Peptide Research and Therapeutics</i> , 1999, 6, 243-245.	0.1	3

#	ARTICLE	IF	CITATIONS
703	Pyrrolidide formation as a side reaction during activation of carboxylic acids by phosphonium salt coupling reagents. <i>International Journal of Peptide Research and Therapeutics</i> , 1999, 6, 243-245.	0.1	6
704	An HPLC-ESMS study on the solid-phase assembly of C-terminal proline peptides. , 1999, 5, 131-140.		39
705	Solid-Phase Synthesis with Tris(alkoxy)benzyl Backbone Amide Linkage (BAL) [â%]. <i>Chemistry - A European Journal</i> , 1999, 5, 2787-2795.	1.7	86
706	Backbone Amide Linker (BAL) Strategy for N ⁹ -Fluorenylmethoxycarbonyl (Fmoc) Solid-Phase Synthesis of Unprotected Peptide-Nitroanilides and Thioesters1. <i>Journal of Organic Chemistry</i> , 1999, 64, 8761-8769.	1.7	149
707	N ⁹ -Alloc temporary protection in solid-phase peptide synthesis. The use of amineâ€borane complexes as allyl group scavengers. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1999, , 2871-2874.	0.9	68
708	Synthesis and Binding Properties of Oligonucleotides Carrying Nuclear Localization Sequences. <i>Bioconjugate Chemistry</i> , 1999, 10, 1005-1012.	1.8	47
709	Title is missing!. <i>International Journal of Peptide Research and Therapeutics</i> , 1998, 5, 247-258.	0.1	0
710	Use of Onium Salt-Based Coupling Reagents in Peptide Synthesis1. <i>Journal of Organic Chemistry</i> , 1998, 63, 9678-9683.	1.7	245
711	Active carbonate resins: Application to the solid-phase synthesis of alcohol, carbamate and cyclic peptides. <i>Tetrahedron</i> , 1998, 54, 10125-10152.	1.0	50
712	â€High-loadâ€ polyethylene glycol-polystyrene (PEG-PS) graft supports for solid-phase synthesis. , 1998, 47, 365-380.		45
713	Reactive intermediates in peptide synthesis: Molecular and crystal structures of HOAt and HOObt, and some ester and amide derivatives of HOBT, HOAt and HOObt. <i>International Journal of Peptide Research and Therapeutics</i> , 1998, 5, 247-258.	0.1	5
714	Solid-phase synthesis of diketopiperazines, useful scaffolds for combinatorial chemistry. <i>Tetrahedron Letters</i> , 1998, 39, 2639-2642.	0.7	58
715	Structural/functional properties of the Glu1â€Ser57 Nâ€terminal fragment of human plasminogen: Conformational characterization and interaction with kringle domains. <i>Protein Science</i> , 1998, 7, 1947-1959.	3.1	16
716	Lysineâ€50 is a likely site for anchoring the plasminogen Nâ€terminal peptide to lysineâ€binding kringles. <i>Protein Science</i> , 1998, 7, 1960-1969.	3.1	37
717	An Easy Entry to a New High-Symmetry, Large Molecular Framework for Molecular Recognition Studies and de Novo Protein Design. Solvent Modulation of the Spontaneous Formation of a Cyclic Monomer, Dimer, or Trimer from a Bis-cysteine Peptide. <i>Journal of the American Chemical Society</i> , 1998, 120, 6639-6650.	6.6	17
718	Backbone Amide Linker (BAL) Strategy for Solid-Phase Synthesis of C-Terminal-Modified and Cyclic Peptides1,2,3. <i>Journal of the American Chemical Society</i> , 1998, 120, 5441-5452.	6.6	292
719	IBTM-Containing Gramicidin S Analogues:â€ Evidence for IBTM as a Suitable Type IIâ€ Î²-Turn Mimetic1,2. <i>Journal of the American Chemical Society</i> , 1997, 119, 10579-10586.	6.6	57
720	Poly(ethylene glycol)-Containing Supports for Solid-Phase Synthesis of Peptides and Combinatorial Organic Libraries. <i>ACS Symposium Series</i> , 1997, , 239-264.	0.5	19

#	ARTICLE	IF	CITATIONS
721	[7] Coupling reagents and activation. <i>Methods in Enzymology</i> , 1997, 289, 104-126.	0.4	91
722	[15] Convergent solid-phase peptide synthesis. <i>Methods in Enzymology</i> , 1997, 289, 313-336.	0.4	33
723	Occurrence and Minimization of Cysteine Racemization during Stepwise Solid-Phase Peptide Synthesis ^{1,2} . <i>Journal of Organic Chemistry</i> , 1997, 62, 4307-4312.	1.7	205
724	Total Synthesis of Dehydrodidemnin B. Use of Uronium and Phosphonium Salt Coupling Reagents in Peptide Synthesis in Solution. <i>Journal of Organic Chemistry</i> , 1997, 62, 354-366.	1.7	86
725	The use of the Nbb-resin for the solid-phase synthesis of peptide alkylesters and alkylamides. Synthesis of leuprolide. <i>Tetrahedron</i> , 1997, 53, 3179-3194.	1.0	20
726	Active carbonate resins for solid-phase synthesis through the anchoring of a hydroxyl function. Synthesis of cyclic and alcohol peptides. <i>Tetrahedron Letters</i> , 1997, 38, 883-886.	0.7	61
727	A new approach to Hmb-backbone protection of peptides: Synthesis and reactivity of N ¹ ±-Fmoc-N ¹ ±-(Hmb)amino acids. <i>Tetrahedron Letters</i> , 1997, 38, 2317-2320.	0.7	34
728	On the use of PyAOP, a phosphonium salt derived from HOAt, in solid-phase peptide synthesis. <i>Tetrahedron Letters</i> , 1997, 38, 4853-4856.	0.7	157
729	Use of Alloc-amino acids in solid-phase peptide synthesis. Tandem deprotection-coupling reactions using neutral conditions. <i>Tetrahedron Letters</i> , 1997, 38, 7275-7278.	0.7	156
730	Preparation and Applications of Xanthenylamide (XAL) Handles for Solid-Phase Synthesis of C-Terminal Peptide Amides under Particularly Mild Conditions ¹⁻³ . <i>Journal of Organic Chemistry</i> , 1996, 61, 6326-6339.	1.7	53
731	Synthesis of derivatives of (2S,4S)-4-hydroxy-2,5-dimethyl-3-oxohexanoic acid, a constituent of the didemnins. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1996, , 1427-1433.	0.9	3
732	On the use of novel coupling reagents for solid-phase peptide synthesis. <i>Techniques in Protein Chemistry</i> , 1996, , 515-523.	0.3	10
733	Use of N-tritylamino acids and PyAOP1 for the suppression of diketopiperazine formation in Fmoc/tBu solid-phase peptide synthesis using alkoxybenzyl ester anchoring linkages. <i>Tetrahedron Letters</i> , 1996, 37, 4195-4198.	0.7	65
734	New carbamate supports for the preparation of 3'-amino-modified oligonucleotides. <i>Bioorganic and Medicinal Chemistry</i> , 1996, 4, 1649-1658.	1.4	21
735	Lips symposia in print and other issues. <i>International Journal of Peptide Research and Therapeutics</i> , 1996, 3, 1-1.	0.1	2
736	3-(1-Piperidinyl)alanine formation during the preparation of C-terminal cysteine peptides with the Fmoc/t-Bu strategy. <i>International Journal of Peptide Research and Therapeutics</i> , 1996, 3, 157-166.	0.1	60
737	A COMPARATIVE STUDY OF SUPPORTS FOR THE SYNTHESIS OF OLIGONUCLEOTIDES WITHOUT USING AMMONIA. <i>Nucleosides & Nucleotides</i> , 1996, 15, 1871-1889.	0.5	21
738	Rearrangement of Glu(OtBu)- and Asp(OtBu)-containing peptides upon fluoride treatment in solid-phase synthesis. <i>International Journal of Peptide Research and Therapeutics</i> , 1995, 1, 213-220.	0.1	13

#	ARTICLE	IF	CITATIONS
739	Synthesis and applications of a new base-labile fluorene derived linker for solid-phase peptide synthesis. <i>Tetrahedron</i> , 1995, 51, 1449-1458.	1.0	41
740	Novel Carboxylic Acid and Carboxamide Protective Groups Based on the Exceptional Stabilization of the Cyclopropylmethyl Cation. <i>Journal of Organic Chemistry</i> , 1995, 60, 7718-7719.	1.7	32
741	Efficiency in Peptide Coupling: 1-Hydroxy-7-azabenzotriazole vs 3,4-Dihydro-3-hydroxy-4-oxo-1,2,3-benzotriazine. <i>Journal of Organic Chemistry</i> , 1995, 60, 3561-3564.	1.7	192
742	Convergent Solid Phase Peptide Synthesis: An Efficient Approach to the Synthesis of Highly Repetitive Protein Domains. <i>Journal of Organic Chemistry</i> , 1995, 60, 7575-7581.	1.7	29
743	Stepwise Automated Solid Phase Synthesis of Naturally Occurring Peptaibols Using Fmoc Amino Acid Fluorides. <i>Journal of Organic Chemistry</i> , 1995, 60, 405-410.	1.7	127
744	S-Phenylacetamidomethyl (Phacm): an orthogonal cysteine protecting group for Boc and Fmoc solid-phase peptide synthesis strategies. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1995, , 1095.	0.9	38
745	Convergent solid-phase peptide synthesis 12. * Chromatographic techniques for the purification of protected peptide segments. <i>International Journal of Peptide and Protein Research</i> , 1995, 46, 119-133.	0.1	7
746	Structural studies of reagents for peptide bond formation: Crystal and molecular structures of HBTU and HATU. <i>International Journal of Peptide Research and Therapeutics</i> , 1994, 1, 57-67.	0.1	86
747	Solid-phase synthesis of peptides using allylic anchoring groups 2. Palladium-catalysed cleavage of Fmoc-protected peptides. <i>Tetrahedron Letters</i> , 1994, 35, 4437-4440.	0.7	30
748	Stepwise solid-phase synthesis of oligonucleotide-peptide hybrids. <i>Tetrahedron Letters</i> , 1994, 35, 2733-2736.	0.7	50
749	Solid-phase N-glycopeptide synthesis using allyl side-chain protected Fmoc-amino acids. <i>Tetrahedron Letters</i> , 1994, 35, 1033-1034.	0.7	42
750	Preparation and applications of polyethylene glycol-polystyrene graft resin supports for solid-phase peptide synthesis. <i>Reactive & Functional Polymers</i> , 1994, 22, 243-258.	0.8	128
751	Racemization studies during solid-phase peptide synthesis using azabenzotriazole-based coupling reagents. <i>Tetrahedron Letters</i> , 1994, 35, 2279-2282.	0.7	199
752	Severe side-reaction in the acidolytic cleavage of a C-terminal Met-containing peptide from the solid support. Formation of the homoserine lactone peptide. <i>Tetrahedron Letters</i> , 1994, 35, 175-178.	0.7	9
753	Solid-phase synthesis of head-to-tail-cyclic peptides via lysine side-chain anchoring. <i>Tetrahedron Letters</i> , 1994, 35, 9633-9636.	0.7	81
754	Formation of Disulfide Bonds in Synthetic Peptides and Proteins. , 1994, 35, 91-170.		109
755	Advantageous applications of azabenzotriazole (triazolopyridine)-based coupling reagents to solid-phase peptide synthesis. <i>Journal of the Chemical Society Chemical Communications</i> , 1994, , 201.	2.0	329
756	Peptides in molecular recognition: synthetic and conformational aspects. <i>Biochemical Society Transactions</i> , 1994, 22, 1045-1048.	1.6	1

#	ARTICLE	IF	CITATIONS
757	Chemical synthesis of a fully active transcriptional repressor protein.. Proceedings of the National Academy of Sciences of the United States of America, 1994, 91, 5178-5182.	3.3	15
758	Solid-Phase Synthesis of Cyclic Peptides. , 1994, , 39-58.		39
759	Antibodies cross-reactive with the scorpion-toxin ii from <i>Androctonus australis Hector</i> elicited in mice by a synthetic peptide. <i>Natural Toxins</i> , 1993, 1, 255-262.	1.0	7
760	Automated Allyl Cleavage for Continuous-Flow Synthesis of Cyclic and Branched Peptides. <i>Analytical Biochemistry</i> , 1993, 212, 303-310.	1.1	128
761	A novel, convenient, three-dimensional orthogonal strategy for solid-phase synthesis of cyclic peptides. <i>Tetrahedron Letters</i> , 1993, 34, 1549-1552.	0.7	250
762	Convergent solid-phase peptide synthesis. XI. Synthesis and purification of protected peptide segments spanning the entire sequence of the uteroglobin monomer using the photolabile nbb-resin.. <i>Tetrahedron</i> , 1993, 49, 10069-10078.	1.0	16
763	Convergent solid-phase peptide synthesis. <i>Tetrahedron</i> , 1993, 49, 11065-11133.	1.0	205
764	The 2,2,4,6,7-pentamethyl-dihydrobenzofuran-5-sulfonyl group (Pbf) as arginine side chain protectant. <i>Tetrahedron Letters</i> , 1993, 34, 7829-7832.	0.7	106
765	Use of a Base-Labile Protected Derivative of 6-Mercaptohexanol for the Preparation of Oligonucleotides Containing a Thiol Group at the 5' End. <i>Nucleosides & Nucleotides</i> , 1993, 12, 993-1005.	0.5	7
766	Design, synthesis, and complexing properties of (1Cys-1'Cys,4Cys-4'Cys)-dithiobis(Ac-L-1Cys-L-Pro-D-Val-L-4Cys-NH ₂). The first example of a new family of ion-binding peptides. <i>Journal of the American Chemical Society</i> , 1993, 115, 11663-11670.	6.6	27
767	Unequivocal synthesis and characterization of a parallel and an antiparallel bis-cystine peptide. <i>Journal of Organic Chemistry</i> , 1993, 58, 6319-6328.	1.7	13
768	Acidolytic cleavage of tris(alkoxy)benzylamide (PAL) -internal reference- amino acyl (IRAA) anchoring linkages: validation of accepted procedures in solid-phase peptide synthesis (SPPS). <i>International Journal of Peptide and Protein Research</i> , 1993, 41, 307-312.	0.1	18
769	Allyl-based orthogonal solid phase peptide synthesis. , 1993, , 191-193.		10
770	Preparation and applications of xanthenylamide (XAL) handles for mild Fmoc solid-phase synthesis of C-terminal peptide amides. , 1993, , 301-304.		0
771	S-2,4,6-trimethoxybenzyl (Tmob): a novel cysteine protecting group for the N.alpha.-(9-fluorenylmethoxycarbonyl) (Fmoc) strategy of peptide synthesis. <i>Journal of Organic Chemistry</i> , 1992, 57, 3013-3018.	1.7	52
772	A synthetic procedure for the preparation of oligonucleotides without using ammonia and its application for the synthesis of oligonucleotides containing O-4-alkyl thymidines.. <i>Tetrahedron</i> , 1992, 48, 4171-4182.	1.0	36
773	-2-(2,4-dinitrophenyl)ethyloxycarbonyl-amino acids, new base labile protected derivatives suitable for solid-phase peptide synthesis.. <i>Tetrahedron Letters</i> , 1992, 33, 4989-4992.	0.7	5
774	Reversible protection of lysine to facilitate the purification of protected peptide segments. <i>Tetrahedron Letters</i> , 1992, 33, 397-400.	0.7	7

#	ARTICLE	IF	CITATIONS
775	S-2-(2,4-dinitrophenyl)ethyl-cysteine: a new derivative for solid-phase peptide synthesis. Tetrahedron Letters, 1992, 33, 2391-2394.	0.7	15
776	A new fluorene-derived anchor for solid-phase synthesis of protected peptides. Tetrahedron Letters, 1992, 33, 1775-1778.	0.7	28
777	Synthesis and ion-binding properties of an immobilized bis-cysteine peptide. Bioorganic and Medicinal Chemistry Letters, 1992, 2, 281-284.	1.0	0
778	A new approach to the solid-phase peptide synthesis of peptide alkyl-amides and esters. Tetrahedron Letters, 1992, 33, 2183-2186.	0.7	11
779	Enzymatic peptide fragment condensation. Choice of reaction media for the synthesis of an insect neuropeptide derivative. Biotechnology Letters, 1992, 6, 69-72.	0.5	3
780	Biopolymer syntheses on novel polyethylene glycol-polystyrene (PEG-PS) graft supports. , 1992, , 603-604.		18
781	Novel cysteine protecting groups for the N ⁺ -9-fluorenylmethoxycarbonyl (Fmoc) strategy of peptide synthesis. , 1992, , 605-606.		1
782	Convergent solid-phase peptide synthesis. , 1992, , 607-608.		1
783	RNA binding characteristics of a 16 kDa glycine-rich protein from maize. Plant Journal, 1992, 2, 999-1003.	2.8	0
784	Synthesis and biological activity of O-glycosylated morphiceptin analogues. Journal of the Chemical Society Perkin Transactions 1, 1991, , 1755-1759.	0.9	36
785	NPE-resin, a new approach to the solid-phase synthesis of protected peptides and oligonucleotides I : Synthesis of the supports and their application to oligonucleotide synthesis.. Tetrahedron Letters, 1991, 32, 1511-1514.	0.7	42
786	NPE-resin, a new approach to the solid-phase synthesis of protected peptides and oligonucleotides II. Synthesis of protected peptides. Tetrahedron Letters, 1991, 32, 1515-1518.	0.7	31
787	Binding and toxicity of apamin. Characterization of the active site. FEBS Journal, 1991, 196, 639-645.	0.2	57
788	Hypersensitive acid-labile (HAL) tris(alkoxy)benzyl ester anchoring for solid-phase synthesis of protected peptide segments. Tetrahedron Letters, 1991, 32, 1015-1018.	0.7	40
789	Solid-phase synthesis of peptides using allylic anchoring groups. An investigation of their palladium-catalysed cleavage. Tetrahedron Letters, 1991, 32, 4207-4210.	0.7	51
790	Convergent solid-phase peptide synthesis. X. Synthesis and purification of protected peptide fragments using the photolabile Nbb-resin. Tetrahedron, 1991, 47, 9867-9880.	1.0	27
791	Synthesis of defined peptide-oligonucleotide hybrids containing a nuclear transport signal sequence.. Tetrahedron, 1991, 47, 4113-4120.	1.0	84
792	Convergent solid-phase peptide synthesis. International Journal of Peptide and Protein Research, 1991, 37, 58-60.	0.1	23

#	ARTICLE	IF	CITATIONS
793	Cyclization of disulfide-containing peptides in solid-phase synthesis ^{>1</sup>. International Journal of Peptide and Protein Research, 1991, 37, 402-413.}	0.1	85
794	Mild orthogonal solid-phase peptide synthesis. , 1991, , 139-142.		3
795	Solid-phase synthesis of new glycosyl enkephalinamides. , 1991, , 416-417.		0
796	Use of BOP reagent for the suppression of diketopiperazine formation in boc/bzl solid-phase peptide synthesis. Tetrahedron Letters, 1990, 31, 7363-7366.	0.7	59
797	Arenesulphonyltriaxolides as condensing reagents in solid phase peptide synthesis. Tetrahedron Letters, 1990, 31, 1915-1918.	0.7	14
798	Solid-Phase Synthesis of Glycopeptide Amides under Mild Conditions: Morphiceptin Analogues. Angewandte Chemie International Edition in English, 1990, 29, 291-292.	4.4	21
799	Festphasen-Synthese von Glycopeptidamiden unter milden Bedingungen: Morphiceptin-Analoga. Angewandte Chemie, 1990, 102, 311-313.	1.6	14
800	Improved method for the synthesis of o-glycosylated fmoc amino acids to be used in solid-phase glycopeptide synthesis (Fmoc = fluoren-9-ylmethoxycarbonyl). Journal of the Chemical Society Chemical Communications, 1990, , 965-967.	2.0	30
801	Preparation and application of the 5-(4-(9-fluorenylmethyloxycarbonyl)aminomethyl-3,5-dimethoxyphenoxy)-valeric acid (PAL) handle for the solid-phase synthesis of C-terminal peptide amides under mild conditions. Journal of Organic Chemistry, 1990, 55, 3730-3743.	1.7	343
802	Solid-phase-mediated peptide heterodisulfide formation. Journal of the American Chemical Society, 1990, 112, 5345-5347.	6.6	21
803	Application of acetamidomethyl and 9-fluorenylmethyl groups for efficient side protection of penicillamine in solid-phase peptide synthesis. International Journal of Peptide and Protein Research, 1990, 35, 434-440.	0.1	11
804	Practical approach to solid-phase synthesis of C-terminal peptide amides under mild conditions based on a photolysable anchoring linkage ^{>1</sup>. International Journal of Peptide and Protein Research, 1990, 36, 31-45.}	0.1	58
805	Orthogonal solid-phase synthesis of human gastrin under mild conditions*. International Journal of Peptide and Protein Research, 1990, 35, 527-538.	0.1	28
806	Solid-phase synthesis of peptides with C-terminal asparagine or glutamine. International Journal of Peptide and Protein Research, 1990, 35, 284-286.	0.1	32
807	Use of polystyrene-1% divinylbenzene and Kel-F-g-styrene for the simultaneous synthesis of peptides. Reactive & Functional Polymers, 1989, 10, 259-268.	0.8	6
808	Quantitative monitoring of carboxyl groups in polymers. Analytica Chimica Acta, 1989, 219, 161-163.	2.6	2
809	Convenient synthesis of a cyclic peptide disulfide: A type II β -turn structural model. Tetrahedron Letters, 1989, 30, 2441-2444.	0.7	35
810	Convergent solid phase peptide synthesis. VII. Good yields in the coupling of protected segments on a solid support. Tetrahedron, 1989, 45, 4637-4648.	1.0	21

#	ARTICLE	IF	CITATIONS
811	Comparative study of supports for solid-phase coupling of protected-peptide segments. <i>Journal of Organic Chemistry</i> , 1989, 54, 360-366.	1.7	51
812	Use of the Npys thiol protection in solid phase peptide synthesis Application to direct peptide-protein conjugation through cysteine residues. <i>International Journal of Peptide and Protein Research</i> , 1989, 34, 124-128.	0.1	47
813	Molecular cloning of cDNAs encoding a putative cell wall protein from <i>Zea mays</i> and immunological identification of related polypeptides. <i>Plant Molecular Biology</i> , 1988, 11, 483-493.	2.0	70
814	Uteroglobulin-like peptide cavities I. Synthesis of antiparallel and parallel dimers of bis-cysteine peptides. <i>Tetrahedron Letters</i> , 1988, 29, 3845-3848.	0.7	34
815	Use of polar picolyl protecting groups in peptide synthesis. <i>Journal of Organic Chemistry</i> , 1988, 53, 5386-5389.	1.7	17
816	On the use of <i>s</i> - <i>t</i> -butylsulphenyl group for protection of cysteine in solid-phase peptide synthesis using fmoc-amino acids. <i>Tetrahedron</i> , 1987, 43, 2675-2680.	1.0	77
817	Convergent solid phase peptide synthesis. v. synthesis of the 1-4, 32-34, and 53-59 protected segments of the toxin ii of <i>androctonus australis hector</i> .. <i>Tetrahedron</i> , 1987, 43, 5961-5971.	1.0	20
818	Convergent solid phase peptide synthesis vi : synthesis by the fmoc procedure with a modified protocol of two protected segments, sequence 5-17 and 18-31 of the neurotoxin ii of the scorpion <i>androctonus australis hector</i> .. <i>Tetrahedron</i> , 1987, 43, 5973-5980.	1.0	15
819	Mild, orthogonal solid-phase peptide synthesis: use of <i>N</i> -dithiasuccinoyl (Dts) amino acids and <i>N</i> -isopropyl(dithio)carbonylproline, together with <i>p</i> -alkoxybenzyl ester anchoring linkages*. <i>International Journal of Peptide and Protein Research</i> , 1987, 30, 177-205.	0.1	36
820	An acid-labile anchoring linkage for solid-phase synthesis of <i>C</i> -terminal peptide amides under mild conditions*. <i>International Journal of Peptide and Protein Research</i> , 1987, 30, 206-216.	0.1	106
821	A convenient general method for synthesis of <i>N</i> - or <i>N</i> -dithiasuccinoyl (Dts) amino acids and dipeptides: application of polyethylene glycol as a carrier for functional purification*. <i>International Journal of Peptide and Protein Research</i> , 1987, 30, 740-783.	0.1	39
822	(<i>S</i>)-9-Fluorenylmethyl-L-cysteine, a useful HF-stable derivative for peptide synthesis. <i>Journal of the Chemical Society Chemical Communications</i> , 1986, , 1501.	2.0	23
823	Synthesis in vitro of a seven amino acid peptide encoded in the leader RNA of Rous sarcoma virus. <i>Journal of Molecular Biology</i> , 1986, 190, 45-57.	2.0	75
824	Three-dimensional orthogonal protection scheme for solid-phase peptide synthesis under mild conditions. <i>Journal of the American Chemical Society</i> , 1985, 107, 4936-4942.	6.6	141
825	Improved approach for anchoring <i>N</i> - or <i>N</i> -fluorenylmethyloxycarbonylamino acids as <i>p</i> -alkoxybenzyl esters in solid-phase peptide synthesis. <i>International Journal of Peptide and Protein Research</i> , 1985, 26, 92-97.	0.1	76
826	Application of <i>N,N</i> -dimethylformamide dineopentyl acetal for efficient anchoring of <i>N</i> -isopropyl- <i>N</i> -fluorenylmethyloxycarbonylamino acids as <i>p</i> -alkoxybenzyl esters in solid-phase peptide synthesis. <i>International Journal of Peptide and Protein Research</i> , 1984, 23, 342-349.	0.1	27
827	Convergent solid phase peptide synthesis. II. Synthesis of the 1-6 apamin protected segment on a NBB-resin. Synthesis of apamin. <i>Tetrahedron</i> , 1982, 38, 1193-1201.	1.0	56
828	Acid-base properties of 4-nitro-l-histidine and related compounds. <i>Bioorganic Chemistry</i> , 1979, 8, 59-67.	2.0	3

#	ARTICLE	IF	CITATIONS
829	The (Classic Concept of) Solid Support. , 0, , 1-14.		1
830	High-Throughput Synthesis of Natural Products. , 0, , 613-640.		1
831	Immobilized Enzymes in Organic Synthesis. , 0, , 365-380.		4
832	CHAPTER 18. Solid-Phase Peptide Synthesis, the State of the Art: Challenges and Opportunities. RSC Drug Discovery Series, 0, , 518-550.	0.2	13
833	Drug discovery: a multifactorial ecosystem. , 0, , 1-5.		0
834	Solid-Phase Peptide Synthesis Using a Four-Dimensional (Safety-Catch) Protecting Group Scheme. Journal of Organic Chemistry, 0, , .	1.7	2
835	2-Methoxy-4-methylsulfinylbenzyl Alcohol as a Safety-Catch Linker for the Fmoc- <i>t</i> -Bu Solid-Phase Peptide Synthesis Strategy. Journal of Organic Chemistry, 0, , .	1.7	4