

Fernando Albericio

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/6676408/fernando-albericio-publications-by-year.pdf>

Version: 2024-04-28

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

815
papers

24,566
citations

72
h-index

119
g-index

942
ext. papers

27,463
ext. citations

5.4
avg, IF

7.35
L-index

#	Paper	IF	Citations
815	The Pharmaceutical Industry in 2021. An Analysis of FDA Drug Approvals from the Perspective of Molecules.. <i>Molecules</i> , 2022 , 27,	4.8	8
814	Linkers: An Assurance for Controlled Delivery of Antibody-Drug Conjugate.. <i>Pharmaceutics</i> , 2022 , 14,	6.4	7
813	Understanding OxymaPure as a Peptide Coupling Additive: A Guide to New Oxyma Derivatives.. <i>ACS Omega</i> , 2022 , 7, 6007-6023	3.9	0
812	2021 FDA TIDES (Peptides and Oligonucleotides) Harvest.. <i>Pharmaceutics</i> , 2022 , 15,	5.2	9
811	s-Triazine: A Multidisciplinary and International Journey. <i>Chemistry Proceedings</i> , 2021 , 3, 53		
810	Minimizing side reactions during amide formation using DIC and oxymapure in solid-phase peptide synthesis. <i>Tetrahedron Letters</i> , 2021 , 85, 153462	2	2
809	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	8.3	3
808	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	3.9	1
807	Improving the Gastrointestinal Stability of Linaclotide. <i>Journal of Medicinal Chemistry</i> , 2021 , 64, 8384-8390,	8.0	5
806	Targeting Energy Expenditure-Drugs for Obesity Treatment. <i>Pharmaceutics</i> , 2021 , 14,	5.2	5
805	An Androsterone-H @C hybrid: Synthesis, Properties and Molecular Docking Simulations with SARS-Cov-2. <i>ChemPlusChem</i> , 2021 , 86, 970-971	2.8	0
804	Scope and Limitations of Barbituric and Thiobarbituric Amino Acid Derivatives as Protecting Groups for Solid-Phase Peptide Synthesis: Towards a Green Protecting Group. <i>ChemistrySelect</i> , 2021 , 6, 6626-6630	1.8	1
803	Super-Cationic Peptide Dendrimers-Synthesis and Evaluation as Antimicrobial Agents. <i>Antibiotics</i> , 2021 , 10,	4.9	2
802	A novel 'smart' 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	6	10
801	Effects of elderflower extract enriched with polyphenols on antioxidant defense of salmon leukocytes. <i>Electronic Journal of Biotechnology</i> , 2021 , 52, 13-20	3.1	2
800	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	3.9	0
799	NOXA upregulation by the prohibitin-binding compound fluorizoline is transcriptionally regulated by integrated stress response-induced ATF3 and ATF4. <i>FEBS Journal</i> , 2021 , 288, 1271-1285	5.7	6

798	Lactoferrin-dual drug nanoconjugate: Synergistic anti-tumor efficacy of docetaxel and the NF- κ B inhibitor celastrol. <i>Materials Science and Engineering C</i> , 2021 , 118, 111422	8.3	13
797	1,3,5-Triazine as core for the preparation of dendrons. <i>Arkivoc</i> , 2021 , 2020, 64-73	0.9	2
796	The Pharmaceutical Industry in 2020. An Analysis of FDA Drug Approvals from the Perspective of Molecules. <i>Molecules</i> , 2021 , 26,	4.8	33
795	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	10	14
794	Green solvents in the biotechnology-based pharmaceutical industry 2021 , 87-104		1
793	Nature-inspired dimerization as a strategy to modulate neuropeptide pharmacology exemplified with vasopressin and oxytocin. <i>Chemical Science</i> , 2021 , 12, 4057-4062	9.4	4
792	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	10	10
791	A native mass spectrometry platform identifies HOP inhibitors that modulate the HSP90-HOP protein-protein interaction. <i>Chemical Communications</i> , 2021 , 57, 10919-10922	5.8	1
790	2020 FDA TIDES (Peptides and Oligonucleotides) Harvest. <i>Pharmaceuticals</i> , 2021 , 14,	5.2	20
789	Propylphosphonic Anhydride (T3P) as Coupling Reagent for Solid-Phase Peptide Synthesis. <i>ChemistrySelect</i> , 2021 , 6, 2649-2657	1.8	3
788	Liquid Phase Peptide Synthesis via One-Pot Nanostar Sieving (PEPSTAR). <i>Angewandte Chemie - International Edition</i> , 2021 , 60, 7786-7795	16.4	8
787	An Androsterone-H @C hybrid: Synthesis, Properties and Molecular Docking Simulations with SARS-Cov-2. <i>ChemPlusChem</i> , 2021 , 86, 972-981	2.8	1
786	The Antiproliferative and Apoptotic Effect of a Novel Synthesized -Triazine Dipeptide Series, and Toxicity Screening in Zebrafish Embryos. <i>Molecules</i> , 2021 , 26,	4.8	3
785	Liquid Phase Peptide Synthesis via One-Pot Nanostar Sieving (PEPSTAR). <i>Angewandte Chemie</i> , 2021 , 133, 7865-7874	3.6	1
784	s-Triazine: A Privileged Structure for Drug Discovery and Bioconjugation. <i>Molecules</i> , 2021 , 26,	4.8	9
783	Rhodiasolv PolarClean is a greener alternative in solid-phase peptide synthesis. <i>Green Chemistry Letters and Reviews</i> , 2021 , 14, 545-550	4.7	2
782	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	1.8	0
781	Tea Bags for Fmoc Solid-Phase Peptide Synthesis: An Example of Circular Economy. <i>Molecules</i> , 2021 , 26,	4.8	2

780	Amide Formation: Choosing the Safer Carbodiimide in Combination with OxymaPure to Avoid HCN Release. <i>Organic Letters</i> , 2021 , 23, 6900-6904	6.2	4
779	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	3	0
778	In vivo 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	7.4	4
777	Protocol for efficient solid-phase synthesis of peptides containing 1-hydroxypyridine-2-one (1,2-HOPO). <i>MethodsX</i> , 2020 , 7, 101082	1.9	1
776	Insights into the chemistry of the amphibactin-metal (M) interaction and its role in antibiotic resistance. <i>Scientific Reports</i> , 2020 , 10, 21049	4.9	0
775	Synthetic peptides to produce antivenoms against the Cys-rich toxins of arachnids. <i>Toxicon: X</i> , 2020 , 6, 100038	2.6	2
774	Enamine Barbiturates and Thiobarbiturates as a New Class of Bacterial Urease Inhibitors. <i>Applied Sciences (Switzerland)</i> , 2020 , 10, 3523	2.6	0
773	Protocol for synthesis of di- and tri-substituted s-triazine derivatives. <i>MethodsX</i> , 2020 , 7, 100825	1.9	1
772	2019 FDA TIDES (Peptides and Oligonucleotides) Harvest. <i>Pharmaceuticals</i> , 2020 , 13,	5.2	29
771	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	5.6	12
770	Synthesis of Stable Cholesteryl-Polyethylene Glycol-Peptide Conjugates with Non-Disperse Polyethylene Glycol Lengths. <i>ACS Omega</i> , 2020 , 5, 5508-5519	3.9	3
769	Breaking a Couple: Disulfide Reducing Agents. <i>ChemBioChem</i> , 2020 , 21, 1947-1954	3.8	6
768	Use of a phosphopeptide as a ligand to purify phospholipase A 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	3.2	2
767	From Ugi Multicomponent Reaction to Linkers for Bioconjugation. <i>ACS Omega</i> , 2020 , 5, 7424-7431	3.9	5
766	Somuncurins: Bioactive Peptides from the Skin of the Endangered Endemic Patagonian Frog. <i>Journal of Natural Products</i> , 2020 , 83, 972-984	4.9	3
765	Crystal Structure and Theoretical Investigation of Thiobarbituric Acid Derivatives as Nonlinear Optical (NLO) Materials. <i>Crystals</i> , 2020 , 10, 442	2.3	1
764	Revisiting NO as Protecting Group of Arginine in Solid-Phase Peptide Synthesis. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	3
763	Barbiturate- and Thiobarbiturate-Based -Triazine Hydrazone Derivatives with Promising Antiproliferative Activities. <i>ACS Omega</i> , 2020 , 5, 15805-15811	3.9	8

762	The Pharmaceutical Industry in 2019. An Analysis of FDA Drug Approvals from the Perspective of Molecules. <i>Molecules</i> , 2020 , 25,	4.8	76
761	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. <i>Journal of Nanobiotechnology</i> , 2020 , 18, 20	9.4	34
760	Greening Fmoc/tBu solid-phase peptide synthesis. <i>Green Chemistry</i> , 2020 , 22, 996-1018	10	44
759	Phenol as a Modulator in the Chemical Reactivity of 2,4,6-Trichloro-1,3,5-triazine: Rules of the Game II. <i>Australian Journal of Chemistry</i> , 2020 , 73, 352	1.2	2
758	Carpino's protecting groups, beyond the Boc and the Fmoc. <i>Peptide Science</i> , 2020 , 112, e24164	3	4
757	Cleaving protected peptides from 2-chlorotrityl chloride resin. Moving away from dichloromethane. <i>Green Chemistry</i> , 2020 , 22, 2840-2845	10	6
756	Solid-Phase Synthesis of Head to Side-Chain Tyr-Cyclodepsipeptides Through a Cyclative Cleavage From Fmoc-MeDbz/MeNbz-resins. <i>Frontiers in Chemistry</i> , 2020 , 8, 298	5	6
755	Naturally Occurring Oxazole-Containing Peptides. <i>Marine Drugs</i> , 2020 , 18,	6	18
754	Successful development of a method for the incorporation of Fmoc-Arg(Pbf)-OH in solid-phase peptide synthesis using N-butylpyrrolidinone (NBP) as solvent. <i>Green Chemistry</i> , 2020 , 22, 3162-3169	10	14
753	OxymaPure Coupling Reagents: Beyond Solid-Phase Peptide Synthesis. <i>Synthesis</i> , 2020 , 52, 3189-3210	2.9	2
752	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	5.1	8
751	Protocol for bevacizumab purification using Ac-PHQGQHIGVSK-agarose. <i>MethodsX</i> , 2020 , 7, 100769	1.9	1
750	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	6.8	13
749	Novel formulation of antimicrobial peptides enhances antimicrobial activity against methicillin-resistant <i>Staphylococcus aureus</i> (MRSA). <i>Amino Acids</i> , 2020 , 52, 1439-1457	3.5	12
748	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	5.1	2
747	Disulfide-Based Protecting Groups for the Cysteine Side Chain. <i>Organic Letters</i> , 2020 , 22, 9644-9647	6.2	2
746	Solid-phase synthesis of peptides containing 1-Hydroxypyridine-2-one (1,2-HOPO). <i>Tetrahedron Letters</i> , 2020 , 61, 152299	2	2
745	N-Butylpyrrolidinone for Solid-Phase Peptide Synthesis is Environmentally Friendlier and Synthetically Better than DMF. <i>ChemSusChem</i> , 2020 , 13, 5288-5294	8.3	13

744	Radical Dendrimers Based on Biocompatible Oligoethylene Glycol Dendrimers as Contrast Agents for MRI. <i>Pharmaceutics</i> , 2020 , 12,	6.4	10
743	Novel 4,6-Disubstituted -Triazin-2-yl Amino Acid Derivatives as Promising Antifungal Agents. <i>Journal of Fungi (Basel, Switzerland)</i> , 2020 , 6,	5.6	4
742	The tea-bag protocol for comparison of Fmoc removal reagents in solid-phase peptide synthesis. <i>Amino Acids</i> , 2020 , 52, 1201-1205	3.5	6
741	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	2	4
740	Gold Nanoparticles Mediate Improved Detection of Amyloid Aggregates by Fluorescence. <i>Nanomaterials</i> , 2020 , 10,	5.4	15
739	Investigating Triorthogonal Chemoselectivity. Effect of Azide Substitution on the Triazine Core. <i>Organic Letters</i> , 2019 , 21, 7888-7892	6.2	7
738	Green Transformation of Solid-Phase Peptide Synthesis. <i>ACS Sustainable Chemistry and Engineering</i> , 2019 , 7, 3671-3683	8.3	52
737	Large-Area Biomolecule Nanopatterns on Diblock Copolymer Surfaces for Cell Adhesion Studies. <i>Nanomaterials</i> , 2019 , 9,	5.4	5
736	Bypassing Osmotic Shock Dilemma in a Polystyrene Resin Using the Green Solvent Cyclopentyl methyl Ether (CPME): A Morphological Perspective. <i>Polymers</i> , 2019 , 11,	4.5	6
735	Optimized Stepwise Synthesis of the API Liraglutide Using BAL Resin and Pseudoprolines. <i>ACS Omega</i> , 2019 , 4, 8674-8680	3.9	4
734	2018 FDA Tides Harvest. <i>Pharmaceutics</i> , 2019 , 12,	5.2	28
733	Carbosilane Dendron-Peptide Nanoconjugates as Antimicrobial Agents. <i>Molecular Pharmaceutics</i> , 2019 , 16, 2661-2674	5.6	19
732	Pseudo-Wang Handle for the Preparation of Fully Protected Peptides. Synthesis of Liraglutide by Fragment Condensation. <i>Organic Letters</i> , 2019 , 21, 2459-2463	6.2	8
731	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 ^{5.1}	5.1	19
730	Troubleshooting When Using Valerolactone (GVL) in Green Solid-Phase Peptide Synthesis. <i>Organic Process Research and Development</i> , 2019 , 23, 1096-1100	3.9	17
729	The Pharmaceutical Industry in 2018. An Analysis of FDA Drug Approvals from the Perspective of Molecules. <i>Molecules</i> , 2019 , 24,	4.8	70
728	Valerolactone (GVL): An eco-friendly anchoring solvent for solid-phase peptide synthesis. <i>Tetrahedron Letters</i> , 2019 , 60, 151058	2	14
727	Identification of Antimicrobial Peptides from the Microalgae (Kylin) Butcher and Bactericidal Activity Improvement. <i>Marine Drugs</i> , 2019 , 17,	6	44

726	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	3.9	5
725	Chemical Modification of Microcin J25 Reveals New Insights on the Stereospecific Requirements for Antimicrobial Activity. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	3
724	Scope and Limitations of γ -Valerolactone (GVL) as a Green Solvent to be Used with Base for Fmoc Removal in Solid Phase Peptide Synthesis. <i>Molecules</i> , 2019 , 24,	4.8	10
723	OctaGel Resin - A New PEG-PS-based Solid Support for Solid-Phase Peptide Synthesis. <i>Letters in Organic Chemistry</i> , 2019 , 16, 935-940	0.6	1
722	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	8.6	3
721	Synthesis and Antimicrobial Activity of a New Series of Thiazolidine-2,4-diones Carboxamide and Amino Acid Derivatives. <i>Molecules</i> , 2019 , 25,	4.8	7
720	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	6.2	3
719	Report of antileishmanial properties of Iberian macroalgae. <i>Natural Product Research</i> , 2019 , 33, 1778-1782	3	3
718	Bacteria Hunt Bacteria through an Intriguing Cyclic Peptide. <i>ChemMedChem</i> , 2019 , 14, 24-51	3.7	6
717	Toward the Synthesis of Phormidolides. <i>ACS Omega</i> , 2018 , 3, 2351-2362	3.9	2
716	Bioconjugation through Mesitylene Thiol Alkylation. <i>Bioconjugate Chemistry</i> , 2018 , 29, 1199-1208	6.3	4
715	Solid-Phase Synthesis of Pyrrole Derivatives through a Multicomponent Reaction Involving Lys-Containing Peptides. <i>ACS Combinatorial Science</i> , 2018 , 20, 187-191	3.9	13
714	1,3,5-Triazino Peptide Derivatives: Synthesis, Characterization, and Preliminary Antileishmanial Activity. <i>ChemMedChem</i> , 2018 , 13, 725-735	3.7	19
713	Application of Decafluorobiphenyl (DFBP) Moiety as a Linker in Bioconjugation. <i>Bioconjugate Chemistry</i> , 2018 , 29, 225-233	6.3	5
712	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	8.2	46
711	3D Electrophoresis-Assisted Lithography (3DEAL): 3D Molecular Printing to Create Functional Patterns and Anisotropic Hydrogels. <i>Advanced Functional Materials</i> , 2018 , 28, 1703014	15.6	8
710	Single step recombinant human growth hormone (rhGH) purification from milk by peptide affinity chromatography. <i>Biotechnology Progress</i> , 2018 , 34, 999-1005	2.8	2
709	Microwave-Assisted Green Solid-Phase Peptide Synthesis Using γ -Valerolactone (GVL) as Solvent. <i>ACS Sustainable Chemistry and Engineering</i> , 2018 , 6, 8034-8039	8.3	45

708	Solid-phase synthesis of homodetic cyclic peptides from Fmoc-MeDbz-resin. <i>Tetrahedron Letters</i> , 2018 , 59, 1779-1782	2	11
707	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	6	9
706	N-methylation in amino acids and peptides: Scope and limitations. <i>Biopolymers</i> , 2018 , 109, e23110	2.2	20
705	Teixobactin as a scaffold for unlimited new antimicrobial peptides: SAR study. <i>Bioorganic and Medicinal Chemistry</i> , 2018 , 26, 2788-2796	3.4	29
704	Antibacterial Activity of Teixobactin Derivatives on Clinically Relevant Bacterial Isolates. <i>Frontiers in Microbiology</i> , 2018 , 9, 1535	5.7	19
703	Further applications of classical amide coupling reagents: Microwave-assisted esterification on solid phase. <i>Journal of Peptide Science</i> , 2018 , 24, e3111	2.1	8
702	Identification of Peptides in Flowers of with Antimicrobial Activity against Aquaculture Pathogens. <i>Molecules</i> , 2018 , 23,	4.8	11
701	Formation of N-terminal 2-dialkyl amino oxazoles from guanidinated derivatives under mild conditions. <i>Organic and Biomolecular Chemistry</i> , 2018 , 16, 5661-5666	3.9	2
700	Exploiting the Thiobarbituric Acid Scaffold for Antibacterial Activity. <i>ChemMedChem</i> , 2018 , 13, 1923-1930	3.7	7
699	The Pharmaceutical Industry in 2017. An Analysis of FDA Drug Approvals from the Perspective of Molecules. <i>Molecules</i> , 2018 , 23,	4.8	34
698	Chemical Synthesis and Functional Analysis of VarvA Cyclotide. <i>Molecules</i> , 2018 , 23,	4.8	6
697	On the Importance of Polyurethane and Polyurea Nanosystems for Future Drug Delivery. <i>Current Drug Delivery</i> , 2018 , 15, 37-43	3.2	10
696	Choosing the Right Coupling Reagent for Peptides: A Twenty-Five-Year Journey. <i>Organic Process Research and Development</i> , 2018 , 22, 760-772	3.9	69
695	Natural Snake Venom Inhibitors and their Pharmaceutical Uses: Challenges and Possibilities. <i>Current Pharmaceutical Design</i> , 2018 , 24, 1737-1747	3.3	3
694	Diethylphosphoryl-OxymaB (DEPO-B) as a Solid Coupling Reagent for Amide Bond Formation. <i>Letters in Organic Chemistry</i> , 2018 , 16, 30-33	0.6	1
693	Improved pharmacokinetic profile of lipophilic anti-cancer drugs using B-targeted polyurethane-polyurea nanoparticles. <i>Nanomedicine: Nanotechnology, Biology, and Medicine</i> , 2018 , 14, 257-267	6	6
692	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	0.8	3
691	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	3.9	26

690	Amphibian Skin Secretions as a Novel Source for the Isolation of Antibacterial Peptides. <i>Molecules</i> , 2018 , 23,	4.8	4
689	Perfluorophenyl Derivatives as Unsymmetrical Linkers for Solid Phase Conjugation. <i>Frontiers in Chemistry</i> , 2018 , 6, 589	5	2
688	Hydroxylamine Derivatives as a New Paradigm in the Search of Antibacterial Agents. <i>ACS Omega</i> , 2018 , 3, 17057-17069	3.9	7
687	A Lasso-Inspired Bicyclic Peptide: Synthesis, Structure and Properties. <i>Chemistry - A European Journal</i> , 2018 , 24, 19250-19257	4.8	4
686	Synthesis, carbonic anhydrase inhibitory activity and antioxidant activity of some 1,3-oxazine derivatives. <i>Drug Development Research</i> , 2018 , 79, 352-361	5.1	7
685	Peptide Ligations by Using Aryloxycarbonyl-o-methylaminoanilides: Chemical Synthesis of Palmitoylated Sonic Hedgehog. <i>Angewandte Chemie - International Edition</i> , 2018 , 57, 16120-16125	16.4	14
684	Peptide Ligations by Using Aryloxycarbonyl-o-methylaminoanilides: Chemical Synthesis of Palmitoylated Sonic Hedgehog. <i>Angewandte Chemie</i> , 2018 , 130, 16352-16357	3.6	5
683	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	5	19
682	2017 FDA Peptide Harvest. <i>Pharmaceuticals</i> , 2018 , 11,	5.2	31
681	Investigation of the Biosynthesis of the Lasso Peptide Chaxapeptin Using an E. coli-Based Production System. <i>Journal of Natural Products</i> , 2018 , 81, 2050-2056	4.9	20
680	One-Pot Peptide Ligation-Oxidative Cyclization Protocol for the Preparation of Short-/Medium-Size Disulfide Cyclopeptides. <i>Organic Letters</i> , 2018 , 20, 4306-4309	6.2	4
679	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	7.9	17
678	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	3.9	38
677	The synthesis of an EDTA-like chelating peptidomimetic building block suitable for solid-phase peptide synthesis. <i>Chemical Communications</i> , 2017 , 53, 2634-2636	5.8	1
676	Microwave-Assisted Synthesis of Antimicrobial Peptides. <i>Methods in Molecular Biology</i> , 2017 , 1548, 51-59.4	5	5
675	Novel Globular Polymeric Supports for Membrane-Enhanced Peptide Synthesis. <i>Macromolecules</i> , 2017 , 50, 1626-1634	5.5	13
674	Tetrahydropyranyl: A Non-aromatic, Mild-Acid-Labile Group for Hydroxyl Protection in Solid-Phase Peptide Synthesis. <i>ChemistryOpen</i> , 2017 , 6, 206-210	2.3	2
673	Understanding Tetrahydropyranyl as a Protecting Group in Peptide Chemistry. <i>ChemistryOpen</i> , 2017 , 6, 168-177	2.3	4

672	Oxidative couplings on tryptophan-based diketopiperazines leading to fused and bridged chemotypes. <i>Chemical Communications</i> , 2017 , 53, 2740-2743	5.8	4
671	Stapled Peptides by Late-Stage C(sp ³)-H Activation. <i>Angewandte Chemie - International Edition</i> , 2017 , 56, 314-318	16.4	113
670	Stapled Peptides by Late-Stage C(sp ³)-H Activation. <i>Angewandte Chemie</i> , 2017 , 129, 320-324	3.6	45
669	Facile solid-phase synthesis of head-side chain cyclothiodipeptides through a cyclative cleavage from MeDbz-resin. <i>Tetrahedron Letters</i> , 2017 , 58, 2788-2791	2	13
668	Role of the Nozaki-Hiyama-Takai-Kishi Reaction in the Synthesis of Natural Products. <i>Chemical Reviews</i> , 2017 , 117, 8420-8446	68.1	82
667	Novel pyrazolyl-s-triazine derivatives, molecular structure and antimicrobial activity. <i>Journal of Molecular Structure</i> , 2017 , 1145, 244-253	3.4	37
666	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	6.6	11
665	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	3.4	25
664	Synthesis, in vitro evaluation, and Ga-radiolabeling of CDP1 toward PET/CT imaging of bacterial infection. <i>Chemical Biology and Drug Design</i> , 2017 , 90, 572-579	2.9	9
663	Intercalative DNA binding of the marine anticancer drug variolin B. <i>Scientific Reports</i> , 2017 , 7, 39680	4.9	13
662	A Trp-BODIPY cyclic peptide for fluorescence labelling of apoptotic bodies. <i>Chemical Communications</i> , 2017 , 53, 945-948	5.8	40
661	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	2.3	5
660	Synthesis, Crystal Structure, DFT Study of m-Methoxy-N ² -(3-Methoxybenzoyl)-N-Phenylbenzohydrazide. <i>Crystals</i> , 2017 , 7, 19	2.3	2
659	Improving gold nanorod delivery to the central nervous system by conjugation to the shuttle Angiopep-2. <i>Nanomedicine</i> , 2017 , 12, 2503-2517	5.6	30
658	Re-evaluating the stability of COMU in different solvents. <i>Journal of Peptide Science</i> , 2017 , 23, 763-768	2.1	12
657	Converting Teixobactin into a Cationic Antimicrobial Peptide (AMP). <i>Journal of Medicinal Chemistry</i> , 2017 , 60, 7476-7482	8.3	37
656	Fmoc-Amox, A Suitable Reagent for the Introduction of Fmoc. <i>Organic Process Research and Development</i> , 2017 , 21, 1533-1541	3.9	3
655	Green solid-phase peptide synthesis 4. γ -Valerolactone and N-formylmorpholine as green solvents for solid phase peptide synthesis. <i>Tetrahedron Letters</i> , 2017 , 58, 2986-2988	2	46

654	Gold nanoparticles as an efficient drug delivery system for GLP-1 peptides. <i>Colloids and Surfaces B: Biointerfaces</i> , 2017 , 158, 25-32	6	28
653	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	18.8	34
652	Sudemycin K: A Synthetic Antitumor Splicing Inhibitor Variant with Improved Activity and Versatile Chemistry. <i>ACS Chemical Biology</i> , 2017 , 12, 163-173	4.9	20
651	Functionalization of CoCr surfaces with cell adhesive peptides to promote HUVECs adhesion and proliferation. <i>Applied Surface Science</i> , 2017 , 393, 82-92	6.7	25
650	Investigation of the N-Terminus Amino Function of Arg-Teixobactin. <i>Molecules</i> , 2017 , 22,	4.8	19
649	Phakellistatins: An Underwater Unsolved Puzzle. <i>Marine Drugs</i> , 2017 , 15,	6	17
648	The Pharmaceutical Industry in 2016. An Analysis of FDA Drug Approvals from a Perspective of the Molecule Type. <i>Molecules</i> , 2017 , 22,	4.8	22
647	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,	4.8	29
646	Structure-Activity Relationship of Arg10-Teixobactin: A Recently Discovered Antimicrobial Peptide. <i>Proceedings (mdpi)</i> , 2017 , 1, 671	0.3	
645	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	2.3	6
644	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	2.3	13
643	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, 724-8	2.3	9
642	Enantioselective Synthesis of the Polyhydroxylated Chain of Oscillariolide and Phormidolides A-C. <i>Organic Letters</i> , 2016 , 18, 4485-7	6.2	7
641	Proximate biochemical composition and mineral content of edible species from the genus <i>Cystoseira</i> in Portugal. <i>Botanica Marina</i> , 2016 ,	1.8	2
640	Short AntiMicrobial Peptides (SAMPs) as a class of extraordinary promising therapeutic agents. <i>Journal of Peptide Science</i> , 2016 , 22, 438-51	2.1	46
639	Peptides conjugated to silver nanoparticles in biomedicine - a "value-added" phenomenon. <i>Biomaterials Science</i> , 2016 , 4, 1713-1725	7.4	25
638	Spacer-free BODIPY fluorogens in antimicrobial peptides for direct imaging of fungal infection in human tissue. <i>Nature Communications</i> , 2016 , 7, 10940	17.4	81
637	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. <i>Chemistry - A European Journal</i> , 2016 , 22, 7033-5	4.8	6

636	Oxyma-T, expanding the arsenal of coupling reagents. <i>Tetrahedron Letters</i> , 2016 , 57, 3523-3525	2	5
635	Synthesis of (E)-4-Bromo-3-methoxybut-3-en-2-one, the Key Fragment in the Polyhydroxylated Chain Common to Oscillariolide and Phormidolides A ₁ . <i>Chemistry - A European Journal</i> , 2016 , 22, 6993-6998	4.8	12
634	One pot synthesis, molecular structure and spectroscopic studies (X-ray, IR, NMR, UV-Vis) of novel 2-(4,6-dimethoxy-1,3,5-triazin-2-yl) amino acid ester derivatives. <i>Spectrochimica Acta - Part A: Molecular and Biomolecular Spectroscopy</i> , 2016 , 159, 184-98	4.4	12
633	Comparative proteomic analysis of growth hormone secretagogue A233 treatment of murine macrophage cells J774A.2 indicates it has a role in antiviral innate response. <i>Biochemistry and Biophysics Reports</i> , 2016 , 5, 379-387	2.2	5
632	CuAAC: An Efficient Click Chemistry Reaction on Solid Phase. <i>ACS Combinatorial Science</i> , 2016 , 18, 1-14	3.9	139
631	An improved and efficient strategy for the total synthesis of a colistin-like peptide. <i>Tetrahedron Letters</i> , 2016 , 57, 1885-1888	2	12
630	Inhibitory effect of short cationic homopeptides against Gram-negative bacteria. <i>Amino Acids</i> , 2016 , 48, 1445-56	3.5	5
629	BbrzSP-32, the first serine protease isolated from Bothrops brazili venom: Purification and characterization. <i>Comparative Biochemistry and Physiology Part A, Molecular & Integrative Physiology</i> , 2016 , 195, 15-25	2.6	17
628	Isololiolide, a carotenoid metabolite isolated from the brown alga <i>Cystoseira tamariscifolia</i> , is cytotoxic and able to induce apoptosis in hepatocarcinoma cells through caspase-3 activation, decreased Bcl-2 levels, increased p53 expression and PARP cleavage. <i>Phytomedicine</i> , 2016 , 23, 550-7	6.5	43
627	Highly chemoselective ligation of thiol- and amino-peptides on a bromomaleimide core. <i>Chemical Communications</i> , 2016 , 52, 2334-7	5.8	8
626	The road to the synthesis of "difficult peptides". <i>Chemical Society Reviews</i> , 2016 , 45, 631-54	58.5	122
625	2-Methyltetrahydrofuran and cyclopentyl methyl ether for green solid-phase peptide synthesis. <i>Amino Acids</i> , 2016 , 48, 419-26	3.5	52
624	Targeting prohibitins induces apoptosis in acute myeloid leukemia cells. <i>Oncotarget</i> , 2016 , 7, 64987-65000	9.0	14
623	Péptidos que atraviesan la membrana celular como potenciales transportadores de fármacos. <i>Revista Bionatura</i> , 2016 , 1,	0.3	2
622	Deprotection Reagents in Fmoc Solid Phase Peptide Synthesis: Moving Away from Piperidine?. <i>Molecules</i> , 2016 , 21,	4.8	48
621	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	3.1	23
620	Galactosidase-A Loaded-Nanoliposomes with Enhanced Enzymatic Activity and Intracellular Penetration. <i>Advanced Healthcare Materials</i> , 2016 , 5, 829-40	10.1	31
619	Enhanced antimicrobial activity of a peptide derived from human lysozyme by arylation of its tryptophan residues. <i>Journal of Peptide Science</i> , 2016 , 22, 123-8	2.1	12

618	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	3.4	28
617	Lysine Scanning of Arg-Teixobactin: Deciphering the Role of Hydrophobic and Hydrophilic Residues. <i>ACS Omega</i> , 2016 , 1, 1262-1265	3.9	46
616	A Facile Synthesis of NODASA-Functionalized Peptide. <i>Synlett</i> , 2016 , 27, 1685-1688	2.2	6
615	Combinatorial Library Screening Coupled to Mass Spectrometry to Identify Valuable Cyclic Peptides. <i>Current Protocols in Chemical Biology</i> , 2016 , 8, 109-130	1.8	5
614	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	8.3	55
613	Synthesis of complex head-to-side-chain cyclodepsipeptides. <i>Nature Protocols</i> , 2016 , 11, 1924-1947	18.8	13
612	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, 13114-9	4.8	51
611	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, 785-93	6.5	6
610	Re-evaluation of the N-terminal substitution and the D-residues of teixobactin. <i>RSC Advances</i> , 2016 , 6, 73827-73829	3.7	33
609	Semipermanent C-terminal carboxylic acid protecting group: application to solubilizing peptides and fragment condensation. <i>Organic Letters</i> , 2015 , 17, 294-7	6.2	15
608	Optimized Microwave Assisted Synthesis of LL37, a Cathelicidin Human Antimicrobial Peptide. <i>International Journal of Peptide Research and Therapeutics</i> , 2015 , 21, 13-20	2.1	7
607	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	5.7	22
606	A solid-phase combinatorial approach for indoloquinolizidine-peptides with high affinity at D(1) and D(2) dopamine receptors. <i>European Journal of Medicinal Chemistry</i> , 2015 , 97, 173-80	6.8	10
605	A new quinoxaline-containing peptide induces apoptosis in cancer cells by autophagy modulation. <i>Chemical Science</i> , 2015 , 6, 4537-4549	9.4	12
604	Methods, setup and safe handling for anhydrous hydrogen fluoride cleavage in Boc solid-phase peptide synthesis. <i>Nature Protocols</i> , 2015 , 10, 1067-83	18.8	36
603	Tetrahydropyranyl, a nonaromatic acid-labile Cys protecting group for Fmoc peptide chemistry. <i>Organic Letters</i> , 2015 , 17, 1680-3	6.2	15
602	Chemical Platforms for Peptide Vaccine Constructs. <i>Advances in Protein Chemistry and Structural Biology</i> , 2015 , 99, 99-130	5.3	4
601	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	3.2	3

600	Hantzsch dihydropyridines: Privileged structures for the formation of well-defined gold nanostars. <i>Journal of Colloid and Interface Science</i> , 2015 , 453, 260-269	9.3	15
599	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	3.9	8
598	EDCI·HCl and Potassium Salts of Oxyma and Oxyma-B as Superior Coupling Cocktails for Peptide Synthesis. <i>European Journal of Organic Chemistry</i> , 2015 , 2015, 3116-3120	3.2	16
597	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, 239-44	4.3	9
596	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	3.9	44
595	Multifunctionalized polyurethane-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	7.3	10
594	BbMP-1, a new metalloproteinase isolated from Bothrops brazili snake venom with in vitro antiplasmodial properties. <i>Toxicon</i> , 2015 , 106, 30-41	2.8	13
593	Single-molecule kinetics and footprinting of DNA bis-intercalation: the paradigmatic case of Thiocoraline. <i>Nucleic Acids Research</i> , 2015 , 43, 2767-79	20.1	26
592	An immunochemical strategy based on peptidoglycan synthetic peptide epitopes to diagnose Staphylococcus aureus infections. <i>Analytica Chimica Acta</i> , 2015 , 889, 203-11	6.6	5
591	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	15.6	47
590	Peptide synthesis beyond DMF: THF and ACN as excellent and friendlier alternatives. <i>Organic and Biomolecular Chemistry</i> , 2015 , 13, 2393-8	3.9	49
589	Addition of Vinylmetallic Reagents to Chiral 2-Formyltetrahydrofuran. <i>European Journal of Organic Chemistry</i> , 2015 , 2015, 235-241	3.2	6
588	Phenolic composition, antioxidant potential and in vitro inhibitory activity of leaves and acorns of Quercus suber on key enzymes relevant for hyperglycemia and Alzheimer's disease. <i>Industrial Crops and Products</i> , 2015 , 64, 45-51	5.9	69
587	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, 40-5	2.1	9
586	Phormidolides B and C, cytotoxic agents from the sea: enantioselective synthesis of the macrocyclic core. <i>Chemistry - A European Journal</i> , 2015 , 21, 150-6	4.8	20
585	β-Ketoamino acid ester derivatives as promising MAO inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015 , 25, 70-4	2.9	13
584	Botryococcus braunii and Nannochloropsis oculata extracts inhibit cholinesterases and protect human dopaminergic SH-SY5Y cells from H2O2-induced cytotoxicity. <i>Journal of Applied Phycology</i> , 2015 , 27, 839-848	3.2	25
583	Installing multifunctionality on titanium with RGD-decorated polyurethane-polyurea roxithromycin loaded nanoparticles: toward new osseointegrative therapies. <i>Advanced Healthcare Materials</i> , 2015 , 4, 1956-60	10.1	20

582	Injectable Hyaluronan Hydrogels with Peptide-Binding Dendrimers Modulate the Controlled Release of BMP-2 and TGF- β . <i>Macromolecular Bioscience</i> , 2015 , 15, 1035-44	5.5	21
581	Review backbone N-modified peptides: How to meet the challenge of secondary amine acylation. <i>Biopolymers</i> , 2015 , 104, 435-52	2.2	17
580	Formylation of electron-rich aromatic rings mediated by dichloromethyl methyl ether and TiCl ₄ : scope and limitations. <i>Molecules</i> , 2015 , 20, 5409-22	4.8	12
579	Synthesis and Preliminary Biological Evaluation of 1,3,5-Triazine Amino Acid Derivatives to Study Their MAO Inhibitors. <i>Molecules</i> , 2015 , 20, 15976-88	4.8	17
578	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	3.5	2
577	New peptide architectures through C-H activation stapling between tryptophan-phenylalanine/tyrosine residues. <i>Nature Communications</i> , 2015 , 6, 7160	17.4	184
576	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, 7197-209	16.4	137
575	Methods for the Peptide Synthesis and Analysis 2015 , 11-73		3
574	The Larock Reaction in the Synthesis of Heterocyclic Compounds. <i>Advances in Heterocyclic Chemistry</i> , 2015 , 116, 1-35	2.4	18
573	Synthesis and Biological Evaluation of a Teixobactin Analogue. <i>Organic Letters</i> , 2015 , 17, 6182-5	6.2	66
572	Stereoselective Allylstannane Addition for a Convergent Synthesis of a Complex Molecule. <i>Organic Letters</i> , 2015 , 17, 6246-9	6.2	5
571	Fatty acid profile of different species of algae of the <i>Cystoseira</i> genus: a nutraceutical perspective. <i>Natural Product Research</i> , 2015 , 29, 1264-70	2.3	25
570	In vitro antioxidant and inhibitory activity of water decoctions of carob tree (<i>Ceratonia siliqua</i> L.) on cholinesterases, α -amylase and α -glucosidase. <i>Natural Product Research</i> , 2015 , 29, 2155-9	2.3	21
569	6-(Bromomaleimido)hexanoic acid as a connector for the construction of multiple branched peptide platforms. <i>Organic Letters</i> , 2015 , 17, 464-7	6.2	5
568	A novel prohibitin-binding compound induces the mitochondrial apoptotic pathway through NOXA and BIM upregulation. <i>Oncotarget</i> , 2015 , 6, 41750-65	3.3	20
567	Immobilized N-chlorosuccinimide as a friendly peptide disulfide-forming reagent. <i>ACS Combinatorial Science</i> , 2014 , 16, 160-3	3.9	10
566	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, 6754-62	16.4	28
565	Thiopeptide engineering: a multidisciplinary effort towards future drugs. <i>Angewandte Chemie - International Edition</i> , 2014 , 53, 6602-16	16.4	70

564	BOP-Oxy, BOP-OBt, and BOP-OAt: novel organophosphinic coupling reagents useful for solution and solid-phase peptide synthesis. <i>Journal of Peptide Science</i> , 2014 , 20, 1-6	2.1	8
563	Solid-phase peptide synthesis (SPPS), C-terminal vs. side-chain anchoring: a reality or a myth. <i>Amino Acids</i> , 2014 , 46, 1827-38	3.5	10
562	Disulfide Formation Strategies in Peptide Synthesis. <i>European Journal of Organic Chemistry</i> , 2014 , 2014, 3519-3530	3.2	64
561	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	3.2	4
560	The potential of N-alkoxymethyl groups as peptide backbone protectants. <i>Tetrahedron Letters</i> , 2014 , 55, 184-188	2	5
559	Multifaceted roles of disulfide bonds. Peptides as therapeutics. <i>Chemical Reviews</i> , 2014 , 114, 901-26	68.1	365
558	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, 15031-9	4.8	14
557	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, 11409-15	4.2	11
556	Immobilized coupling reagents: synthesis of amides/peptides. <i>ACS Combinatorial Science</i> , 2014 , 16, 579-601	9.1	20
555	Proline N-oxides: modulators of the 3D conformation of linear peptides through "NO-turns". <i>Organic and Biomolecular Chemistry</i> , 2014 , 12, 4479-90	3.9	11
554	Synthesis of cyclohexapeptides as antimalarial and anti-trypanosomal agents. <i>MedChemComm</i> , 2014 , 5, 1309-1316	5	12
553	Tackling lipophilicity of peptide drugs: replacement of the backbone N-methyl group of cilengitide by N-oligoethylene glycol (N-OEG) chains. <i>Bioconjugate Chemistry</i> , 2014 , 25, 11-7	6.3	14
552	Polythiazole linkers as functional rigid connectors: a new RGD cyclopeptide with enhanced integrin selectivity. <i>Chemical Science</i> , 2014 , 5, 3929	9.4	8
551	Multivalent dendrimers presenting spatially controlled clusters of binding epitopes in thermoresponsive hyaluronan hydrogels. <i>Acta Biomaterialia</i> , 2014 , 10, 4340-50	10.8	21
550	A Trifluorinated Thiazoline Scaffold Leading to Pro-apoptotic Agents Targeting Prohibitins. <i>Angewandte Chemie</i> , 2014 , 126, 10314-10318	3.6	1
549	Dissecting the structure of thiopeptides: assessment of thiazoline and tail moieties of baringolin and antibacterial activity optimization. <i>Journal of Medicinal Chemistry</i> , 2014 , 57, 4185-95	8.3	20
548	Novel peptide-based platform for the dual presentation of biologically active peptide motifs on biomaterials. <i>ACS Applied Materials & Interfaces</i> , 2014 , 6, 6525-36	9.5	56
547	A trifluorinated thiazoline scaffold leading to pro-apoptotic agents targeting prohibitins. <i>Angewandte Chemie - International Edition</i> , 2014 , 53, 10150-4	16.4	25

546	Facile and mild synthesis of linear and cyclic peptides via thioesters. <i>Organic Letters</i> , 2014 , 16, 3922-5	6.2	12
545	High Control, Fast Growth OEG-Based Dendron Synthesis via a Sequential Two-Step Process of Copper-Free Diazo Transfer and Click Chemistry. <i>Macromolecules</i> , 2014 , 47, 2585-2591	5.5	3
544	Thioester Bonds of Thiocoraline Can Be Replaced with NMe-Amide Bridges without Affecting Its DNA-Binding Properties. <i>ACS Medicinal Chemistry Letters</i> , 2014 , 5, 45-50	4.3	5
543	Selective Formation of a Z-trisubstituted double bond using a 1-(tert-butyl)tetrazolyl sulfone. <i>Journal of Organic Chemistry</i> , 2014 , 79, 10648-54	4.2	8
542	Oxyma-B, an excellent racemization suppressor for peptide synthesis. <i>Organic and Biomolecular Chemistry</i> , 2014 , 12, 8379-85	3.9	22
541	Microreactors for peptide synthesis: looking through the eyes of twenty first century !!! <i>Amino Acids</i> , 2014 , 46, 2091-104	3.5	12
540	Linear versus branched poly-lysine/arginine as polarity enhancer tags. <i>Organic and Biomolecular Chemistry</i> , 2014 , 12, 7194-6	3.9	6
539	Antimicrobial peptides from skin secretions of <i>Hypsiboas pulchellus</i> (Anura: Hylidae). <i>Journal of Natural Products</i> , 2014 , 77, 831-41	4.9	21
538	Controlling multivalency and multimodality: up to pentamodal dendritic platforms based on diethylenetriaminepentaacetic acid cores. <i>Organic Letters</i> , 2014 , 16, 1318-21	6.2	13
537	Cysteine pseudoprolines for thiol protection and peptide macrocyclization enhancement in Fmoc-based solid-phase peptide synthesis. <i>Organic Letters</i> , 2014 , 16, 1772-5	6.2	19
536	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, 428-47	3.9	23
535	Amphiphilic cationic carbosilane-PEG dendrimers: synthesis and applications in gene therapy. <i>European Journal of Medicinal Chemistry</i> , 2014 , 76, 43-52	6.8	33
534	Morphological characterization of fullerene-androsterone conjugates. <i>Beilstein Journal of Nanotechnology</i> , 2014 , 5, 374-9	3	6
533	TOMBU and COMBU as Novel Uronium-type peptide coupling reagents derived from Oxyma-B. <i>Molecules</i> , 2014 , 19, 18953-65	4.8	9
532	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, 2568-72 ^{2.9}		10
531	Alkylation of histidine residues of <i>Bothrops jararacussu</i> venom proteins and isolated phospholipases A2: a biotechnological tool to improve the production of antibodies. <i>BioMed Research International</i> , 2014 , 2014, 981923	3	2
530	Thiopeptide antibiotics: retrospective and recent advances. <i>Marine Drugs</i> , 2014 , 12, 317-51	6	122
529	A novel phospholipase A2 (D49) from the venom of the <i>Crotalus oreganus abyssus</i> (North American Grand canyon rattlesnake). <i>BioMed Research International</i> , 2014 , 2014, 654170	3	8

528	Engineering von Thiopeptiden: ein multidisziplinärer Weg zu neuen Wirkstoffen. <i>Angewandte Chemie</i> , 2014 , 126, 6720-6735	3.6	10
527	"Clicking" Porphyrins to Magnetic Nanoparticles for Photodynamic Therapy. <i>ChemPlusChem</i> , 2014 , 79, 90-98	2.8	22
526	Gold nanoparticles for photothermally controlled drug release. <i>Nanomedicine</i> , 2014 , 9, 2023-39	5.6	32
525	Peptide affinity chromatography based on combinatorial strategies for protein purification. <i>Methods in Molecular Biology</i> , 2014 , 1129, 277-302	1.4	13
524	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	3.7	10
523	Isolation and biochemical characterization of a new thrombin-like serine protease from Bothrops pirajai snake venom. <i>BioMed Research International</i> , 2014 , 2014, 595186	3	11
522	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, 435-49	5.6	24
521	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	6.8	30
520	Covalent immobilization of hLf1-11 peptide on a titanium surface reduces bacterial adhesion and biofilm formation. <i>Acta Biomaterialia</i> , 2014 , 10, 3522-34	10.8	104
519	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	3.2	49
518	Structural glance into a novel anti-staphylococcal peptide. <i>Biopolymers</i> , 2014 , 102, 49-57	2.2	5
517	Chiral thiazoline and thiazole building blocks for the synthesis of peptide-derived natural products. <i>Current Topics in Medicinal Chemistry</i> , 2014 , 14, 1244-56	3	12
516	Low-epimerization Peptide Bond Formation with Oxyma Pure: Preparation of Z-L-Phg-Val-OMe 2014 , 306-315		1
515	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, 8129-35	4.2	91
514	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	4.1	10
513	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	5.9	25
512	N-chlorosuccinimide, an efficient peptide disulfide bond-forming reagent in aqueous solution. <i>RSC Advances</i> , 2013 , 3, 14277	3.7	15
511	Synthesis and NMR elucidation of pentacycloundecane-derived hydroxy acid peptides as potential anti-HIV-1 agents. <i>Structural Chemistry</i> , 2013 , 24, 1461-1471	1.8	3

510	Friendly strategy to prepare encoded one bead-one compound cyclic peptide library. <i>ACS Combinatorial Science</i> , 2013 , 15, 525-9	3.9	11
509	The first total synthesis of the cyclodepsipeptide pipercolidepsin A. <i>Nature Communications</i> , 2013 , 4, 2352-7.4	7.4	42
508	RADA-16: A Tough Peptide [Strategies for Synthesis and Purification. <i>European Journal of Organic Chemistry</i> , 2013 , 2013, 5871-5878	3.2	8
507	The backbone N-(4-azidobutyl) linker for the preparation of peptide chimera. <i>Organic Letters</i> , 2013 , 15, 4572-5	6.2	11
506	Inhibitory effect of short cationic homopeptides against gram-positive bacteria. <i>Journal of Peptide Science</i> , 2013 , 19, 792-800	2.1	22
505	"Head-to-side-chain" cyclodepsipeptides of marine origin. <i>Marine Drugs</i> , 2013 , 11, 1693-717	6	38
504	Rescuing biological activity from synthetic phakellistatin 19. <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 9780-8	8.3	24
503	Multifunctional nanovesicle-bioactive conjugates prepared by a one-step scalable method using CO ₂ -expanded solvents. <i>Nano Letters</i> , 2013 , 13, 3766-74	11.5	31
502	K-Oxyma: a Strong Acylation-Promoting, 2-CTC Resin-Friendly Coupling Additive. <i>European Journal of Organic Chemistry</i> , 2013 , 2013, 6372-6378	3.2	23
501	From 2,6-Dichloronicotinic Acid to Thiopeptide Cores. <i>European Journal of Organic Chemistry</i> , 2013 , 2013, 6404-6419	3.2	5
500	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	3.7	3
499	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	5.8	15
498	Synthesis and biological evaluation of a post-synthetically modified Trp-based diketopiperazine. <i>MedChemComm</i> , 2013 , 4, 1171	5	14
497	N-Chlorosuccinimide, an efficient reagent for on-resin disulfide formation in solid-phase peptide synthesis. <i>Organic Letters</i> , 2013 , 15, 616-9	6.2	35
496	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	3.2	2
495	Use of an internal reference for the quantitative HPLC-UV analysis of solid-phase reactions: a case study of 2-chlorotrityl chloride resin. <i>ACS Combinatorial Science</i> , 2013 , 15, 229-34	3.9	6
494	Tetrahydrofuran-containing macrolides: a fascinating gift from the deep sea. <i>Chemical Reviews</i> , 2013 , 113, 4567-610	68.1	226
493	Stable conjugates of peptides with gold nanorods for biomedical applications with reduced effects on cell viability. <i>ACS Applied Materials & Interfaces</i> , 2013 , 5, 4076-85	9.5	59

- 492 Handles for Fmoc solid-phase synthesis of protected peptides. *ACS Combinatorial Science*, **2013**, 15, 217-38 53
- 491 Marine Peptides and Proteins with Cytotoxic and Antitumoral Properties **2013**, 407-430
- 490 Enzyme-labile protecting groups for the synthesis of natural products: solid-phase synthesis of thiocoraline. *Angewandte Chemie - International Edition*, **2013**, 52, 5726-30 16.4 17
- 489 Biocompatible, multifunctional, and well-defined OEG-based dendritic platforms for biomedical applications. *Organic and Biomolecular Chemistry*, **2013**, 11, 4109-21 3.9 13
- 488 COMU: scope and limitations of the latest innovation in peptide acyl transfer reagents. *Journal of Peptide Science*, **2013**, 19, 408-14 2.1 30
- 487 Electrostatic binding and hydrophobic collapse of peptide-nucleic acid aggregates quantified using force spectroscopy. *ACS Nano*, **2013**, 7, 5102-13 16.7 23
- 486 Orthogonal chemistry for the synthesis of thiocoraline-triostin hybrids. Exploring their structure-activity relationship. *Journal of Medicinal Chemistry*, **2013**, 56, 5587-600 8.3 21
- 485 Wang linker free of side reactions. *Organic Letters*, **2013**, 15, 246-9 6.2 14
- 484 Imidazole-1-sulfonyl azide-based diazo-transfer reaction for the preparation of azido solid supports for solid-phase synthesis. *ACS Combinatorial Science*, **2013**, 15, 331-4 3.9 18
- 483 Polyproline α -DEG Co-Oligomeric Dendrimers: A Family of Highly Branched Polyproline Macromolecules. *European Journal of Organic Chemistry*, **2013**, 2013, 8279-8287 3.2
- 482 Total Synthesis and Stereochemical Assignment of Baringolin. *Angewandte Chemie*, **2013**, 125, 7972-7975 6 9
- 481 Enzyme-Labile Protecting Groups for the Synthesis of Natural Products: Solid-Phase Synthesis of Thiocoraline. *Angewandte Chemie*, **2013**, 125, 5838-5842 3.6 5
- 480 Total synthesis and stereochemical assignment of baringolin. *Angewandte Chemie - International Edition*, **2013**, 52, 7818-21 16.4 33
- 479 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. *Molecules*, **2013**, 18, 14747-59 4.8 16
- 478 Understanding acid lability of cysteine protecting groups. *Molecules*, **2013**, 18, 5155-62 4.8 11
- 477 Neurotoxicity of prion peptides mimicking the central domain of the cellular prion protein. *PLoS ONE*, **2013**, 8, e70881 3.7 17
- 476 Liquid phase organic synthesis of 3,5-disubstituted 1,3,5-thiadiazinane-2-thione derivatives on polyethylene glycol (PEG) support. *Arkivoc*, **2013**, 2012, 326-338 0.9 3
- 475 Cell adhesion and focal contact formation on linear RGD molecular gradients: study of non-linear concentration dependence effects. *Nanomedicine: Nanotechnology, Biology, and Medicine*, **2012**, 8, 432-9⁶ 34

474	Cell-penetrating peptide/antimicrobial undecapeptide conjugates with anticancer activity. <i>Tetrahedron</i> , 2012 , 68, 4406-4412	2.4	11
473	Cyanoacetamide-based oxime carbonates: an efficient, simple alternative for the introduction of Fmoc with minimal dipeptide formation. <i>Tetrahedron</i> , 2012 , 68, 3056-3062	2.4	10
472	Orthogonal Protecting Groups in the Synthesis of Tryptophanyl-Hexahydropyrroloindoles. <i>European Journal of Organic Chemistry</i> , 2012 , 2012, 67-73	3.2	10
471	Myoblast cell interaction with polydopamine coated liposomes. <i>Biointerphases</i> , 2012 , 7, 8	1.8	38
470	Trimethoxyphenylthio as a highly labile replacement for tert-butylthio cysteine protection in Fmoc solid phase synthesis. <i>Organic Letters</i> , 2012 , 14, 5468-71	6.2	43
469	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, 2313-5	5.8	9
468	Eco-friendly combination of the immobilized PGA enzyme and the S-Phacm protecting group for the synthesis of Cys-containing peptides. <i>Chemistry - A European Journal</i> , 2012 , 18, 16166-76	4.8	22
467	H-bonding promotion of peptide solubility and cyclization by fluorinated alcohols. <i>RSC Advances</i> , 2012 , 2, 2729	3.7	12
466	Introducing an Asp-Pro linker in the synthesis of random one-bead-one-compound hexapeptide libraries compatible with ESI-MS analysis. <i>ACS Combinatorial Science</i> , 2012 , 14, 145-9	3.9	5
465	Synthesis and in vivo evaluation of the biodistribution of a 18F-labeled conjugate gold-nanoparticle-peptide with potential biomedical application. <i>Bioconjugate Chemistry</i> , 2012 , 23, 399-408	6.3	82
464	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. <i>Botanica Marina</i> , 2012 , 55,	1.8	25
463	Efficient amino-proline-derived cell penetrating peptide-superparamagnetic iron oxide nanoparticle conjugates via aniline-catalyzed oxime chemistry as bimodal imaging nanoagents. <i>Chemical Communications</i> , 2012 , 48, 5322-4	5.8	20
462	Surface-adhered composite poly(vinyl alcohol) physical hydrogels: polymersome-aided delivery of therapeutic small molecules. <i>Advanced Healthcare Materials</i> , 2012 , 1, 791-5	10.1	34
461	Resin-to-Resin Transfer Reactions (RRTR) via Sonogashira Coupling 2012 , 59-66		
460	Rapid and high-yielding cysteine labelling of peptides with N-succinimidyl 4-[18F]fluorobenzoate. <i>Chemical Communications</i> , 2012 , 48, 6118-20	5.8	11
459	Acid-labile Cys-protecting groups for the Fmoc/tBu strategy: filling the gap. <i>Organic Letters</i> , 2012 , 14, 5472-5	6.2	29
458	Vascular effects and electrolyte homeostasis of the natriuretic peptide isolated from <i>Crotalus oreganus abyssus</i> (North American Grand Canyon rattlesnake) venom. <i>Peptides</i> , 2012 , 36, 206-12	3.8	14
457	Targeting nanoparticles to dendritic cells for immunotherapy. <i>Methods in Enzymology</i> , 2012 , 509, 143-63	1.7	90

456	Thiadiazines, N,N-heterocycles of biological relevance. <i>Molecules</i> , 2012 , 17, 7612-28	4.8	16
455	Cancer prognostics by direct detection of p53-antibodies on gold surfaces by impedance measurements. <i>Small</i> , 2012 , 8, 2106-15	11	19
454	Biosensors: Cancer Prognostics by Direct Detection of p53-Antibodies on Gold Surfaces by Impedance Measurements (Small 13/2012). <i>Small</i> , 2012 , 8, 1962-1962	11	
453	Solid-phase synthesis of NMe-IB-01212, a highly N-methylated cyclic peptide. <i>Organic Letters</i> , 2012 , 14, 612-5	6.2	29
452	Screening of N-alkyl-cyanoacetamido oximes as substitutes for N-hydroxysuccinimide. <i>ChemistryOpen</i> , 2012 , 1, 147-52	2.3	13
451	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. <i>Biopolymers</i> , 2012 , 98, 89-97	2.2	37
450	Microalgae of different phyla display antioxidant, metal chelating and acetylcholinesterase inhibitory activities. <i>Food Chemistry</i> , 2012 , 131, 134-140	8.5	68
449	Synthesis and Thermal Properties of Novel Polyamides Containing α -Amino Acid Moieties: Structure-Property Relationship. <i>Journal of Macromolecular Science - Pure and Applied Chemistry</i> , 2012 , 49, 41-54	2.2	5
448	Enhancing immunogenicity and cross-reactivity of HIV-1 antigens by in vivo targeting to dendritic cells. <i>Nanomedicine</i> , 2012 , 7, 1591-610	5.6	5
447	Drug Delivery: Surface-Adhered Composite Poly(Vinyl Alcohol) Physical Hydrogels: Polymersome-Aided Delivery of Therapeutic Small Molecules (Adv. Healthcare Mater. 6/2012). <i>Advanced Healthcare Materials</i> , 2012 , 1, 790-790	10.1	2
446	Gene Promoters and Transcription Control Regions as Therapeutic Targets 2012 , 327-350		
445	Oxime-based carbonates as useful reagents for both N-protection and peptide coupling. <i>Molecules</i> , 2012 , 17, 14361-76	4.8	1
444	Polymers and drug delivery systems. <i>Current Drug Delivery</i> , 2012 , 9, 367-94	3.2	174
443	Affinity chromatography based on a combinatorial strategy for erythropoietin purification. <i>ACS Combinatorial Science</i> , 2011 , 13, 251-8	3.9	25
442	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, 7005-12	5.7	43
441	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, 4506-13	4.2	19
440	Targeting nanosystems to human DCs via Fc receptor as an effective strategy to deliver antigen for immunotherapy. <i>Molecular Pharmaceutics</i> , 2011 , 8, 104-16	5.6	72
439	Total synthesis of aeruginazole A. <i>Organic Letters</i> , 2011 , 13, 4648-51	6.2	15

438	Acridine and quindoline oligomers linked through a 4-aminoproline backbone prefer G-quadruplex structures. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2011 , 1810, 769-76	4	11
437	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-6-chloro-5-formyl-2-methyl-1,4-dihydropyridine-3-carboxylate. <i>Journal of Heterocyclic Chemistry</i> , 2011 , 47, 2199-2207	4.8	7
436	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, 852-8	3.8	18
435	Solid-phase synthesis of a library of amphipatic hydantoins. Discovery of new hits for TRPV1 blockade. <i>ACS Combinatorial Science</i> , 2011 , 13, 458-65	3.9	9
434	Synthesis and Aminolysis of 2,4-Dinitrophenyl and 5-NitropyridineN-Hydroxy Oxime Derivatives. <i>Bulletin of the Chemical Society of Japan</i> , 2011 , 84, 633-639	5.1	6
433	Aspartimide formation in peptide chemistry: occurrence, prevention strategies and the role of N-hydroxylamines. <i>Tetrahedron</i> , 2011 , 67, 8595-8606	2.4	63
432	Highly efficient, multigram and enantiopure synthesis of (S)-2-(2,4?-bithiazol-2-yl)pyrrolidine. <i>Tetrahedron Letters</i> , 2011 , 52, 5435-5437	2	8
431	Leishmania mexicana: LACK (Leishmania homolog of receptors for activated C-kinase) is a plasminogen binding protein. <i>Experimental Parasitology</i> , 2011 , 127, 752-61	2.1	15
430	Peptide coupling reagents, more than a letter soup. <i>Chemical Reviews</i> , 2011 , 111, 6557-602	68.1	716
429	Functionalization of gold surfaces: recent developments and applications. <i>Journal of Materials Science</i> , 2011 , 46, 7643-7648	4.3	19
428	Phytochemical profile, antioxidant and cytotoxic activities of the carob tree (Ceratonia siliqua L.) germ flour extracts. <i>Plant Foods for Human Nutrition</i> , 2011 , 66, 78-84	3.9	47
427	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. <i>International Journal of Peptide Research and Therapeutics</i> , 2011 , 17, 225-230	2.1	2
426	Total synthesis of a depsidomycin analogue by convergent solid-phase peptide synthesis and macrolactonization strategy for antitubercular activity. <i>Journal of Peptide Science</i> , 2011 , 17, 683-9	2.1	22
425	Fmoc Methodology: Cleavage from the Resin and Final Deprotection 2011 , 349-369		3
424	Peptide-Coupling Reagents 2011 , 407-444		2
423	Optimized Fmoc solid-phase synthesis of the cysteine-rich peptide linaclootide. <i>Biopolymers</i> , 2011 , 96, 69-80	2.2	40
422	Structure, bioactivity and synthesis of natural products with hexahydropyrrolo[2,3-b]indole. <i>Chemistry - A European Journal</i> , 2011 , 17, 1388-408	4.8	367
421	Progress on lamellarins. <i>MedChemComm</i> , 2011 , 2, 689-697	5	69

4 ²⁰	Study of various presentation forms for a peptide mimetic of Neisseria meningitidis serogroup B capsular polysaccharide. <i>Bioconjugate Chemistry</i> , 2011 , 22, 33-41	6.3	3
4 ¹⁹	Trivalent PEGylated platform for the conjugation of bioactive compounds. <i>Bioconjugate Chemistry</i> , 2011 , 22, 2172-8	6.3	1
4 ¹⁸	Biotin ergopeptide probes for dopamine receptors. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 1080-90	8.3	13
4 ¹⁷	TRPV1 modulators: structure-activity relationships using a rational combinatorial approach. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011 , 21, 3541-5	2.9	6
4 ¹⁶	Solid-phase synthesis of new Trp(Nps)-containing dipeptide derivatives as TRPV1 channel blockers. <i>Molecules</i> , 2010 , 15, 4924-33	4.8	1
4 ¹⁵	Synthesis of 2-(4,6-dimethoxy-1,3,5-triazin-2-yloxyimino) derivatives: application in solution peptide synthesis. <i>Molecules</i> , 2010 , 15, 9403-17	4.8	6
4 ¹⁴	A hybrid indoloquinolizidine peptide as allosteric modulator of dopamine D1 receptors. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2010 , 332, 876-85	4.7	11
4 ¹³	The Sea as a Source of New Drugs 2010 , 237-249		2
4 ¹²	Isolation, structural assignment, and total synthesis of barmumycin. <i>Journal of Organic Chemistry</i> , 2010 , 75, 8508-15	4.2	28
4 ¹¹	Engineering advanced capsosomes: maximizing the number of subcompartments, cargo retention, and temperature-triggered reaction. <i>ACS Nano</i> , 2010 , 4, 1351-61	16.7	129
4 ¹⁰	N-Hydroxylamines for Peptide Synthesis 2010 ,		1
4 ⁰⁹	Total regioselective control of tartaric acid. <i>Journal of Organic Chemistry</i> , 2010 , 75, 5746-9	4.2	3
4 ⁰⁸	The antitumoral depsipeptide IB-01212 kills Leishmania through an apoptosis-like process involving intracellular targets. <i>Molecular Pharmaceutics</i> , 2010 , 7, 1608-17	5.6	20
4 ⁰⁷	Improving the brain delivery of gold nanoparticles by conjugation with an amphipathic peptide. <i>Nanomedicine</i> , 2010 , 5, 897-913	5.6	85
4 ⁰⁶	PyOxP and PyOxB: the Oxyma-based novel family of phosphonium salts. <i>Organic and Biomolecular Chemistry</i> , 2010 , 8, 3665-73	3.9	33
4 ⁰⁵	Amphiphilic peptides and their cross-disciplinary role as building blocks for nanoscience. <i>Chemical Society Reviews</i> , 2010 , 39, 241-63	58.5	222
4 ⁰⁴	Multifunctionalized gold nanoparticles with peptides targeted to gastrin-releasing peptide receptor of a tumor cell line. <i>Bioconjugate Chemistry</i> , 2010 , 21, 1070-8	6.3	60
4 ⁰³	A novel dipeptidomimetic containing a cyclic threonine. <i>Chemical Communications</i> , 2010 , 46, 1266-8	5.8	11

402	Synthesis of orthogonally protected L-threo-beta-ethoxyasparagine. <i>Amino Acids</i> , 2010 , 39, 161-5	3.5	5
401	Targeted PLGA nano- but not microparticles specifically deliver antigen to human dendritic cells via DC-SIGN in vitro. <i>Journal of Controlled Release</i> , 2010 , 144, 118-26	11.7	218
400	Solid-Phase Synthesis of Aza-Kahalalide F Analogues: (2R,3R)-2-Amino-3-azidobutanoic Acid as Precursor of the Aza-Threonine. <i>European Journal of Organic Chemistry</i> , 2010 , 2010, 2536-2543	3.2	12
399	Oxime Carbonates: Novel Reagents for the Introduction of Fmoc and Alloc Protecting Groups, Free of Side Reactions. <i>European Journal of Organic Chemistry</i> , 2010 , 2010, 3275-3280	3.2	14
398	2,2,4,6,7-Pentamethyl-2,3-dihydrobenzofuran-5-methyl (Pbfm) as an Alternative to the Trityl Group for the Side-Chain Protection of Cysteine and Asparagine/Glutamine. <i>European Journal of Organic Chemistry</i> , 2010 , 2010, 3631-3640	3.2	11
397	A Novel Family of Onium Salts Based Upon Isonitroso Meldrum's Acid Proves Useful as Peptide Coupling Reagents. <i>European Journal of Organic Chemistry</i> , 2010 , 2010, 3641-3649	3.2	27
396	Streamlined Access to Functionalized Chromenes and Quinolines using Domino Reactions of Salicylic Aldehydes and Methyl 4-Chloro-2-butynoate. <i>European Journal of Organic Chemistry</i> , 2010 , 2010, 5373-5379	3.2	16
395	Capsosomes with Multilayered Subcompartments: Assembly and Loading with Hydrophobic Cargo. <i>Advanced Functional Materials</i> , 2010 , 20, 59-66	15.6	106
394	Postsynthetic modification of peptides: chemoselective C-arylation of tryptophan residues. <i>Chemistry - A European Journal</i> , 2010 , 16, 1124-7	4.8	138
393	Amide-to-Ester Substitution Allows Fine-Tuning of the Cyclopeptide Conformational Ensemble. <i>Angewandte Chemie</i> , 2010 , 122, 2792-2797	3.6	7
392	Amide-to-ester substitution allows fine-tuning of the cyclopeptide conformational ensemble. <i>Angewandte Chemie - International Edition</i> , 2010 , 49, 2732-7	16.4	15
391	Microwave assisted SPPS of amylin and its toxicity of the pure product to RIN-5F cells. <i>Biopolymers</i> , 2010 , 94, 323-30	2.2	16
390	Sample preparation for sequencing hits from one-bead-one-peptide combinatorial libraries by matrix-assisted laser desorption/ionization time-of-flight mass spectrometry. <i>Analytical Biochemistry</i> , 2010 , 400, 295-7	3.1	20
389	COMU: a third generation of uronium-type coupling reagents. <i>Journal of Peptide Science</i> , 2010 , 16, 6-9	2.1	73
388	A convenient microwave-enhanced solid-phase synthesis of short chain N-methyl-rich peptides. <i>Journal of Peptide Science</i> , 2010 , 16, 136-40	2.1	27
387	Improved antimicrobial activity of h-lysozyme (107-115) by rational Ala substitution. <i>Journal of Peptide Science</i> , 2010 , 16, 424-9	2.1	7
386	ChemMatrix(®) for complex peptides and combinatorial chemistry. <i>Journal of Peptide Science</i> , 2010 , 16, 675-8	2.1	48
385	Synthesis and Aminolysis of N,N-Diethyl Carbamic Ester of HOBt Derivatives. <i>Bulletin of the Korean Chemical Society</i> , 2010 , 31, 75-81	1.2	7

384	Solid-phase synthesis of peptides with C-terminal asparagine or glutamine. <i>International Journal of Peptide and Protein Research</i> , 2009 , 35, 284-286		29
383	Convergent solid-phase peptide synthesis. <i>International Journal of Peptide and Protein Research</i> , 2009 , 37, 58-60		21
382	Oxyma: an efficient additive for peptide synthesis to replace the benzotriazole-based HOBT and HOAt with a lower risk of explosion. <i>Chemistry - A European Journal</i> , 2009 , 15, 9394-403	4.8	268
381	COMU: a safer and more effective replacement for benzotriazole-based uronium coupling reagents. <i>Chemistry - A European Journal</i> , 2009 , 15, 9404-16	4.8	219
380	Structure-activity relationships of SSAO/VAP-1 arylalkylamine-based substrates. <i>ChemMedChem</i> , 2009 , 4, 495-503	3.7	13
379	Indoloquinolizidine-peptide hybrids as multiple agonists for D1 and D2 dopamine receptors. <i>ChemMedChem</i> , 2009 , 4, 1514-22	3.7	14
378	Synthesis and Application of N-Hydroxylamine Derivatives as Potential Replacements for HOBT. <i>European Journal of Organic Chemistry</i> , 2009 , 2009, 1499-1501	3.2	22
377	Oxathiocoraline: Lessons to be Learned from the Synthesis of Complex N-Methylated Depsipeptides. <i>European Journal of Organic Chemistry</i> , 2009 , 2009, 2957-2974	3.2	13
376	Siamese Depsipeptides: Constrained Bicyclic Architectures. <i>Angewandte Chemie</i> , 2009 , 121, 8716-8719	3.6	2
375	Enhanced microwave-assisted method for on-bead disulfide bond formation: synthesis of alpha-conotoxin MII. <i>Biopolymers</i> , 2009 , 92, 23-34	2.2	27
374	Design and facile solid-phase synthesis of peptide-based LPS-inhibitors containing PEG-like functionalities. <i>Biopolymers</i> , 2009 , 92, 508-17	2.2	4
373	Optimized Fmoc solid-phase synthesis of Thymosin alpha1 by side-chain anchoring onto a PEG resin. <i>Biopolymers</i> , 2009 , 92, 565-72	2.2	9
372	Solid-phase peptide synthesis using acetonitrile as a solvent in combination with PEG-based resins. <i>Journal of Peptide Science</i> , 2009 , 15, 629-33	2.1	40
371	Optical tweezers study of topoisomerase inhibition. <i>Small</i> , 2009 , 5, 1269-72	11	5
370	Microwave irradiation and COMU: a potent combination for solid-phase peptide synthesis. <i>Tetrahedron Letters</i> , 2009 , 50, 6200-6202	2	44
369	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, 1139-45	3.4	54
368	Amino acid-protecting groups. <i>Chemical Reviews</i> , 2009 , 109, 2455-504	68.1	535
367	Siamese depsipeptides: constrained bicyclic architectures. <i>Angewandte Chemie - International Edition</i> , 2009 , 48, 8564-7	16.4	5

366	Total synthesis and antiproliferative activity screening of (+/-)-aplicyanins A, B and E and related analogues. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 6217-23	8.3	32
365	Lamellarin D bioconjugates II: Synthesis and cellular internalization of dendrimer and nuclear location signal derivatives. <i>Bioconjugate Chemistry</i> , 2009 , 20, 1112-21	6.3	23
364	Solid-phase synthesis of chiral bicyclic guanidinium oligomers. <i>ACS Combinatorial Science</i> , 2009 , 11, 410-21	6	
363	Lamellarin D bioconjugates I: Synthesis and cellular internalization of PEG-derivatives. <i>Bioconjugate Chemistry</i> , 2009 , 20, 1100-11	6.3	22
362	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, 146-50		40
361	Adenosine A2A receptor-antagonist/dopamine D2 receptor-agonist bivalent ligands as pharmacological tools to detect A2A-D2 receptor heteromers. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 5590-602	8.3	116
360	Conjugation of Kahalalide F with gold nanoparticles to enhance in vitro antitumoral activity. <i>Bioconjugate Chemistry</i> , 2009 , 20, 138-46	6.3	67
359	Antioxidant activity and in vitro inhibition of tumor cell growth by leaf extracts from the carob tree (<i>Ceratonia siliqua</i>). <i>Pharmaceutical Biology</i> , 2009 , 47, 721-728	3.8	18
358	NMe amide as a synthetic surrogate for the thioester moiety in thiocoraline. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 834-9	8.3	32
357	Manufacturing peptides as active pharmaceutical ingredients. <i>Future Medicinal Chemistry</i> , 2009 , 1, 361-77.1	7.1	116
356	Synthesis of the pyrrolo[2,3-c]carbazole core of the dictyodendrins. <i>Organic and Biomolecular Chemistry</i> , 2009 , 7, 860-2	3.9	33
355	Kahalalide F, an antitumor depsipeptide in clinical trials, and its analogues as effective antileishmanial agents. <i>Molecular Pharmaceutics</i> , 2009 , 6, 813-24	5.6	35
354	1,2-Dimethylindole-3-sulfonyl (MIS) as protecting group for the side chain of arginine. <i>Organic and Biomolecular Chemistry</i> , 2009 , 7, 2565-9	3.9	10
353	Solid-phase peptide synthesis in water using microwave-assisted heating. <i>Organic Letters</i> , 2009 , 11, 4488-91	9.1	61
352	N,N,N',N'-tetramethylchloroformamidinium hexafluorophosphate (TCFH), a powerful coupling reagent for bioconjugation. <i>Bioconjugate Chemistry</i> , 2008 , 19, 1968-71	6.3	14
351	THAL, a sterically unhindered linker for the solid-phase synthesis of acid-sensitive protected peptide acids. <i>Journal of Organic Chemistry</i> , 2008 , 73, 7342-4	4.2	4
350	Morpholine-based immonium and halogenoamidinium salts as coupling reagents in Peptide synthesis1. <i>Journal of Organic Chemistry</i> , 2008 , 73, 2731-7	4.2	55
349	Structure-activity relationship of kahalalide F synthetic analogues. <i>Journal of Medicinal Chemistry</i> , 2008 , 51, 4920-31	8.3	32

348	A novel protecting/activating strategy for beta-hydroxy acids and its use in convergent peptide synthesis. <i>Journal of Organic Chemistry</i> , 2008 , 73, 2311-4	4.2	10
347	Synthesis and antitumor activity of mechercharmycin A analogues. <i>Journal of Medicinal Chemistry</i> , 2008 , 51, 5722-30	8.3	21
346	Solid-phase combinatorial synthesis of a lysyl-tRNA synthetase (LysRS) inhibitory library. <i>ACS Combinatorial Science</i> , 2008 , 10, 391-400		10
345	Cysteine-S-trityl a key derivative to prepare N-methyl cysteines. <i>ACS Combinatorial Science</i> , 2008 , 10, 69-78		13
344	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	0.2	1
343	Asymmetric Synthesis of α -Unsubstituted β -Hydroxy Acids. <i>Current Organic Synthesis</i> , 2008 , 5, 151-161	1.9	8
342	2008 ,		27
341	Phenyl-EDOTn derivatives as super acid labile carboxylic acid protecting groups for peptide synthesis. <i>Tetrahedron Letters</i> , 2008 , 49, 3304-3307	2	9
340	Solid Phase Preparation of 1,3-Disubstituted Indazole derivatives. <i>QSAR and Combinatorial Science</i> , 2008 , 27, 1267-1273		2
339	Protection by conformationally restricted mobility: first solid-phase synthesis of triostin A. <i>Chemistry - A European Journal</i> , 2008 , 14, 4475-8	4.8	14
338	Conformationally restricted hydantoin-based peptidomimetics as inhibitors of caspase-3 with basic groups allowed at the S3 enzyme subsite. <i>ChemMedChem</i> , 2008 , 3, 979-85	3.7	10
337	Use of N-methylpiperazine for the preparation of piperazine-based unsymmetrical bis-ureas as anti-HIV agents. <i>ChemMedChem</i> , 2008 , 3, 1034-7	3.7	8
336	Nanostructure formation enhances the activity of LPS-neutralizing peptides. <i>ChemMedChem</i> , 2008 , 3, 1748-55	3.7	11
335	Synthesis of Natural Product Derivatives Containing 2,4-Concatenated Oxazoles. <i>European Journal of Organic Chemistry</i> , 2008 , 2008, 3389-3396	3.2	19
334	EDOTn and MIM, new peptide backbone protecting groups. <i>Biopolymers</i> , 2008 , 90, 444-9	2.2	21
333	Synthesis of oligonucleotide derivatives using ChemMatrix supports. <i>Chemistry and Biodiversity</i> , 2008 , 5, 209-18	2.5	8
332	Design, synthesis and antiproliferative properties of oligomers with chromophore units linked by amide backbones. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 2440-4	2.9	13
331	Solid-phase synthesis of oligomers carrying several chromophore units linked by phosphodiester backbones. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 2306-10	2.9	9

330	Stereomeric studies on the oxidation and alkylation of 4-thiazolidinones. <i>Tetrahedron Letters</i> , 2008 , 49, 1569-1572	2	12
329	[[Cu(pzPh)(Opo)] ₂ (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	2.7	28
328	Design and synthesis of FAJANU: a de novo C(2) symmetric cyclopeptide family. <i>Journal of Medicinal Chemistry</i> , 2008 , 51, 3194-202	8.3	9
327	Trishomocubane amino acid as a beta-turn scaffold. <i>Chemical Biology and Drug Design</i> , 2008 , 71, 125-30	2.9	20
326	Tiratricol neutralizes bacterial endotoxins and reduces lipopolysaccharide-induced TNF-alpha production in the cell. <i>Chemical Biology and Drug Design</i> , 2008 , 72, 320-8	2.9	3
325	A nonacid degradable linker for solid-phase synthesis. <i>Organic Letters</i> , 2007 , 9, 4319-22	6.2	13
324	Smallest peptoids with antiproliferative activity on human neoplastic cells. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 2443-9	8.3	16
323	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, 1429-32	6.2	12
322	Novel ergopeptides as dual ligands for adenosine and dopamine receptors. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 3062-9	8.3	32
321	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, 6133-43	8.3	17
320	From the one-bead-one-compound concept to one-bead-one-reactor. <i>ACS Combinatorial Science</i> , 2007 , 9, 395-8		9
319	Solid-phase synthesis of sulfamate peptidomimetics. <i>ACS Combinatorial Science</i> , 2007 , 9, 501-6		4
318	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, 1110-4	3.8	23
317	Fmoc-2-mercaptobenzothiazole, for the introduction of the Fmoc moiety free of side-reactions. <i>Biopolymers</i> , 2007 , 88, 733-7	2.2	30
316	Simple machine-assisted protocol for solid-phase synthesis of depsipeptides. <i>Biopolymers</i> , 2007 , 88, 823-82		6
315	Amide-to-ester substitution in coiled coils: the effect of removing hydrogen bonds on protein structure. <i>Angewandte Chemie - International Edition</i> , 2007 , 46, 7766-9	16.4	36
314	Estersubstitutionen in Helicalen Coiled-Coil-Peptiden: Effekt der Eliminierung von Wasserstoffbrücken auf die Struktur von Proteinen. <i>Angewandte Chemie</i> , 2007 , 119, 7912-7916	3.6	12
313	Synthesis of a 24-Membered Cyclic Peptide-Biphenyl Hybrid. <i>European Journal of Organic Chemistry</i> , 2007 , 2007, 1301-1308	3.2	17

312	Regioselective Monobromination of Free and Protected Phenols. <i>European Journal of Organic Chemistry</i> , 2007 , 2007, 1921-1924	3.2	16
311	Inhibition of VAP1: quickly gaining ground as an anti-inflammatory therapy. <i>ChemMedChem</i> , 2007 , 2, 173-4	3.7	12
310	Synthesis and structure-activity relationship of cytotoxic marine cyclodepsipeptide IB-01212 analogues. <i>ChemMedChem</i> , 2007 , 2, 1076-84	3.7	8
309	Formation of dihydrouracils via cyclization of N-substituted 3-thioureidopropanoic acids and facile desulfurization. <i>Tetrahedron</i> , 2007 , 63, 8949-8953	2.4	0
308	Preparation of penta-azole containing cyclopeptides: challenges in macrocyclization. <i>Tetrahedron</i> , 2007 , 63, 9862-9870	2.4	23
307	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-one peptide combinatorial libraries. <i>Analytical Biochemistry</i> , 2007 , 370, 215-22	3.1	24
306	Chlorotriyl Chloride (CTC) Resin as a Reusable Carboxyl Protecting Group. <i>QSAR and Combinatorial Science</i> , 2007 , 26, 1027-1035		25
305	Understanding the mechanism of action of the novel SSAO substrate (C ₇ NH ₁₀) ₆ (V ₁₀ O ₂₈).2H ₂ O, a prodrug of peroxovanadate insulin mimetics. <i>Chemical Biology and Drug Design</i> , 2007 , 69, 423-8	2.9	36
304	Chemical Synthesis of 19F-labeled HIV-1 Protease using Fmoc-Chemistry and ChemMatrix Resin. <i>International Journal of Peptide Research and Therapeutics</i> , 2007 , 13, 221-227	2.1	16
303	Beyond Azathiocoraline: Synthesis of Analogues. <i>International Journal of Peptide Research and Therapeutics</i> , 2007 , 13, 295-306	2.1	3
302	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	2.1	2
301	Partially Fluorinated Heterocycles from 4,4-Bis(trifluoromethyl)-hetero-1,3-dienes via C-B Bond Activation: Synthesis of 2-Fluoro-3-(trifluoromethyl)furans. <i>Monatshefte für Chemie</i> , 2007 , 138, 227-236	1.4	7
300	Enolase as a plasminogen binding protein in <i>Leishmania mexicana</i> . <i>Parasitology Research</i> , 2007 , 101, 1511-6	1.6	81
299	Oral insulin-mimetic compounds that act independently of insulin. <i>Diabetes</i> , 2007 , 56, 486-93	0.9	50
298	Novel Synthesis of Arylethynyl Heterocycles. <i>Synthesis</i> , 2007 , 2007, 1559-1565	2.9	4
297	New developments in the synthesis of oligonucleotide-peptide conjugates. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2007 , 26, 963-7	1.4	3
296	Advances in solid-phase cycloadditions for heterocyclic synthesis. <i>ACS Combinatorial Science</i> , 2007 , 9, 521-65		32
295	Peptides and metallic nanoparticles for biomedical applications. <i>Nanomedicine</i> , 2007 , 2, 287-306	5.6	109

294	Synthesis of IB-01211, a cyclic peptide containing 2,4-concatenated thia- and oxazoles, via Hantzsch macrocyclization. <i>Organic Letters</i> , 2007 , 9, 809-11	6.2	40
293	Solid-phase synthesis of oxathiocoraline by a key intermolecular disulfide dimer. <i>Journal of the American Chemical Society</i> , 2007 , 129, 5322-3	16.4	45
292	Novel proton acceptor immonium-type coupling reagents: application in solution and solid-phase peptide synthesis. <i>Organic Letters</i> , 2007 , 9, 4475-7	6.2	36
291	Cell-penetrating proline-rich peptidomimetics. <i>Methods in Molecular Biology</i> , 2007 , 386, 241-67	1.4	4
290	The synthesis of naturally occurring peptides and their analogs. <i>Current Opinion in Drug Discovery & Development</i> , 2007 , 10, 768-83		15
289	Total solid-phase synthesis of the azathiocoraline class of symmetric bicyclic peptides. <i>Chemistry - A European Journal</i> , 2006 , 12, 9001-9	4.8	27
288	The synergy of ChemMatrix resin and pseudoproline building blocks renders RANTES, a complex aggregated chemokine. <i>Biopolymers</i> , 2006 , 84, 566-75	2.2	49
287	Design and Synthesis of New Immonium-Type Coupling Reagents. <i>European Journal of Organic Chemistry</i> , 2006 , 2006, 1563-1573	3.2	22
286	p-Nitrobenzyloxycarbonyl (pNZ) as an Alternative to Fmoc for the Protection of Amines in Solid-Phase Peptide Synthesis 2006 , 116-117		
285	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. <i>Heterocycles</i> , 2006 , 69, 569	0.8	9
284	Role of the Acid Group in the Pictet-Spengler Reaction of β -Amino Acids. <i>Synlett</i> , 2006 , 2006, 1903-1907	2.2	10
283	Solid-phase chemistry in the total synthesis of non-peptidic natural products. <i>Mini-Reviews in Medicinal Chemistry</i> , 2006 , 6, 11-25	3.2	15
282	Convergent approaches for the synthesis of the antitumoral peptide, Kahalalide F. Study of orthogonal protecting groups. <i>Journal of Organic Chemistry</i> , 2006 , 71, 7196-204	4.2	22
281	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, 3335-8	4.2	37
280	Total solid-phase synthesis of marine cyclodepsipeptide IB-01212. <i>Journal of Organic Chemistry</i> , 2006 , 71, 3339-44	4.2	18
279	ChemMatrix, a poly(ethylene glycol)-based support for the solid-phase synthesis of complex peptides. <i>ACS Combinatorial Science</i> , 2006 , 8, 213-20		213
278	New Nomenclature for Complex Cyclopeptides 2006 , 142-143		
277	New efficient substrates for semicarbazide-sensitive amine oxidase/VAP-1 enzyme: analysis by SARs and computational docking. <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 6197-208	8.3	23

- 276 Hexafluoroacetone as protecting and activating reagent: new routes to amino, hydroxy, and mercapto acids and their application for peptide and glyco- and depsipeptide modification. *Chemical Reviews*, **2006**, 106, 4728-46 68.1 30
- 275 Synthesis and structure-activity relationship study of potent cytotoxic analogues of the marine alkaloid Lamellarin D. *Journal of Medicinal Chemistry*, **2006**, 49, 3257-68 8.3 93
- 274 Enhancing Atom Economy of SPS: Recoverable and Reusable Building Blocks for Depsipeptide Synthesis **2006**, 108-109
- 273 Total Solid Phase Synthesis of a Marine Cyclodepsipeptide IB-01212 **2006**, 210-211
- 272 Chlorotriyl Chloride (CTC) Resin as a Convenient Reusable Protecting Group **2006**, 220-221 2
- 271 Solid-Phase Peptide Synthesis Using ChemMatrix[®], a Polyethylenglycol (PEG)-based Solid **2006**, 114-115 1
- 270 1-Hydroxy-6,7-dimethoxy-8-nitro-1,2,3,4-tetrahydroisoquinoline. *Acta Crystallographica Section E: Structure Reports Online*, **2006**, 62, o2285-o2287
- 269 Solid-Phase N-Electrophilic Amination of 2-Aminopyridines: Preparation of 2-Substituted-[1,2,4]triazolo[1,5-a]pyridine Derivatives. *QSAR and Combinatorial Science*, **2006**, 25, 961-965 6
- 268 Solid-Phase Synthesis and Structural Study of Substituted 1,4,5,6-Tetrahydro-6-oxopyridine-3-carboxylic Acids. *QSAR and Combinatorial Science*, **2006**, 25, 921-927 9
- 267 Design of a minimized cyclic tetrapeptide that neutralizes bacterial endotoxins. *Journal of Peptide Science*, **2006**, 12, 491-6 2.1 14
- 266 Facile solid-phase synthesis of biotinylated alkyl thiols. *Tetrahedron*, **2006**, 62, 6876-6881 2.4 18
- 265 Sulfoxidations in the solid phase. *Tetrahedron: Asymmetry*, **2006**, 17, 3327-3331 4
- 264 A convenient semicarbazide resin for the solid-phase synthesis of peptide ketones and aldehydes. *Tetrahedron Letters*, **2006**, 47, 1657-1661 2 16
- 263 Homologation of β -hydroxy acids to α -unsubstituted β -hydroxy carboxamides via Arndt-Eistert reaction. *Tetrahedron Letters*, **2006**, 47, 4557-4560 2 7
- 262 Microwave-assisted synthesis of 1,3-dihydro-[1,2,5]thiadiazolo[3,4-b]pyrazine-2,2-dioxides. *Tetrahedron Letters*, **2006**, 47, 8603-8606 2 9
- 261 Synthesis of alpha-trifluoromethyl alpha-amino acids with aromatic, heteroaromatic and ferrocenyl subunits in the side chain. *Amino Acids*, **2006**, 31, 55-62 3.5 8
- 260 Synthetic Approaches to Disulfide-free Circular Bovine Pancreatic Trypsin Inhibitor (c-BPTI) Analogues. *International Journal of Peptide Research and Therapeutics*, **2006**, 12, 93-104 2.1 2
- 259 Domino reactions with fluorinated five-membered heterocycles. alpha-Trifluoromethyl alpha-amino acids with unsaturated side-chains. *Amino Acids*, **2006**, 31, 427-33 3.5 4

258	Solid-phase synthesis and characterization of N-methyl-rich peptides. <i>Chemical Biology and Drug Design</i> , 2005 , 65, 153-66		94
257	Backbone amide linker strategies for the solid-phase synthesis of C-terminal modified peptides. <i>Methods in Molecular Biology</i> , 2005 , 298, 195-208	1.4	7
256	Cell-penetrating cis-gamma-amino-L-proline-derived peptides. <i>Journal of the American Chemical Society</i> , 2005 , 127, 9459-68	16.4	90
255	A straightforward synthesis of 5'-peptide oligonucleotide conjugates using N(alpha)-Fmoc-protected amino acids. <i>Organic Letters</i> , 2005 , 7, 4349-52	6.2	26
254	Preparation of de novo globular proteins based on proline dendrimers. <i>Journal of Organic Chemistry</i> , 2005 , 70, 6274-81	4.2	21
253	A new strategy for solid-phase depsipeptide synthesis using recoverable building blocks. <i>Organic Letters</i> , 2005 , 7, 597-600	6.2	20
252	Peptide and amide bond-containing dendrimers. <i>Chemical Reviews</i> , 2005 , 105, 1663-81	68.1	296
251	Modular total synthesis of lamellarin D. <i>Journal of Organic Chemistry</i> , 2005 , 70, 8231-4	4.2	94
250	Evaluation of solution and solid-phase approaches to the synthesis of libraries of alpha,alpha-disubstituted-alpha-acylaminoketones. <i>ACS Combinatorial Science</i> , 2005 , 7, 843-63		10
249	Suzuki coupling reaction for the solid-phase preparation of 5-substituted nicotinic acid derivatives. <i>Tetrahedron Letters</i> , 2005 , 46, 581-585	2	24
248	An efficient strategy for the preparation of one-bead-one-peptide libraries on a new biocompatible solid support. <i>Tetrahedron Letters</i> , 2005 , 46, 1561-1564	2	39
247	5,6-Dihydropyrrolo[2,1-b]isoquinolines as scaffolds for synthesis of lamellarin analogues. <i>Tetrahedron Letters</i> , 2005 , 46, 2041-2044	2	39
246	N-[Chloro(dimethylamino)methylene]-N-methylmethanaminium chloride (TMUCl Cl), the reagent of choice for the solid-phase synthesis of anilides. <i>Tetrahedron Letters</i> , 2005 , 46, 5383-5386	2	12
245	NO as temporary guanidino-protecting group provides efficient access to Pbf-protected argininic acid. <i>Tetrahedron Letters</i> , 2005 , 46, 6733-6735	2	6
244	Avoiding pyran ring opening during palladium acetate catalyzed C-glycosidation of peracetylated glycals. <i>Tetrahedron Letters</i> , 2005 , 46, 7271-7274	2	22
243	Semipermanent p-nitrobenzyloxycarbonyl (pNZ) protection of Orn and Lys side chains: prevention of undesired Fmoc removal and application to the synthesis of cyclic peptides. <i>Tetrahedron Letters</i> , 2005 , 46, 7733-7736	2	11
242	Use of p-nitrobenzyloxycarbonyl (pNZ) as a permanent protecting group in the synthesis of Kahalalide F analogs. <i>Tetrahedron Letters</i> , 2005 , 46, 7737-7741	2	16
241	A new approach to 3-hydroxyquinoline-2-carboxylic acid. <i>Tetrahedron</i> , 2005 , 61, 1407-1411	2.4	10

240	Combinatorial approaches towards the discovery of new trypsin inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005 , 15, 1659-64	2.9	17
239	Multicomponent reactions with dihydroazines: efficient synthesis of a diverse set of pyrido-fused tetrahydroquinolines. <i>ACS Combinatorial Science</i> , 2005 , 7, 33-41		45
238	Abbreviated nomenclature for cyclic and branched homo- and hetero-detic peptides. <i>Chemical Biology and Drug Design</i> , 2005 , 65, 550-5		43
237	p-Nitrobenzyloxycarbonyl (pNBZ) 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	3.2	43
236	Synthesis and screening of a small library of proline-based biodendrimers for use as delivery agents. <i>Biopolymers</i> , 2005 , 80, 800-14	2.2	27
235	Hexafluoroacetone as a Protecting and Activating Reagent. N- and O-Glycosylation of Isoleucine and Isocysteine. <i>Monatshefte für Chemie</i> , 2005 , 136, 577-595	1.4	4
234	Orthogonally Protected, Carboxy-Activated L-Homoleucine, 2-Methyl-L-homoleucine, and Homocysteine Derivatives. New Building Blocks for Peptide and Dipeptide Modification. <i>Monatshefte für Chemie</i> , 2005 , 136, 763-776	1.4	4
233	Domino Reactions with Fluorinated Five-membered Heterocycles [Syntheses of Trifluoromethyl Substituted Butenolides and β -Ketoacids. <i>Monatshefte für Chemie</i> , 2005 , 136, 1763-1779	1.4	13
232	Solid-Phase Preparation of a Library Based on a Phenylalanine Scaffold. <i>QSAR and Combinatorial Science</i> , 2005 , 24, 913-922		9
231	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	2.1	17
230	Chapter 1 Lamellarins: Isolation, activity and synthesis. <i>Progress in Heterocyclic Chemistry</i> , 2005 , 16, 1-26	0.8	20
229	Directly Linked Polyazoles: Important Moieties in Natural Products. <i>Synthesis</i> , 2005 , 2005, 1907-1922	2.9	114
228	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	0.6	3
227	Application of hexafluoroacetone as protecting and activating reagent in solid phase peptide and dipeptide synthesis. <i>Arkivoc</i> , 2005 , 2005, 191-199	0.9	10
226	Re-Evaluation of a Solid-Phase Hydantoin Synthesis. <i>Letters in Organic Chemistry</i> , 2004 , 1, 224-226	0.6	20
225	From production of peptides in milligram amounts for research to multi-tons quantities for drugs of the future. <i>Current Pharmaceutical Biotechnology</i> , 2004 , 5, 29-43	2.6	142
224	Inhibition of beta-amyloid toxicity by short peptides containing N-methyl amino acids. <i>Chemical Biology and Drug Design</i> , 2004 , 63, 324-8		44
223	One-Pot Preparation of N-Carbamate Protected Amino Acids via the Azide. <i>Organic Process Research and Development</i> , 2004 , 8, 920-924	3.9	25

222	Synthesis of Polyheterocyclic Nitrogen-Containing Marine Natural Products. <i>Monatshefte für Chemie</i> , 2004 , 135, 615-627	1.4	39
221	Hexafluoroacetone as a Protecting and Activating Reagent. Regioselective Esterification of Aspartic, Malic, and Thiomalic Acid. <i>Monatshefte für Chemie</i> , 2004 , 135, 1427-1443	1.4	5
220	Gaining diversity in solid-phase synthesis by modulation of cleavage conditions from hydroxymethyl-based supports. Application to lamellarin synthesis. <i>Tetrahedron</i> , 2004 , 60, 8669-8675	2.4	23
219	A Combination of Different Spectroscopic Techniques to Monitor the <i>In situ</i> Solid-phase Synthesis of Organic Molecules. <i>QSAR and Combinatorial Science</i> , 2004 , 23, 61-68		5
218	A Re-evaluation of the Use of Rink, BAL, and PAL Resins and Linkers. <i>QSAR and Combinatorial Science</i> , 2004 , 23, 145-152		33
217	Solid-phase synthesis of second-generation polyproline dendrimers. <i>Biopolymers</i> , 2004 , 76, 283-97	2.2	24
216	Synthetic Circularized Analogues of Bovine Pancreatic Trypsin Inhibitor. <i>European Journal of Organic Chemistry</i> , 2004 , 2004, 4541-4544	3.2	4
215	Synthesis of Polyheterocyclic Nitrogen-Containing Marine Natural Products.. <i>ChemInform</i> , 2004 , 35, no		1
214	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	2	3
213	Solid-phase synthesis of lamellarins Q and O. <i>Tetrahedron</i> , 2004 , 60, 8659-8668	2.4	46
212	Developments in peptide and amide synthesis. <i>Current Opinion in Chemical Biology</i> , 2004 , 8, 211-21	9.7	208
211	Solid-phase combinatorial synthesis of peptide-biphenyl hybrids as calpain inhibitors. <i>Organic Letters</i> , 2004 , 6, 4089-92	6.2	21
210	A comparative study of different presentation strategies for an HIV peptide immunogen. <i>Bioconjugate Chemistry</i> , 2004 , 15, 112-20	6.3	45
209	A new class of foldamers based on cis-gamma-amino-L-proline. <i>Journal of the American Chemical Society</i> , 2004 , 126, 6048-57	16.4	92
208	Solid-phase syntheses of furopyridine and furoquinoline systems. <i>Organic Letters</i> , 2004 , 6, 1405-8	6.2	34
207	Monoclonal antibody purification by affinity chromatography with ligands derived from the screening of peptide combinatorial libraries. <i>Biotechnology Letters</i> , 2003 , 25, 1545-8	3	29
206	Synthesis and antifungal activity of an acivicine-based dipeptide library. <i>International Journal of Peptide Research and Therapeutics</i> , 2003 , 10, 645-653		2
205	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, 1096-103	4.8	6

204	Solid-phase synthesis of C-terminal modified peptides. <i>Biopolymers</i> , 2003 , 71, 454-77	2.2	56
203	Saturated resins or stress of the resin. <i>Tetrahedron Letters</i> , 2003 , 44, 1751-1754	2	14
202	o-Formylation of electron-rich phenols with dichloromethyl methyl ether and TiCl ₄ . <i>Tetrahedron Letters</i> , 2003 , 44, 4961-4963	2	19
201	Solid-phase synthesis: a linker for side-chain anchoring of arginine. <i>Tetrahedron Letters</i> , 2003 , 44, 5319-5321		23
200	Synthesis of variolin B. <i>Tetrahedron Letters</i> , 2003 , 44, 6191-6194	2	18
199	BAL resin for the preparation of secondary amines. <i>Tetrahedron Letters</i> , 2003 , 44, 6907-6910	2	18
198	Synthesis and SAR of alpha-acylaminoketone ligands for control of gene expression. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2003 , 13, 475-8	2.9	43
197	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		17
196	Solid-phase total synthesis of the pentacyclic system lamellarins U and L. <i>Organic Letters</i> , 2003 , 5, 2959-62		70
195	Solution structure of the antitumor candidate trunkamide A by 2D NMR and restrained simulated annealing methods. <i>Journal of Organic Chemistry</i> , 2003 , 68, 211-5	4.2	31
194	Bicyclic homodetic peptide libraries: comparison of synthetic strategies for their solid-phase synthesis. <i>ACS Combinatorial Science</i> , 2003 , 5, 760-8		22
193	Synthesis and NMR structure of p41icf, a potent inhibitor of human cathepsin L. <i>Journal of the American Chemical Society</i> , 2003 , 125, 1508-17	16.4	21
192	Tentoxin as a scaffold for drug discovery. Total solid-phase synthesis of tentoxin and a library of analogues. <i>Organic Letters</i> , 2003 , 5, 2115-8	6.2	19
191	Qualitative colorimetric tests for solid phase synthesis. <i>Methods in Enzymology</i> , 2003 , 369, 21-35	1.7	27
190	Total syntheses of variolin B and deoxyvariolin B. <i>Journal of Organic Chemistry</i> , 2003 , 68, 10020-9	4.2	47
189	Fmoc solid-phase synthesis of peptide thioesters by masking as trithioortho esters. <i>Organic Letters</i> , 2003 , 5, 2951-3	6.2	101
188	Exploring solid-phase approaches for the preparation of new beta-lactams from amino acids. <i>Molecular Diversity</i> , 2003 , 6, 75-84	3.1	11
187	¹ H NMR spectroscopy with internal and external standards for the quantification of libraries. <i>Molecular Diversity</i> , 2003 , 6, 165-8	3.1	5

186	Small molecules targeting the vanilloid receptor complex as drugs for inflammatory pain. <i>Drugs of the Future</i> , 2003 , 28, 787	2.3	13
185	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	3.2	15
184	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	3.2	24
183	Synthesis of peptides containing α -ketoamide hydroamino acids. Scope and limitations. <i>International Journal of Peptide Research and Therapeutics</i> , 2002 , 9, 135-141		2
182	Practical protocols for stepwise solid-phase synthesis of cysteine-containing peptides. <i>Chemical Biology and Drug Design</i> , 2002 , 60, 292-9		61
181	Structural, kinetic and cytotoxicity aspects of 12-28 beta-amyloid protein fragment: a reappraisal. <i>Journal of Peptide Science</i> , 2002 , 8, 578-88	2.1	21
180	Four-dimensional orthogonal solid-phase synthesis of new scaffolds based on cyclic tetra-peptides. <i>Tetrahedron Letters</i> , 2002 , 43, 2029-2032	2	19
179	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	2	16
178	Undesired removal of the Fmoc group by the free amino function of a lysine residue. <i>Tetrahedron Letters</i> , 2002 , 43, 7813-7815	2	27
177	Solid phase synthesis of α -acylamino- β -disubstituted ketones. <i>Tetrahedron Letters</i> , 2002 , 43, 7491-7494	2	8
176	Combined solid phase and solution synthesis of a library of β -disubstituted- α -acylamino ketones. <i>Tetrahedron Letters</i> , 2002 , 43, 7495-7498	2	11
175	Synthesis of Fmoc-protected amino ketones bearing tert-butyl based side-chain protecting groups. <i>Tetrahedron Letters</i> , 2002 , 43, 7499-7502	2	14
174	Synthesis of peptides containing α -ketoamide hydroamino acids. Scope and limitations. <i>International Journal of Peptide Research and Therapeutics</i> , 2002 , 9, 135-141		6
173	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, 863-72	3.8	4
172	Peptide dendrimers based on polyproline helices. <i>Journal of the American Chemical Society</i> , 2002 , 124, 8876-83	16.4	104
171	Synthesis and applications of a bis-sulfonyl handle for solid-phase synthesis of peptides 2002 , 307-308		
170	Solid-phase peptide synthesis in the N->C direction 2002 , 78-79		
169	Backbone Amide Linker (BAL) methodology to accommodate C-terminal hindered, unreactive, and/or sensitive modifications 2002 , 102-103		

- 168 Disulfide Bond Based Self-Assembly of Peptides Leading To Spheroidal Cyclic Trimers **2002**, 243-256
- 167 Backbone amide linker (BAL) for solid-phase synthesis of 2,5-piperazinediones (DKP), useful scaffolds for combinatorial chemistry **2002**, 37-39
- 166 Backbone Amide Linker (BAL) Strategy for Solid-Phase Synthesis **2001**, 121-138 3
- 165 Solid-phase peptide synthesis using N ϵ -trityl-amino acids. *International Journal of Peptide Research and Therapeutics*, **2001**, 8, 331-338
- 164 A useful and sensitive color test to monitor aldehydes on solid-phase. *Tetrahedron Letters*, **2001**, 42, 6691-6693 2 24
- 163 Solid-phase syntheses of constrained RGD scaffolds and their binding to the $\alpha_5\beta_1$ integrin receptor. *Tetrahedron Letters*, **2001**, 42, 7387-7391 2 28
- 162 2-Mercaptopyridine-1-oxide-based peptide coupling reagents. *Tetrahedron*, **2001**, 57, 9607-9613 2.4 21
- 161 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 2 23
- 160 Inhibition of HIV-2(ROD) replication in a lymphoblastoid cell line by the alpha1-antitrypsin Portland variant (alpha1-PDX) and the decRVKRCmk peptide: comparison with HIV-1(LAI). *Microbes and Infection*, **2001**, 3, 1073-84 9.3 3
- 159 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, 115-20 2.1 19
- 158 Solid-Phase Synthesis of Peptides Containing β -Didehydroamino Acids. *European Journal of Organic Chemistry*, **2001**, 2001, 45-48 3.2 8
- 157 Solid-phase peptide synthesis using N ϵ -trityl-amino acids. *International Journal of Peptide Research and Therapeutics*, **2001**, 8, 331-338 8
- 156 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 1.7 17
- 155 Solid-phase total synthesis of trunkamide A(1). *Journal of Organic Chemistry*, **2001**, 66, 7568-74 4.2 41
- 154 Synthesis and structure determination of kahalalide F (1,2). *Journal of the American Chemical Society*, **2001**, 123, 11398-401 16.4 103
- 153 Synthesis of a sulfahydantoin library. *ACS Combinatorial Science*, **2001**, 3, 290-300 22
- 152 Proline: A Key Building Block in *De novo* Designed Peptide Molecules **2001**, 432-434
- 151 Design, synthesis, and conformational analysis of azacycloalkane amino acids as conformationally constrained probes for mimicry of peptide secondary structures. *Biopolymers*, **2000**, 55, 101-22 2.2 99

150	Orthogonal protecting groups for N(alpha)-amino and C-terminal carboxyl functions in solid-phase peptide synthesis. <i>Biopolymers</i> , 2000 , 55, 123-39	2.2	76
149	An efficient solid-phase strategy for the construction of chemokines. <i>Journal of Peptide Science</i> , 2000 , 6, 512-8	2.1	13
148	Solid phase synthesis of sulfahydantoin. <i>Tetrahedron Letters</i> , 2000 , 41, 3161-3163	2	24
147	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	2	30
146	Synthesis of aspartimide-free protected peptides on base-labile functionalized resins. <i>Tetrahedron Letters</i> , 2000 , 41, 8093-8096	2	9
145	Kahalalide B. Synthesis of a natural cyclodepsipeptide. <i>Tetrahedron Letters</i> , 2000 , 41, 9765-9769	2	21
144	Nsc and Fmoc Nalpha-amino protection for solid-phase peptide synthesis: a parallel study. <i>Chemical Biology and Drug Design</i> , 2000 , 56, 63-9		17
143	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		2
142	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		
141	Disulfide Bonded Cyclic Peptide Dimers and Trimers: An Easy Entry to High Symmetry Peptide Frameworks. <i>Synlett</i> , 2000 , 2000, 172-181	2.2	7
140	Solid-phase peptide synthesis in the reverse (N --> C) direction. <i>Organic Letters</i> , 2000 , 2, 1815-7	6.2	50
139	Substituted guanidines: introducing diversity in combinatorial chemistry. <i>Organic Letters</i> , 2000 , 2, 3539-42		31
138	Understanding the structure/reactivity of aminium/uronium salts as coupling reagents in peptide synthesis. <i>Tetrahedron Letters</i> , 1999 , 40, 2641-2644	2	9
137	Useful scaffolds and handles for creating diversity in the preparation of chemical libraries. <i>Reactive and Functional Polymers</i> , 1999 , 41, 103-110	4.6	10
136	Continuous-flow solid-phase peptide synthesis using polystyrene resins. <i>Chemical Biology and Drug Design</i> , 1999 , 53, 682-3		5
135	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		3
134	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		4
133	An HPLC-ESMS study on the solid-phase assembly of C-terminal proline peptides. <i>Journal of Peptide Science</i> , 1999 , 5, 131-40	2.1	32

132	Solid-Phase Synthesis with Tris(alkoxy)benzyl Backbone Amide Linkage (BAL) <i>Chemistry - A European Journal</i> , 1999 , 5, 2787-2795	4.8	76
131	Backbone Amide Linker (BAL) Strategy for N(alpha)-(9-Fluorenylmethoxycarbonyl (Fmoc) Solid-Phase Synthesis of Unprotected Peptide p-Nitroanilides and Thioesters(1). <i>Journal of Organic Chemistry</i> , 1999 , 64, 8761-8769	4.2	140
130	N-Boc temporary protection in solid-phase peptide synthesis. The use of amine-borane complexes as allyl group scavengers. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1999 , 2871-2874		63
129	Synthesis and binding properties of oligonucleotides carrying nuclear localization sequences. <i>Bioconjugate Chemistry</i> , 1999 , 10, 1005-12	6.3	35
128	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		
127	Use of Onium Salt-Based Coupling Reagents in Peptide Synthesis ¹ . <i>Journal of Organic Chemistry</i> , 1998 , 63, 9678-9683	4.2	206
126	Active carbonate resins: Application to the solid-phase synthesis of alcohol, carbamate and cyclic peptides. <i>Tetrahedron</i> , 1998 , 54, 10125-10152	2.4	47
125	"High-load" polyethylene glycol-polystyrene (PEG-PS) graft supports for solid-phase synthesis. <i>Biopolymers</i> , 1998 , 47, 365-80	2.2	33
124	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		3
123	Solid-phase synthesis of diketopiperazines, useful scaffolds for combinatorial chemistry. <i>Tetrahedron Letters</i> , 1998 , 39, 2639-2642	2	56
122	Structural/functional properties of the Glu1-HSer57 N-terminal fragment of human plasminogen: conformational characterization and interaction with kringle domains. <i>Protein Science</i> , 1998 , 7, 1947-59	6.3	16
121	Lysine-50 is a likely site for anchoring the plasminogen N-terminal peptide to lysine-binding kringles. <i>Protein Science</i> , 1998 , 7, 1960-9	6.3	35
120	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	16.4	12
119	Backbone Amide Linker (BAL) Strategy for Solid-Phase Synthesis of C-Terminal-Modified and Cyclic Peptides ^{1,2,3} . <i>Journal of the American Chemical Society</i> , 1998 , 120, 5441-5452	16.4	273
118	IBTM-Containing Gramicidin S Analogues: Evidence for IBTM as a Suitable Type III Turn Mimetic ^{1,2} . <i>Journal of the American Chemical Society</i> , 1997 , 119, 10579-10586	16.4	50
117	Poly(ethylene glycol)-Containing Supports for Solid-Phase Synthesis of Peptides and Combinatorial Organic Libraries. <i>ACS Symposium Series</i> , 1997 , 239-264	0.4	14
116	Coupling reagents and activation. <i>Methods in Enzymology</i> , 1997 , 289, 104-26	1.7	72
115	Convergent solid-phase peptide synthesis. <i>Methods in Enzymology</i> , 1997 , 289, 313-36	1.7	23

114	Occurrence and Minimization of Cysteine Racemization during Stepwise Solid-Phase Peptide Synthesis(1),(2). <i>Journal of Organic Chemistry</i> , 1997 , 62, 4307-4312	4.2	182
113	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	4.2	70
112	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	2.4	15
111	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	2	55
110	A new approach to Hmb-backbone protection of peptides: Synthesis and reactivity of N ^F Fmoc-N ^H (Hmb)amino acids. <i>Tetrahedron Letters</i> , 1997 , 38, 2317-2320	2	30
109	On the use of PyAOP, a phosphonium salt derived from HOAt, in solid-phase peptide synthesis. <i>Tetrahedron Letters</i> , 1997 , 38, 4853-4856	2	137
108	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	2	137
107	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	4.2	43
106	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		3
105	On the use of novel coupling reagents for solid-phase peptide synthesis. <i>Techniques in Protein Chemistry</i> , 1996 , 515-523		9
104	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	2	56
103	New carbamate supports for the preparation of 3'-amino-modified oligonucleotides. <i>Bioorganic and Medicinal Chemistry</i> , 1996 , 4, 1649-58	3.4	19
102	3-(1-Piperidiny)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		52
101	A COMPARATIVE STUDY OF SUPPORTS FOR THE SYNTHESIS OF OLIGONUCLEOTIDES WITHOUT USING AMMONIA. <i>Nucleosides & Nucleotides</i> , 1996 , 15, 1871-1889		17
100	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, 119-33		4
99	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		13
98	Synthesis and applications of a new base-labile fluorene derived linker for solid-phase peptide synthesis. <i>Tetrahedron</i> , 1995 , 51, 1449-1458	2.4	33
97	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	4.2	23

96	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	4.2	176
95	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	4.2	23
94	Stepwise Automated Solid Phase Synthesis of Naturally Occurring Peptaibols Using Fmoc Amino Acid Fluorides. <i>Journal of Organic Chemistry</i> , 1995 , 60, 405-410	4.2	113
93	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		32
92	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		76
91	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	2	27
90	Stepwise solid-phase synthesis of oligonucleotide-peptide hybrids. <i>Tetrahedron Letters</i> , 1994 , 35, 2733-2736		45
89	Solid-phase N-glycopeptide synthesis using allyl side-chain protected Fmoc-amino acids. <i>Tetrahedron Letters</i> , 1994 , 35, 1033-1034	2	37
88	Preparation and applications of polyethylene glycol-polystyrene graft resin supports for solid-phase peptide synthesis. <i>Reactive & Functional Polymers</i> , 1994 , 22, 243-258		118
87	Racemization studies during solid-phase peptide synthesis using azabenzotriazole-based coupling reagents. <i>Tetrahedron Letters</i> , 1994 , 35, 2279-2282	2	172
86	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	2	8
85	Solid-phase synthesis of bead-to-tail cyclic peptides via lysine side-chain anchoring. <i>Tetrahedron Letters</i> , 1994 , 35, 9633-9636	2	76
84	Formation of disulfide bonds in synthetic peptides and proteins. <i>Methods in Molecular Biology</i> , 1994 , 35, 91-169	1.4	87
83	Advantageous applications of azabenzotriazole (triazolopyridine)-based coupling reagents to solid-phase peptide synthesis. <i>Journal of the Chemical Society Chemical Communications</i> , 1994 , 201		292
82	Peptides in molecular recognition: synthetic and conformational aspects. <i>Biochemical Society Transactions</i> , 1994 , 22, 1045-8	5.1	1
81	Chemical synthesis of a fully active transcriptional repressor protein. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1994 , 91, 5178-82	11.5	15
80	Solid-Phase Synthesis of Cyclic Peptides 1994 , 39-58		31
79	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, 307-12		14

78	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		6
77	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	16.4	23
76	Unequivocal synthesis and characterization of a parallel and an antiparallel bis-cystine peptide. <i>Journal of Organic Chemistry</i> , 1993 , 58, 6319-6328	4.2	9
75	Antibodies cross-reactive with the scorpion-toxin II from <i>Androctonus australis</i> Hector elicited in mice by a synthetic peptide. <i>Natural Toxins</i> , 1993 , 1, 255-62		7
74	Automated allyl cleavage for continuous-flow synthesis of cyclic and branched peptides. <i>Analytical Biochemistry</i> , 1993 , 212, 303-10	3.1	123
73	A novel, convenient, three-dimensional orthogonal strategy for solid-phase synthesis of cyclic peptides. <i>Tetrahedron Letters</i> , 1993 , 34, 1549-1552	2	223
72	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	2.4	14
71	Convergent solid-phase peptide synthesis. <i>Tetrahedron</i> , 1993 , 49, 11065-11133	2.4	151
70	The 2,2,4,6,7-pentamethyldihydrobenzofuran-5-sulfonyl group (Pbf) as arginine side chain protectant. <i>Tetrahedron Letters</i> , 1993 , 34, 7829-7832	2	91
69	Novel polyethylene glycol-polystyrene (PEG-PS) graft supports for solid-phase peptide synthesis 1993 , 267-268		16
68	Allyl-based orthogonal solid phase peptide synthesis 1993 , 191-193		10
67	Strategies and tactics for the solid-phase synthesis of cystine-containing peptides 1993 , 19-23		1
66	Preparation and applications of xanthenylamide (XAL) handles for mild Fmoc solid-phase synthesis of C-terminal peptide amides 1993 , 301-304		
65	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	4.2	45
64	A synthetic procedure for the preparation of oligonucleotides without using ammonia and its application for the synthesis of oligonucleotides containing 0-4-alkyl thymidines.. <i>Tetrahedron</i> , 1992 , 48, 4171-4182	2.4	31
63	-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	2	4
62	Reversible protection of lysine to facilitate the purification of protected peptide segments. <i>Tetrahedron Letters</i> , 1992 , 33, 397-400	2	6
61	S-2-(2,4-dinitrophenyl)ethyl--cysteine: a new derivative for solid-phase peptide synthesis. <i>Tetrahedron Letters</i> , 1992 , 33, 2391-2394	2	12

60	A new fluorene-derived anchor for solid-phase synthesis of protected peptides. <i>Tetrahedron Letters</i> , 1992 , 33, 1775-1778	2	26
59	Synthesis and ion-binding properties of an immobilized bis-cysteine peptide. <i>Bioorganic and Medicinal Chemistry Letters</i> , 1992 , 2, 281-284	2.9	
58	A new approach to the solid-phase peptide synthesis of peptide alkyl-amides and esters. <i>Tetrahedron Letters</i> , 1992 , 33, 2183-2186	2	7
57	Enzymatic peptide fragment condensation. Choice of reaction media for the synthesis of an insect neuropeptide derivative. <i>Biotechnology Letters</i> , 1992 , 6, 69-72		2
56	Synthesis and applications of XAL, a new acid-labile handle for solid-phase synthesis of peptide amides 1992 , 601-602		10
55	Biopolymer syntheses on novel polyethylene glycol-polystyrene (PEG-PS) graft supports 1992 , 603-604		16
54	Novel cysteine protecting groups for the N ϵ -fluorenylmethyloxycarbonyl (Fmoc) strategy of peptide synthesis 1992 , 605-606		1
53	Convergent solid-phase peptide synthesis 1992 , 607-608		1
52	RNA binding characteristics of a 16 kDa glycine-rich protein from maize. <i>Plant Journal</i> , 1992 , 2, 999-1003	6.9	
51	Cyclization of disulfide-containing peptides in solid-phase synthesis. <i>International Journal of Peptide and Protein Research</i> , 1991 , 37, 402-13		70
50	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.. <i>Tetrahedron Letters</i> , 1991 , 32, 1511-1514	2	37
49	NPE-resin, a new approach to the solid-phase synthesis of protected peptides and oligonucleotides II. Synthesis of protected peptides. <i>Tetrahedron Letters</i> , 1991 , 32, 1515-1518	2	25
48	Binding and toxicity of apamin. Characterization of the active site. <i>FEBS Journal</i> , 1991 , 196, 639-45		49
47	Hypersensitive acid-labile (HAL) tris(alkoxy)benzyl ester anchoring for solid-phase synthesis of protected peptide segments. <i>Tetrahedron Letters</i> , 1991 , 32, 1015-1018	2	34
46	Solid-phase synthesis of peptides using allylic anchoring groups. An investigation of their palladium-catalysed cleavage. <i>Tetrahedron Letters</i> , 1991 , 32, 4207-4210	2	48
45	Convergent solid-phase peptide synthesis. X. Synthesis and purification of protected peptide fragments using the photolabile Nbb-resin. <i>Tetrahedron</i> , 1991 , 47, 9867-9880	2.4	21
44	Synthesis of defined peptide-oligonucleotide hybrids containing a nuclear transport signal sequence.. <i>Tetrahedron</i> , 1991 , 47, 4113-4120	2.4	78
43	Synthesis and biological activity of O-glycosylated morphiceptin analogues. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1991 , 1755-1759		34

42	Mild orthogonal solid-phase peptide synthesis 1991 , 139-142		1
41	Solid-phase synthesis of new glycosyl enkephalinamides 1991 , 416-417		
40	Application of acetamidomethyl and 9-fluorenylmethyl groups for efficient side protection of penicillamine in solid-phase peptide synthesis. <i>International Journal of Peptide and Protein Research</i> , 1990 , 35, 434-40		8
39	Practical approach to solid-phase synthesis of C-terminal peptide amides under mild conditions based on a photolysable anchoring linkage. <i>International Journal of Peptide and Protein Research</i> , 1990 , 36, 31-45		44
38	Orthogonal solid-phase synthesis of human gastrin-I under mild conditions. <i>International Journal of Peptide and Protein Research</i> , 1990 , 35, 527-38		21
37	Use of BOP reagent for the suppression of diketopiperazine formation in boc/bzl solid-phase peptide synthesis. <i>Tetrahedron Letters</i> , 1990 , 31, 7363-7366	2	50
36	Arenesulphonyltriazolides as condensing reagents in solid phase peptide synthesis. <i>Tetrahedron Letters</i> , 1990 , 31, 1915-1918	2	12
35	Solid-Phase Synthesis of Glycopeptide Amides under Mild Conditions: Morphiceptin Analogues. <i>Angewandte Chemie International Edition in English</i> , 1990 , 29, 291-292		20
34	Festphasen-Synthese von Glycopeptidamiden unter milden Bedingungen: Morphiceptin-Analoga. <i>Angewandte Chemie</i> , 1990 , 102, 311-313	3.6	12
33	Improved method for the synthesis of o-glycosylated fmoc amino acids to be used in solid-phase glycopeptide synthesis (Fmoc = fluoren-9-ylmethoxycarbonyl). <i>Journal of the Chemical Society Chemical Communications</i> , 1990 , 965-967		26
32	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. <i>Journal of Organic Chemistry</i> , 1990 , 55, 3730-3743	4.2	310
31	Solid-phase-mediated peptide heterodisulfide formation. <i>Journal of the American Chemical Society</i> , 1990 , 112, 5345-5347	16.4	18
30	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, 124-8		41
29	Use of polystyrene-1% divinylbenzene and Kel-F-g-styrene for the simultaneous synthesis of peptides. <i>Reactive & Functional Polymers</i> , 1989 , 10, 259-268		6
28	Quantitative monitoring of carboxyl groups in polymers. <i>Analytica Chimica Acta</i> , 1989 , 219, 161-163	6.6	2
27	Convenient synthesis of a cyclic peptide disulfide: A type II Eurn structural model. <i>Tetrahedron Letters</i> , 1989 , 30, 2441-2444	2	31
26	Convergent solid phase peptide synthesis. VII. Good yields in the coupling of protected segments on a solid support. <i>Tetrahedron</i> , 1989 , 45, 4637-4648	2.4	21
25	Comparative study of supports for solid-phase coupling of protected-peptide segments. <i>Journal of Organic Chemistry</i> , 1989 , 54, 360-366	4.2	46

24	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, 483-93	4.6	64
23	Uteroglobin-like peptide cavities I. Synthesis of antiparallel and parallel dimers of bis-cysteine peptides. <i>Tetrahedron Letters</i> , 1988 , 29, 3845-3848	2	30
22	Use of polar picolyl protecting groups in peptide synthesis. <i>Journal of Organic Chemistry</i> , 1988 , 53, 5386-5389	4.389	13
21	Mild, orthogonal solid-phase peptide synthesis: use of N alpha-dithiasuccinoyl (Dts) amino acids and N-(iso-propylidithio)carbonylproline, together with p-alkoxybenzyl ester anchoring linkages. <i>International Journal of Peptide and Protein Research</i> , 1987 , 30, 177-205		26
20	An acid-labile anchoring linkage for solid-phase synthesis of C-terminal peptide amides under mild conditions. <i>International Journal of Peptide and Protein Research</i> , 1987 , 30, 206-16		95
19	A convenient general method for synthesis of N alpha- or N omega-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, 740-83		28
18	On the use of s-t-butylsulphenyl group for protection of cysteine in solid-phase peptide synthesis using fmoc-amino acids. <i>Tetrahedron</i> , 1987 , 43, 2675-2680	2.4	70
17	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	2.4	19
16	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	2.4	14
15	(S)-9-Fluorenylmethyl-L-cysteine, a useful HF-stable derivative for peptide synthesis. <i>Journal of the Chemical Society Chemical Communications</i> , 1986 , 1501		20
14	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	6.5	58
13	Improved approach for anchoring N alpha-9-fluorenylmethyloxycarbonylamino acids as p-alkoxybenzyl esters in solid-phase peptide synthesis. <i>International Journal of Peptide and Protein Research</i> , 1985 , 26, 92-7		66
12	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	16.4	117
11	Application of N,N-dimethylformamide dineopentyl acetal for efficient anchoring of N alpha-9-fluorenylmethyloxycarbonylamino acids as p-alkoxybenzyl esters in solid-phase peptide synthesis. <i>International Journal of Peptide and Protein Research</i> , 1984 , 23, 342-9		21
10	Convergent solid phase peptide synthesis. II. Synthesis of the 1B apamin protected segment on a NBB-resin. Synthesis of apamin. <i>Tetrahedron</i> , 1982 , 38, 1193-1201	2.4	46
9	Acid-base properties of 4-nitro-l-histidine and related compounds. <i>Bioorganic Chemistry</i> , 1979 , 8, 59-67	5.1	3
8	The (Classic Concept of) Solid Support1-14		0
7	High-Throughput Synthesis of Natural Products613-640		0

6 Peptides493-527

5 Coupling and Introducing Building Block Reagents141-181

0

4 Immobilized Enzymes in Organic Synthesis365-380

1

3 Acid-Labile Resins381-416

2 Base/Nucleophile-Labile Resins417-436

1

1 CHAPTER 18:Solid-Phase Peptide Synthesis, the State of the Art: Challenges and Opportunities.
RSC Drug Discovery Series,518-550

0.6 10