Antonio Guarna

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/10899101/publications.pdf

Version: 2024-02-01

142 papers

3,573 citations

147801 31 h-index 206112 48 g-index

174 all docs

174 docs citations

174 times ranked

2876 citing authors

#	Article	IF	CITATIONS
1	A New Family of Cinchona-Derived Amino Phosphine Precatalysts: Application to the Highly Enantioand Diastereoselective Silver-Catalyzed Isocyanoacetate Aldol Reaction. Journal of the American Chemical Society, 2011, 133, 1710-1713.	13.7	225
2	1,3-Aminoalcohols by reductive cleavage of isoxazolidines with molybdenum hexacarbonyl. Tetrahedron Letters, 1990, 31, 3351-3354.	1.4	209
3	Rearrangement of isoxazoline-5-spiro derivatives. 2. Synthesis and rearrangement of tetrahydroisoxazole-5-spirocyclopropanes. Preparation of precursors of quinolizine, isoquinoline, and indole alkaloids. Journal of Organic Chemistry, 1988, 53, 2430-2434.	3.2	82
4	New Synthetic Approach to Cyclopenta-Fused Heterocycles Based upon a Mild Nazarov Reactionâ€. Journal of Organic Chemistry, 2003, 68, 9728-9741.	3.2	78
5	Suzuki Reaction of Vinyl Triflates from Six- and Seven-MemberedN-Alkoxycarbonyl Lactams with Boronic Acids and Esters. Journal of Organic Chemistry, 2001, 66, 2459-2465.	3.2	77
6	Organocatalytic Diastereo- and Enantioselective Michael Addition Reactions of 5-Aryl-1,3-dioxolan-4-ones. Organic Letters, 2007, 9, 2107-2110.	4.6	72
7	Preparation and Suzukiâ^'Miyaura Coupling Reactions of Tetrahydropyridine-2-boronic Acid Pinacol Esters. Journal of Organic Chemistry, 2005, 70, 7324-7330.	3.2	64
8	19-Nor-10-azasteroids:  A Novel Class of Inhibitors for Human Steroid 5α-Reductases 1 and 2. Journal of Medicinal Chemistry, 1997, 40, 1112-1129.	6.4	58
9	Rearrangement of isoxazoline-5-spiro derivatives. 5. Diastereofacial selectivity in the cycloaddition of substituted five-membered cyclic nitrones and methylenecyclopropanes. Stereoselective synthesis of 3,5-substituted indolizidinones. Journal of Organic Chemistry, 1990, 55, 1762-1767.	3.2	57
10	New Synthesis of Azaheterocycles by Rearrangement of Isoxazoline-5-spirocycloalkane Compounds. Synlett, 1993, 1993, 1-8.	1.8	54
11	Preparation and Cycloaddition Reactions of Enantiopure 2-(N-Acylamino)-1,3-dienes for the Synthesis of Octahydroquinoline Derivatives. Journal of Organic Chemistry, 2003, 68, 6360-6368.	3.2	53
12	Selective non-steroidal inhibitors of 5α-reductase type 1. Journal of Steroid Biochemistry and Molecular Biology, 2004, 88, 1-16.	2.5	52
13	Synthesis and Conformational Analysis of Small Peptides Containing 6-Endo-BT(t)L Scaffolds as Reverse Turn Mimetics. Journal of Organic Chemistry, 2002, 67, 7483-7492.	3.2	51
14	New Synthetic Approach to Cyclopenta-Fused Heterocycles Based upon a Mild Nazarov Reaction. 2. Further Studies on the Torquoselectivity. Journal of Organic Chemistry, 2004, 69, 7705-7709.	3.2	51
15	Remote Stereocontrol in the Nazarov Reaction:Â A New Approach to the Core of Roseophilin. Journal of Organic Chemistry, 2005, 70, 4542-4545.	3.2	51
16	Redirection of allergen-specific TH2 responses by a modified adenine through Toll-like receptor 7 interaction and IL-12/IFN release. Journal of Allergy and Clinical Immunology, 2006, 118, 511-517.	2.9	50
17	Click-Chemistry-Derived Triazole Ligands of Arginineâ^'Glycineâ^'Aspartate (RGD) Integrins with a Broad Capacity To Inhibit Adhesion of Melanoma Cells and Both in Vitro and in Vivo Angiogenesis. Journal of Medicinal Chemistry, 2010, 53, 7119-7128.	6.4	49
18	The Lewis Acid-Catalyzed Nazarov Reaction of 2-(N-Methoxycarbonylamino)-1,4-pentadien-3-ones. Organic Letters, 2006, 8, 781-784.	4.6	44

#	Article	IF	CITATIONS
19	Synthesis and Reactivity of Bicycles Derived from Tartaric Acid and α-Amino Acids: A Novel Class of Conformationally Constrained Dipeptide Isosteres Based upon Enantiopure 3-Aza-6,8-dioxabicyclo[3.2.1]octane-7-carboxylic Acid. Journal of Organic Chemistry, 1999, 64, 7347-7364.	3.2	43
20	Density Functional Studies on the Nazarov Reaction Involving Cyclic Systems. Chemistry - A European Journal, 2006, 12, 2836-2845.	3.3	42
21	A new synthesis of (2S)-4-oxopipecolic acid by thermal rearrangement of enantiopure spirocyclopropaneisoxazolidine. Tetrahedron Letters, 1996, 37, 4205-4208.	1.4	41
22	Cleavage of Isoxazolines with Tricarbonyltris(acetonitrile)molybdenum and Silica Gel. Synthesis of 1-(2-Oxoalkyl)cyclopropanols from Isoxazoline-5-spirocyclopropanes. Synthesis, 1989, 1989, 175-178.	2.3	39
23	Rearrangement of isoxazoline-5-spiro derivatives. Part 7. Thermal rearrangement of 4,5-dihydro and tetrahydroisoxazole-5-spirocyclobutanes to azepin-4-one derivatives. Tetrahedron, 1992, 48, 5283-5300.	1.9	38
24	Convenient Route to Enantiopure Fmoc-Protected Morpholine-3-carboxylic Acid. Journal of Organic Chemistry, 2007, 72, 4254-4257.	3.2	36
25	The isoxazoline-5-spirocyclopropane route to $(\hat{A}\pm)$ -Pumiliotoxin C. Tetrahedron Letters, 1992, 33, 6697-6700.	1.4	35
26	Carbonylative Suzuki–Miyaura Coupling Reaction of Lactam-, Lactone-, and Thiolactone-Derived Enol Triflates for the Synthesis of Unsymmetrical Dienones. European Journal of Organic Chemistry, 2007, 2007, 2152-2163.	2.4	34
27	The TLR7 Ligand 9-Benzyl-2-Butoxy-8-Hydroxy Adenine Inhibits IL-17 Response by Eliciting IL-10 and IL-10–Inducing Cytokines. Journal of Immunology, 2011, 186, 4707-4715.	0.8	34
28	Rearrangement of isoxazoline-5-spiro derivatives. 1. Synthesis of 4,5-dihydroisoxazole-5-spirocyclopropanes and their rearrangement to 5,6-dihydro-4-pyridones. Journal of Organic Chemistry, 1988, 53, 2426-2429.	3.2	33
29	Stereoselective cyclopropanation of serine- and threonine-derived oxazines to access new morpholine-based scaffolds. Organic and Biomolecular Chemistry, 2008, 6, 3328.	2.8	33
30	Diversityâ€Oriented Synthesis of Morpholineâ€Containing Molecular Scaffolds. Chemistry - A European Journal, 2009, 15, 7871-7875.	3.3	33
31	Enantiodivergent Chemoenzymatic Synthesis of 4â€Hydroxypiperidine Alkaloids. European Journal of Organic Chemistry, 2010, 2010, 5831-5840.	2.4	33
32	Diversity-Oriented Synthesis as a Tool for Chemical Genetics. Molecules, 2014, 19, 16506-16528.	3.8	32
33	Benzo[c]quinolizin-3-ones:  A Novel Class of Potent and Selective Nonsteroidal Inhibitors of Human Steroid 5α-Reductase 1. Journal of Medicinal Chemistry, 2000, 43, 3718-3735.	6.4	31
34	Synthesis of \hat{l} ±-Acyl-Functionalized Azacycles by Pd-Catalyzed Cross-Coupling Reactions of \hat{l} ±-Alkoxyboronates with Lactam-Derived Vinyl Triflates. Journal of Organic Chemistry, 2002, 67, 7144-7146.	3.2	31
35	Skeletal Diversity from Carbohydrates: Use of Mannose for the Diversity-Oriented Synthesis of Polyhydroxylated Compounds. Journal of Organic Chemistry, 2015, 80, 2182-2191.	3.2	30
36	The chemistry of fulminic acid revised. Tetrahedron, 1985, 41, 5181-5185.	1.9	29

#	Article	IF	CITATIONS
37	Rearrangement of isoxazoline-5-spiro derivatives. part 4. Synthesis of medium size benzofused azaheterocycles. Tetrahedron, 1989, 45, 5917-5924.	1.9	29
38	Microbial biotransformations in water/organic solvent system. Enantioselective reduction of aromatic \hat{l}^2 - and \hat{l}^3 -nitroketones. Tetrahedron: Asymmetry, 1998, 9, 1389-1394.	1.8	29
39	Identification of Inhibitors of Drug-Resistant <i>Candida albicans</i> Strains from a Library of Bicyclic Peptidomimetic Compounds. Journal of Medicinal Chemistry, 2010, 53, 2502-2509.	6.4	29
40	Enantiospecific synthesis of 3-aza-6,8-dioxa-bicyclo[3.2.1]octane carboxylic acids from erythrose. Tetrahedron, 2003, 59, 5251-5258.	1.9	28
41	New Evidence of Similarity between Human and Plant Steroid Metabolism: 5α-Reductase Activity in Solanum malacoxylon. Endocrinology, 2003, 144, 220-229.	2.8	28
42	Synthesis, Biological Activity, and Three-Dimensional Quantitative Structureâ^'Activity Relationship Model for a Series of Benzo[c]quinolizin-3-ones, Nonsteroidal Inhibitors of Human Steroid 5α-Reductase 1. Journal of Medicinal Chemistry, 2004, 47, 3546-3560.	6.4	28
43	Baker's yeast reduction of prochiral \hat{I}^3 -nitroketones. II.1 straightforward enantioselective synthesis of 2,7-dimethyl-1,6-dioxaspiro[4.4]nonanes. Tetrahedron: Asymmetry, 1995, 6, 2971-2976.	1.8	27
44	19-Nor-10-azasteroids. 5.1A Synthetic Strategy for the Preparation of (+)-17-(3-Pyridyl)-(5 \hat{l}^2)-10-azaestra-1,16-dien-3-one, a Novel Potential Inhibitor for Human Cytochrome P45017 \hat{l} ±(17 \hat{l} ±-Hydroxylase/C17,20-lyase). Journal of Organic Chemistry, 1999, 64, 4985-4989.	3.2	27
45	Modification of the Aza-Robinson Annulation for the Synthesis of 4-Methyl-Benzo[c]quinolizin-3-ones, Potent Inhibitors of Steroid 5î±-Reductase 1. Journal of Organic Chemistry, 2000, 65, 8093-8095.	3.2	27
46	Pd(0)-Catalyzed Cross-Coupling Reactions of Boron Derivatives with a Lactam-Derived N-Boc Enol Triflate. Organic Letters, 2000, 2, 1241-1242.	4.6	27
47	Skeletal diversity by sequential one-pot and stepwise routes using morpholine ester scaffolds. Tetrahedron Letters, 2010, 51, 6282-6285.	1.4	27
48	19-Nor-10-azasteroids, a New Class of Steroid 5î±-Reductase Inhibitors. 2. X-ray Structure, Molecular Modeling, Conformational Analysis of 19-Nor-10-azasteroids and Comparison with 4-Azasteroids and 6-Azasteroids. Journal of Medicinal Chemistry, 1997, 40, 3466-3477.	6.4	26
49	Enantioselective addition of diethylzinc to aldehydes using 1,4-aminoalcohols as chiral ligands. Tetrahedron: Asymmetry, 2004, 15, 1319-1324.	1.8	26
50	¹²⁵ I-Radiolabeled Morpholine-Containing Arginine–Glycine–Aspartate (RGD) Ligand of α _v β ₃ Integrin As a Molecular Imaging Probe for Angiogenesis. Journal of Medicinal Chemistry, 2012, 55, 5024-5033.	6.4	26
51	Trimethylsilanecarbonitrile Oxide. Synthesis, 1982, 1982, 719-721.	2.3	25
52	Morpholine-based RGD-cyclopentapeptides as $\hat{l}\pm v\hat{l}^23/\hat{l}\pm v\hat{l}^25$ integrin ligands: Role of configuration towards receptor binding affinity. Bioorganic and Medicinal Chemistry, 2009, 17, 1542-1549.	3.0	25
53	Baker's yeast reduction of prochiral \hat{l}^3 -nitroketones: Enantioselective synthesis of (S)-4-nitroalcohols. Tetrahedron, 1995, 51, 1775-1788.	1.9	24
54	A Concise Route to 19-Nor-10-azasteroids, a New Class of Steroid $5\hat{l}_{\pm}$ -Reductase Inhibitors. 3.1 Synthesis of (+)-19-Nor-10-azatestosterone and (+)-17 \hat{l}^2 -(Acetyloxy)-($5\hat{l}^2$)-10-azaestr-1-en-3-one. Journal of Organic Chemistry, 1998, 63, 4111-4115.	3.2	24

#	Article	IF	CITATIONS
55	Synthesis of a new enantiopure bicyclic $^{\hat{j}}$ \hat{l} -amino acid (BTKa) derived from tartaric acid and \hat{l} ±-amino acetophenone. Tetrahedron, 2002, 58, 9865-9870.	1.9	24
56	Modified Adenine (9-Benzyl-2-Butoxy-8-Hydroxyadenine) Redirects Th2-Mediated Murine Lung Inflammation by Triggering TLR7. Journal of Immunology, 2009, 182, 880-889.	0.8	24
57	Novel small molecules for the treatment of infections caused by∢i>Candida albicans∢/i>: a patent review (2002 – 2010). Expert Opinion on Therapeutic Patents, 2011, 21, 381-397.	5.0	24
58	Stereoselective Meisenheimer rearrangement using BTAa's as chiral auxiliaries. Tetrahedron: Asymmetry, 2000, 11, 4227-4238.	1.8	23
59	Effect of C-ring modifications in benzo[c]quinolizin-3-ones, new selective inhibitors of human 5α-reductase 1. Bioorganic and Medicinal Chemistry, 2001, 9, 1385-1393.	3.0	22
60	Synthesis of benzo[c]quinolizin-3-ones: Selective non-steroidal inhibitors of steroid 5α-reductase 1. Bioorganic and Medicinal Chemistry Letters, 1998, 8, 2871-2876.	2.2	21
61	Oligomers of Enantiopure Bicyclic \hat{I}^3/\hat{I} -Amino Acids (BTAa). 1. Synthesis and Conformational Analysis of 3-Aza-6,8-dioxabicyclo [3.2.1] octane-7-carboxylic Acid Oligomers (PolyBTG). Organic Letters, 2000, 2, 3987-3990.	4.6	21
62	Synthesis of bicyclic molecular scaffolds (BTAa): An investigation towards new selective MMP-12 inhibitors. Bioorganic and Medicinal Chemistry, 2006, 14, 7392-7403.	3.0	21
63	Nitrile oxides cycloadditions to cinnamaldehyde. Facile dehydrogenation of 4â€formylâ€4,5â€dihydroisoxazoles. Journal of Heterocyclic Chemistry, 1983, 20, 1505-1507.	2.6	20
64	N-Bridgehead polycyclic compounds by sequential rearrangement-annulation of isoxazoline-5-spirocyclopropanes. 6. A general synthetic method for 5,6-dihydro-7(8H)- and 2,3,5,6-tetrahydro-7(1H)-indolizinones. Journal of Organic Chemistry, 1992, 57, 4206-4211.	3.2	20
65	Novel inhibitors of 5î±-reductase. Expert Opinion on Therapeutic Patents, 2002, 12, 201-215.	5.0	20
66	Selectivity of Daucus carota roots and baker $\hat{a} \in \mathbb{N}$ s yeast in the enantioselective reduction of \hat{l}^3 -nitroketones. Tetrahedron: Asymmetry, 2005, 16, 1479-1483.	1.8	20
67	Parallel Synthesis of an Amide Library Based on the 6,8-Dioxa-3-azabicyclo[3.2.1]octane Scaffold by Direct Aminolysis of Methyl Esters. ACS Combinatorial Science, 2007, 9, 454-461.	3.3	20
68	Evaluation of stereochemically dense morpholine-based scaffolds as proline surrogates in \hat{l}^2 -turn peptides. Organic and Biomolecular Chemistry, 2010, 8, 916-924.	2.8	20
69	New synthesis of azepin-4-ones by flash vacuum thermolysis of dihydro and tetrahydroisoxazole-5-spirocyclobutane derivatives. Tetrahedron Letters, 1986, 27, 5271-5274.	1.4	19
70	Enantioselective synthesis of indolizine derivatives by rearrangement-cyclization of isoxazoline-5-spirocyclopropanes. Tetrahedron, 1993, 49, 10629-10642.	1.9	19
71	Synthesis of 8-chloro-benzo[c]quinolizin-3-ones as potent and selective inhibitors of human steroid 51±-reductase 1. Bioorganic and Medicinal Chemistry Letters, 2000, 10, 353-356.	2.2	19
72	N-Substituent effects on the diethylzinc addition to benzaldehyde catalysed by bicyclic 1,4-amino alcohols. Tetrahedron: Asymmetry, 2009, 20, 340-350.	1.8	19

#	Article	IF	Citations
73	5α-Reductase Inhibitors, Chemical and Clinical Models. Steroids, 1998, 63, 355-361.	1.8	18
74	Introduction of the new dipeptide isostere 7-endo-BtA as reverse turn inducer in a Bowman-Birk proteinase inhibitor. Bioorganic and Medicinal Chemistry, 2001, 9, 1625-1632.	3.0	18
75	A new bicyclic proline-mimetic amino acid. Tetrahedron Letters, 2003, 44, 3489-3492.	1.4	18
76	Peptidomimetics as protein arginine deiminase 4 (PAD4) inhibitors. Journal of Enzyme Inhibition and Medicinal Chemistry, 2015, 30, 466-471.	5.2	18
77	Behaviour of nitrile oxides towards nucleophiles. Part II. Substituent effect on the rate of dimerisation of aromatic nitrile oxides to 3,6-diaryl-1,4,2,5-dioxadiazines. Journal of the Chemical Society Perkin Transactions II, 1976, , 626.	0.9	17
78	5î±-Reductase activity in Lycopersicon esculentum: Cloning and functional characterization of LeDET2 and evidence of the presence of two isoenzymes. Journal of Steroid Biochemistry and Molecular Biology, 2005, 96, 287-299.	2.5	17
79	A Short and Convenient Synthesis of Enantiopure cis- and trans-4-Hydroxypipecolic Acid. Synthesis, 2009, 2009, 3611-3616.	2.3	17
80	Cyclopropane Pipecolic Acids as Templates for Linear and Cyclic Peptidomimetics: Application in the Synthesis of an Argâ€Glyâ€Asp (RGD)â€Containing Peptide as an α _v β ₃ Integrin Ligand Chemistry - A European Journal, 2014, 20, 11187-11203.	. 3.3	17
81	Regioselectivity in the 1,3-dipolar cycloaddition of nitrile oxides to alkylidenecyclopropanes Tetrahedron Letters, 1987, 28, 3845-3848.	1.4	16
82	Synthesis of a new 1,4-aminoalcohol and its use as catalyst in the enantioselective addition of organozinc to aldehydes. Tetrahedron: Asymmetry, 2006, 17, 1409-1414.	1.8	16
83	Stereoselective Synthesis of (2 <i>S</i> ,4 <i>R</i>)â€4â€Hydroxypipecolic Acid. European Journal of Organic Chemistry, 2008, 2008, 524-531.	2.4	16
84	Diastereoselective Synthesis of Highly Constrained Spiroâ€Î²â€Lactams by the Staudinger Reaction Using an Unsymmetrical Bicyclic Ketene. European Journal of Organic Chemistry, 2007, 2007, 4594-4599.	2.4	15
85	Chemical genetics approach to identify new small molecule modulators of cell growth by phenotypic screening of Saccharomyces cerevisiae strains with a library of morpholine-derived compounds. Organic and Biomolecular Chemistry, 2010, 8, 5552.	2.8	15
86	Synthesis and Rearrangement of Cycloadducts from Trimethylsilanecarbonitrile Oxide. Heterocycles, 1983, 20, 511.	0.7	15
87	Configurationally driven folding of model tetrapeptides containing <scp>L</scp> â€or <scp>D</scp> â€morpholineâ€3â€carboxylic acids as βâ€turn nucleators. Chirality, 2009, 21, 584-594.	2.6	14
88	Rearrangement of isoxazoline-5-spiro derivatives. Part 3. Indolizine, quinolizine and pyrido[1,2-a]azepine derivatives by sequential rearrangement-annulation. Journal of the Chemical Society Perkin Transactions 1, 1989, , 1253.	0.9	13
89	A Systems Biology Approach to Dissection of the Effects of Small Bicyclic Peptidomimetics on a Panel of Saccharomyces cerevisiae Mutants. Journal of Biological Chemistry, 2010, 285, 23477-23485.	3.4	13
90	Diastereodivergent Synthesis of 4â€Hydroxyâ€2,3â€methanopipecolic Acid Derivatives as Conformationally Constrained Homoserine Analogues. European Journal of Organic Chemistry, 2011, 2011, 6544-6552.	2.4	13

#	Article	IF	CITATIONS
91	One-pot sequential Ti-/Cu-catalysis for tandem amidation/Ullmann-type cyclization: synthesis of model benzodiazepine(di)ones promoted by microwave irradiation. Organic and Biomolecular Chemistry, 2012, 10, 2780.	2.8	13
92	A novel allergen-adjuvant conjugate suitable for specific immunotherapy of respiratory allergy. Journal of Allergy and Clinical Immunology, 2013, 132, 84-92.e6.	2.9	13
93	A lactam-derived vinyl boronate as a stable and crystalline reagent for the synthesis of 2-substituted piperidines by Pd-catalyzed coupling reactions. Tetrahedron Letters, 2004, 45, 5271-5274.	1.4	12
94	Synthesis and Conformational Analysis of Constrained \hat{l}^2 -Turn Mimetics Incorporating a Bicyclic Turn Inducer by Use of the Petasis Three-Component Reaction on Solid Phase. European Journal of Organic Chemistry, 2007, 2007, 1659-1668.	2.4	12
95	Radiosynthesis and micro-SPECT analysis of triazole-based RGD integrin ligands as non-peptide molecular imaging probes for angiogenesis. Bioorganic and Medicinal Chemistry, 2015, 23, 1112-1122.	3.0	12
96	Simple in situ preparation of fulmimic acid. Tetrahedron Letters, 1983, 24, 1815-1816.	1.4	11
97	Sequential rearrangement-annulation of isoxazoline-5-spirocyclopropanes. Total Synthesis of $(\hat{A}\pm)$ \hat{i} "9(11) -19-Nor-10-Aza-Testosterone Tetrahedron Letters, 1991, 32, 6395-6398.	1.4	11
98	A Short and Efficient Route to Enantiopure 3,5-Diarylpyrrolizidines. Journal of Organic Chemistry, 1999, 64, 1727-1732.	3.2	11
99	Synthesis and activity of 8-substituted benzo[c]quinolizin-3-ones as dual inhibitors of human 51±-reductases 1 and 2. Bioorganic and Medicinal Chemistry Letters, 2005, 15, 145-148.	2.2	11
100	Cyclic DGR-peptidomimetic containing a bicyclic reverse turn inducer as a selective $\hat{l}\pm v\hat{l}^2$ 5 integrin ligand. Amino Acids, 2010, 38, 329-337.	2.7	11
101	Bicyclic peptidomimetics targeting secreted aspartic protease 2 (SAP2) from Candida albicans reveal a constrained inhibitory chemotype. Bioorganic and Medicinal Chemistry, 2012, 20, 7206-7213.	3.0	11
102	d-Proline-based peptidomimetic inhibitors of anthrax lethal factor. European Journal of Medicinal Chemistry, 2012, 56, 96-107.	5.5	11
103	Insight into the structural similarity between HIV protease and secreted aspartic protease-2 and binding mode analysis of HIV- <i>Candida albicans</i> inhibitors. Journal of Enzyme Inhibition and Medicinal Chemistry, 2013, 28, 936-943.	5.2	11
104	Insight to the binding mode of triazole RGD-peptidomimetics to integrin-rich cancer cells by NMR and molecular modeling. Bioorganic and Medicinal Chemistry, 2016, 24, 989-994.	3.0	11
105	Behaviour of nitrile oxides towards nucleophiles. Part IV. Heteromacrocycles from acetonitrile oxide. Journal of the Chemical Society Perkin Transactions 1, 1976, , 1827.	0.9	10
106	Reactions of organomercury fulminates with acetylene derivatives. Journal of Organometallic Chemistry, 1984, 269, 115-121.	1.8	10
107	Stereoselectivity in the TiCl4-catalysed reaction of Danishefsky's diene with a N-(acyloxy)iminium ion: Synthesis of $5\hat{l}^2$ versus $5\hat{l}^2$ \hat{l}^3 1(2)-19-Nor-10-azasteroids. 4. Tetrahedron, 1998, 54, 11589-11596.	1.9	10
108	Synthesis and preliminary biological characterization of a new potential 125I-Radioligand for dopamine and serotonin receptors. Bioorganic and Medicinal Chemistry, 2001, 9, 3197-3206.	3.0	10

7

#	Article	IF	CITATIONS
109	Synthesis of 17β-N-Substituted 19-Nor-10-azasteroids as Inhibitors of Human 5α-Reductases I and II. Bioorganic and Medicinal Chemistry, 2002, 10, 3455-3461.	3.0	10
110	Solvent-Dependent Conformational Behaviour of Model Tetrapeptides Containing a Bicyclic Proline Mimetic. European Journal of Organic Chemistry, 2004, 2004, 4621-4627.	2.4	10
111	Synthesis of a constrained tricyclic scaffold based on trans-4-hydroxy-l-proline. Tetrahedron Letters, 2005, 46, 7813-7816.	1.4	10
112	Synthesis of Glycidol- and Sugar-Derived Bicyclic \hat{l}^2 - and $\hat{l}^3\hat{l}^4$ -Amino Acids for Peptidomimetic Design. European Journal of Organic Chemistry, 2005, 2005, 4372-4381.	2.4	10
113	Identification of constrained peptidomimetic chemotypes as HIV protease inhibitors. European Journal of Medicinal Chemistry, 2014, 84, 444-453.	5 . 5	10
114	Synthesis of enantiopure 2,7-diaryl-1,6-dioxaspiro [4.4] nonanes via enantioselective reduction of prochiral \hat{l}^3 -nitroketones by diisopinocampheylchloroborane (DIP-C1 \hat{a} ,¢). Tetrahedron: Asymmetry, 1996, 7, 1929-1942.	1.8	9
115	Multinuclear magnetic resonance study of organomercury fulminates. The structure of mercury fulminate in solution. Magnetic Resonance in Chemistry, 1984, 22, 372-375.	0.7	8
116	Condensation Product between (R,R)-Tartaric Acid and a L-Phenylalanine Derivative as a New Molecular Scaffold. Archiv Der Pharmazie, 1997, 330, 201-202.	4.1	8
117	Synthesis of new molecular scaffolds: 3-aza-7,9-dioxa-bicyclo[4.2.1]nonane (8-exo BTKa) and 3-aza-8,10-dioxa-bicyclo[5.2.1]decane (9-exo BTKa) carboxylic acids. Tetrahedron, 2004, 60, 2583-2591.	1.9	8
118	Synthesis and conformational studies of a hybrid β-alanine–morpholine tetramer. Tetrahedron, 2012, 68, 9701-9705.	1.9	8
119	A multinuclear magnetic resonance study of nitrile oxides. Journal of Magnetic Resonance, 1982, 50, 64-70.	0.5	7
120	Bicyclic Compounds Derived from Tartaric Acid and \hat{l} ±-Amino Acids (BTAas): Synthesis of New Molecular Scaffolds Derived from the Combination of (R,R)-Tartaric Acid and L-Serine. European Journal of Organic Chemistry, 2002, 2002, 873-880.	2.4	7
121	Evaluation of efficacy, pharmacokinetics and tolerability of peptidomimetic aspartic proteinase inhibitors as cream formulation in experimental vaginal candidiasis. Journal of Pharmacy and Pharmacology, 2014, 66, 1094-1101.	2.4	7
122	A study of ad-proline peptidomimetic inhibitor of melanoma and endothelial cell invasion through activity towards MMP-2 and MMP-9. MedChemComm, 2015, 6, 277-282.	3.4	7
123	3-Aza-6,8-dioxabicyclo[3.2.1]octanes as new enantiopure heteroatom-rich tropane-like ligands of human dopamine transporter. Bioorganic and Medicinal Chemistry, 2006, 14, 5110-5120.	3.0	6
124	Role of Sideâ€Chain Bioisosteres in Determining the Binding Affinity of Click Chemistry Derived RGD Peptidomimetics to α _v β ₃ Integrin. European Journal of Organic Chemistry, 2014, 2014, 7595-7604.	2.4	6
125	Combination of click chemistry and sulfonamides to develop three-armed triazole compounds. Tetrahedron, 2014, 70, 5439-5449.	1.9	6
126	Deciphering the mechanism of action of 089, a compound impairing the fungal cell cycle. Scientific Reports, 2018, 8, 5964.	3.3	6

#	Article	IF	CITATIONS
127	Reactivity in the gas phase. Behaviour of isoxazoles under negative ion chemical ionization conditions. Organic Mass Spectrometry, 1989, 24, 490-496.	1.3	5
128	Chemical genetics approach to drug discovery by diversity-oriented synthesis (DOS) of peptidomimetics. Pure and Applied Chemistry, 2011, 83, 687-698.	1.9	5
129	Synthesis of a chemiluminescent probe useful for the purification of steroid 5α-reductase. Bioorganic and Medicinal Chemistry Letters, 1996, 6, 1997-2002.	2.2	4
130	A solid-phase approach towards the development of 3-aza-6,8-dioxabicyclo[3.2.1]octane scaffolds. Molecular Diversity, 2000, 6, 245-250.	3.9	4
131	Neat reaction of carboxylic acid methyl esters and amines for efficient parallel synthesis of scaffold amide libraries. Comptes Rendus Chimie, 2003, 6, 631-633.	0.5	4
132	Asymmetric hydrogenation of prochiral \hat{I}^3 -nitroketones by ruthenium complexes. Journal of Molecular Catalysis A, 1996, 110, 129-134.	4.8	3
133	Synthesis and Reactivity of Bicycles Derived from Tartaric Acid and α-Amino Acids: A Novel Class of Conformationally Constrained Dipeptide Isosteres Based upon Enantiopure 3-Aza-6,8-dioxabicyclo[3.2.1]octane-7-carboxylic Acid Journal of Organic Chemistry, 2000, 65, 4782-4782.	3.2	3
134	3-Aza-8,10-dioxa-bicyclo[5.2.1]decane (9-exo BTKa) carboxylic acid as a new reverse turn inducer: synthesis and conformational analysis of a model peptide. Tetrahedron, 2006, 62, 1575-1582.	1.9	2
135	One-Pot Pictet-Spengler Reaction and Esterification for the Preparation of a Key Tadalafil Synthetic Intermediate. Letters in Organic Chemistry, 2010, 7, 311-313.	0.5	1
136	New Synthetic Approach to Cyclopenta-Fused Heterocycles Based upon a Mild Nazarov Reaction ChemInform, 2004, 35, no.	0.0	0
137	Synthesis of New Molecular Scaffolds: 3-Aza-7,9-dioxa-bicyclo[4.2.1]nonane (8-exo BTKa) and 3-Aza-8,10-dioxa-bicyclo[5.2.1]decane (9-exo BTKa) Carboxylic Acids ChemInform, 2004, 35, no.	0.0	0
138	Enantioselective Addition of Diethylzinc to Aldehydes Using 1,4-Aminoalcohols as Chiral Ligands ChemInform, 2004, 35, no.	0.0	0
139	A Lactam-Derived Vinyl Boronate as a Stable and Crystalline Reagent for the Synthesis of 2-Substituted Piperidines by Pd-Catalyzed Coupling Reactions ChemInform, 2004, 35, no.	0.0	0
140	New Synthetic Approach to Cyclopenta-Fused Heterocycles Based upon a Mild Nazarov Reaction. Part 2. Further Studies on the Torquoselectivity ChemInform, 2005, 36, no.	0.0	0
141	Synthesis and Activity of 8-Substituted Benzo[c]quinolizin-3-ones (I) as Dual Inhibitors of Human 5?-Reductases 1 and 2 ChemInform, 2005, 36, no.	0.0	0
142	Preparation and Suzuki—Miyaura Coupling Reactions of Tetrahydropyridine-2-boronic Acid Pinacol Esters ChemInform, 2006, 37, no.	0.0	0