Alessandra Tolomelli

List of Publications by Year
in descending order

109	2,415	186265	289244
papers	citations	28 h-index	40 g-index

133	133	133	2244
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Peptides and Peptidomimetics in Medicine, Surgery and Biotechnology. Current Medicinal Chemistry, 2006, 13, 2449-2466.	2.4	148
2	A Straightforward Method for the Synthesis of Alkylidene and Arylidene Malonates Through Proline-Catalyzed Knoevenagel Condensation. Synthetic Communications, 2003, 33, 1587-1594.	2.1	88
3	Green Solvent Mixtures for Solid-Phase Peptide Synthesis: A Dimethylformamide-Free Highly Efficient Synthesis of Pharmaceutical-Grade Peptides. ACS Sustainable Chemistry and Engineering, 2019, 7, 12867-12877.	6.7	69
4	A Stereoselective Synthesis of (2R,3S)-N-Benzoylphenylisoserine Methyl Ester. Journal of Organic Chemistry, 1998, 63, 2351-2353.	3.2	66
5	Therapeutic Peptides Targeting PPI in Clinical Development: Overview, Mechanism of Action and Perspectives. Frontiers in Molecular Biosciences, 2021, 8, 697586.	3.5	64
6	Sustainability in peptide chemistry: current synthesis and purification technologies and future challenges. Green Chemistry, 2022, 24, 975-1020.	9.0	57
7	Recent Advances in the Investigation of the Bioactive Conformation of Peptides Active at the μ-opioid Receptor. Conformational Analysis of Endomorphins. Current Topics in Medicinal Chemistry, 2004, 4, 105-121.	2.1	53
8	Unusual Amino Acids: Synthesis and Introduction into Naturally Occurring Peptides and Biologically Active Analogues. Mini-Reviews in Medicinal Chemistry, 2006, 6, 293-304.	2.4	51
9	Enzymatic Resolution of Î±-Alkyl Î²-Amino Acids Using Immobilized Penicillin G Acylase. Journal of Organic Chemistry, 1996, 61, 8651-8654.	3.2	48
10	Synthesis and Evaluation of the Affinity toward Î¼-Opioid Receptors of Atypical, Lipophilic Ligands Based on the Sequencec[-Tyr-Pro-Trp-Phe-Gly-]. Journal of Medicinal Chemistry, 2004, 47, 5198-5203.	6.4	47
11	Novel Ligands Targeting Î±4Î²1 Integrin: Therapeutic Applications and Perspectives. Frontiers in Chemistry, 2019, 7, 489.	3.6	46
12	Stability against enzymatic hydrolysis of endomorphin-1 analogues containing Î²-proline. Organic and Biomolecular Chemistry, 2003, 1, 1498-1502.	2.8	41
13	Ring expansion of N-acyl aziridine-2-imides to oxazoline-4-imides, useful precursors of pure Î²-Hydroxy Î±-aminoacids. Tetrahedron Letters, 1997, 38, 6953-6956.	1.4	40
14	Highly Diastereoselective Allylic Azide Formation and Isomerization. Synthesis of 3(2â€˜-Amino)-Î²-lactams. Organic Letters, 2005, 7, 533-536.	4.6	40
15	Targeting integrins Î±vÎ²3 and Î±5Î²1 with new Î²-lactam derivatives. European Journal of Medicinal Chemistry, 2014, 83, 284-293.	5.5	40
16	Two-dimensional UV spectroscopy: a new insight into the structure and dynamics of biomolecules. Chemical Science, 2019, 10, 9907-9921.	7.4	40
17	Conjugate addition of hydroxylamino derivatives to alkylidene malonates in the presence of chiral Lewis acids. Tetrahedron: Asymmetry, 2001, 12, 2395-2398.	1.8	37
18	Antinociception by a peripherally administered novel endomorphin-1 analogue containing Î²-proline. European Journal of Pharmacology, 2003, 469, 89-95.	3.5	37

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19	Synthesis of Dehydro-Î²-amino esters via Highly Regioselective Amination of Allylic Carbonates. Organic Letters, 2008, 10, 2425-2428.	4.6	36
20	A Practical Method for the Synthesis of Î²-Amino Î±-Hydroxy Acids. Synthesis of Enantiomerically Pure Hydroxyaspartic Acid and Isoserine. Synlett, 1999, 1999, 1727-1730.	1.8	34
21	Peripheral antinociceptive effects of the cyclic endomorphin-1 analog c[YpwFG] in a mouse visceral pain model. Peptides, 2010, 31, 2135-2140.	2.4	34
22	Molecular Docking of Opiates and Opioid Peptides, a Tool for the Design of Selective Agonists and Antagonists, and for the Investigation of Atypical Ligand-Receptor Interactions. Current Medicinal Chemistry, 2012, 19, 1587-1601.	2.4	31
23	Synthesis of Aziridine-2,2-dicarboxylates via 1,4-Addition ofN,O-(Bistrimethylsilyl)hydroxylamine to Î±,Î²-Unsaturated Malonates. Journal of Organic Chemistry, 2001, 66, 8657-8660.	3.2	29
24	Investigation of the interaction between the atypical agonist c[YpwFG] and MOR. FEBS Journal, 2008, 275, 2315-2337.	4.7	29
25	Antiangiogenic Effect of Dual/Selective Î±₅Î²₁/Î±_vÎ²₃ Integrin Antagonists Designed on Partially Modified Retro-Inverso Cyclotetrapeptide Mimetics. Journal of Medicinal Chemistry, 2010, 53, 106-118.	6.4	29
26	Can Integrin Agonists Have Cards to Play against Cancer? A Literature Survey of Small Molecules Integrin Activators. Cancers, 2017, 9, 78.	3.7	29
27	Asymmetric synthesis of 5-isopropyl-oxazoline-4-imide as syn-hydroxyleucine precursor. Tetrahedron: Asymmetry, 2001, 12, 563-569.	1.8	28
28	Controlled Solid Phase Peptide Bond Formation Using N-Carboxyanhydrides and PEG Resins in Water.. ACS Sustainable Chemistry and Engineering, 2013, 1, 566-569.	6.7	28
29	Spectroscopic fingerprints of DNA/RNA pyrimidine nucleobases in third-order nonlinear electronic spectra. Theoretical Chemistry Accounts, 2016, 135, 1.	1.4	28
30	A New Diastereoselective Synthesis ofanti-Î±-Alkyl Î±-Hydroxy Î²-Amino Acids. European Journal of Organic Chemistry, 1999, 1999, 155-161.	2.4	27
31	Inhibition of cancer cell adhesion by heterochiral Pro-containing RGD mimetics. Bioorganic and Medicinal Chemistry Letters, 2007, 17, 2329-2333.	2.2	27
32	Integrin Ligands with Î±/Î²â€Hybrid Peptide Structure: Design, Bioactivity, and Conformational Aspects. Medicinal Research Reviews, 2016, 36, 389-424.	10.5	27
33	NMR Investigations on Boron Complexes in the Conjugate Addition on Î±,Î²-Unsaturated Imides. Organic Letters, 2001, 3, 1165-1167.	4.6	26
34	Microwave-assisted ring expansion of N-acetyl 3â€²-unsubstituted aziridine in the presence of Lewis acids. Tetrahedron, 2001, 57, 2807-2812.	1.9	26
35	Fast Heckâ€“Cassarâ€“Sonogashira (HCS) Reactions in Green Solvents. Organic Letters, 2020, 22, 3969-3973.	4.6	26
36	A New Selective Synthesis of the Ile-allo-Thr-Gly Tripeptide Fragment of Lysobactin. Organic Letters, 2000, 2, 1105-1107.	4.6	25

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37	The Inverse Typeâ€…II Î²â€Turn on <scp>D</scp>â€Trpâ€Phe, a Pharmacophoric Motif for MOR Agonists. ChemMedChem, 2011, 6, 1640-1653.	3.2	25
38	Diastereoselective synthesis of 3â€²-unsubstituted N-BOC-aziridine from a readily available chiral Î±,Î²-unsaturated imide. Tetrahedron, 1998, 54, 8217-8222.	1.9	23
39	Expedient synthesis of pseudo-Pro-containing peptides: towards constrained peptidomimetics and foldamers. Organic and Biomolecular Chemistry, 2012, 10, 2307.	2.8	23
40	Synthesis of optically pure threonine-containing dipeptides by regio- and stereo-controlled ring expansion of aziridine-2-imide derivatives. Chemical Communications, 1999, , 167-168.	4.1	22
41	Development of Isoxazolineâ€Containing Peptidomimetics as Dual Î±_vÎ²₃ and Î±₅Î²₁ Integrin Ligands. ChemMedChem, 2011, 6, 2264-2272.	3.2	22
42	Formation of Aziridine-2-amides through 5-Halo-6-methylperhydropyrimidin-4-ones. A Route to Enantiopurel- andd-Threonine andallo-Threonine. Journal of Organic Chemistry, 1998, 63, 3458-3462.	3.2	21
43	Asymmetric synthesis of syn hydroxyphenylalanine via aziridine ring expansion to an oxazoline. Tetrahedron Letters, 1999, 40, 8261-8264.	1.4	21
44	Enantioselective synthesis of aziridine 2,2-dicarboxylates. Part I: Copper(II)-bisoxazoline complex-catalysed Michael reaction on alkylidene malonates. Tetrahedron: Asymmetry, 2002, 13, 1407-1410.	1.8	21
45	Dehydro-Î²-proline Containing Î±₄Î²₁ Integrin Antagonists: Stereochemical Recognition in Ligandâ€“Receptor Interplay. ACS Medicinal Chemistry Letters, 2015, 6, 701-706.	2.8	21
46	Steps towards sustainable solid phase peptide synthesis: use and recovery of N-octyl pyrrolidone. Green Chemistry, 2021, 23, 4095-4106.	9.0	21
47	Palladium Catalyst Recycling for Heckâ€Cassarâ€Sonogashira Crossâ€Coupling Reactions in Green Solvent/Base Blend. ChemSusChem, 2021, 14, 2591-2600.	6.8	21
48	Î±-Bromo-Î²,Î³-unsaturated ketenes for the synthesis of Î±-benzylamino-Î²,Î³-unsaturated acids. Tetrahedron: Asymmetry, 2004, 15, 593-601.	1.8	20
49	Synthesis and biological evaluation of non-peptide Î±vÎ²3/Î±5Î²1 integrin dual antagonists containing 5,6-dihydropyridin-2-one scaffolds. Bioorganic and Medicinal Chemistry, 2007, 15, 7380-7390.	3.0	20
50	Synthesis and biological evaluation of unprecedented classes of spiro-Î²-lactams and azido-Î²-lactams as acyl-CoA:cholesterol acyltransferase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2007, 17, 1946-1950.	2.2	20
51	New isoxazolidinone and 3,4-dehydro-Î²-proline derivatives as antibacterial agents and MAO-inhibitors: A complex balance between two activities. European Journal of Medicinal Chemistry, 2016, 124, 906-919.	5.5	20
52	Asymmetric 1,4 addition of Grignard reagents to chiral Î±,Î²-unsaturated esters in the presence of Lewis acids. Tetrahedron, 1999, 55, 6231-6242.	1.9	19
53	A Microwave-Enhanced, Lewis Acid-Catalyzed Synthesis of 1,3-Dioxolanes and Oxazolines from Epoxides. Advanced Synthesis and Catalysis, 2007, 349, 1256-1264.	4.3	19
54	Modulation of Î±vÎ²3- and Î±5Î²1-integrin-mediated adhesion by dehydro-Î²-amino acids containing peptidomimetics. European Journal of Medicinal Chemistry, 2013, 66, 258-268.	5.5	19

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55	An improved microwave assisted protocol for Yonemitsu-type trimolecular condensation. Tetrahedron, 2014, 70, 6781-6788.	1.9	19
56	Synthesis of the Phenylserineâ"Leucine Dipeptide Fragment Present in the Antibiotic Lysobactin from an Aziridine-2-imide Precursor. European Journal of Organic Chemistry, 2000, 2000, 2489-2494.	2.4	18
57	Stable and Biocompatible Monodispersion of C₆₀ in Water by Peptides. Bioconjugate Chemistry, 2019, 30, 808-814.	3.6	18
58	Integrin-mediated adhesive properties of neutrophils are reduced by hyperbaric oxygen therapy in patients with chronic non-healing wound. PLoS ONE, 2020, 15, e0237746.	2.5	18
59	Synthesis of Four-Membered Ring Spiro-Î²-lactams by Epoxide Ring-Opening. European Journal of Organic Chemistry, 2007, 2007, 3199-3205.	2.4	17
60	Opioid Activity Profiles of Oversimplified Peptides Lacking in the Protonable N-Terminus. Journal of Medicinal Chemistry, 2012, 55, 10292-10296.	6.4	17
61	A translation of the twelve principles of green chemistry to guide the development of cross-coupling reactions. Catalysis Today, 2022, 397-399, 265-271.	4.4	17
62	Synthesis of enantiomerically pure syn and anti Î±-hydroxy Î²-amino acids through diastereoselective hydroxylation of perhydropyrimidin-4-ones. Tetrahedron, 1995, 51, 11831-11840.	1.9	16
63	Dipeptides containing D-serine or D-isoserine from the same (R)-aziridine-2-imide by a simple reversal of the synthetic procedure. Tetrahedron, 1999, 55, 15151-15158.	1.9	16
64	Practical synthesis of 3-bromo-5,6-dihydropyridin-2-ones via Î²,Î³-unsaturated Î±-bromo-ketene/imine cycloaddition. Tetrahedron, 2004, 60, 5031-5040.	1.9	16
65	Convenient Synthesis of the Antibiotic Linezolid via an Oxazolidineâ€2,4â€dione Intermediate Derived from the Chiral Building Block Isoserine. European Journal of Organic Chemistry, 2014, 2014, 7614-7620.	2.4	16
66	Synthesis and assay of retro-Î±4Î²1 integrin-targeting motifs. European Journal of Medicinal Chemistry, 2014, 73, 225-232.	5.5	16
67	Synthesis and Biological Evaluation of Azido- and Aziridino-hydroxyl-Î²-lactams through Stereo- and Regioselective Epoxide Ring Opening. Journal of Organic Chemistry, 2006, 71, 9229-9232.	3.2	15
68	Replacing piperidine in solid phase peptide synthesis: effective Fmoc removal by alternative bases. Green Chemistry, 2021, 23, 8096-8107.	9.0	15
69	Synthesis and Conformational Analysis of Cyclotetrapeptide Mimetic Î²â€Turn Templates and Validation as 3D Scaffolds. ChemMedChem, 2009, 4, 517-523.	3.2	13
70	Topological Exploration of Cyclic Endomorphin-1 Analogues, Structurally Defined Models for Investigating the Bioactive Conformation of MOR Agonists. Protein and Peptide Letters, 2007, 14, 51-56.	0.9	12
71	Lewis Acid Induced Highly Regioselective Synthesis of a New Class of Substituted Isoxazolidines. Synlett, 2008, 2008, 2605-2608.	1.8	12
72	A Nonclassical Stereoselective Semi-Synthesis of Drospirenone via Cross-Metathesis Reaction. Synthesis, 2008, 2008, 3801-3804.	2.3	12

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73	A convenient synthesis of functionalized isoxazolines and related 5-hydroxyisoxazolidine-4-carboxylates. Tetrahedron, 2009, 65, 2478-2483.	1.9	12
74	A simple route towards peptide analogues containing substituted (S)- or (R)-tryptophans. Tetrahedron Letters, 2010, 51, 2576-2579.	1.4	12
75	Enantioselective synthesis of aziridine 2,2-dicarboxylates. Part II: Determination of the absolute configuration. Tetrahedron: Asymmetry, 2002, 13, 1411-1415.	1.8	11
76	Enzymatic resolution of ethyl 3-hydroxy-2(1â€²substituted-methylidene)-butyrate by Pseudomonas cepacia lipase catalyzed acetylation. Tetrahedron: Asymmetry, 2007, 18, 2227-2232.	1.8	11
77	Synthesis of Ethyl 5â€Hydroxyisoxazolidineâ€4â€carboxylates via Michael Addition/Intramolecular Hemiketalisation. European Journal of Organic Chemistry, 2008, 2008, 6119-6127.	2.4	11
78	In-peptide synthesis of di-oxazolidinone and dehydroamino acidâ€“oxazolidinone motifs as Î²-turn inducers. Organic and Biomolecular Chemistry, 2013, 11, 4316.	2.8	11
79	Dehydroâ€Î²â€amino Acid Containing Peptides as Promising Sequences for Drug Development. European Journal of Organic Chemistry, 2009, 2009, 5991-5997.	2.4	10
80	Highly Regio- and Diastereoselective Palladium-Catalyzed Allylic Substitution. Synthesis of 3-(2-Aminobutylidene)- 4-arylazetidin-2-ones. Advanced Synthesis and Catalysis, 2005, 347, 833-838.	4.3	9
81	Cyclotetrapeptide Mimics Based on a 13â€Membered, Partially Modified Retroâ€Inverso Structure. European Journal of Organic Chemistry, 2008, 2008, 729-735.	2.4	9
82	A straightforward route to enantiopure 2-substituted-3,4-dehydro-Î²-proline via ring closing metathesis. Amino Acids, 2011, 41, 575-586.	2.7	9
83	Synthesis of Constrained Peptidomimetics Containing 2-Oxo-1,3-oxazolidine-4-carboxylic Acids. European Journal of Organic Chemistry, 2011, 2011, n/a-n/a.	2.4	8
84	Synthesis of Enantiopure Isosteres of Amino Acids Containing a Quaternary Stereocenter: Experimental and Computational Evaluation of a Novel Class of Atropisomers. European Journal of Organic Chemistry, 2018, 2018, 6524-6536.	2.4	8
85	Introduction of hydroxyl- or keto- functionalities in azetidin-2-ones side chain via allylic bromide rearrangement, followed by supported reagent substitution. Arkivoc, 2005, 2005, 136-152.	0.5	8
86	Conformational analysis and Î¼-opioid receptor affinity of short peptides, endomorphin models in a low polarity solvent. Organic and Biomolecular Chemistry, 2003, 1, 3010-3014.	2.8	7
87	Synthesis of enantiomerically pure Î±vÎ²3 integrin ligands based on a 5,6-dihydropyridin-2-one scaffold. Tetrahedron: Asymmetry, 2006, 17, 167-170.	1.8	7
88	Synthesis of chiral non-racemic intermediates and Arg-Gly-Asp mimetics by CaLB-catalyzed resolution. Tetrahedron: Asymmetry, 2010, 21, 96-102.	1.8	7
89	Novel insights into the chemistry of an old medicine: A general degradative pathway for penicillins from a piperacillin/tazobactam stability study. European Journal of Pharmaceutical Sciences, 2019, 136, 104957.	4.0	7
90	A Review of Strategies for the Development of Alkyl Prolines in Drug Discovery. Current Bioactive Compounds, 2016, 12, 146-160.	0.5	7

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91	Exploring the reactivity of alkylidene malonamides: synthesis of polyfunctionalized isoxazolidinones, aziridines and oxazolines.. Arkivoc, 2012, 2012, 196-209.	0.5	7
92	Cyclopeptide Analogs for Generating New Molecular and 3D Diversity. Combinatorial Chemistry and High Throughput Screening, 2009, 12, 929-939.	1.1	6
93	Fluorene benzothiadiazole co-oligomer based aqueous self-assembled nanoparticles. RSC Advances, 2020, 10, 444-450.	3.6	6
94	Aziridines and Oxazolines: Valuable Intermediates in the Synthesis of Unusual Amino Acids. ChemInform, 2004, 35, no.	0.0	5
95	The Cycloaddition Reaction Between Î±â€Bromo Vinylketenes and Imines: A Combined Experimental and Theoretical Study. Advanced Synthesis and Catalysis, 2008, 350, 2261-2273.	4.3	5
96	Oneâ€Pot Twoâ€Step Microwaveâ€Assisted Synthesis of Alkylidene Acetoacetamido Esters, Useful Intermediates for Î²â€Dehydropeptides. European Journal of Organic Chemistry, 2016, 2016, 3217-3222.	2.4	5
97	Highly stable atropisomers by electrophilic amination of a chiral Î³-lactam within the synthesis of an elusive conformationally restricted analogue of Î±-methylhomoserine. Amino Acids, 2016, 48, 461-478.	2.7	5
98	Lipase-mediated kinetic resolution of allylic(hydroxymethyl)methylenecyclopentane building blocks. Tetrahedron: Asymmetry, 2000, 11, 1289-1294.	1.8	4
99	Highly regio- and stereoselective palladium-catalyzed allylic carbonate amination. A practical route to dehydro-Î²-amino esters. Tetrahedron, 2010, 66, 4994-4999.	1.9	4
100	Synthesis of Î±/Î² dipeptides containing linear or cyclic Î±-dehydro-Î²-amino acids as scaffolds for bioactive compounds. Amino Acids, 2019, 51, 1475-1483.	2.7	4
101	Side chain effect in the modulation of Î±vÎ²3/Î±5Î²1 integrin activity via clickable isoxazoline-RGD-mimetics: development of molecular delivery systems. Scientific Reports, 2020, 10, 7410.	3.3	4
102	Ampicillin sodium: Isolation, identification and synthesis of the last unknown impurity after 60 years of clinical use. Journal of Pharmaceutical and Biomedical Analysis, 2020, 191, 113584.	2.8	2
103	Modulation of Î±vÎ²3- and Î±5Î²1-integrin-mediated adhesion by dehydro-Î²-amino acids containing peptidomimetics. , 2013, 66, 258-258.		1
104	Practical Synthesis of 3-Bromo-5,6-dihydropyridin-2-ones via Î²,Î³-Unsaturated Î±-Bromo-ketene/imine Cycloaddition.. ChemInform, 2004, 35, no.	0.0	0
105	Highly Diastereoselective Allylic Azide Formation and Isomerization. Synthesis of 3(2â€²-Amino)-Î²-lactams.. ChemInform, 2005, 36, no.	0.0	0
106	Zinc Metal-Promoted Nucleophilic Addition of Azetidin-2-ones to Aldehydes and Nitriles. Synthesis, 2005, 2005, 61-70.	2.3	0
107	An Investigation of the Reactivity of MCPBA and Î±-Bromoalkenes under ÂTraditional or Microwave-Assisted Conditions: Selective Formation of ÂEpoxides or Allylic Bromides. Synlett, 2005, 2005, 2204-2208.	1.8	0
108	A Novel Family of Minimal PMRI Cyclic Peptides as Versatile Scaffolds for Generating New Molecular Topology. Medicinal Chemistry, 2006, 2, 395-400.	1.5	0

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109	Asymmetric Synthesis of Three- and Four-Membered Ring Heterocycles. , 0, , 1-50.		0