Alessandra Tolomelli

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/9193114/publications.pdf

Version: 2024-02-01

133

all docs

109 2,415 28
papers citations h-index

citations h-index g-index

133 133 2244
docs citations times ranked citing authors

289244

40

#	Article	IF	CITATIONS
1	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
2	Sustainability in peptide chemistry: current synthesis and purification technologies and future challenges. Green Chemistry, 2022, 24, 975-1020.	9.0	57
3	Steps towards sustainable solid phase peptide synthesis: use and recovery of <i>N</i> -octyl pyrrolidone. Green Chemistry, 2021, 23, 4095-4106.	9.0	21
4	Replacing piperidine in solid phase peptide synthesis: effective Fmoc removal by alternative bases. Green Chemistry, 2021, 23, 8096-8107.	9.0	15
5	Palladium Catalyst Recycling for Heckâ€Cassarâ€Sonogashira Crossâ€Coupling Reactions in Green Solvent/Base Blend. ChemSusChem, 2021, 14, 2591-2600.	6.8	21
6	Therapeutic Peptides Targeting PPI in Clinical Development: Overview, Mechanism of Action and Perspectives. Frontiers in Molecular Biosciences, 2021, 8, 697586.	3.5	64
7	Fluorene benzothiadiazole co-oligomer based aqueous self-assembled nanoparticles. RSC Advances, 2020, 10, 444-450.	3.6	6
8	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
9	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
10	Side chain effect in the modulation of $\hat{l}\pm v\hat{l}^23/\hat{l}\pm 5\hat{l}^21$ integrin activity via clickable isoxazoline-RGD-mimetics: development of molecular delivery systems. Scientific Reports, 2020, 10, 7410.	3.3	4
11	Fast Heck–Cassar–Sonogashira (HCS) Reactions in Green Solvents. Organic Letters, 2020, 22, 3969-3973.	4.6	26
12	Novel Ligands Targeting $\hat{l}\pm4\hat{l}^21$ Integrin: Therapeutic Applications and Perspectives. Frontiers in Chemistry, 2019, 7, 489.	3.6	46
13	Synthesis of $\hat{l}\pm /\hat{l}^2$ dipeptides containing linear or cyclic $\hat{l}\pm$ -dehydro- \hat{l}^2 -amino acids as scaffolds for bioactive compounds. Amino Acids, 2019, 51, 1475-1483.	2.7	4
14	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
15	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
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	Stable and Biocompatible Monodispersion of C ₆₀ in Water by Peptides. Bioconjugate Chemistry, 2019, 30, 808-814.	3.6	18
18	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

#	Article	IF	CITATIONS
19	Can Integrin Agonists Have Cards to Play against Cancer? A Literature Survey of Small Molecules Integrin Activators. Cancers, 2017, 9, 78.	3.7	29
20	Integrin Ligands with α/βâ€Hybrid Peptide Structure: Design, Bioactivity, and Conformational Aspects. Medicinal Research Reviews, 2016, 36, 389-424.	10.5	27
21	Oneâ€Pot Twoâ€Step Microwaveâ€Assisted Synthesis of Alkylidene Acetoacetamido Esters, Useful Intermediates for βâ€Dehydropeptides. European Journal of Organic Chemistry, 2016, 2016, 3217-3222.	2.4	5
22	Spectroscopic fingerprints of DNA/RNA pyrimidine nucleobases in third-order nonlinear electronic spectra. Theoretical Chemistry Accounts, 2016, 135, 1.	1.4	28
23	New isoxazolidinone and 3,4-dehydro- \hat{l}^2 -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
24	Highly stable atropisomers by electrophilic amination of a chiral \hat{I}^3 -lactam within the synthesis of an elusive conformationally restricted analogue of \hat{I}^{\pm} -methylhomoserine. Amino Acids, 2016, 48, 461-478.	2.7	5
25	A Review of Strategies for the Development of Alkyl Prolines in Drug Discovery. Current Bioactive Compounds, 2016, 12, 146-160.	0.5	7
26	Dehydro-β-proline Containing α ₄ β ₁ Integrin Antagonists: Stereochemical Recognition in Ligand–Receptor Interplay. ACS Medicinal Chemistry Letters, 2015, 6, 701-706.	2.8	21
27	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
28	Targeting integrins $\hat{l}\pm\hat{vl}^2$ 3 and $\hat{l}\pm5\hat{l}^2$ 1 with new \hat{l}^2 -lactam derivatives. European Journal of Medicinal Chemistry, 2014, 83, 284-293.	5.5	40
29	An improved microwave assisted protocol for Yonemitsu-type trimolecular condensation. Tetrahedron, 2014, 70, 6781-6788.	1.9	19
30	Synthesis and assay of retro- $\hat{l}\pm4\hat{l}^21$ integrin-targeting motifs. European Journal of Medicinal Chemistry, 2014, 73, 225-232.	5.5	16
31	Controlled Solid Phase Peptide Bond Formation Using $\langle i \rangle N \langle j \rangle$ -Carboxyanhydrides and PEG Resins in Water ACS Sustainable Chemistry and Engineering, 2013, 1, 566-569.	6.7	28
32	In-peptide synthesis of di-oxazolidinone and dehydroamino acid–oxazolidinone motifs as β-turn inducers. Organic and Biomolecular Chemistry, 2013, 11, 4316.	2.8	11
33	Modulation of $\hat{l}\pm\hat{vl}^2$ 3- and $\hat{l}\pm\hat{5l}^2$ 1-integrin-mediated adhesion by dehydro- \hat{l}^2 -amino acids containing peptidomimetics. European Journal of Medicinal Chemistry, 2013, 66, 258-268.	5.5	19
34	Modulation of $\hat{l}\pm v\hat{l}^2$ 3- and $\hat{l}\pm 5\hat{l}^2$ 1-integrin-mediated adhesion by dehydro- \hat{l}^2 -amino acids containing peptidomimetics. , 2013, 66, 258-258.		1
35	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
36	Opioid Activity Profiles of Oversimplified Peptides Lacking in the Protonable N-Terminus. Journal of Medicinal Chemistry, 2012, 55, 10292-10296.	6.4	17

#	Article	IF	Citations
37	Expedient synthesis of pseudo-Pro-containing peptides: towards constrained peptidomimetics and foldamers. Organic and Biomolecular Chemistry, 2012, 10, 2307.	2.8	23
38	Exploring the reactivity of alkylidene malonamides: synthesis of polyfunctionalized isoxazolidinones, aziridines and oxazolines Arkivoc, 2012, 2012, 196-209.	0.5	7
39	A straightforward route to enantiopure 2-substituted-3,4-dehydro- \hat{l}^2 -proline via ring closing metathesis. Amino Acids, 2011, 41, 575-586.	2.7	9
40	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
41	The Inverse Type Il βâ€Turn on <scp>D</scp> â€Trpâ€Phe, a Pharmacophoric Motif for MOR Agonists. ChemMedChem, 2011, 6, 1640-1653.	3.2	25
42	Development of Isoxazolineâ€Containing Peptidomimetics as Dual α _v β ₃ and α ₅ β ₁ Integrin Ligands. ChemMedChem, 2011, 6, 2264-2272.	3.2	22
43	Highly regio- and stereoselective palladium-catalyzed allylic carbonate amination. A practical route to dehydro-l ² -amino esters. Tetrahedron, 2010, 66, 4994-4999.	1.9	4
44	Synthesis of chiral non-racemic intermediates and Arg-Gly-Asp mimetics by CaLB-catalyzed resolution. Tetrahedron: Asymmetry, 2010, 21, 96-102.	1.8	7
45	A simple route towards peptide analogues containing substituted (S)- or (R)-tryptophans. Tetrahedron Letters, 2010, 51, 2576-2579.	1.4	12
46	Antiangiogenic Effect of Dual/Selective $\hat{l}\pm < sub>5 < / sub>\hat{l}^2 < sub>1 < / sub>/\hat{l}\pm < sub>v < / sub>\hat{l}^2 < sub>3 < / sub> Integrin Antagonists Designed on Partially Modified Retro-Inverso Cyclotetrapeptide Mimetics. Journal of Medicinal Chemistry, 2010, 53, 106-118.$	6.4	29
47	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
48	Synthesis and Conformational Analysis of Cyclotetrapeptide Mimetic βâ€∓urn Templates and Validation as 3D Scaffolds. ChemMedChem, 2009, 4, 517-523.	3.2	13
49	Dehydroâ€Î²â€amino Acid Containing Peptides as Promising Sequences for Drug Development. European Journal of Organic Chemistry, 2009, 2009, 5991-5997.	2.4	10
50	A convenient synthesis of functionalized isoxazolines and related 5-hydroxyisoxazolidine-4-carboxylates. Tetrahedron, 2009, 65, 2478-2483.	1.9	12
51	Cyclopeptide Analogs for Generating New Molecular and 3D Diversity. Combinatorial Chemistry and High Throughput Screening, 2009, 12, 929-939.	1.1	6
52	Cyclotetrapeptide Mimics Based on a 13â€Membered, Partially Modified Retroâ€Inverso Structure. European Journal of Organic Chemistry, 2008, 2008, 729-735.	2.4	9
53	Synthesis of Ethyl 5â€Hydroxyisoxazolidineâ€4â€carboxylates via Michael Addition/Intramolecular Hemiketalisation. European Journal of Organic Chemistry, 2008, 2008, 6119-6127.	2.4	11
54	The Cycloaddition Reaction Between αâ€Bromo Vinylketenes and Imines: A Combined Experimental and Theoretical Study. Advanced Synthesis and Catalysis, 2008, 350, 2261-2273.	4.3	5

#	Article	IF	CITATIONS
55	Investigation of the interaction between the atypical agonist c[YpwFG] and MOR. FEBS Journal, 2008, 275, 2315-2337.	4.7	29
56	Synthesis of Dehydro-β-amino esters <i>via</i> Highly Regioselective Amination of Allylic Carbonates. Organic Letters, 2008, 10, 2425-2428.	4.6	36
57	Lewis Acid Induced Highly Regioselective Synthesis of a New Class of Substituted Isoxazolidines. Synlett, 2008, 2008, 2605-2608.	1.8	12
58	A Nonclassical Stereoselective Semi-Synthesis of Drospirenone via Cross-Metathesis Reaction. Synthesis, 2008, 2008, 3801-3804.	2.3	12
59	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
60	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
61	Synthesis of Four-Membered Ring Spiro-β-lactams by Epoxide Ring-Opening. European Journal of Organic Chemistry, 2007, 2007, 3199-3205.	2.4	17
62	Enzymatic resolution of ethyl 3-hydroxy-2(1′substituted-methylidene)-butyrate by Pseudomonas cepacia lipase catalyzed acetylation. Tetrahedron: Asymmetry, 2007, 18, 2227-2232.	1.8	11
63	Synthesis and biological evaluation of non-peptide $\hat{l}\pm v\hat{l}^23/\hat{l}\pm 5\hat{l}^21$ integrin dual antagonists containing 5,6-dihydropyridin-2-one scaffolds. Bioorganic and Medicinal Chemistry, 2007, 15, 7380-7390.	3.0	20
64	Synthesis and biological evaluation of unprecedented classes of spiro- \hat{l}^2 -lactams and azido- \hat{l}^2 -lactams as acyl-CoA:cholesterol acyltransferase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2007, 17, 1946-1950.	2.2	20
65	Inhibition of cancer cell adhesion by heterochiral Pro-containing RGD mimetics. Bioorganic and Medicinal Chemistry Letters, 2007, 17, 2329-2333.	2.2	27
66	Synthesis and Biological Evaluation of Azido- and Aziridino-hydroxyl- \hat{l}^2 -lactams through Stereo- and Regioselective Epoxide Ring Opening. Journal of Organic Chemistry, 2006, 71, 9229-9232.	3.2	15
67	Synthesis of enantiomerically pure $\hat{l}\pm v\hat{l}^23$ integrin ligands based on a 5,6-dihydropyridin-2-one scaffold. Tetrahedron: Asymmetry, 2006, 17, 167-170.	1.8	7
68	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
69	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
70	Peptides and Peptidomimetics in Medicine, Surgery and Biotechnology. Current Medicinal Chemistry, 2006, 13, 2449-2466.	2.4	148
71	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
72	Highly Diastereoselective Allylic Azide Formation and Isomerization. Synthesis of 3(2′-Amino)-β-lactams ChemInform, 2005, 36, no.	0.0	0

#	Article	IF	CITATIONS
73	Zinc Metal-Promoted Nucleophilic Addition of Azetidin-2-ones to Aldehydes and Nitriles. Synthesis, 2005, 2005, 61-70.	2.3	O
74	An Investigation of the Reactivity of MCPBA and α-Bromoalkenes under ÂTraditional or Microwave-Assisted Conditions: Selective Formation of ÂEpoxides or Allylic Bromides. Synlett, 2005, 2005, 2204-2208.	1.8	0
75	Highly Diastereoselective Allylic Azide Formation and Isomerization. Synthesis of 3(2â€~-Amino)-β-lactams. Organic Letters, 2005, 7, 533-536.	4.6	40
76	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
77	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
78	Practical synthesis of 3-bromo-5,6-dihydropyridin-2-ones via \hat{l}^2 , \hat{l}^3 -unsaturated $\hat{l}\pm$ -bromo-ketene/imine cycloaddition. Tetrahedron, 2004, 60, 5031-5040.	1.9	16
79	\hat{l}_{\pm} -Bromo- \hat{l}_{-}^2 , \hat{l}_{-}^3 -unsaturated ketenes for the synthesis of \hat{l}_{\pm} -benzylamino- \hat{l}_{-}^2 , \hat{l}_{-}^3 -unsaturated acids. Tetrahedron: Asymmetry, 2004, 15, 593-601.	1.8	20
80	Aziridines and Oxazolines: Valuable Intermediates in the Synthesis of Unusual Amino Acids. ChemInform, 2004, 35, no.	0.0	5
81	Practical Synthesis of 3-Bromo-5,6-dihydropyridin-2-ones via \hat{l}^2 , \hat{l}^3 -Unsaturated $\hat{l}\pm$ -Bromo-ketene/imine Cycloaddition ChemInform, 2004, 35, no.	0.0	0
82	Synthesis and Evaluation of the Affinity toward $\hat{l}\frac{1}{4}$ -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
83	Antinociception by a peripherally administered novel endomorphin-1 analogue containing \hat{l}^2 -proline. European Journal of Pharmacology, 2003, 469, 89-95.	3.5	37
84	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
85	Conformational analysis and $\hat{l}\frac{1}{4}$ -opioid receptor affinity of short peptides, endomorphin models in a low polarity solvent. Organic and Biomolecular Chemistry, 2003, 1, 3010-3014.	2.8	7
86	Stability against enzymatic hydrolysis of endomorphin-1 analogues containing \hat{l}^2 -proline. Organic and Biomolecular Chemistry, 2003, 1, 1498-1502.	2.8	41
87	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
88	Enantioselective synthesis of aziridine 2,2-dicarboxylates. Part II: Determination of the absolute configuration. Tetrahedron: Asymmetry, 2002, 13, 1411-1415.	1.8	11
89	Synthesis of Aziridine-2,2-dicarboxylates via 1,4-Addition of N,O-(Bistrimethylsilyl) hydroxylamine to $\hat{1}\pm,\hat{1}^2$ -Unsaturated Malonates. Journal of Organic Chemistry, 2001, 66, 8657-8660.	3.2	29
90	NMR Investigations on Boron Complexes in the Conjugate Addition on \hat{l}_{\pm}, \hat{l}^2 -Unsaturated Imides. Organic Letters, 2001, 3, 1165-1167.	4.6	26

#	Article	IF	Citations
91	Asymmetric synthesis of 5-isopropyl-oxazoline-4-imide as syn-hydroxyleucine precursor. Tetrahedron: Asymmetry, 2001, 12, 563-569.	1.8	28
92	Conjugate addition of hydroxylamino derivatives to alkylidene malonates in the presence of chiral Lewis acids. Tetrahedron: Asymmetry, 2001, 12, 2395-2398.	1.8	37
93	Microwave-assisted ring expansion of N-acetyl 3′-unsubstituted aziridine in the presence of Lewis acids. Tetrahedron, 2001, 57, 2807-2812.	1.9	26
94	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
95	Lipase-mediated kinetic resolution of allylic(hydroxymethyl)methylenecyclopentane building blocks. Tetrahedron: Asymmetry, 2000, 11, 1289-1294.	1.8	4
96	A New Selective Synthesis of the Ile-allo-Thr-Gly Tripeptide Fragment of Lysobactin. Organic Letters, 2000, 2, 1105-1107.	4.6	25
97	A Practical Method for the Synthesis of \hat{l}^2 -Amino \hat{l}_\pm -Hydroxy Acids. Synthesis of Enantiomerically Pure Hydroxyaspartic Acid and Isoserine. Synlett, 1999, 1999, 1727-1730.	1.8	34
98	Asymmetric synthesis of syn hydroxyphenylalanine via aziridine ring expansion to an oxazoline. Tetrahedron Letters, 1999, 40, 8261-8264.	1.4	21
99	Asymmetric 1,4 addition of Grignard reagents to chiral $\hat{l}\pm,\hat{l}^2$ -unsaturated esters in the presence of Lewis acids. Tetrahedron, 1999, 55, 6231-6242.	1.9	19
100	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
101	A New Diastereoselective Synthesis ofanti-α-Alkyl α-Hydroxy β-Amino Acids. European Journal of Organic Chemistry, 1999, 1999, 155-161.	2.4	27
102	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
103	Diastereoselective synthesis of $3\hat{a}\in^2$ -unsubstituted N-BOC-aziridine from a readily available chiral $\hat{l}\pm,\hat{l}^2$ -unsaturated imide. Tetrahedron, 1998, 54, 8217-8222.	1.9	23
104	A Stereoselective Synthesis of (2R,3S)-N-Benzoylphenylisoserine Methyl Ester. Journal of Organic Chemistry, 1998, 63, 2351-2353.	3.2	66
105	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
106	Ring expansion of N-acyl aziridine-2-imides to oxazoline-4-imides, useful precursors of pure \hat{l}^2 -Hydroxy \hat{l}_2 -aminoacids. Tetrahedron Letters, 1997, 38, 6953-6956.	1.4	40
107	Enzymatic Resolution of α-Alkyl β-Amino Acids Using Immobilized Penicillin G Acylase. Journal of Organic Chemistry, 1996, 61, 8651-8654.	3.2	48
108	Synthesis of enantiomerically pure syn and anti \hat{l}_{\pm} -hydroxy \hat{l}^2 -amino acids through diastereoselective hydroxylation of perhydropyrimidin-4-ones. Tetrahedron, 1995, 51, 11831-11840.	1.9	16

ARTICLE IF CITATIONS

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