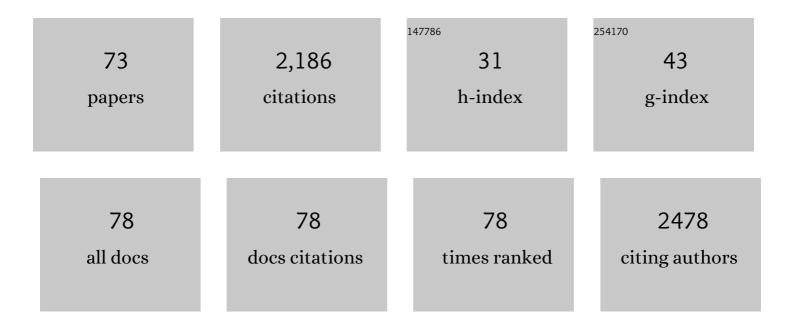
Leonardo Manzoni

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	N-Trichloroethoxycarbonyl-glucosamine derivatives as glycosyl donors. Carbohydrate Research, 1996, 296, 135-147.	2.3	153
2	Cyclic RGD Functionalized Gold Nanoparticles for Tumor Targeting. Bioconjugate Chemistry, 2011, 22, 664-672.	3.6	82
3	Biological and molecular properties of a new αvβ3/αvβ5 integrin antagonist. Molecular Cancer Therapeutics, 2005, 4, 1670-1680.	4.1	75
4	Novel SMAC-mimetics synergistically stimulate melanoma cell death in combination with TRAIL and Bortezomib. British Journal of Cancer, 2010, 102, 1707-1716.	6.4	70
5	Regulation of HuR structure and function by dihydrotanshinone-I. Nucleic Acids Research, 2017, 45, 9514-9527.	14.5	64
6	Targeting integrins: Insights into structure and activity of cyclic RGD pentapeptide mimics containing azabicycloalkane amino acids. Bioorganic and Medicinal Chemistry, 2006, 14, 169-180.	3.0	61
7	Cyclic RGDâ€Peptidomimetics Containing Bifunctional Diketopiperazine Scaffolds as New Potent Integrin Ligands. Chemistry - A European Journal, 2009, 15, 12184-12188.	3.3	58
8	Synthesis of Gd and ⁶⁸ Ga Complexes in Conjugation with a Conformationally Optimized RGD Sequence as Potential MRI and PET Tumorâ€Imaging Probes. ChemMedChem, 2012, 7, 1084-1093.	3.2	53
9	Designing Smac-mimetics as antagonists of XIAP, cIAP1, and cIAP2. Biochemical and Biophysical Research Communications, 2009, 378, 162-167.	2.1	50
10	Potent Integrin Antagonists from a Small Library of RGD-Including Cyclic Pseudopeptides. Organic Letters, 2001, 3, 1001-1004.	4.6	49
11	Conformationally constrained dipeptides: Synthesis of 7,5- and 6,5-fused bicyclic lactams by stereoselective radical cyclizations. Tetrahedron Letters, 1995, 36, 625-628.	1.4	46
12	Rapid synthesis of oligosaccharides using an anomeric fluorous silyl protecting groupElectronic supplementary information (ESI) available: experimental data. See http://www.rsc.org/suppdata/cc/b3/b311448a/. Chemical Communications, 2003, , 2930.	4.1	45
13	Cyclic RGDâ€Containing Functionalized Azabicycloalkane Peptides as Potent Integrin Antagonists for Tumor Targeting. ChemMedChem, 2009, 4, 615-632.	3.2	44
14	Conformational Analysis of Azabicycloalkane Amino Acid Scaffolds as Reverse-Turn Inducer Dipeptide Mimics. European Journal of Organic Chemistry, 2000, 2000, 2563-2569.	2.4	43
15	Froc:  A New Fluorous Protective Group for Peptide and Oligosaccharide Synthesisâ€. Organic Letters, 2006, 8, 955-957.	4.6	42
16	Stereoselective Synthesis of Conformationally Constrained Cyclohexanediols:  A Set of Molecular Scaffolds for the Synthesis of Glycomimetics. Journal of Organic Chemistry, 2001, 66, 6209-6216.	3.2	41
17	Design, Synthesis, and Biological Evaluation of Novel cRGD–Paclitaxel Conjugates for Integrin-Assisted Drug Delivery. Bioconjugate Chemistry, 2012, 23, 1610-1622.	3.6	41
18	Synthesis of new bicyclic lactam peptidomimetics by ring-closing metathesis reactions. Tetrahedron, 2003, 59, 4501-4513.	1.9	40

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#	Article	IF	CITATIONS
19	Targeting the X-Linked Inhibitor of Apoptosis Protein through 4-Substituted Azabicyclo[5.3.0]alkane Smac Mimetics. Structure, Activity, and Recognition Principles. Journal of Molecular Biology, 2008, 384, 673-689.	4.2	40
20	Structural Basis for Bivalent Smac-Mimetics Recognition in the IAP Protein Family. Journal of Molecular Biology, 2009, 392, 630-644.	4.2	40
21	Synthesis of N -acetylglucosamine containing Lewis A and Lewis X building blocks based on N -tetrachlorophthaloyl protection—synthesis of Lewis X pentasaccharide. Carbohydrate Research, 1998, 310, 157-171.	2.3	39
22	Practical stereoselective synthesis of conformationally constrained unnatural proline-based amino acids and peptidomimetics. Tetrahedron, 2001, 57, 6463-6473.	1.9	39
23	Functionalized Azabicycloalkane Amino Acids by Nitrone 1,3-Dipolar Intramolecular Cycloaddition. Journal of Organic Chemistry, 2005, 70, 4124-4132.	3.2	39
24	A new optical imaging probe targeting <i>α</i> _V <i>β</i> ₃ integrin in glioblastoma xenografts. Contrast Media and Molecular Imaging, 2011, 6, 449-458.	0.8	39
25	Interfering with HuR–RNA Interaction: Design, Synthesis and Biological Characterization of Tanshinone Mimics as Novel, Effective HuR Inhibitors. Journal of Medicinal Chemistry, 2018, 61, 1483-1498.	6.4	39
26	MicroPET/CT imaging of αvl ² 3 integrin via a novel 68Ga-NOTA-RGD peptidomimetic conjugate in rat myocardial infarction. European Journal of Nuclear Medicine and Molecular Imaging, 2013, 40, 1265-1274.	6.4	38
27	Rational design, synthesis and characterization of potent, non-peptidic Smac mimics/XIAP inhibitors as proapoptotic agents for cancer therapy. Bioorganic and Medicinal Chemistry, 2009, 17, 5834-5856.	3.0	36
28	Synthesis of the Lewis a Trisaccharide Based on an Anomeric Silyl Fluorous Tag. Organic Letters, 2004, 6, 4195-4198.	4.6	35
29	Synthesis of 7,5-fused bicyclic lactams by stereoselective radical cyclization. Tetrahedron Letters, 1994, 35, 4031-4034.	1.4	34
30	A Practical Way to 2,5-Disubstituted Pyrrolidine Derivatives. Synlett, 1996, 1996, 441-443.	1.8	34
31	The first asymmetric synthesis of enantiopure .alphasulfenyl dithioacetals and .alphasulfenyl aldehydes. Journal of Organic Chemistry, 1993, 58, 3165-3168.	3.2	33
32	Integrin-Targeted Peptide- and Peptidomimetic-Drug Conjugates for the Treatment of Tumors. Recent Patents on Anti-Cancer Drug Discovery, 2017, 12, 148-168.	1.6	33
33	Mimics of ganglioside GM1 as cholera toxin ligands: replacement of the GalNAc residueElectronic supplementary information (ESI) available: synthetic details, product characterisations and full NOE contact list. See http://www.rsc.org/suppdata/ob/b2/b210503a/. Organic and Biomolecular Chemistry, 2003. 1. 785-792.	2.8	31
34	Novel second mitochondria-derived activator of caspases (Smac) mimetic compounds sensitize human leukemic cell lines to conventional chemotherapeutic drug-induced and death receptor-mediated apoptosis. Investigational New Drugs, 2011, 29, 1264-1275.	2.6	31
35	Dimeric Smac mimetics/IAP inhibitors as in vivo-active pro-apoptotic agents. Part II: Structural and biological characterization. Bioorganic and Medicinal Chemistry, 2012, 20, 6709-6723.	3.0	29
36	Synthesis of Lewis a and Lewis X Pentasaccharides Based on N-Trichloroethoxycarbonyl Protection. Journal of Carbohydrate Chemistry, 1998, 17, 739-758.	1.1	28

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37	Synthesis of Azabicycloalkane Amino Acid Scaffolds as Reverse-Turn Inducer Dipeptide Mimics. European Journal of Organic Chemistry, 2000, 2000, 2571-2581.	2.4	27
38	A Potent Integrin Antagonist from a Small Library of Cyclic RGD Pentapeptide Mimics Including Benzyl‧ubstituted Azabicycloalkane Amino Acids. ChemMedChem, 2008, 3, 1589-1603.	3.2	27
39	Diastereoselective addition of metal-coordinated and â€~naked' tri-sec-butylborohydrides to a norephedrine-derived 2-acetyloxazolidine. Journal of the Chemical Society Chemical Communications, 1992, , 1027-1029.	2.0	26
40	Functionalized Cyclic RGD Peptidomimetics: Conjugable ligands for α _v l² ₃ Receptor Imaging. Bioconjugate Chemistry, 2009, 20, 1611-1617.	3.6	26
41	The first example of ring-closing olefin metathesis of dehydroamino acids: an application to the synthesis of azabicyclo[X.Y.0]alkanes. Tetrahedron Letters, 2004, 45, 2623-2625.	1.4	25
42	Enhancement of the Uptake and Cytotoxic Activity of Doxorubicin in Cancer Cells by Novel cRGD-Semipeptide-Anchoring Liposomes. Molecular Pharmaceutics, 2014, 11, 2280-2293.	4.6	25
43	Synthesis of substituted conformationally constrained 6,5- and 7,5-fused bicyclic lactams as dipeptide mimics. Tetrahedron, 2003, 59, 6241-6250.	1.9	22
44	Design, synthesis and biological evaluation of novel dimeric and tetrameric cRGD–paclitaxel conjugates for integrin-assisted drug delivery. Organic and Biomolecular Chemistry, 2015, 13, 7530-7541.	2.8	22
45	Click chemistry to functionalise peptidomimetics. Tetrahedron Letters, 2006, 47, 3697-3700.	1.4	20
46	Homo- and heterodimeric Smac mimetics/IAP inhibitors as in vivo-active pro-apoptotic agents. Part I: Synthesis. Bioorganic and Medicinal Chemistry, 2012, 20, 6687-6708.	3.0	20
47	Synthesis of spiroazabicycloalkane amino acid scaffolds as reverse-turn inducer dipeptide mimics. Tetrahedron, 2001, 57, 249-255.	1.9	19
48	Synthesis and biological evaluation of dual action <i>cyclo</i> -RGD/SMAC mimetic conjugates targeting α _v β ₃ /α _v β ₅ integrins and IAP proteins. Organic and Biomolecular Chemistry, 2014, 12, 3288-3302.	2.8	19
49	Cyclic RGD Peptides Containing Azabicycloalkane Reverse-Turn Mimics. Helvetica Chimica Acta, 2002, 85, 4353-4368.	1.6	18
50	Synthesis of some oligopyridine–galactose conjugates and their metal complexes: a simple entry to multivalent sugar ligands. Tetrahedron, 2005, 61, 10048-10060.	1.9	18
51	4-Aminoproline-based arginine-glycine-aspartate integrin binders with exposed ligation points: practical in-solution synthesis, conjugation and binding affinity evaluation. Organic and Biomolecular Chemistry, 2009, 7, 4924.	2.8	18
52	Investigating the Interaction of Cyclic RGD Peptidomimetics with $\hat{I}\pm V\hat{I}^26$ Integrin by Biochemical and Molecular Docking Studies. Cancers, 2017, 9, 128.	3.7	18
53	Solid-Phase Synthesis of Peptides Containing Reverse-Turn Mimetic Bicyclic Lactams. , 1999, 1999, 379-388.		17
54	Computational design of novel peptidomimetic inhibitors of cadherin homophilic interactions. Organic and Biomolecular Chemistry, 2015, 13, 2570-2573.	2.8	16

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55	Iron Oxide-Gold Core-Shell Nanoparticles as Multimodal Imaging Contrast Agent. IEEE Sensors Journal, 2013, 13, 2341-2347.	4.7	15
56	Stereoselective synthesis of 6,5-bicyclic reverse-turn peptidomimetics. Tetrahedron, 1998, 54, 5325-5336.	1.9	14
57	Diastereoselective addition of metal-coordinated and "naked―nucleophilic reagents to norephedrine derived 2-acyl-N-tosyl-oxazolidines. Tetrahedron, 1997, 53, 1759-1776.	1.9	12
58	Synthesis of a Pseudo Tetrasaccharide Mimic of Ganglioside GM1. European Journal of Organic Chemistry, 1999, 1999, 1311-1317.	2.4	12
59	Diastereoselective Addition of Organometallic Reagents to Nor-Ephedrine-Derived 2-Acyl-N-Tosyl-Oxazolidines. Synlett, 1995, 1995, 71-73.	1.8	11
60	Nonpeptide Integrin Antagonists: RGD Mimetics Incorporating Substituted Azabicycloalkanes as Amino Acid Replacements. European Journal of Organic Chemistry, 2007, 2007, 1309-1317.	2.4	10
61	Stereoselective synthesis of Cα-tetrasubstituted azabicyclo[X.3.0]alkane amino acids. Tetrahedron Letters, 2004, 45, 6311-6315.	1.4	7
62	Dual action Smac mimetics–zinc chelators as pro-apoptotic antitumoral agents. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 4613-4619.	2.2	6
63	Synthesis of Functionalized Azabicycloalkane Amino Acids as Dipeptide Mimics. Synthesis, 2006, 2006, 1133-1140.	2.3	5
64	Design, Synthesis, Conformational Analysis and Application of AzabicycloÂalkane Amino Acids as Constrained Dipeptide Mimics. Synlett, 2004, 2004, 1449-1471.	1.8	4
65	New potent α _v î² ₃ integrin ligands based on azabicycloalkane (î³,α)-dipeptide mimics. Organic and Biomolecular Chemistry, 2016, 14, 3221-3233.	2.8	4
66	4-Connected azabicyclo[5.3.0]decane Smac mimetics-Zn 2+ chelators as dual action antitumoral agents. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 2336-2344.	2.2	4
67	Asymmetric Synthesis of Enantiopure α-Sulfenyl Dithioacetals and α-Sulfenyl Aldehydes. Phosphorus, Sulfur and Silicon and the Related Elements, 1993, 74, 381-382.	1.6	3
68	Synthesis of Conformationally Restricted and Optically Pure Analogues of Serine-Proline Dipeptide via Aldol Condensation. Synthesis, 2004, 2004, 353-358.	2.3	1
69	Molecular Targeting of Imaging and Drug Delivery Probes in Atherosclerosis. Annual Reports in Medicinal Chemistry, 2013, 48, 105-118.	0.9	1
70	Stereoselective Synthesis of a Functionalized 2-Oxo-1-azabicyclo[5.3.0]alkane as a Potential Scaffold for Targeted Chemotherapy Strategies. Synthesis, 2003, 2003, 2363-2367.	2.3	0
71	The First Example of Ring-Closing Olefin Metathesis of Dehydroamino Acids: An Application to the Synthesis of Azabicyclo[X.Y.0]alkanes ChemInform, 2004, 35, no.	0.0	0
72	Characterization of iron oxide-gold core-shell multifunctional nanoparticles in biomedical imaging. , 2011, , .		0

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73	Bisphosphonate-functionalized cyclic Arg-Gly-Asp peptidomimetics. Arkivoc, 2013, 2013, 185-200.	0.5	0