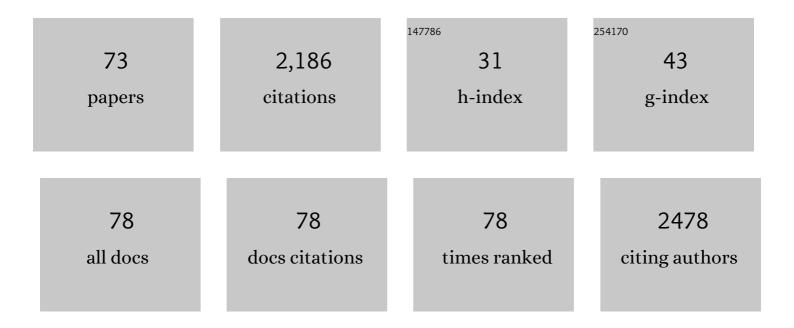
Leonardo Manzoni

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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|----|---|-----|-----------|
| 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 |