Daniela Arosio

List of Publications by Year in descending order

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83 papers 2,467 citations

147801 31 h-index 233421 45 g-index

85 all docs

85 docs citations

85 times ranked 3270 citing authors

#	Article	IF	CITATIONS
1	An inhibitor of tau hyperphosphorylation prevents severe motor impairments in tau transgenic mice. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 9673-9678.	7.1	206
2	Advancement in integrin facilitated drug delivery. Advanced Drug Delivery Reviews, 2016, 97, 111-143.	13.7	128
3	Cyclic RGD Functionalized Gold Nanoparticles for Tumor Targeting. Bioconjugate Chemistry, 2011, 22, 664-672.	3.6	82
4	A Synthetic Divalent Cholera Toxin Glycocalix[4]arene Ligand Having Higher Affinity than Natural GM1 Oligosaccharide. Journal of the American Chemical Society, 2005, 127, 3660-3661.	13.7	79
5	Synthesis and cholera toxin binding properties of multivalent GM1 mimicsElectronic supplementary information (ESI) available: characterization of the polyvalent compounds? imide by-products. See http://www.rsc.org/suppdata/ob/b4/b405344c/. Organic and Biomolecular Chemistry, 2004, 2, 2113.	2.8	77
6	Intramolecular Carbohydrate-Aromatic Interactions and Intermolecular van der Waals Interactions Enhance the Molecular Recognition Ability of GM1 Glycomimetics for Cholera Toxin. Chemistry - A European Journal, 2004, 10, 4395-4406.	3.3	69
7	Synthesis and Biological Evaluation (in Vitro and in Vivo) of Cyclic Arginine–Glycine–Aspartate (RGD) Peptidomimetic–Paclitaxel Conjugates Targeting Integrin α _V β ₃ . Journal of Medicinal Chemistry, 2012, 55, 10460-10474.	6.4	68
8	Dihydrotanshinone-I interferes with the RNA-binding activity of HuR affecting its post-transcriptional function. Scientific Reports, 2015, 5, 16478.	3.3	65
9	Cyclic RGD Peptidomimetics Containing Bifunctional Diketopiperazine Scaffolds as New Potent Integrin Ligands. Chemistry - A European Journal, 2012, 18, 6195-6207.	3.3	62
10	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
11	Designing Smac-mimetics as antagonists of XIAP, cIAP1, and cIAP2. Biochemical and Biophysical Research Communications, 2009, 378, 162-167.	2.1	50
12	Effective Targeting of DC-SIGN by \hat{l}_{\pm} -Fucosylamide Functionalized Gold Nanoparticles. Bioconjugate Chemistry, 2014, 25, 2244-2251.	3.6	50
13	Synthesis and Biological Evaluation of RGD Peptidomimetic–Paclitaxel Conjugates Bearing Lysosomally Cleavable Linkers. Chemistry - A European Journal, 2015, 21, 6921-6929.	3 . 3	48
14	Cyclic RGDâ€Containing Functionalized Azabicycloalkane Peptides as Potent Integrin Antagonists for Tumor Targeting. ChemMedChem, 2009, 4, 615-632.	3.2	44
15	Kiss and Run: Promoting Effective and Targeted Cellular Uptake of a Drug Delivery Vehicle Composed of an Integrin-Targeting Diketopiperazine Peptidomimetic and a Cell-Penetrating Peptide. Bioconjugate Chemistry, 2019, 30, 2011-2022.	3.6	44
16	Retromer stabilization results in neuroprotection in a model of Amyotrophic Lateral Sclerosis. Nature Communications, 2020, 11, 3848.	12.8	44
17	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
18	Design, Synthesis, and Biological Evaluation of Novel cRGDâ€"Paclitaxel Conjugates for Integrin-Assisted Drug Delivery. Bioconjugate Chemistry, 2012, 23, 1610-1622.	3.6	41

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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	Functionalized Azabicycloalkane Amino Acids by Nitrone 1,3-Dipolar Intramolecular Cycloaddition. Journal of Organic Chemistry, 2005, 70, 4124-4132.	3.2	39
21	A new optical imaging probe targeting $\langle i \rangle \hat{l} \pm \langle i \rangle \langle sub \rangle V \langle sub \rangle \langle i \rangle \hat{l}^2 \langle i \rangle \langle sub \rangle 3 \langle sub \rangle$ integrin in glioblastoma xenografts. Contrast Media and Molecular Imaging, 2011, 6, 449-458.	0.8	39
22	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
23	Solid phase immunoadsorption for therapeutic and analytical studies on neuropathy-associated anti-GM1 antibodies. Glycobiology, 2007, 17, 294-303.	2.5	38
24	MicroPET/CT imaging of $\hat{l}\pm v\hat{l}^2$ 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
25	HuR/ELAVL1 drives malignant peripheral nerve sheath tumor growth and metastasis. Journal of Clinical Investigation, 2020, 130, 3848-3864.	8.2	38
26	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
27	Synthesis of Novel c(AmpRGD)–Sunitinib Dual Conjugates as Molecular Tools Targeting the α _v β ₃ Integrin/VEGFR2 Couple and Impairing Tumor-Associated Angiogenesis. Journal of Medicinal Chemistry, 2017, 60, 248-262.	6.4	36
28	Integrin-Mediated Drug Delivery in Cancer and Cardiovascular Diseases with Peptide-Functionalized Nanoparticles. Current Medicinal Chemistry, 2012, 19, 3128-3151.	2.4	34
29	Cyclic <i>iso</i> DGR and RGD Peptidomimetics Containing Bifunctional Diketopiperazine Scaffolds are Integrin Antagonists. Chemistry - A European Journal, 2015, 21, 6265-6271.	3.3	33
30	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
31	Improved synthesis of both enantiomers of trans-cyclohex-4-ene-1,2-dicarboxylic acid. Tetrahedron: Asymmetry, 1999, 10, 3403-3407.	1.8	32
32	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
33	Synthesis and biological evaluation of RGD and isoDGR peptidomimetic-α-amanitin conjugates for tumor-targeting. Beilstein Journal of Organic Chemistry, 2018, 14, 407-415.	2.2	30
34	Neutrophil Elastase Promotes Linker Cleavage and Paclitaxel Release from an Integrinâ€Targeted Conjugate. Chemistry - A European Journal, 2019, 25, 1696-1700.	3.3	29
35	Cyclic <i>iso</i> DGR Peptidomimetics as Lowâ€Nanomolar α _v β ₃ Integrin Ligands. Chemistry - A European Journal, 2013, 19, 3563-3567.	3.3	28
36	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

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37	Multivalency Increases the Binding Strength of RGD Peptidomimeticâ€Paclitaxel Conjugates to Integrin α _V β ₃ . Chemistry - A European Journal, 2017, 23, 14410-14415.	3.3	27
38	Regiospecific Synthesis of Mono-N-substituted Indolopyrrolocarbazoles. Organic Letters, 2005, 7, 4573-4576.	4.6	26
39	Functionalized Cyclic RGD Peptidomimetics: Conjugable ligands for \hat{l}_{s} sub> \hat{l}_{s} sub	3. 6	26
40	Synthesis and Biological Evaluation of RGD and <i>iso</i> DGR–Monomethyl Auristatin Conjugates Targeting Integrin α _V β ₃ . ChemMedChem, 2019, 14, 938-942.	3.2	26
41	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
42	Synthesis, Characterization, and Biological Evaluation of a Dualâ€Action Ligand Targeting α _v β ₃ Integrin and VEGF Receptors. ChemistryOpen, 2015, 4, 633-641.	1.9	25
43	Mimicking gangliosides by design: mimics of GM1 headgroup. Neurochemical Research, 2002, 27, 539-545.	3.3	23
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	Insights into the Binding of Cyclic RGD Peptidomimetics to \hat{l}_{\pm} ₅ \hat{l}^{2} ₁ Integrin by using Live-Cell NMR And Computational Studies. ChemistryOpen, 2017, 6, 128-136.	1.9	21
46	Click chemistry to functionalise peptidomimetics. Tetrahedron Letters, 2006, 47, 3697-3700.	1.4	20
47	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
48	Synthesis and biological evaluation of dual action <i>cyclo</i> -RGD/SMAC mimetic conjugates targeting \hat{l}_{\pm} _v \hat{l}_{\pm} _{\hat{l}_{\pm}_{\hat}}	2.8	19
49	Investigating the Interaction of Cyclic RGD Peptidomimetics with $\hat{l}\pm\hat{Vl^2}$ 6 Integrin by Biochemical and Molecular Docking Studies. Cancers, 2017, 9, 128.	3.7	18
50	Intracisternal delivery of PEG-coated gold nanoparticles results in high brain penetrance and long-lasting stability. Journal of Nanobiotechnology, 2019, 17, 49.	9.1	18
51	Ganglioside GM1 mimics: lipophilic substituents improve affinity for cholera toxin. Bioorganic and Medicinal Chemistry Letters, 2003, 13, 3831-3834.	2.2	17
52	Tumor Targeting with an <i>iso</i> DGR–Drug Conjugate. Chemistry - A European Journal, 2017, 23, 7910-7914.	3.3	17
53	Conjugates of Cryptophycin and RGD or <i>iso</i> DGR Peptidomimetics for Targeted Drug Delivery. ChemistryOpen, 2019, 8, 737-742.	1.9	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	Synthesis and Biological Evaluation of Paclitaxel Conjugates Involving Linkers Cleavable by Lysosomal Enzymes and α _V β ₃ â€Integrin Ligands for Tumor Targeting. European Journal of Organic Chemistry, 2018, 2018, 2902-2909.	2.4	16
56	Trehalose-based neuroprotective autophagy inducers. Bioorganic and Medicinal Chemistry Letters, 2021, 40, 127929.	2.2	16
57	Interfering with the Tumor–Immune Interface: Making Way for Triazine-Based Small Molecules as Novel PD-L1 Inhibitors. Journal of Medicinal Chemistry, 2021, 64, 16020-16045.	6.4	16
58	Iron Oxide-Gold Core-Shell Nanoparticles as Multimodal Imaging Contrast Agent. IEEE Sensors Journal, 2013, 13, 2341-2347.	4.7	15
59	Targeting Integrin \hat{l}_{\pm} _V \hat{l}^2 ₃ with Theranostic RGD-Camptothecin Conjugates Bearing a Disulfide Linker: Biological Evaluation Reveals a Complex Scenario. ChemistrySelect, 2017, 2, 4759-4766.	1.5	14
60	Nanolipid-Trehalose Conjugates and Nano-Assemblies as Putative Autophagy Inducers. Pharmaceutics, 2019, 11, 422.	4.5	14
61	\hat{l}^2 -Glucuronidase triggers extracellular MMAE release from an integrin-targeted conjugate. Organic and Biomolecular Chemistry, 2019, 17, 4705-4710.	2.8	14
62	Synthesis and preclinical evaluation of a novel, selective ¹¹¹ In-labelled aminoproline-RGD-peptide for non-invasive melanoma tumor imaging. MedChemComm, 2015, 6, 2175-2183.	3.4	11
63	HuR-targeted agents: An insight into medicinal chemistry, biophysical, computational studies and pharmacological effects on cancer models. Advanced Drug Delivery Reviews, 2022, 181, 114088.	13.7	11
64	Rational Design of Antiangiogenic Helical Oligopeptides Targeting the Vascular Endothelial Growth Factor Receptors. Frontiers in Chemistry, 2019, 7, 170.	3.6	10
65	Multimeric Presentation of RGD Peptidomimetics Enhances Integrin Binding and Tumor Cell Uptake. Chemistry - A European Journal, 2020, 26, 7492-7496.	3.3	10
66	Rational design, synthesis and characterization of potent, drug-like monomeric Smac mimetics as pro-apoptotic anticancer agents. Bioorganic and Medicinal Chemistry Letters, 2012, 22, 2204-2208.	2.2	7
67	Shifting Towards α _V β ₆ Integrin Ligands Using Novel Aminoprolineâ€Based Cyclic Peptidomimetics. Chemistry - A European Journal, 2020, 26, 13468-13475.	3.3	7
68	Squalene-Based Nano-Assemblies Improve the Pro-Autophagic Activity of Trehalose. Pharmaceutics, 2022, 14, 862.	4.5	7
69	Dual action Smac mimetics–zinc chelators as pro-apoptotic antitumoral agents. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 4613-4619.	2.2	6
70	Synthesis of non glycosidic nucleobase-sugar mimetics. Comptes Rendus Chimie, 2010, 13, 1284-1300.	0.5	5
71	SPION-Smac mimetic nano-conjugates: Putative pro-apoptotic agents in oncology. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 2374-2378.	2.2	5
72	Cyclic RGD and isoDGR Integrin Ligands Containing cis-2-amino-1-cyclopentanecarboxylic (cis-Î ² -ACPC) Scaffolds. Molecules, 2020, 25, 5966.	3.8	5

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73	New potent $\hat{l}\pm < \text{sub}>v < / \text{sub}> \hat{l}^2 < \text{sub}> 3 < / \text{sub}> integrin ligands based on azabicycloalkane (\hat{l}^3, \hat{l}\pm)-dipeptide mimics. Organic and Biomolecular Chemistry, 2016, 14, 3221-3233.$	2.8	4
74	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
75	Stereodivergent synthesis of 5-aminopipecolic acids and application in the preparation of a cyclic RGD peptidomimetic as a nanomolar \hat{l}_{\pm} _{\hat{l}^{2}_{\hat{s} integrin ligand. Organic and Biomolecular Chemistry, 2018, 16, 3402-3414.}}	2.8	4
76	A dimeric bicyclic RGD ligand displays enhanced integrin binding affinity and strong biological effects on U-373 MG glioblastoma cells. Organic and Biomolecular Chemistry, 2019, 17, 8913-8917.	2.8	4
77	Small molecules as pro-apoptotic anticancer agents. Pharmaceutical Patent Analyst, 2012, 1, 483-505.	1.1	3
78	Synthesis and Characterization of Novel Mono- and Bis-Guanyl Hydrazones as Potent and Selective ASIC1 Inhibitors Able to Reduce Brain Ischemic Insult. Journal of Medicinal Chemistry, 2021, 64, 8333-8353.	6.4	3
79	Solid-phase synthesis of combinatorial libraries based on enatiomerically pure (1S,2S,4R,5S)-4,5-dihydroxycyclohexan-1,2-dicarboxylic acid scaffolds. Il Farmaco, 2002, 57, 861-864.	0.9	1
80	Molecular Targeting of Imaging and Drug Delivery Probes in Atherosclerosis. Annual Reports in Medicinal Chemistry, 2013, 48, 105-118.	0.9	1
81	Characterization of iron oxide-gold core-shell multifunctional nanoparticles in biomedical imaging. , 2011, , .		O
82	Bisphosphonate-functionalized cyclic Arg-Gly-Asp peptidomimetics. Arkivoc, 2013, 2013, 185-200.	0.5	0
83	Frontispiece: Multivalency Increases the Binding Strength of RGD Peptidomimeticâ€Paclitaxel Conjugates to Integrin α _V β ₃ . Chemistry - A European Journal, 2017, 23, .	3.3	0