

Cesare Gennari

List of Publications by Year in descending order

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228
papers

8,410
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272
times ranked

4551
citing authors

#	ARTICLE	IF	CITATIONS
1	Direct synthesis of Z-unsaturated esters. A useful modification of the horner-emmons olefination.. Tetrahedron Letters, 1983, 24, 4405-4408.	1.4	1,024
2	Combinatorial Libraries of Chiral Ligands for Enantioselective Catalysis. Chemical Reviews, 2003, 103, 3071-3100.	47.7	271
3	Asymmetric electrophilic amination: synthesis of .alpha.-amino and .alpha.-hydrazino acids with high optical purity. Journal of the American Chemical Society, 1986, 108, 6394-6395.	13.7	217
4	Synthesis of Sulfonamido-Pseudopeptides: New Chiral Unnatural Oligomers. Angewandte Chemie International Edition in English, 1994, 33, 2067-2069.	4.4	142
5	The rational design and systematic analysis of asymmetric aldol reactions using enol borinates: Applications of transition state computer modelling. Tetrahedron: Asymmetry, 1995, 6, 2613-2636.	1.8	115
6	Supramolecular ligand-ligand and ligand-substrate interactions for highly selective transition metal catalysis. Dalton Transactions, 2011, 40, 4355.	3.3	115
7	Investigation of a New Family of Chiral Ligands for Enantioselective Catalysis via Parallel Synthesis and High-Throughput Screening. Journal of Organic Chemistry, 1998, 63, 5312-5313.	3.2	114
8	Asymmetric synthesis of trans-Î²-lactams through TiCl4-mediated addition to imines.. Tetrahedron Letters, 1987, 28, 227-230.	1.4	111
9	Conformational Preferences of Peptides Containing Reverse-Turn Mimetic Bicyclic Lactams: Inverse Î³-Turns versus Type-IIâ€² Î²-Turns â€“ Insights into Î²-Hairpin Stability. , 1999, 1999, 389-400.		92
10	Lewis acid mediated aldol condensations using thioester silyl ketene acetals. Tetrahedron, 1986, 42, 893-909.	1.9	87
11	Rh-Catalyzed Asymmetric Hydrogenation of Prochiral Olefins with a Dynamic Library of Chiral TROPOS Phosphorus Ligands. Chemistry - A European Journal, 2005, 11, 6701-6717.	3.3	86
12	Effect of Ligands and Additives on the Palladium-Promoted Carbonylative Coupling of Vinyl Stannanes and Electron-Poor Enol Triflates. Journal of Organic Chemistry, 2000, 65, 6254-6256.	3.2	85
13	Titanium tetrachloride-mediated enantioselective synthesis of trans Î²-lactones. Tetrahedron, 1992, 48, 5557-5564.	1.9	81
14	A Catalytic and Enantioselective Desymmetrization of meso Cyclic Allylic Bisdiethylphosphates with Organozinc Reagents. Angewandte Chemie - International Edition, 2003, 42, 234-236.	13.8	81
15	Enantioselective synthesis of anti-.alpha.-methyl-.beta.-hydroxy esters through titanium tetrachloride-mediated aldol condensation. Journal of the American Chemical Society, 1985, 107, 5812-5813.	13.7	79
16	Discovery of a New Efficient Chiral Ligand for Copper-Catalyzed Enantioselective Michael Additions by High-Throughput Screening of a Parallel Library. Angewandte Chemie - International Edition, 2000, 39, 916-918.	13.8	79
17	Rh-Catalyzed Enantioselective Conjugate Addition of Arylboronic Acids with a Dynamic Library of Chiral Tropos Phosphorus Ligands. Chemistry - A European Journal, 2007, 13, 1547-1558.	3.3	73
18	A Practical Approach to the Resolution of Racemic N-Benzyl Î±-Amino Acids by Liquid-Liquid Extraction with a Lipophilic Chiral Salenâ€“Cobalt(III) Complex. Angewandte Chemie - International Edition, 2006, 45, 2449-2453.	13.8	70

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19	Transition-state modeling of the aldol reaction of boron enolates: a force field approach. <i>Journal of Organic Chemistry</i> , 1990, 55, 3576-3581.	3.2	69
20	Synthesis and Biological Evaluation (in Vitro and in Vivo) of Cyclic Arginine-Glycine-Aspartate (RGD) Peptidomimetic-Paclitaxel Conjugates Targeting Integrin $\alpha_5\beta_3$. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 10460-10474.	6.4	68
21	Innovative Linker Strategies for Tumor-Targeted Drug Conjugates. <i>Chemistry - A European Journal</i> , 2019, 25, 14740-14757.	3.3	68
22	Synthetic Receptors Based on Vinylogous Sulfonyl Peptides. <i>Angewandte Chemie International Edition in English</i> , 1995, 34, 1765-1768.	4.4	66
23	Combinatorial Libraries: Studies in Molecular Recognition and the Quest for New Catalysts. <i>Liebigs Annalen</i> , 1997, 1997, 637-647.	0.8	66
24	Titanium chloride-mediated reactions of the silyl ketene acetals derived from N-methylephedrine esters: an asymmetric variant of the Mukaiyama reaction. <i>Journal of Organic Chemistry</i> , 1987, 52, 2754-2760.	3.2	64
25	Improved enantioselective synthesis of anti-1-methyl-2-hydroxyesters through TiCl_4 - PPh_3 mediated aldol condensation. <i>Tetrahedron Letters</i> , 1986, 27, 1735-1738.	1.4	63
26	Cyclic RGD Peptidomimetics Containing Bifunctional Diketopiperazine Scaffolds as New Potent Integrin Ligands. <i>Chemistry - A European Journal</i> , 2012, 18, 6195-6207.	3.3	62
27	A new method for the solution and solid phase synthesis of chiral 2-sulfonopeptides under mild conditions. <i>Tetrahedron Letters</i> , 1996, 37, 8589-8592.	1.4	61
28	High diastereoface selection in an ester enolate addition to α -alkoxy aldehydes: stereoselective synthesis of α -methylene- β -hydroxy- γ -alkoxy esters. <i>Journal of Organic Chemistry</i> , 1984, 49, 3784-3790.	3.2	60
29	Discovery of a New Efficient Chiral Ligand for Copper-Catalyzed Enantioselective Michael Additions by High-Throughput Screening of a Parallel Library. <i>Chemistry - A European Journal</i> , 2001, 7, 2628-2634.	3.3	59
30	Cyclic RGD Peptidomimetics Containing Bifunctional Diketopiperazine Scaffolds as New Potent Integrin Ligands. <i>Chemistry - A European Journal</i> , 2009, 15, 12184-12188.	3.3	58
31	Rhodium-Catalyzed Asymmetric Hydrogenation of Olefins with PhthalaPhos, a New Class of Chiral Supramolecular Ligands. <i>Chemistry - A European Journal</i> , 2012, 18, 1383-1400.	3.3	57
32	Chiral (Cyclopentadienone)iron Complexes for the Catalytic Asymmetric Hydrogenation of Ketones. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 1887-1893.	2.4	56
33	The rational design of highly stereoselective boron enolates using transition-state computer modeling: a novel, asymmetric anti aldol reaction for ketones. <i>Journal of Organic Chemistry</i> , 1992, 57, 5173-5177.	3.2	55
34	Hydrogen-Bonding Donor/Acceptor Scales in 2-Sulfonamidopeptides. <i>Chemistry - A European Journal</i> , 1998, 4, 1924-1931.	3.3	55
35	Stereoselectivity of intramolecular nitrile oxide cycloadditions to Z and E chiral alkenes. <i>Journal of Organic Chemistry</i> , 1987, 52, 4674-4681.	3.2	54
36	A trifunctional steroid-based scaffold for combinatorial chemistry. <i>Tetrahedron Letters</i> , 1999, 40, 2849-2852.	1.4	52

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37	Ureas: New efficient Lewis base catalysts for the allylation of aldehydes. <i>Tetrahedron Letters</i> , 1999, 40, 3633-3634.	1.4	52
38	Origins of β -face selectivity in the aldol reactions of chiral E-enol borinates: a computational study using transition state modelling. <i>Tetrahedron</i> , 1993, 49, 685-696.	1.9	51
39	Solid-Phase Synthesis of Vinylogous Sulfonyl Peptides. <i>Angewandte Chemie International Edition in English</i> , 1995, 34, 1763-1765.	4.4	51
40	Diastereofacial selectivity in the aldol reactions of chiral β -methyl aldehydes: a computer modelling approach. <i>Tetrahedron</i> , 1992, 48, 4439-4458.	1.9	50
41	Conformational Studies of Chiral Vinylogous Sulfonamidopeptides. <i>Chemistry - A European Journal</i> , 1996, 2, 644-655.	3.3	50
42	PhthalaPhos: Chiral Supramolecular Ligands for Enantioselective Rhodium-Catalyzed Hydrogenation Reactions. <i>Angewandte Chemie - International Edition</i> , 2010, 49, 6633-6637.	13.8	50
43	Effects of allylic and homoallylic substituents on the ring closing metathesis reaction used to synthesise simplified eleuthesides. <i>Tetrahedron Letters</i> , 2003, 44, 7913-7919.	1.4	49
44	A Formal Total Synthesis of Eleutherobin Using the Ring-Closing Metathesis (RCM) Reaction of a Densely Functionalized Diene as the Key Step: Investigation of the Unusual Kinetically Controlled RCM Stereochemistry. <i>Chemistry - A European Journal</i> , 2006, 12, 51-62.	3.3	49
45	Synthetic opportunities offered by anti α -methylene- β -hydroxy- γ -alkoxy esters: stereoselective reactions at the double bond. <i>Journal of Organic Chemistry</i> , 1985, 50, 4442-4447.	3.2	48
46	Asymmetric synthesis of functionalized β -amino- γ -hydroxy acids via chiral norephedrine-derived oxazolidines. <i>Tetrahedron</i> , 1988, 44, 5563-5572.	1.9	48
47	Reagent Control in the Aldol Addition Reaction of Chiral Boron Enolates with Chiral α -Amino Aldehydes. Total Synthesis of (3S,4S)-Statine. <i>Journal of Organic Chemistry</i> , 1995, 60, 6248-6249.	3.2	48
48	Synthesis and Biological Evaluation of RGD Peptidomimetic-Paclitaxel Conjugates Bearing Lysosomally Cleavable Linkers. <i>Chemistry - A European Journal</i> , 2015, 21, 6921-6929.	3.3	48
49	Synthesis and Conformational Studies of Peptidomimetics Containing a New Bifunctional Diketopiperazine Scaffold Acting as a β -Hairpin Inducer. <i>Journal of Organic Chemistry</i> , 2008, 73, 652-660.	3.2	47
50	Origins of stereoselectivity in chiral boron enolate aldol reactions: A computational study using transition state modellings. <i>Tetrahedron</i> , 1991, 47, 3471-3484.	1.9	46
51	Copper Phosphoramidite-Catalyzed Enantioselective Desymmetrization of meso-Cyclic Allylic Bisdiethyl Phosphates. <i>Organic Letters</i> , 2003, 5, 4493-4496.	4.6	46
52	A Formal Total Synthesis of Eleutherobin Through an Unprecedented Kinetically Controlled Ring-Closing-Metathesis Reaction of a Densely Functionalized Diene. <i>Angewandte Chemie - International Edition</i> , 2005, 44, 588-591.	13.8	46
53	Enantioselective conjugate addition of phenylboronic acid to enones catalysed by a chiral tropos/atropos rhodium complex at the coalescence temperature. <i>Chemical Communications</i> , 2005, , 5281.	4.1	46
54	Chelation-controlled enantioselective synthesis of key intermediates for the preparation of carbapenem antibiotics PS-5 and 1. β -methyl-PS-5. <i>Journal of Organic Chemistry</i> , 1988, 53, 4015-4021.	3.2	45

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55	Bifunctional 2,5-Diketopiperazines as Rigid Three-Dimensional Scaffolds in Receptors and Peptidomimetics. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 217-228.	2.4	45
56	Synthesis of BINOL-Derived (Cyclopentadienone)iron Complexes and Their Application in the Catalytic Asymmetric Hydrogenation of Ketones. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 5526-5536.	2.4	45
57	γ -Integrin-Targeted Peptide/Peptidomimetic-Drug Conjugates: In-Depth Analysis of the Linker Technology. <i>Current Topics in Medicinal Chemistry</i> , 2015, 16, 314-329.	2.1	44
58	Optimization of New Chiral Ligands for the Copper-Catalysed Enantioselective Conjugate Addition of Et ₂ Zn to Nitroolefins by High-Throughput Screening of a Parallel Library. <i>European Journal of Organic Chemistry</i> , 2001, 2001, 803-807.	2.4	43
59	A Straightforward Total Synthesis of Chaetominine. <i>Chemistry - A European Journal</i> , 2009, 15, 7922-7929.	3.3	43
60	Efficient Synthesis of Amines by Iron-Catalyzed C=N Transfer Hydrogenation and C=O Reductive Amination. <i>Advanced Synthesis and Catalysis</i> , 2018, 360, 1054-1059.	4.3	43
61	Highly enantio- and diastereoselective boron aldol reactions of α -heterosubstituted thioacetates with aldehydes and silyl imines. <i>Tetrahedron</i> , 1997, 53, 5909-5924.	1.9	42
62	Chelation controlled aldol additions of the enolsilane derived from tert-butyl thioacetate : a stereoselective approach to 1 ² -methylthienamycin. <i>Tetrahedron</i> , 1988, 44, 5965-5974.	1.9	41
63	TiCl ₄ -mediated reactions of silyl ketene acetals derived from n-methylephedrine esters: asymmetric synthesis of β -lactams. <i>Tetrahedron</i> , 1988, 44, 4221-4232.	1.9	41
64	Synthetic studies on sarcodictyins and eleutherobin: Synthesis of fully functionalized cyclization precursors. <i>Tetrahedron Letters</i> , 1999, 40, 153-156.	1.4	41
65	Synthesis of novel simplified sarcodictyin/eleutherobin analogs with potent microtubule-stabilizing activity, using ring closing metathesis as the key-step. <i>Tetrahedron</i> , 2003, 59, 8803-8820.	1.9	41
66	Stereoselective aldol condensations via alkenyloxy dialkoxyboranes: synthetic applications using thioesters. <i>Tetrahedron</i> , 1984, 40, 4059-4065.	1.9	40
67	Asymmetric Synthesis with Enol Ethers. , 1991, , 629-660.		40
68	Taxol Semisynthesis: A Highly Enantio- and Diastereoselective Synthesis of the Side Chain and a New Method for Ester Formation at C-13 Using Thioesters. <i>Journal of Organic Chemistry</i> , 1997, 62, 4746-4755.	3.2	40
69	Rh-catalysed asymmetric hydrogenations with a dynamic library of chiral tropos phosphorus-ligands. <i>Tetrahedron Letters</i> , 2004, 45, 6859-6862.	1.4	40
70	Natural products with taxol-like anti-tumor activity: Synthetic approaches to eleutherobin and dictyostatin. <i>Pure and Applied Chemistry</i> , 2007, 79, 173-180.	1.9	40
71	Asymmetric dihydroxylations via chiral oxazolidines. <i>Tetrahedron Letters</i> , 1985, 26, 5459-5462.	1.4	39
72	Computer-Assisted Design of Chiral Boron Enolates: A Novel, Highly Enantioselective Aldol Reaction for Thioacetates and Thiopropionates. <i>Angewandte Chemie International Edition in English</i> , 1993, 32, 1618-1621.	4.4	39

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73	2-Benzoylamino-2-deoxy-2-hydroxymethyl-D-hexono-1,4-lactones: synthesis from D-fructose and utilization in the total synthesis of thermozyomicidin (myriocin). <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1983, , 1613.	0.9	38
74	Synthetic studies on the sarcodictyins: synthesis of fully functionalized cyclization precursors. <i>Tetrahedron</i> , 2001, 57, 8531-8542.	1.9	38
75	Resolution of Racemic <i>N</i> -Benzyl $\hat{\pm}$ -Amino Acids by Liquid-Liquid Extraction: A Practical Method Using a Lipophilic Chiral Cobalt(III) Salen Complex and Mechanistic Studies. <i>European Journal of Organic Chemistry</i> , 2008, 2008, 1253-1264.	2.4	38
76	Stereocontrolled synthesis of polyketide libraries: Boron-mediated aldol reactions with aldehydes on solid support. <i>Tetrahedron</i> , 1998, 54, 14999-15016.	1.9	37
77	Synthesis and Screening of New Chiral Ligands for the Copper-Catalysed Enantioselective Allylic Substitution. <i>Helvetica Chimica Acta</i> , 2002, 85, 3388-3399.	1.6	37
78	Theoretical studies of stereoselective aldol condensations. <i>Journal of Organic Chemistry</i> , 1986, 51, 612-616.	3.2	36
79	1,4-addition to $\hat{\pm}$, $\hat{1}^2$ -unsaturated carbonyl compounds bearing a $\hat{1}^3$ -stereocenter: A molecular mechanics model for steric interactions in the transition state.. <i>Tetrahedron: Asymmetry</i> , 1990, 1, 21-32.	1.8	36
80	Combinations of Acidic and Basic Monodentate Binaphtholic Phosphites as Supramolecular Bidentate Ligands for Enantioselective Rh-catalyzed Hydrogenations. <i>European Journal of Organic Chemistry</i> , 2009, 2009, 2539-2547.	2.4	36
81	Chiral acyl anion and enolonium ion equivalents. Asymmetric synthesis of $\hat{\pm}$ -methoxy-aldehydes. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1981, , 1278-1283.	0.9	35
82	Total synthesis of (+)-thermozyomicidin (myriocin) from D-fructose. <i>Journal of the Chemical Society Chemical Communications</i> , 1982, , 488.	2.0	35
83	Fast Cyclization of a Proline-derived Self-immolative Spacer Improves the Efficacy of Carbamate Prodrugs. <i>Angewandte Chemie - International Edition</i> , 2020, 59, 4176-4181.	13.8	35
84	Chiral acyl anion equivalents: asymmetric synthesis of $\hat{\pm}$ -methoxytolualdehyde. <i>Journal of the Chemical Society Chemical Communications</i> , 1979, , 591-592.	2.0	34
85	Synthesis of [Bis(hexamethylene)cyclopentadienone]iron Tricarbonyl and its Application to the Catalytic Reduction of C=O Bonds. <i>ChemCatChem</i> , 2017, 9, 1461-1468.	3.7	34
86	Stereoselective aldol condensations via enolboronates.. <i>Tetrahedron Letters</i> , 1984, 25, 2283-2286.	1.4	33
87	High diastereoselectivity in lewis acid mediated aldol condensations using thioester silyl ketene acetals.. <i>Tetrahedron Letters</i> , 1985, 26, 797-800.	1.4	33
88	Synthesis of a simplified sarcodictyin analogue which retains microtubule stabilising properties. <i>Tetrahedron Letters</i> , 2001, 42, 9187-9190.	1.4	33
89	Cyclative cleavage via solid-phase supported stabilized sulfur ylides: synthesis of macrocyclic lactones. <i>Tetrahedron Letters</i> , 2002, 43, 761-766.	1.4	33
90	A Library Approach to the Development of BenzaPhos: Highly Efficient Chiral Supramolecular Ligands for Asymmetric Hydrogenation. <i>Chemistry - A European Journal</i> , 2012, 18, 10368-10381.	3.3	33

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91	Cyclic <i>iso</i> -DGR and RGD Peptidomimetics Containing Bifunctional Diketopiperazine Scaffolds are Integrin Antagonists. <i>Chemistry - A European Journal</i> , 2015, 21, 6265-6271.	3.3	33
92	Stereoselective radical-mediated cyclization of norephedrine derived $\hat{\pm}$ -iodoamides: synthesis of enantiopure pyrrolidines and transition state modelling. <i>Tetrahedron</i> , 1992, 48, 3945-3960.	1.9	32
93	Origins of stereoselectivity in the addition of allyl- and crotylboronates to aldehydes : the development and application of a force field model of the transition state. <i>Tetrahedron</i> , 1994, 50, 8815-8826.	1.9	32
94	A carbonylative cross-coupling strategy to the total synthesis of the sarcodictyins: preliminary studies and synthesis of a cyclization precursor. <i>Tetrahedron Letters</i> , 2001, 42, 7421-7425.	1.4	32
95	Stereospecific Synthesis of Chiral $\hat{\pm}$ -Sulfinylhydrazones. <i>Synthesis</i> , 1982, 1982, 829-831.	2.3	30
96	Auxiliary structure and asymmetric induction in the Mukaiyama-aldol reactions of chiral silyl ketene acetals. <i>Tetrahedron Letters</i> , 1989, 30, 5163-5166.	1.4	30
97	Synthesis and biological evaluation of RGD and isoDGR peptidomimetic- $\hat{\pm}$ -amanitin conjugates for tumor-targeting. <i>Beilstein Journal of Organic Chemistry</i> , 2018, 14, 407-415.	2.2	30
98	Origins of stereoselectivity in the addition of chiral allyl- and crotylboranes to aldehydes: the development and application of a force field model of the transition state. <i>Journal of Organic Chemistry</i> , 1993, 58, 1711-1718.	3.2	29
99	Combination of a binaphthol-derived phosphite and a C1-symmetric phosphinamine generates heteroleptic catalysts in Rh- and Pd-mediated reactions. <i>Chemical Communications</i> , 2009, , 3539.	4.1	29
100	Asymmetric Hydrogenation of 3-Substituted Pyridinium Salts. <i>Chemistry - A European Journal</i> , 2016, 22, 9528-9532.	3.3	29
101	Neutrophil Elastase Promotes Linker Cleavage and Paclitaxel Release from an Integrin-Targeted Conjugate. <i>Chemistry - A European Journal</i> , 2019, 25, 1696-1700.	3.3	29
102	Enantioselective cyanosilylation of aldehydes catalysed by a diastereomeric mixture of atropisomeric thioureas. <i>Tetrahedron: Asymmetry</i> , 2006, 17, 999-1006.	1.8	28
103	Highly enantioselective Rh-catalyzed hydrogenations with heterocombinations of pentafluorobenzyl- and methoxybenzyl-derived binaphthyl phosphites. <i>Tetrahedron Letters</i> , 2008, 49, 755-759.	1.4	28
104	Cyclic <i>iso</i> -DGR Peptidomimetics as Low-Nanomolar $\hat{\pm}$ -Integrin Ligands. <i>Chemistry - A European Journal</i> , 2013, 19, 3563-3567.	3.3	28
105	Recent Catalytic Applications of (Cyclopentadienone)iron Complexes. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 3192-3205.	2.4	28
106	Stereoselective aldol condensations via alkenyloxy dialkoxyboranes : mechanistic and stereochemical details. <i>Tetrahedron</i> , 1984, 40, 4051-4058.	1.9	27
107	Stereoselective aldol additions to $\hat{\pm}$ -alkoxy aldehydes using thioester silyl ketene acetals,. <i>Tetrahedron Letters</i> , 1985, 26, 2373-2376.	1.4	27
108	Boron aldol reaction of $\hat{\pm}$ -halosubstituted thioacetates with silyl imines: A highly enantio- and diastereoselective synthesis of aziridines. <i>Tetrahedron Letters</i> , 1996, 37, 3747-3750.	1.4	27

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109	Expanding the Catalytic Scope of (Cyclopentadienone)iron Complexes to the Hydrogenation of Activated Esters to Alcohols. <i>ChemCatChem</i> , 2016, 8, 3431-3435.	3.7	27
110	Multivalency Increases the Binding Strength of RGD Peptidomimeticâ€Paclitaxel Conjugates to Integrin $\alpha_5\beta_1$. <i>Chemistry - A European Journal</i> , 2017, 23, 14410-14415.	3.3	27
111	Stereoselective aldol reactions using tlc14 as stereochemical template. <i>Tetrahedron Letters</i> , 1985, 26, 4129-4132.	1.4	26
112	Synthesis of novel, simplified, C-7 substituted eleutheside analogues with potent microtubule-stabilizing activity. <i>Tetrahedron</i> , 2005, 61, 2123-2139.	1.9	26
113	Rhodium-catalyzed asymmetric reactions with a dynamic library of chiral tropos phosphorus ligands. <i>Pure and Applied Chemistry</i> , 2006, 78, 303-310.	1.9	26
114	Bifunctional 2,5â€Diketopiperazines as Efficient Organocatalysts for the Enantioselective Conjugate Addition of Aldehydes to Nitroolefins. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 5599-5607.	2.4	26
115	Synthesis and Biological Evaluation of RGD and <i>iso</i> -DGRâ€Monomethyl Auristatin Conjugates Targeting Integrin $\alpha_5\beta_1$. <i>ChemMedChem</i> , 2019, 14, 938-942.	3.2	26
116	Stereoselective radical-mediated cyclization of norephedrine derived o-bromobenzamides: Enantioselective synthesis of 4-substituted 1,2,3,4-tetrahydroisoquinolines. <i>Tetrahedron: Asymmetry</i> , 1993, 4, 273-280.	1.8	25
117	Copper-Catalysed, Enantioselective Desymmetrisation of meso Cyclic Allylic Bis(diethyl phosphates) with Organozinc Reagents. <i>European Journal of Organic Chemistry</i> , 2005, 2005, 895-906.	2.4	25
118	Bicyclic carbohydrate-derived scaffolds for combinatorial libraries. <i>Bioorganic and Medicinal Chemistry</i> , 2006, 14, 3349-3367.	3.0	25
119	Synthesis, Characterization, and Biological Evaluation of a Dualâ€Action Ligand Targeting $\alpha_5\beta_1$ Integrin and VEGF Receptors. <i>ChemistryOpen</i> , 2015, 4, 633-641.	1.9	25
120	Chiral acyl anion equivalents: asymmetric synthesis of 11-deoxy-ent-prostaglandin intermediates. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1981, , 1284.	0.9	24
121	Enolboronates: New practical reagents for regioselective aldol condensations.. <i>Tetrahedron Letters</i> , 1984, 25, 2279-2282.	1.4	24
122	Synthesis of novel simplified eleutheside analogues with potent microtubule-stabilizing activity, using ring-closing metathesis as the key-step. <i>Tetrahedron Letters</i> , 2003, 44, 681-684.	1.4	24
123	Synthese von Sulfonamidâ€Pseudopeptiden: neue chirale synthetische Oligomere. <i>Angewandte Chemie</i> , 1994, 106, 2181-2183.	2.0	23
124	Title is missing!. <i>Angewandte Chemie</i> , 2003, 115, 244-246.	2.0	23
125	A highly stereoselective synthesis of the C10â€C23 fragment of (â€)-dictyostatin. <i>Chemical Communications</i> , 2007, , 4271.	4.1	23
126	Asymmetric synthesis of 34--substituted β -lactams via chiral norephedrine-derived oxazolidines.. <i>Tetrahedron</i> , 1989, 45, 7397-7404.	1.9	22

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127	A computational study of the 1,4-addition of lithium enolates to conjugated carbonyl compounds. <i>Journal of Organic Chemistry</i> , 1992, 57, 7029-7034.	3.2	22
128	Semisynthesis of Taxol: A Highly Enantio- and Diastereoselective Synthesis of the Side Chain and a New Method for Ester Formation at C13 Using Thioesters. <i>Angewandte Chemie International Edition in English</i> , 1996, 35, 1723-1725.	4.4	22
129	Determination of the binding epitope of RGD-peptidomimetics to $\alpha_5\beta_1$ and $\alpha_5\beta_3$ integrin-rich intact cells by NMR and computational studies. <i>Organic and Biomolecular Chemistry</i> , 2013, 11, 3886.	2.8	22
130	Chiral Formyl-Group Equivalents: Conjugate Addition to α,β -Unsaturated Ketones. <i>Synthesis</i> , 1981, 1981, 74-76.	2.3	21
131	Biosynthesis of citrinin and synthesis of its biogenetic precursors. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1981, , 2594.	0.9	21
132	Reagent control in the aldol addition reaction of chiral boron enolates with chiral aldehydes. Total synthesis of (3S,4S)-Statine. <i>Tetrahedron</i> , 1997, 53, 5593-5608.	1.9	21
133	Copper catalysed 1,4-addition of organozinc reagents to α,β -unsaturated carbonyl compounds: a mechanistic investigation. <i>Journal of Organometallic Chemistry</i> , 2004, 689, 2169-2176.	1.8	21
134	Synthesis of the C15-C23 fragment of dictyostatin using a highly stereoselective Carreira alkylation. <i>Tetrahedron</i> , 2007, 63, 5873-5878.	1.9	21
135	Insights into the Binding of Cyclic RGD Peptidomimetics to $\alpha_5\beta_1$ Integrin by using Live-Cell NMR And Computational Studies. <i>ChemistryOpen</i> , 2017, 6, 128-136.	1.9	21
136	Reagent control in the aldol addition reaction of chiral boron enolates with chiral aldehydes. <i>Tetrahedron Letters</i> , 1994, 35, 4623-4626.	1.4	20
137	Biosynthetic origin and revised structure of ascochitine, a phytotoxic fungal metabolite. Incorporation of [1- ¹³ C]- and [1,2- ¹³ C ₂]-acetates and [Me- ¹³ C]methionine. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1980, , 675.	0.9	19
138	6-Farnesyl-5,7-dihydroxy-4-methylphthalide oxidation mechanism in mycophenolic acid biosynthesis. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1982, , 365.	0.9	19
139	Total Synthesis of (+)- ϵ -Helianane and (+)- ϵ -Chloro- ϵ -Helianane through Stereoselective Aromatic Claisen Rearrangement. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 6794-6801.	2.4	19
140	Synthesis and biological evaluation of dual action <i>cyclo</i> -RGD/SMAC mimetic conjugates targeting $\alpha_v\beta_3$ / $\alpha_v\beta_5$ integrins and IAP proteins. <i>Organic and Biomolecular Chemistry</i> , 2014, 12, 3288-3302.	2.8	19
141	(E,E)-10-(1,3-Dihydro-4,6-dihydroxy-7-methyl-3-oxoisobenzofuran-5-yl)4,8-dimethyldeca-4,8-dienoic acid: total synthesis and role in mycophenolic acid biosynthesis. <i>Journal of the Chemical Society Chemical Communications</i> , 1979, , 1021.	2.0	18
142	Developing a force field for the transition state of the aldol reaction of enolborinates: Evaluation of the use of fixed point charges. <i>Tetrahedron</i> , 1992, 48, 4183-4192.	1.9	18
143	A Mixed Ligand Approach for the Asymmetric Hydrogenation of 2ϵ -Substituted Pyridinium Salts. <i>Advanced Synthesis and Catalysis</i> , 2016, 358, 2589-2593.	4.3	18
144	Investigating the Interaction of Cyclic RGD Peptidomimetics with $\alpha_5\beta_6$ Integrin by Biochemical and Molecular Docking Studies. <i>Cancers</i> , 2017, 9, 128.	3.7	18

#	ARTICLE	IF	CITATIONS
145	Acetogenin synthesis. Organocopper reagents, anions of 1,3-dithians and of protected cyanohydrins as intermediates in ketide side-chain synthesis. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1980, , 136.	0.9	17
146	A highly enantio- and diastereoselective aldol reaction for $\hat{1}\pm$ -heterosubstituted thioacetates. <i>Tetrahedron Letters</i> , 1994, 35, 4857-4860.	1.4	17
147	Computer-assisted design of chiral boron enolates: The role of ate complexes in determining aldol stereoselectivity.. <i>Tetrahedron</i> , 1994, 50, 1227-1242.	1.9	17
148	Synthetische Rezeptoren aus vinylogen Sulfonylpeptiden. <i>Angewandte Chemie</i> , 1995, 107, 1894-1896.	2.0	17
149	Synthesis of Chiral Vinylogous Sulfonamidopeptides (vs-Peptides). , 1998, 1998, 945-959.		17
150	Synthesis of Combinatorial Libraries of Vinylogous Sulfonamidopeptides (vs-Peptides). , 1998, 1998, 2437-2449.		17
151	Solid-Phase Synthesis of Peptides Containing Reverse-Turn Mimetic Bicyclic Lactams. , 1999, 1999, 379-388.		17
152	Enantioselective binding of dipeptides using acyclic receptors. <i>Chemical Communications</i> , 2001, , 1358-1359.	4.1	17
153	Tumor Targeting with an <i>iso</i> -DGR "Drug Conjugate. <i>Chemistry - A European Journal</i> , 2017, 23, 7910-7914.	3.3	17
154	Conjugates of Cryptophycin and RGD or <i>iso</i> -DGR Peptidomimetics for Targeted Drug Delivery. <i>ChemistryOpen</i> , 2019, 8, 737-742.	1.9	17
155	Regiodivergent Reductive Opening of Epoxides by Catalytic Hydrogenation Promoted by a (Cyclopentadienone)iron Complex. <i>ACS Catalysis</i> , 2022, 12, 235-246.	11.2	17
156	Lewis acid promoted aldol additions of $\hat{1}\pm$ -thiosilylketeneacetals to $\hat{1}\pm$ -alkoxy aldehydes: diastereoselective synthesis of $\hat{1}\pm$ -methylene- $\hat{1}^2$ -hydroxy- $\hat{1}\pm$ -alkoxy esters.. <i>Tetrahedron Letters</i> , 1985, 26, 6509-6512.	1.4	16
157	Stereoselective radical-mediated cyclization of norephedrine derived $\hat{1}\pm$ -iodoamides: Experiments and TS-modelling. <i>Tetrahedron: Asymmetry</i> , 1991, 2, 793-796.	1.8	16
158	Rationally Designed Bicyclic Lactams Control Different Turn Motifs and Folding Patterns in Hexapeptide Mimics. , 2000, 2000, 695-699.		16
159	Highly Stereoselective Total Synthesis of (+)- $\hat{1}\pm$ -epi- $\hat{1}\pm$ -dictyostatin and ($\hat{1}\pm$)- $\hat{1}\pm$ -dictyostatin. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 2643-2661.	2.4	16
160	Assisted Tandem Catalysis: Metathesis Followed by Asymmetric Hydrogenation from a Single Ruthenium Source. <i>Advanced Synthesis and Catalysis</i> , 2015, 357, 2223-2228.	4.3	16
161	Synthesis of a $\hat{1}\pm$ -vinyltetrahydrocarbazole by Palladium-Catalyzed Asymmetric Allylic Alkylation of Indole-Containing Allylic Carbonates. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 6669-6678.	2.4	16
162	Synthesis and Biological Evaluation of Paclitaxel Conjugates Involving Linkers Cleavable by Lysosomal Enzymes and $\hat{1}\pm$ - $\hat{1}^2$ - $\hat{1}^3$ -Integrin Ligands for Tumor Targeting. <i>European Journal of Organic Chemistry</i> , 2018, 2018, 2902-2909.	2.4	16

#	ARTICLE	IF	CITATIONS
163	Hydrogen-Borrowing Amination of Secondary Alcohols Promoted by a (Cyclopentadienone)iron Complex. <i>Synthesis</i> , 2019, 51, 3545-3555.	2.3	15
164	Chiral (cyclopentadienone)iron complexes with a stereogenic plane as pre-catalysts for the asymmetric hydrogenation of polar double bonds. <i>Tetrahedron</i> , 2019, 75, 1415-1424.	1.9	15
165	Peptide bond formation using an enzyme mimicking approach. <i>Tetrahedron</i> , 1990, 46, 7289-7300.	1.9	14
166	A computational study of the 1,4-addition of lithium enolates to conjugated carbonyl compounds.. <i>Tetrahedron Letters</i> , 1991, 32, 823-826.	1.4	14
167	Computational studies on the aldol-type addition of boron enolates to imines: An ab-initio approach. <i>Tetrahedron</i> , 1997, 53, 7705-7714.	1.9	14
168	Efficient resolution of racemic N-benzyl β -amino acids by iterative liquid-liquid extraction with a chiral (salen)cobalt(III) complex as enantioselective selector. <i>Organic and Biomolecular Chemistry</i> , 2007, 5, 3464.	2.8	14
169	Targeting Integrin α _V β ₃ with Theranostic RGD-Camptothecin Conjugates Bearing a Disulfide Linker: Biological Evaluation Reveals a Complex Scenario. <i>ChemistrySelect</i> , 2017, 2, 4759-4766.	1.5	14
170	Use of the Trost Ligand in the Ruthenium-Catalyzed Asymmetric Hydrogenation of Ketones. <i>ChemCatChem</i> , 2017, 9, 3125-3130.	3.7	14
171	β -Glucuronidase triggers extracellular MMAE release from an integrin-targeted conjugate. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 4705-4710.	2.8	14
172	Enantioselective Synthesis of (-)-(R)-5-Hydroxy-1-(4-hydroxy-3-methoxyphenyl)-3-decanone [(-)-(R)-[6]-Gingerol]. <i>Synthesis</i> , 1984, 1984, 702-703.	2.3	13
173	A Modular Approach to a New Class of Monodentate Chiral Phosphorus Ligands and Their Application in Enantioselective Copper-Catalysed Conjugate Additions of Diethylzinc to Cyclohexenone. <i>European Journal of Organic Chemistry</i> , 2004, 2004, 3557-3565.	2.4	13
174	Chiral β -sulphinyl hydrazones as effective reagents for stereoselective aldol-type condensation. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1985, 251-254.	0.9	12
175	Festphasensynthese von vinylogen Sulfonylpeptiden. <i>Angewandte Chemie</i> , 1995, 107, 1892-1893.	2.0	12
176	Mechanistic insights from ab initio calculations on a nitrogen analogue of the boron-mediated aldol reaction. <i>Tetrahedron</i> , 1995, 51, 4853-4866.	1.9	12
177	Improving C=N Bond Reductions with (Cyclopentadienone)iron Complexes: Scope and Limitations. <i>European Journal of Organic Chemistry</i> , 2019, 2019, 647-654.	2.4	12
178	Acceleration of hemiacetal cleavage through hydrogen bonding: a new synthetic catalyst with balanced conformational flexibility and preorganization. <i>Journal of Organic Chemistry</i> , 1991, 56, 3201-3203.	3.2	11
179	Synthesis of tertiary acylolins using acyl anions formed from 1,3-dithians. Total synthesis of (A \pm)-(2E)-1,7-dihydroxy-3,7,11-trimethyl-dodeca-2,10-dien-6-one. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1978, 1036-1041.	0.9	10
180	Biosynthesis of ascochitine and synthesis of its biogenetic precursors. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1980, 2549.	0.9	10

#	ARTICLE	IF	CITATIONS
181	Computerunterstütztes Design von chiralen Borenlolaten: Eine neue, hoch enantioselektive Aldolreaktion für Thioacetate und Thiopropionate. <i>Angewandte Chemie</i> , 1993, 105, 1717-1719.	2.0	10
182	Rationally designed chiral enol borinates: Powerful reagents for the stereoselective synthesis of natural products. <i>Pure and Applied Chemistry</i> , 1997, 69, 507-512.	1.9	10
183	Rational Design of Antiangiogenic Helical Oligopeptides Targeting the Vascular Endothelial Growth Factor Receptors. <i>Frontiers in Chemistry</i> , 2019, 7, 170.	3.6	10
184	The Importance of Detail: How Differences in Ligand Structures Determine Distinct Functional Responses in Integrin $\alpha 5 \beta 3$. <i>Chemistry - A European Journal</i> , 2019, 25, 5959-5970.	3.3	10
185	Multimeric Presentation of RGD Peptidomimetics Enhances Integrin Binding and Tumor Cell Uptake. <i>Chemistry - A European Journal</i> , 2020, 26, 7492-7496.	3.3	10
186	Biosynthesis of cochliquinones. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1980, , 2686.	0.9	9
187	Stereoselective synthesis of α -methylene- β -hydroxy- γ -alkoxy esters from α -alkoxy aldehydes. <i>Journal of the Chemical Society Chemical Communications</i> , 1983, , 1112-1113.	2.0	9
188	Novel reverse-turn mimics inhibit farnesyl transferase. <i>Bioorganic and Medicinal Chemistry Letters</i> , 1999, 9, 847-852.	2.2	9
189	Synthesis, Conformational Studies and Binding Properties of Acyclic Receptors for N-Protected Amino Acids and Dipeptides. <i>European Journal of Organic Chemistry</i> , 2001, 2001, 4625.	2.4	9
190	A Highly Stereoselective Total Synthesis of (+)- <i>epi</i> -Dictyostatin. <i>European Journal of Organic Chemistry</i> , 2010, 2010, 5767-5771.	2.4	9
191	Selective O-acylation of unprotected N-benzylserine methyl ester and O,N-acyl transfer in the formation of cyclo[Asp-Ser] diketopiperazines. <i>Tetrahedron</i> , 2010, 66, 9528-9531.	1.9	9
192	Biosynthesis of mycophenolic acid. Oxidation of 6-farnesyl-5,7-dihydroxy-4-methylphthalide in a cell-free preparation from <i>Penicillium brevicompactum</i> . <i>Journal of the Chemical Society Chemical Communications</i> , 1978, , 434.	2.0	8
193	Detection of one symmetrical precursor during the biosynthesis of the fungal metabolite austdiol using [1,2- ^{13}C]acetate and [Me- ^{13}C]methionine. <i>Journal of the Chemical Society Chemical Communications</i> , 1981, , 575.	2.0	8
194	Asymmetric Transfer Hydrogenation of Ketones with Modified Grubbs Metathesis Catalysts: On the Way to a Tandem Process. <i>Advanced Synthesis and Catalysis</i> , 2016, 358, 515-519.	4.3	8
195	Fast Cyclization of a Proline-Derived Self-Immolative Spacer Improves the Efficacy of Carbamate Prodrugs. <i>Angewandte Chemie</i> , 2020, 132, 4205-4210.	2.0	8
196	Biosynthesis of citrinin. <i>Journal of the Chemical Society Chemical Communications</i> , 1980, , 1132.	2.0	7
197	Enantioselective Rh-Catalyzed Addition of Arylboronic Acids to N-Tosylarylimines. <i>Synlett</i> , 2007, 2007, 2213-2216.	1.8	7
198	Chiral (salen)Co(III)(N-benzyl-L-serine)-derived phosphites: monodentate P-ligands for enantioselective catalytic applications. <i>Tetrahedron: Asymmetry</i> , 2009, 20, 1185-1190.	1.8	7

#	ARTICLE	IF	CITATIONS
199	A trifunctional self-immolative spacer enables drug release with two non-sequential enzymatic cleavages. <i>Chemical Communications</i> , 2021, 57, 7778-7781.	4.1	7
200	Stereoselective aldol reactions of $\hat{\text{I}}^3$ -thiobutyrolactone: The benzaldehyde anomaly. <i>Tetrahedron Letters</i> , 1990, 31, 2453-2456.	1.4	6
201	Semisyntese von Taxol: eine hoehenantio- und diastereoselektive Synthese der Seitenkette und eine neue Methode zur Esterbildung an C13 unter Verwendung von Thioestern. <i>Angewandte Chemie</i> , 1996, 108, 1809-1812.	2.0	6
202	Development and Biochemical Characterization of Self-Immolative Linker Containing GnRH-III-Drug Conjugates. <i>International Journal of Molecular Sciences</i> , 2022, 23, 5071.	4.1	6
203	Peptide bond formation using an enzyme mimicking approach. <i>Tetrahedron Letters</i> , 1990, 31, 2929-2932.	1.4	5
204	Advanced Pyrrolidine-Carbamate Self-Immolative Spacer with Tertiary Amine Handle Induces Superfast Cyclative Drug Release. <i>ChemMedChem</i> , 2022, 17, .	3.2	5
205	Biosynthesis of cochlioquinones. <i>Journal of the Chemical Society Chemical Communications</i> , 1978, , 679.	2.0	4
206	Biosynthesis of ascochitine: incorporation studies with advanced precursors. <i>Journal of the Chemical Society Chemical Communications</i> , 1979, , 492.	2.0	4
207	A Practical Synthesis of the C1-C9 Fragment of Dictyostatin. <i>Synthesis</i> , 2008, 2008, 2158-2162.	2.3	4
208	Enantioselective synthesis of 1-vinyltetrahydroisoquinolines through palladium-catalysed intramolecular allylic amidation with chiral PhthalaPhos ligands. <i>Tetrahedron: Asymmetry</i> , 2014, 25, 844-850.	1.8	4
209	A dimeric bicyclic RGD ligand displays enhanced integrin binding affinity and strong biological effects on U-373 MG glioblastoma cells. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 8913-8917.	2.8	4
210	Riding the Wave of Monodentate Ligand Revival: From the A/B Concept to Noncovalent Interactions. <i>Chemical Record</i> , 2016, 16, 2544-2560.	5.8	3
211	Computer-Assisted Design and Synthetic Applications of Chiral Enol Borinates: Novel, Highly Enantioselective Aldol Reagents. <i>Journal of the Brazilian Chemical Society</i> , 1998, 9, .	0.6	3
212	Biosynthesis of austdiol and synthesis of a deuterium labelled biogenetic precursor. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1983, , 2745.	0.9	2
213	Stereoselectivity in (Z)-Vinylmetal Additions to the Dictyostatin C1-C9 $\hat{\text{I}}^2$ -Silyloxy Aldehyde. <i>European Journal of Organic Chemistry</i> , 2012, 2012, 144-153.	2.4	2
214	The Italian Chemical Society Is 100 Years Old (Eur. J. Org. Chem. 19/2009). <i>European Journal of Organic Chemistry</i> , 2009, 2009, 3095-3097.	2.4	1
215	Functionalized 2-Hydroxybenzaldehyde-PEG Modules as Portable Tags for the Engagement of Protein Lysine $\hat{\mu}$ -Amino Groups. <i>European Journal of Organic Chemistry</i> , 2021, 2021, 1763-1767.	2.4	1
216	Hydrogen-Bonding Donor/Acceptor Scales in -Sulfonamidopeptides. <i>Chemistry - A European Journal</i> , 1998, 4, 1924-1931.	3.3	1

#	ARTICLE	IF	CITATIONS
217	Origins of stereoselectivity in the addition of allyl- and crotylboronates to aldehydes: The development and application of a force field model of the transition state. AIP Conference Proceedings, 1995, , .	0.4	0
218	Synthesis and Screening of New Chiral Ligands for the Copper-Catalyzed Enantioselective Allylic Substitution.. ChemInform, 2003, 34, no.	0.0	0
219	A Catalytic and Enantioselective Desymmetrization of meso Cyclic Allylic Bisdiethylphosphates with Organozinc Reagents.. ChemInform, 2003, 34, no.	0.0	0
220	Combinatorial Libraries of Chiral Ligands for Enantioselective Catalysis. ChemInform, 2003, 34, no.	0.0	0
221	Copper Phosphoramidite-Catalyzed Enantioselective Desymmetrization of meso-Cyclic Allylic Bisdiethyl Phosphates. ChemInform, 2004, 35, no.	0.0	0
222	Copper-Catalyzed, Enantioselective Desymmetrization of meso Cyclic Allylic Bis(diethyl phosphates) with Organozinc Reagents.. ChemInform, 2005, 36, no.	0.0	0
223	The Italian Chemical Society Is 100 Years Old (Eur. J. Org. Chem. 18/2009). European Journal of Inorganic Chemistry, 2009, 2009, 2567-2569.	2.0	0
224	Inside Cover: Cyclic RGD Peptidomimetics Containing Bifunctional Diketopiperazine Scaffolds as New Potent Integrin Ligands (Chem. Eur. J. 20/2012). Chemistry - A European Journal, 2012, 18, 6106-6106.	3.3	0
225	Frontispiece: Multivalency Increases the Binding Strength of RGD Peptidomimetic Paclitaxel Conjugates to Integrin $\alpha_5\beta_1$. Chemistry - A European Journal, 2017, 23, .	3.3	0
226	Frontispiece: Innovative Linker Strategies for Tumor-Targeted Drug Conjugates. Chemistry - A European Journal, 2019, 25, .	3.3	0
227	TUMOR TARGETING WITH INTEGRIN LIGAND - DRUG CONJUGATES. Istituto Lombardo - Accademia Di Scienze E Lettere - Rendiconti Di Scienze, 2020, , .	0.0	0
228	CONFORMATIONAL CONTROL IN NATURAL PRODUCTS SYNTHESIS. , 1984, , 199-209.		0