

Mikael Bols

List of Publications by Year in descending order

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times ranked

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citing authors

#	ARTICLE	IF	CITATIONS
1	Taming of the DIBAL Promoted Debenzylation of α -Cyclodextrin. Kinetics, Substituent Effects and Efficient Synthesis of Lings Tetrol**. Chemistry - A European Journal, 2022, 28, e202200564.	1.7	4
2	Silylated Sugars – Synthesis and Properties. Synlett, 2022, 33, 415-428.	1.0	1
3	Attachment of cyclodextrin acids to PEGA resin and study of binding with fluorescence microscopy. Bioorganic and Medicinal Chemistry Letters, 2021, 43, 128060.	1.0	2
4	Imino- and Azasugar Protonation Inside Human Acid α -Glucosidase, the Enzyme that is Defective in Gaucher Disease. Angewandte Chemie, 2020, 132, 10552-10555.	1.6	6
5	Synthesis of Isofagomine Derivatives as New Fluorescence pH Indicators/Glycosidase Inhibitors. European Journal of Organic Chemistry, 2020, 2020, 3989-3996.	1.2	4
6	Imino- and Azasugar Protonation Inside Human Acid α -Glucosidase, the Enzyme that is Defective in Gaucher Disease. Angewandte Chemie - International Edition, 2020, 59, 10466-10469.	7.2	16
7	Design and Combinatorial Development of Shield-1 Peptide Mimetics Binding to Destabilized FKBP12. ACS Combinatorial Science, 2020, 22, 156-164.	3.8	4
8	Synthesis of Shld Derivatives, Their Binding to the Destabilizing Domain, and Influence on Protein Accumulation in Transgenic Plants. Journal of Medicinal Chemistry, 2019, 62, 5191-5216.	2.9	5
9	Zinc and Copper Complexes of Methylated α - and Tetraaminocyclodextrins. European Journal of Organic Chemistry, 2019, 2019, 1083-1091.	1.2	5
10	An Inexpensive and Scalable Synthesis of Shld. Journal of Organic Chemistry, 2018, 83, 6050-6055.	1.7	11
11	On the nature of the electronic effect of multiple hydroxyl groups in the 6-membered ring – the effects are additive but steric hindrance plays a role too. Organic and Biomolecular Chemistry, 2017, 15, 1164-1173.	1.5	11
12	Glycosyl Fluorides as Intermediates in $\text{BF}_3 \cdot \text{OEt}_2$ -Promoted Glycosylation with Trichloroacetimidates. European Journal of Organic Chemistry, 2017, 2017, 1281-1284.	1.2	9
13	Conformationally superarmed S-ethyl glycosyl donors as effective building blocks for chemoselective oligosaccharide synthesis in one pot. Organic and Biomolecular Chemistry, 2017, 15, 559-563.	1.5	12
14	Determination of protonation states of iminosugar-enzyme complexes using photoinduced electron transfer. Chemical Science, 2017, 8, 7383-7393.	3.7	17
15	Artificial Metallooxidases from Cyclodextrin Diacids. Chemistry - A European Journal, 2017, 23, 13766-13775.	1.7	21
16	Silyl-protective groups influencing the reactivity and selectivity in glycosylations. Beilstein Journal of Organic Chemistry, 2017, 13, 93-105.	1.3	59
17	C-H Functionalization on Carbohydrates. European Journal of Organic Chemistry, 2016, 2016, 2740-2756.	1.2	34
18	Selenoureido-iminosugars: A new family of multitarget drugs. European Journal of Medicinal Chemistry, 2016, 123, 155-160.	2.6	27

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19	Super arming of a glycosyl donor using a molecular lever. <i>Tetrahedron Letters</i> , 2016, 57, 35-38.	0.7	5
20	Exploring the relationship between the conformation and pK _a : can a pK _a value be used to determine the conformational equilibrium?. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 3116-3121.	1.5	13
21	Î ² -Mannosylation with 4,6-benzylidene protected mannosyl donors without preactivation. <i>Chemical Communications</i> , 2015, 51, 13283-13285.	2.2	36
22	Mechanisms of Glycosylation Reactions Studied by Low-Temperature Nuclear Magnetic Resonance. <i>Chemical Reviews</i> , 2015, 115, 4963-5013.	23.0	142
23	Synthesis of <sc>l</sc>-Hexoses. <i>Chemical Reviews</i> , 2015, 115, 3615-3676.	23.0	68
24	A fluorescence study of isofagomine protonation in Î ² -glucosidase. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 6562-6566.	1.5	10
25	Synthesis of All Eight <sc>L</sc>-Glycopyranosyl Donors Using C₆H Activation. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 13889-13893.	7.2	31
26	Synthesis of All Eight Stereoisomeric 6-Deoxy-<sc>L</sc>-Hexopyranosyl Donors - Trends in Using Stereoselective Reductions or Mitsunobu Epimerizations. <i>European Journal of Organic Chemistry</i> , 2014, 2014, 7924-7939.	1.2	34
27	Total Synthesis of Five Lipoteichoic acids of <i>Clostridium difficile</i>. <i>Chemistry - A European Journal</i> , 2014, 20, 13511-13516.	1.7	8
28	Î ² -Selective Mannosylation with a 4,6-Silylene-Tethered Thiomannosyl Donor. <i>Organic Letters</i> , 2014, 16, 1116-1119.	2.4	67
29	Cyclodextrin-based artificial oxidases with high rate accelerations and selectivity. <i>Tetrahedron Letters</i> , 2014, 55, 2304-2307.	0.7	7
30	Superarming of Glycosyl Donors by Combined Neighboring and Conformational Effects. <i>Organic Letters</i> , 2013, 15, 4904-4907.	2.4	26
31	Enzyme inhibition by iminosugars: Analysis and insight into the glycosidase-iminosugar dependency of pH. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 4755-4761.	1.4	15
32	Influence of O6 in Mannosylations Using Benzylidene Protected Donors: Stereoelectronic or Conformational Effects?. <i>Journal of Organic Chemistry</i> , 2013, 78, 2191-2205.	1.7	41
33	Artificial enzyme activity from cyclodextrins with cyanohydrins on the secondary rim. <i>Tetrahedron Letters</i> , 2013, 54, 2458-2461.	0.7	11
34	Conformationally Armed 3,6-Tethered Glycosyl Donors: Synthesis, Conformation, Reactivity, and Selectivity. <i>Journal of Organic Chemistry</i> , 2013, 78, 7234-7248.	1.7	56
35	Synthesis and Thermotropic Phase Behavior of Four Glycoglycerolipids. <i>Molecules</i> , 2013, 18, 13546-13573.	1.7	6
36	Total synthesis of biologically active lipoteichoic acids. <i>Arkivoc</i> , 2013, 2013, 249-275.	0.3	0

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37	Easy Access to α -Mannosides and α -Galactosides by Using $C_{12}H_{10}$ Activation of the Corresponding 6-Deoxysugars. <i>Angewandte Chemie - International Edition</i> , 2012, 51, 12285-12288.	7.2	50
38	Synthesis of Tin-Containing Cyclodextrins as Potential Enzyme Models. <i>European Journal of Organic Chemistry</i> , 2012, 2012, 6383-6389.	1.2	10
39	Artificial enzymes based on cyclodextrin with phenol as the catalytic group. <i>Tetrahedron Letters</i> , 2012, 53, 5023-5026.	0.7	8
40	Two Diastereomeric Artificial Enzymes with Different Catalytic Activity. <i>European Journal of Organic Chemistry</i> , 2012, 2012, 5366-5372.	1.2	7
41	Rhamnosylation: Diastereoselectivity of Conformationally Armed Donors. <i>Journal of Organic Chemistry</i> , 2012, 77, 5559-5568.	1.7	40
42	Synthesis of Cyclodextrins with Carboxylic Acids at the Secondary Rim and an Evaluation of Their Properties as Chemzymes for Glycoside Hydrolysis. <i>European Journal of Organic Chemistry</i> , 2012, 2012, 4063-4070.	1.2	10
43	β -Cyclodextrin as a mimetic of the natural GFP-chromophore environment. <i>Tetrahedron Letters</i> , 2012, 53, 973-976.	0.7	11
44	Conversion of d-glucose into 5-hydroxymethylfurfural (HMF) using zeolite in [Bmim]Cl or tetrabutylammonium chloride (TBAC)/CrCl ₂ . <i>Tetrahedron Letters</i> , 2012, 53, 983-985.	0.7	70
45	A uronic acid analogue of isofagomine lactam as a nanomolar glucuronidase inhibitor. <i>Tetrahedron Letters</i> , 2012, 53, 2045-2047.	0.7	4
46	An Isofagomine Analogue with an Amidine at the Pseudoanomeric Position. <i>Organic Letters</i> , 2011, 13, 2908-2911.	2.4	15
47	The Influence of Neighboring Group Participation on the Hydrolysis of 2-O-Substituted Methyl Glucopyranosides. <i>Organic Letters</i> , 2011, 13, 5956-5959.	2.4	32
48	A study of anhydrocelluloses " Is a cellulose structure with residues in a 1C ₄ -conformation more prone to hydrolysis?. <i>Organic and Biomolecular Chemistry</i> , 2011, 9, 7525.	1.5	8
49	Simple cyclodextrin aldehydes as excellent artificial oxidases. <i>Journal of Inclusion Phenomena and Macrocyclic Chemistry</i> , 2011, 69, 397-402.	1.6	10
50	Substrate structure governs maximum rate of catalysis exerted by cyclodextrin oxidase chemzymes. <i>Journal of Inclusion Phenomena and Macrocyclic Chemistry</i> , 2011, 69, 417-423.	1.6	2
51	Regioselective difunctionalization of cyclodextrins. <i>Journal of Inclusion Phenomena and Macrocyclic Chemistry</i> , 2011, 69, 425-431.	1.6	6
52	Synthesis of 5-Bromomethylfurfural from Cellulose as a Potential Intermediate for Biofuel. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 1266-1270.	1.2	43
53	Cyclodextrin Ketones with the Catalytic Group at the Secondary Rim and Their Effectiveness in Enzyme-Like Epoxidation of Stilbenes. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 2339-2345.	1.2	16
54	Recognition of Peptides by Cyclodextrin Trimers. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 5279-5290.	1.2	22

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55	3-Deoxyglucosone is an Intermediate in the Formation of Furfurals from D-Glucose.. ChemSusChem, 2011, 4, 1049-1051.	3.6	83
56	Direct Experimental Evidence for the High Chemical Reactivity of 2,5- and 2,6-Xylopyranosides Adopting a ^{2,5} B Conformation in Glycosyl Transfer. Chemistry - A European Journal, 2011, 17, 7345-7356.	1.7	14
57	Quantifying the Electronic Effects of Carbohydrate Hydroxy Groups by Using Aminosugar Models. Chemistry - A European Journal, 2011, 17, 7080-7086.	1.7	42
58	Glycosyl donors in "unusual" conformations influence on reactivity and selectivity. Comptes Rendus Chimie, 2011, 14, 17-43.	0.2	66
59	Synthesis of monofluorinated isofagomine analogues and evaluation as glycosidase inhibitors. Journal of Fluorine Chemistry, 2011, 132, 838-845.	0.9	15
60	Amino-Acetone-Bridged Cyclodextrins - Artificial Alcohol Oxidases. European Journal of Organic Chemistry, 2010, 2010, 157-167.	1.2	15
61	The Grignard Reaction of Cyclodextrin-Aldehydes Revisited: A Study of the Stereoselectivity Upon Addition of Organometallic Reagents to Aldehydes and Ketones. European Journal of Organic Chemistry, 2010, 2010, 3883-3896.	1.2	13
62	Substantial Spatial Flexibility and Hydrogen Bonding within the Catalysis Exerted by Cyclodextrin Artificial Glycosidases. European Journal of Organic Chemistry, 2010, 2010, 3487-3500.	1.2	11
63	Quantifying Electronic Effects of Common Carbohydrate Protecting Groups in a Piperidine Model System. Chemistry - A European Journal, 2010, 16, 13982-13994.	1.7	48
64	Binding and Orientation of Tricyclic Antidepressants within the Central Substrate Site of the Human Serotonin Transporter. Journal of Biological Chemistry, 2010, 285, 8363-8374.	1.6	85
65	Cyclodextrins as Supramolecular Organo-Catalysts. Current Organic Chemistry, 2010, 14, 1380-1398.	0.9	34
66	First Step in Chemical Preparation of Metal Nanogaps Bridged by Thiol End-Capped Molecular Wires. Journal of Physical Chemistry B, 2010, 114, 11771-11777.	1.2	9
67	Substrate control through per-O-methylation of cyclodextrin acids. Chemical Communications, 2010, 46, 7769.	2.2	7
68	Cyclodextrin derivatives that display Enzyme Catalysis. Trends in Glycoscience and Glycotechnology, 2009, 21, 309-323.	0.0	24
69	Cyclodextrin Aldehydes are Oxidase Mimics. ChemBioChem, 2009, 10, 2494-2503.	1.3	27
70	Synthesis and biological evaluation of potent glycosidase inhibitors: 4-deoxy-4,4-difluoroisofagomine and analogues. Tetrahedron, 2009, 65, 3717-3727.	1.0	17
71	An extended study of dimeric phenyl tropanes. Bioorganic and Medicinal Chemistry, 2009, 17, 4900-4909.	1.4	11
72	Selective Discrimination of Cyclodextrin Diols Using Cyclic Sulfates. Organic Letters, 2009, 11, 1983-1985.	2.4	29

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73	Difluoromethylenated polyhydroxylated pyrrolidines: facile synthesis, crystal structure and biological evaluation. <i>Future Medicinal Chemistry</i> , 2009, 1, 991-997.	1.1	2
74	Cyclodextrin ketones as oxidation catalysts: Investigation of bridged derivatives. <i>Organic and Biomolecular Chemistry</i> , 2009, 7, 933.	1.5	20
75	Artificial enzymes, "Chemzymes": current state and perspectives. <i>Applied Microbiology and Biotechnology</i> , 2008, 81, 1-11.	1.7	113
76	Hydrolysis of Toxic Natural Glucosides Catalyzed by Cyclodextrin Dicyanohydrins. <i>European Journal of Organic Chemistry</i> , 2008, 2008, 745-752.	1.2	23
77	Effective synthesis of negatively charged cyclodextrins. Selective access to phosphate cyclodextrins. <i>Tetrahedron</i> , 2008, 64, 7587-7593.	1.0	12
78	Binding of Serotonin to the Human Serotonin Transporter. Molecular Modeling and Experimental Validation. <i>Journal of the American Chemical Society</i> , 2008, 130, 3853-3865.	6.6	123
79	Conformationally armed glycosyl donors: reactivity quantification, new donors and one pot reactions. <i>Chemical Communications</i> , 2008, , 2465.	2.2	77
80	"Super Armed" Glycosyl Donors: A Conformational Arming of Thioglycosides by Silylation. <i>Journal of the American Chemical Society</i> , 2007, 129, 9222-9235.	6.6	168
81	Anomer-Selective Glycosidase Inhibition by 2-N-Alkylated 1-Azafagomines. <i>ChemBioChem</i> , 2007, 8, 657-661.	1.3	14
82	Going to Extremes: "Super" Armed Glycosyl Donors in Glycosylation Chemistry. <i>Chemistry - A European Journal</i> , 2007, 13, 7576-7582.	1.7	85
83	Synthesis of Some Trifluoromethylated Cyclodextrin Derivatives and Analysis of Their Properties as Artificial Glycosidases and Oxidases. <i>European Journal of Organic Chemistry</i> , 2007, 2007, 704-710.	1.2	48
84	Active Site Protonation of 1-Azafagomine in Glucosidases Studied by Solid-State NMR Spectroscopy. <i>European Journal of Organic Chemistry</i> , 2007, 2007, 1735-1742.	1.2	8
85	New cup-shaped β -cyclodextrin derivatives and a study of their catalytic properties in oxidation reactions. <i>Tetrahedron</i> , 2007, 63, 8872-8880.	1.0	22
86	Synthesis, inhibition and binding of simple non-nitrogen inhibitors of monoamine transporters. <i>Bioorganic and Medicinal Chemistry</i> , 2007, 15, 4159-4174.	1.4	20
87	QSAR studies and pharmacophore identification for arylsubstituted cycloalkenecarboxylic acid methyl esters with affinity for the human dopamine transporter. <i>Bioorganic and Medicinal Chemistry</i> , 2007, 15, 5262-5274.	1.4	11
88	High dopamine transporter selectivity can be displayed by remarkably simple non-nitrogen containing inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007, 17, 6019-6025.	1.0	1
89	Stereoelectronic Substituent Effects. <i>Accounts of Chemical Research</i> , 2006, 39, 259-265.	7.6	139
90	Synthesis and Biological Evaluation of Glycosidase Inhibitors: "gem-Difluoromethylenated Nojirimycin Analogues. <i>Journal of Medicinal Chemistry</i> , 2006, 49, 2989-2997.	2.9	45

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91	Cyclodextrin derivatives with cyanohydrin and carboxylate groups as artificial glycosidases. Canadian Journal of Chemistry, 2006, 84, 650-658.	0.6	19
92	Very High Rate Enhancement of Benzyl Alcohol Oxidation by an Artificial Enzyme. Angewandte Chemie - International Edition, 2006, 45, 4590-4593.	7.2	86
93	The role of the active site Zn in the catalytic mechanism of the GH38 Golgi α -mannosidase II: Implications from noeuromycin inhibition. Biocatalysis and Biotransformation, 2006, 24, 55-61.	1.1	13
94	Stereochemical substituent effects: investigation of the cyano, amide and carboxylate group. Tetrahedron, 2005, 61, 115-122.	1.0	11
95	Artificial Epoxidase II. Synthesis of Cyclodextrin Ketoesters and Epoxidation of Alkenes. European Journal of Organic Chemistry, 2005, 2005, 2734-2739.	1.2	32
96	Artificial Glycosyl Phosphorylases. Chemistry - A European Journal, 2005, 11, 5094-5101.	1.7	40
97	Safe Radical Azidonation Using Polystyrene Supported Diazidoiodate(I).. ChemInform, 2005, 36, no.	0.1	0
98	Cyclodextrins Containing an Acetone Bridge. Synthesis and Study as Epoxidation Catalysts.. ChemInform, 2005, 36, no.	0.1	0
99	Radical Substitution with Azide: TMSN ₃ Ph(OAc) ₂ as a Substitute of IN ₃ .. ChemInform, 2005, 36, no.	0.1	0
100	Safe radical azidonation using polystyrene supported diazidoiodate(I). Tetrahedron, 2005, 61, 123-127.	1.0	33
101	Supramolecular Oxidation of Anilines Using Hydrogen Peroxide as Stoichiometric Oxidant. Journal of the American Chemical Society, 2005, 127, 17578-17579.	6.6	62
102	Four Orders of Magnitude Rate Increase in Artificial Enzyme-Catalyzed Aryl Glycoside Hydrolysis. Journal of Organic Chemistry, 2005, 70, 7217-7226.	1.7	45
103	Radical substitution with azide: TMSN ₃ Ph(OAc) ₂ as a substitute of IN ₃ . Organic and Biomolecular Chemistry, 2005, 3, 816-822.	1.5	92
104	Remarkable Supramolecular Catalysis of Glycoside Hydrolysis by a Cyclodextrin Cyanohydrin. Journal of the American Chemical Society, 2005, 127, 3238-3239.	6.6	110
105	On the electronic effects of OH groups. Synthesis and investigation of tetrahydroxylated azabicycloheptanes. Organic and Biomolecular Chemistry, 2005, 3, 1514.	1.5	8
106	Benzyl 4-C-nitrosomethyl- β -D-arabinopyranoside. Acta Crystallographica Section E: Structure Reports Online, 2004, 60, o660-o661.	0.2	0
107	The C-4 Configuration as a Probe for the Study of Glycosidation Reactions. European Journal of Organic Chemistry, 2004, 2004, 323-329.	1.2	25
108	Radical Azidonation of Aldehydes.. ChemInform, 2004, 35, no.	0.1	0

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109	An artificial enzyme that catalyzes hydrolysis of aryl glycosides. <i>Tetrahedron Letters</i> , 2004, 45, 8709-8711.	0.7	44
110	Conformational Effects on Glycoside Reactivity: A Study of the High Reactive Conformer of Glucose. <i>Journal of the American Chemical Society</i> , 2004, 126, 12374-12385.	6.6	82
111	Two- and Three Dimensional Combinatorial Chemistry from Multicomponent Grignard Reagents. <i>ACS Combinatorial Science</i> , 2004, 6, 509-519.	3.3	13
112	Synthesis and Chemistry of Noeuromycin and Isofagomine Analogues. <i>Journal of Carbohydrate Chemistry</i> , 2004, 23, 223-238.	0.4	13
113	Cyclodextrins containing an acetone bridge. Synthesis and study as epoxidation catalysts. <i>Organic and Biomolecular Chemistry</i> , 2004, 2, 3476.	1.5	46
114	Combinatorial synthesis of benzotropine libraries and their evaluation as monoamine transporter inhibitors. <i>Organic and Biomolecular Chemistry</i> , 2004, 2, 2861-2869.	1.5	24
115	The Disarming Effect of the 4,6-Acetal Group on Glycoside Reactivity: A Torsional or Electronic?. <i>Journal of the American Chemical Society</i> , 2004, 126, 9205-9213.	6.6	223
116	Synthesis of furan 4- ϵ -thio-C-nucleosides, their methylsulfonium and sulfoxide derivatives. Evaluation as glycosidase inhibitors. <i>Tetrahedron</i> , 2003, 59, 2801-2809.	1.0	11
117	Steric Effects Are Not the Cause of the Rate Difference in Hydrolysis of Stereoisomeric Glycosides. <i>Organic Letters</i> , 2003, 5, 3419-3421.	2.4	62
118	Radical Azidonation of Aldehydes. <i>Journal of Organic Chemistry</i> , 2003, 68, 9453-9455.	1.7	85
119	Isofagomine lactams, synthesis and enzyme inhibition. <i>Organic and Biomolecular Chemistry</i> , 2003, 1, 282-287.	1.5	30
120	Aziridines as a structural motif to conformational restriction of azasugars. <i>Organic and Biomolecular Chemistry</i> , 2003, 1, 478-482.	1.5	13
121	An Azido-Hanessian Reaction. <i>Synlett</i> , 2002, 2002, 1111-1112.	1.0	22
122	Equatorial Contra Axial Polar Substituents. The Relation of a Chemical Reaction to Stereochemical Substituent Constants. <i>Journal of Organic Chemistry</i> , 2002, 67, 8970-8974.	1.7	48
123	Recent Developments of Transition-State Analogue Glycosidase Inhibitors of Non-Natural Product Origin. <i>Chemical Reviews</i> , 2002, 102, 515-554.	23.0	673
124	Unusual hydrogen-bonding differences in stereoisomeric 6-C-alkylated cyclodextrins Electronic supplementary information (ESI) available: IR spectra of compounds 5 and 7 in CH ₂ Cl ₂ solution. See http://www.rsc.org/suppdata/p1/b2/b207033m/ . <i>Journal of the Chemical Society, Perkin Transactions 1</i> , 2002, , 2880-2885.	1.3	10
125	Investigation of the base promoted tandem syn-elimination- ϵ -Favorskii rearrangement of levoglucosan sulfonates. <i>Journal of the Chemical Society, Perkin Transactions 1</i> , 2002, , 1297-1301.	1.3	7
126	Synthesis and evaluation of sulfonium analogues of isofucofagomine as glycosidase inhibitors. <i>Journal of the Chemical Society, Perkin Transactions 1</i> , 2002, , 1242-1246.	1.3	14

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127	Synthesis and investigation of L-fuco- and D-glucurono-azafagomine. <i>Journal of the Chemical Society, Perkin Transactions 1</i> , 2002, , 1190-1198.	1.3	37
128	Solution-phase combinatorial synthesis using multicomponent Grignard reagents. <i>Journal of the Chemical Society, Perkin Transactions 1</i> , 2002, , 503-508.	1.3	10
129	Substitution of $\hat{1}\pm$ -azido ethers with Grignard reagents and its use in combinatorial chemistry. <i>Journal of the Chemical Society, Perkin Transactions 1</i> , 2002, , 509-512.	1.3	14
130	Stereoelectronic Substituent Effects in Polyhydroxylated Piperidines and Hexahydropyridazines. <i>Chemistry - A European Journal</i> , 2002, 8, 1218.	1.7	85
131	A bioisosteric oligosaccharide mimetic based on isofagomine-type monomers. <i>Journal of the Chemical Society, Perkin Transactions 1</i> , 2001, , 2764-2773.	1.3	4
132	Isogalactofagomine lactam. A neutral nanomolar galactosidase inhibitor. <i>Journal of the Chemical Society, Perkin Transactions 1</i> , 2001, , 1584-1585.	1.3	13
133	Noeuromycin, 1A Glycosyl Cation Mimic that Strongly Inhibits Glycosidases. <i>Journal of the American Chemical Society</i> , 2001, 123, 5116-5117.	6.6	95
134	Garner's aldehyde. <i>Journal of the Chemical Society, Perkin Transactions 1</i> , 2001, , 2136-2157.	1.3	128
135	Synthesis of 1-azagalactofagomine, a potent galactosidase inhibitor. <i>Journal of the Chemical Society, Perkin Transactions 1</i> , 2001, , 905-909.	1.3	32
136	Accurate Determination of Rate Constants of Very Slow, Tight-Binding Competitive Inhibitors by Numerical Solution of Differential Equations, Independently of Precise Knowledge of the Enzyme Concentration. <i>Analytical Biochemistry</i> , 2001, 295, 186-193.	1.1	6
137	Synthesis of 5-Azacastanospermine, a Conformationally Restricted Azafagomine Analogue. <i>Chemistry - A European Journal</i> , 2001, 7, 2324-2331.	1.7	18
138	Efficient Synthesis of Isofagomine and Noeuromycin. <i>Chemistry - A European Journal</i> , 2001, 7, 3744-3747.	1.7	43
139	Radical Azidation of Benzylic Positions with Iodonium Azide. <i>Angewandte Chemie - International Edition</i> , 2001, 40, 623-625.	7.2	57
140	A Free-Energy Relationship between the Rate of Acidic Hydrolysis of Glycosides and the pKa of Isofagomines. <i>Angewandte Chemie - International Edition</i> , 2001, 40, 3447-3449.	7.2	63
141	Specific Glycosidase Activity Isolated from a Random Phage Display Antibody Library. <i>Biotechnology Progress</i> , 2001, 17, 197-202.	1.3	5
142	Slow inhibition of almond $\hat{1}2$ -glucosidase by azasugars: determination of activation energies for slow binding. <i>BBA - Proteins and Proteomics</i> , 2001, 1545, 207-215.	2.1	10
143	Synthesis of an $\hat{1}\epsilon^2$ -Azasugar Analogue of Maltose. <i>Synthesis</i> , 2001, 2001, 0339-0342.	1.2	8
144	Radical Azidation of Benzylic Positions with Iodonium Azide. <i>Angewandte Chemie - International Edition</i> , 2001, 40, 623-625.	7.2	3

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145	Investigation of the slow inhibition of almond Î²-glucosidase and yeast isomaltase by 1-azasugar inhibitors: evidence for the "direct binding" model. <i>Biochemical Journal</i> , 2000, 349, 211.	1.7	17
146	Investigation of the slow inhibition of almond Î²-glucosidase and yeast isomaltase by 1-azasugar inhibitors: evidence for the "direct binding" model. <i>Biochemical Journal</i> , 2000, 349, 211-215.	1.7	25
147	Enantiospecific Synthesis of 1-Azafagomine. <i>Chemistry - A European Journal</i> , 2000, 6, 278-287.	1.7	63
148	A New Method for the Deprotection of Benzyl Ethers or the Selective Protection of Alcohols. <i>Chemistry - A European Journal</i> , 2000, 6, 1140-1146.	1.7	15
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