Yuichi Yoshimura

List of Publications by Year in descending order

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257429 330122 1,907 94 24 37 citations h-index g-index papers 113 113 113 1301 docs citations times ranked citing authors all docs

#	Article	lF	Citations
1	A Novel Synthesis of $2\hat{a}\in -M$ odified $2\hat{a}\in -D$ eoxy- $4\hat{a}\in -T$ thiocytidines from d-Glucose 1. Journal of Organic Chemistry, 1997, 62, 3140-3152.	3.2	102
2	A Novel Synthesis of New Antineoplastic 2â€~-Deoxy-2â€~-substituted-4â€~-thiocytidines. Journal of Organic Chemistry, 1996, 61, 822-823.	3.2	81
3	α-1- <i>C</i> -Butyl-1,4-dideoxy-1,4-imino- <scp>l</scp> -arabinitol as a Second-Generation Iminosugar-Based Oral α-Glucosidase Inhibitor for Improving Postprandial Hyperglycemia. Journal of Medicinal Chemistry, 2012, 55, 10347-10362.	6.4	72
4	A Facile, Alternative Synthesis of 4â€~-Thioarabinonucleosides and Their Biological Activities. Journal of Medicinal Chemistry, 1997, 40, 2177-2183.	6.4	62
5	Antitumor activity of a novel orally effective nucleoside, 1 -(2-deoxy-2-fluoro-4-thio- \hat{l}^2 -d-arabinofuranosyl)cytosine. Cancer Letters, 1998, 129, 103-110.	7.2	59
6	Acceleration Effect of an Allylic Hydroxy Group on Ringâ€Closing Enyne Metathesis of Terminal Alkynes: Scope, Application, and Mechanistic Insights. Chemistry - A European Journal, 2008, 14, 10762-10771.	3.3	56
7	Nucleosides and Nucleotides. 175. Structural Requirements of the Sugar Moiety for the Antitumor Activities of New Nucleoside Antimetabolites, 1-(3- <i>C</i> -Ethynyl-β- <scp>d</scp> - <i>r</i> >i>io>ipo-pentofuranosyl)cytosine and -uracil. Journal of Medicinal Chemistry, 1998, 41, 2892-2902.	6.4	55
8	Docking study and biological evaluation of pyrrolidine-based iminosugars as pharmacological chaperones for Gaucher disease. Organic and Biomolecular Chemistry, 2016, 14, 1039-1048.	2.8	46
9	Synthesis of Novel Iso-4â€~-thionucleosides Using the Mitsunobu Reaction. Journal of Organic Chemistry, 1998, 63, 6891-6899.	3.2	44
10	Synthesis of l-enantiomers of 4′-thioarabinofuranosyl pyrimidine nucleosides. Bioorganic and Medicinal Chemistry Letters, 1998, 8, 989-992.	2.2	42
11	The synthesis and biological evaluation of 1-C-alkyl-l-arabinoiminofuranoses, a novel class of \hat{l}_{\pm} -glucosidase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 738-741.	2.2	39
12	Synthesis and biological activities of 2′-deoxy-2′-fluoro-4′-thioarabinofuranosylpyrimidine and -purine nucleosides. Bioorganic and Medicinal Chemistry, 2000, 8, 1545-1558.	3.0	38
13	Synthesis of both enantiomers of hydroxypipecolic acid derivatives equivalent to 5-azapyranuronic acids and evaluation of their inhibitory activities against glycosidases. Bioorganic and Medicinal Chemistry, 2008, 16, 8273-8286.	3.0	38
14	Stereoselective Synthesis of the β-Anomer of 4'-Thionucleosides Based on Electrophilic Glycosidation to 4-Thiofuranoid Glycals. Journal of Organic Chemistry, 2002, 67, 5919-5927.	3.2	37
15	Nucleosides and Nucleotides. 108. Synthesis and Optical Properties of Syn-Fixed Carbon-Bridged Pyrimidine Cyclonuclesides Chemical and Pharmaceutical Bulletin, 1992, 40, 1761-1769.	1.3	36
16	An Alternative Synthesis of the Antineoplastic Nucleoside 4â€~-ThioFAC and Its Application to the Synthesis of 4â€~-ThioFAG and 4â€~-Thiocytarazid. Journal of Organic Chemistry, 1999, 64, 7912-7920.	3.2	35
17	A practical synthesis of 4′-thioribonucleosides. Tetrahedron Letters, 2006, 47, 591-594.	1.4	31
18	Recent Advances in the Synthesis of Conformationally Locked Nucleosides and Their Success in Probing the Critical Question of Conformational Preferences by Their Biological Targets. Nucleosides, Nucleotides and Nucleic Acids, 2003, 22, 547-557.	1.1	30

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19	Nucleosides and nucleotides. 134. Synthesis of $2\hat{a}\in^2$ -C-alkynyl- $2\hat{a}\in^2$ -deoxy- $1\cdot\hat{l}^2$ -D-arabinofuranosylpyrimidines via radical deoxygenation of tert-propargyl alcohols in the sugar moiety. Tetrahedron, 1994, 50, 10397-10406.	1.9	29
20	Synthesis of 6,3'-methanocytidine,6,3'-methanouridine, and their 2'-deoxyribonucleosides (nucleosides) Tj ETQq0	0 _{1.3} rgBT	/Overlock 10
21	Nucleosides and nucleotides. 102. Stereoselective radical deoxygenation of tert-propargyl alcohols in sugar moiety of pyrimidine nucleosides: synthesis of $2\hat{a}\in^2$ -C-alkynyl- $2\hat{a}\in^2$ -deoxy- $1-\hat{l}^2$ -d-arabinofuranosylpyrimidines. Tetrahedron Letters, 1991, 32, 6003-6006.	1.4	27
22	Design and synthesis of a novel ring-expanded 4′-thio-apio-nucleoside derivatives. Tetrahedron Letters, 2007, 48, 4519-4522.	1.4	26
23	SYNTHESIS AND BIOLOGICAL ACTIVITY OF BRANCHED CHAIN-SUGAR NUCLEOSIDES1. Nucleosides, Nucleotides and Nucleic Acids, 1989, 8, 743-752.	1.1	25
24	Synthesis of 1-(5,6-dihydro-2H-thiopyran-2-yl)uracil by a Pummerer-type thioglycosylation reaction: the regioselectivity of allylic substitution. Tetrahedron, 2009, 65, 9091-9102.	1.9	25
25	Homeostatic and pathogenic roles of <scp>GM</scp> 3 ganglioside molecular species in <scp>TLR</scp> 4 signaling in obesity. EMBO Journal, 2020, 39, e101732.	7.8	25
26	Nucleosides and Nucleotides. 104. Radical and Palladium-Catalyzed Deoxygenation of the Allylic Alcohol Systems in the Sugar Moiety of Pyrimidine Nucleosides $\hat{A}_{,1}$, Nucleosides & Nucleotides, 1992, 11, 197-226.	0.5	24
27	Synthesis and Biological Evaluation of $1\hat{a}\in^2-\langle i\rangle C\langle i\rangle$ -Cyano-Pyrimidine Nucleosides. Nucleosides & Nucleotides, 1996, 15, 305-324.	0.5	24
28	Enantioselective Synthesis of Bicyclo[3.1.0]hexane Carbocyclic Nucleosides via a Lipase-Catalyzed Asymmetric Acetylation. Characterization of an Unusual Acetal Byproduct. Journal of Organic Chemistry, 2002, 67, 5938-5945.	3.2	24
29	Novel Stereoselective Entry to 2â€~1̂²-Carbon-Substituted 2â€~-Deoxy-4â€~-thionucleosides from 4-Thiofuranoid Glycals. Organic Letters, 2004, 6, 2645-2648.	4.6	24
30	Docking and SAR studies of d- and l-isofagomine isomers as human \hat{l}^2 -glucocerebrosidase inhibitors. Bioorganic and Medicinal Chemistry, 2011, 19, 3558-3568.	3.0	24
31	Rationale of 5-(125)I-iodo-4'-thio-2'-deoxyuridine as a potential iodinated proliferation marker. Journal of Nuclear Medicine, 2002, 43, 1218-26.	5.0	24
32	An alternative synthesis of antineoplastic 4′-thiocytidine analogue 4′-thioFAC. Tetrahedron Letters, 1999, 40, 1937-1940.	1.4	23
33	New Synthesis of (\hat{A}_{\pm}) -Isonucleosides. Organic Letters, 2006, 8, 6015-6018.	4.6	23
34	Asymmetric synthesis of 2,5-disubstituted 3-hydroxypyrrolidines based on stereodivergent intramolecular iridium-catalyzed allylic aminations. Organic and Biomolecular Chemistry, 2014, 12, 1983.	2.8	23
35	Development of radioiodinated nucleoside analogs for imaging tissue proliferation: comparisons of six 5-iodonucleosides. Nuclear Medicine and Biology, 2003, 30, 687-696.	0.6	22
36	A New Entry to Carbocyclic Nucleosides: Oxidative Coupling Reaction of Cycloalkenylsilanes with a Nucleobase Mediated by Hypervalent Iodine Reagent. Organic Letters, 2008, 10, 3449-3452.	4.6	22

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37	Nucleosides and nucleotides. LXXXVIII. Synthesis of 6,6'-cyclo-5',6'-dideoxy-1-(.BETAD-allofuranosyl)cytosine and related nucleosides Chemical and Pharmaceutical Bulletin, 1989, 37, 660-664.	1.3	21
38	Asymmetric Synthesis of All Stereoisomers of Isofagomine Using [2,3]-Wittig Rearrangement. Heterocycles, 2007, 72, 633.	0.7	21
39	Nucleosides and nucleotides. 120. Stereoselective radical deoxygenation of tert-alcohols in the sugar moiety of nucleosides: synthesis of 2′,3′-dideoxy-2′-C-methyl2′-C-ethynyl-β-d-threo-pentofuranosyl pyrimidines and adenine as potential antiviral and antitumor agents. Tetrahedron, 1993, 49, 8513-8528.	1.9	20
40	Synthesis of $6,5\hat{a}\in^2$ -C-Cyclouridine by a Novel Tandem Radical 1,6-Hydrogen Transfer and Cyclization Reaction. Synlett, 2007, 2007, 0111-0114.	1.8	20
41	A New Preparation of HomochiralN-Protected 5-Hydroxy-3- piperidenes, Promising Chiral Building Blocks, by Palladium- Catalyzed Deracemization of Their Alkyl Carbonates. Advanced Synthesis and Catalysis, 2007, 349, 685-693.	4.3	20
42	Nucleosides & nucleotides. 118. Synthesis of oligonucleotides containing a novel $2\hat{a}\in^2$ -deoxyuridine analogue that carries an aminoalkyl tether at $1\hat{a}\in^2$ -position; stabilization of duplex formation by an intercalatin group accommodated in the minor groove. Bioorganic and Medicinal Chemistry Letters, 1993, 3, 615-618.	2.2	19
43	Synthesis of 1-(2-Deoxy-2-C-fluoromethyl- \hat{l}^2 -D-arabinofuranosyl)cytosine as a potential antineoplastic agent. Bioorganic and Medicinal Chemistry Letters, 1994, 4, 721-724.	2.2	19
44	Synthesis of all stereoisomers of 3-hydroxypipecolic acid and 3-hydroxy-4,5-dehydropipecolic acid and their evaluation as glycosidase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2008, 18, 1810-1813.	2.2	19
45	Design and Synthesis of Isonucleosides Constructed on a 2-Oxa-6-thiabicyclo[3.2.0]heptane Scaffold. Journal of Organic Chemistry, 2010, 75, 4161-4171.	3.2	16
46	Synthesis and Antiviral Activities of 5-Substituted 1-(2-Deoxy-2-C-methylene-4-thio- \hat{l}^2 -D-erythro-pentofuranosyl)uracilst. Nucleosides, Nucleotides and Nucleic Acids, 1998, 17, 65-79.	1.1	15
47	Lithiation at the 6-Position of Uridine with Lithium Hexamethyldisilazide:  Crucial Role of Temporary Silylation. Organic Letters, 2004, 6, 1793-1795.	4.6	15
48	A Facile Synthesis of Fully Protected meso-Diaminopimelic Acid (DAP) and Its Application to the Preparation of Lipophilic N-Acyl iE-DAP. Molecules, 2013, 18, 1162-1173.	3.8	15
49	Anti-herpesvirus activity profile of 4′-thioarabinofuranosyl purine and uracil nucleosides and activity of 1-β-d-2′-fluoro-4′-thioarabinofuranosyl guanine and 2,6-diaminopurine against clinical isolates of human cytomegalovirus. Antiviral Research, 1998, 39, 129-137.	4.1	14
50	Nucleosides and Nucleotides. 177. 9-(6,7-Dideoxy- $\hat{1}^2$ -d-allo-hept-5- ynofuranosyl)adenine: \hat{A} A Selective and Potent Ligand for P3Purinoceptor-like Protein1. Journal of Medicinal Chemistry, 1998, 41, 2676-2678.	6.4	14
51	Comparison of 1-(2-deoxy-2-fluoro-4-thio- \hat{l}^2 -d-arabinofuranosyl)cytosine with gemcitabine in its antitumor activity. Cancer Letters, 1999, 144, 177-182.	7.2	14
52	Synthesis of $6,1\hat{a}\in^2$ -propanouridine, fixed in syn-conformation by a spiro-carbon bridge. Tetrahedron Letters, 1991, 32, 4549-4552.	1.4	13
53	Nucleosides and nucleotides. 142. an alternative synthesis of and its antiviral activity. Bioorganic and Medicinal Chemistry Letters, 1995, 5, 1685-1688.	2.2	13
54	Synthesis of 5-thiodidehydropyranylcytosine derivatives as potential anti-HIV agents. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 3313-3316.	2.2	12

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55	The Antitumor Mechanism of 1-(2-Deoxy-2-fluoro-4-thio- \hat{l}^2 -D-arabinofuranosyl)-cytosine: Effects of Its Triphosphate on Mammalian DNA Polymerases. Japanese Journal of Cancer Research, 2001, 92, 562-567.	1.7	11
56	Synthesis of Various Heterocycles Having a Dienamide Moiety by Ring-Closing Metathesis of Ene-ynamides. Synthesis, 2018, 50, 3467-3486.	2.3	11
57	Synthesis and biological evaluation of α-1-C-4′-arylbutyl-l-arabinoiminofuranoses, a new class of α-glucosidase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 3298-3301.	2.2	10
58	Strategy for Designing Selective Lysosomal Acid \hat{l}_{\pm} -Glucosidase Inhibitors: Binding Orientation and Influence on Selectivity. Molecules, 2020, 25, 2843.	3.8	10
59	Synthesis and Conformational Studies ofO5′, 6-Methanouridine—A New Type of Pyrimidine Cyclonucleoside. Nucleosides & Nucleotides, 1992, 11, 615-635.	0.5	9
60	Comparison of the Selectivity of Antiâ€Varicellaâ€Zoster Virus Nucleoside Analogues. Microbiology and Immunology, 1995, 39, 201-206.	1.4	9
61	Synthesis of (2'S)-1-(2-C-Azidomethyl-2-deoxy and) Tj ETQq1 1 0.784314 rgBT /Overlock 10 Tf 50 507 Td (2-C-C 1995, 14, 427-429.	yanomethy 1.1	vl-2-deoxy-Î ² - 9
62	A New Route to N1-Substituted Uracil Derivatives Using Hypervalent Iodine. Synthesis, 2012, 44, 1163-1170.	2.3	9
63	ASYMMETRIC SYNTHESIS OF 1-ALKYL-2-DEOXYIMINOFURANOSES VIA THE IRIDIUM-CATALYZED INTRAMOLECULAR CYCLIZATION OF AN ALLYLIC CARBONATE. Heterocycles, 2012, 86, 1401.	0.7	9
64	Asymmetric Synthesis of 2-Propylisofagomine Using Allylic Hydroxy Group Accelerated Ring-Closing Enyne Metathesis. Heterocycles, 2012, 84, 929.	0.7	9
65	Recent Advances in Cyclonucleosides: C-Cyclonucleosides and Spore Photoproducts in Damaged DNA. Molecules, 2012, 17, 11630-11654.	3.8	9
66	Design and synthesis of a nucleoside and a phosphonate analogue constructed on a branched-threo-tetrofuranose skeleton. Tetrahedron Letters, 2013, 54, 3949-3952.	1.4	9
67	Synthesis and Biological Activities of 2′-Modified 4′-Thionucleosides. Nucleosides & Nucleotides, 1997, 16, 1103-1106.	0.5	8
68	A straightforward stereoselective synthesis of meso-, (S,S)- and (R,R)-2,6-diaminopimelic acids from cis-1,4-diacetoxycyclohept-2-ene. Bioorganic and Medicinal Chemistry Letters, 2007, 17, 5894-5896.	2.2	8
69	An Access to the \hat{I}^2 -Anomer of $4\hat{a}\in \hat{I}^2$ -Thio-C-ribonucleosides: Hydroboration of $1-\langle i\rangle C \langle i\rangle$ -Aryl- or $1-\langle i\rangle C \langle i\rangle$ -Heteroaryl-4-thiofuranoid Glycals and Its Regiochemical Outcome. Journal of Organic Chemistry, 2011, 76, 8658-8669.	3.2	8
70	Palladium-Catalyzed Three-Component Coupling of Ynamides. Organic Letters, 2020, 22, 5299-5303.	4.6	8
71	Design of New Types of Antitumor Nucleosides: The Synthesis and Antitumor Activity of $2\hat{a}\in^2$ -Deoxy-($2\hat{a}\in^2$ -C-Substituted)Cytidines. , 1993, , 1-22.		8

Alternative synthesis of 2'-deoxy-6,2'-methano-pyrimidine nucleosides (nucleosides and nucleotides.) Tj ETQq0 0 0 rgBT /Overlock 10 Tf

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73	Palladium-Catalyzed Regioselective Hydroarylation of Ynamides with Aryl Iodides: Easy Synthesis of Various Substituted Enamides Containing Stilbene Derivatives. Synlett, 2017, 28, 2135-2138.	1.8	7
74	Development of a Glycosylation Reaction: A Key to Accessing Structurally Unique Nucleosides. Heterocycles, 2017, 94, 1625.	0.7	7
75	Synthesis of $4\hat{a}\in^2$ -Thionucleosides as Antitumor and Antiviral Agents. Chemical and Pharmaceutical Bulletin, 2018, 66, 139-146.	1.3	7
76	Glycosylation reactions mediated by hypervalent iodine: application to the synthesis of nucleosides and carbohydrates. Beilstein Journal of Organic Chemistry, 2018, 14, 1595-1618.	2.2	7
77	Development of Stereoselective Synthesis of Biologically Active Nitrogen-heterocyclic Compounds: Applications for Syntheses of Natural Product and Organocatalyst. Yuki Gosei Kagaku Kyokaishi/Journal of Synthetic Organic Chemistry, 2016, 74, 335-349.	0.1	7
78	In Vitro and in Vivo Antitumor Activity of a Novel Nucleoside, 4'-Thio-2'-deoxy-2'-methylidenecytidine Biological and Pharmaceutical Bulletin, 1996, 19, 1311-1315.	1.4	6
79	Inhibitory Effects of 9-(4-ThioBETAD-ribo-pentofuranosyl)guanine on Tumor Growth and Angiogenesis. Biological and Pharmaceutical Bulletin, 2004, 27, 520-523.	1.4	6
80	Chemoselective O-tert-butoxycarbonylation of phenols using 6,7-dimethoxyisoquinoline as a novel organocatalyst. Tetrahedron Letters, 2010, 51, 6915-6917.	1.4	6
81	Synthesis of a Dihydropyranonucleoside Using an Oxidative Glycosylation ReactionÂ-Mediated by Hypervalent Iodine. Synthesis, 2014, 46, 879-886.	2.3	6
82	Synthesis and Properties of 4′-ThioLNA/BNA. Organic Letters, 2021, 23, 4062-4066.	4.6	6
83	Modified 3-Hydroxypipecolic Acid Derivatives as an Organocatalyst. Heterocycles, 2009, 77, 635.	0.7	6
84	Synthetic Studies on 2′-Substituted-4′-thiocytidine Derivatives as Antineoplastic Agents. Nucleosides & Nucleotides, 1999, 18, 815-820.	0.5	5
85	Synthesis of 6,3′-Methanothymidine from a Ribofuranos-3-Ulose and 2,4-Dimethoxy-5,6-Dimethylpyrimidine. Nucleosides & Nucleotides, 1988, 7, 409-416.	0.5	4
86	Synthesis and antiviral evaluation of α-d-2′,3′-didehydro-2′,3′-dideoxy-3′-C-hydroxymethyl nucleosi Bioorganic and Medicinal Chemistry Letters, 2010, 20, 6013-6016.	des 2.2	4
87	Catalytic asymmetric synthesis of stereoisomers of 1-C-n-butyl-LABs for the SAR study of \hat{l}_{\pm} -glucosidase inhibition. Tetrahedron, 2019, 75, 2866-2876.	1.9	4
88	Construction of an Isonucleoside on a 2,6-Dioxobicyclo [3.2.0]-heptane Skeleton. Molecules, 2015, 20, 4623-4634.	3.8	3
89	Concise Syntheses of Violaceoids A and C. Chemical and Pharmaceutical Bulletin, 2021, 69, 232-235.	1.3	3
90	Synthesis and evaluation of trypanocidal activity of derivatives of naturally occurring 2,5-diphenyloxazoles. Bioorganic and Medicinal Chemistry, 2021, 42, 116253.	3.0	3

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91	Development of Glycoside Bond Formation Reactions and Their Applications to the Synthesis of Novel Biologically Active Nucleosides. Yuki Gosei Kagaku Kyokaishi/Journal of Synthetic Organic Chemistry, 2009, 67, 798-808.	0.1	2
92	Practical Synthesis of 4′â€Thioribonucleosides from <scp>L</scp> â€Arabinose via Novel Reductive Ringâ€Contraction Reaction and Pummererâ€Type Thioglycosylation. Current Protocols in Nucleic Acid Chemistry, 2017, 71, 1.43.1-1.43.12.	0.5	1
93	Synthesis of 2′-aminouridine derivatives as an organocatalyst for Diels-Alder reaction. Nucleosides, Nucleotides and Nucleic Acids, 2020, 39, 365-383.	1.1	1
94	Study on the reaction mechanism of C-6 lithiation of pyrimidine nucleosides by using lithium hexamethyldisilazide as a base. Nucleic Acids Symposium Series, 2003, 3, 17-18.	0.3	0