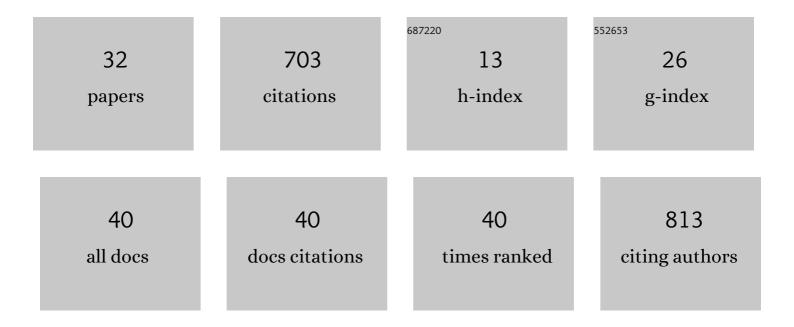
Yoshihiro Natori

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Asymmetric Synthesis of Neolignans (â^')- <i>epi</i> -Conocarpan and (+)-Conocarpan via Rh(II)-Catalyzed Câ^'H Insertion Process and Revision of the Absolute Configuration of (â^')- <i>epi</i> -Conocarpan. Journal of Organic Chemistry, 2009, 74, 4418-4421.	1.7	86
2	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.	2.9	72
3	Catalytic enantioselective C–H functionalization of indoles with α-diazopropionates using chiral dirhodium(II) carboxylates: asymmetric synthesis of the (+)-α-methyl-3-indolylacetic acid fragment of acremoauxin A. Tetrahedron: Asymmetry, 2011, 22, 907-915.	1.8	62
4	Conjugate vaccine produces long-lasting attenuation of fentanyl vs. food choice and blocks expression of opioid withdrawal-induced increases in fentanyl choice in rats. Neuropsychopharmacology, 2019, 44, 1681-1689.	2.8	56
5	Docking study and biological evaluation of pyrrolidine-based iminosugars as pharmacological chaperones for Gaucher disease. Organic and Biomolecular Chemistry, 2016, 14, 1039-1048.	1.5	46
6	Efficacious Vaccine against Heroin Contaminated with Fentanyl. ACS Chemical Neuroscience, 2018, 9, 1269-1275.	1.7	44
7	The synthesis and biological evaluation of 1-C-alkyl-l-arabinoiminofuranoses, a novel class of α-glucosidase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 738-741.	1.0	39
8	Catalytic asymmetric synthesis of (â^')-E-Î'-viniferin via an intramolecular C–H insertion of diaryldiazomethane using Rh2(S-TFPTTL)4. Tetrahedron Letters, 2015, 56, 4324-4327.	0.7	31
9	Improved Admixture Vaccine of Fentanyl and Heroin Hapten Immunoconjugates: Antinociceptive Evaluation of Fentanyl-Contaminated Heroin. ACS Omega, 2018, 3, 11537-11543.	1.6	31
10	Enantioselective Synthesis of 3-Arylindan-1-ones via Intramolecular C-H Insertion Reactions of α-Diazo-β-Ketoesters Catalyzed by Chiral Dirhodium(II) Carboxylates. Heterocycles, 2006, 70, 635.	0.4	26
11	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.	3.5	25
12	Asymmetric synthesis of 2,5-disubstituted 3-hydroxypyrrolidines based on stereodivergent intramolecular iridium-catalyzed allylic aminations. Organic and Biomolecular Chemistry, 2014, 12, 1983.	1.5	23
13	A chemically contiguous hapten approach for a heroin–fentanyl vaccine. Beilstein Journal of Organic Chemistry, 2019, 15, 1020-1031.	1.3	22
14	A Highly Efficacious Carfentanil Vaccine That Blunts Opioid-Induced Antinociception and Respiratory Depression. ACS Chemical Biology, 2021, 16, 277-282.	1.6	16
15	Synthesis of 5-thiodidehydropyranylcytosine derivatives as potential anti-HIV agents. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 3313-3316.	1.0	12
16	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.	1.0	10
17	Strategy for Designing Selective Lysosomal Acid α-Glucosidase Inhibitors: Binding Orientation and Influence on Selectivity. Molecules, 2020, 25, 2843.	1.7	10
18	A New Route to N1-Substituted Uracil Derivatives Using Hypervalent Iodine. Synthesis, 2012, 44, 1163-1170.	1.2	9

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19	ASYMMETRIC SYNTHESIS OF 1-ALKYL-2-DEOXYIMINOFURANOSES VIA THE IRIDIUM-CATALYZED INTRAMOLECULAR CYCLIZATION OF AN ALLYLIC CARBONATE. Heterocycles, 2012, 86, 1401.	0.4	9
20	Design and synthesis of a nucleoside and a phosphonate analogue constructed on a branched-threo-tetrofuranose skeleton. Tetrahedron Letters, 2013, 54, 3949-3952.	0.7	9
21	Efficient Syntheses of Cocaine Vaccines and Their <i>in Vivo</i> Evaluation. ACS Medicinal Chemistry Letters, 2018, 9, 411-416.	1.3	9
22	Palladium-Catalyzed Three-Component Coupling of Ynamides. Organic Letters, 2020, 22, 5299-5303.	2.4	8
23	Palladium-Catalyzed Regioselective Hydroarylation of Ynamides with Aryl Iodides: Easy Synthesis of Various Substituted Enamides Containing Stilbene Derivatives. Synlett, 2017, 28, 2135-2138.	1.0	7
24	Synthesis of 4′-Thionucleosides as Antitumor and Antiviral Agents. Chemical and Pharmaceutical Bulletin, 2018, 66, 139-146.	0.6	7
25	Glycosylation reactions mediated by hypervalent iodine: application to the synthesis of nucleosides and carbohydrates. Beilstein Journal of Organic Chemistry, 2018, 14, 1595-1618.	1.3	7
26	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.	0.0	7
27	Synthesis of a Dihydropyranonucleoside Using an Oxidative Glycosylation ReactionÂ-Mediated by Hypervalent Iodine. Synthesis, 2014, 46, 879-886.	1.2	6
28	Synthesis and Properties of 4′-ThioLNA/BNA. Organic Letters, 2021, 23, 4062-4066.	2.4	6
29	Catalytic asymmetric synthesis of stereoisomers of 1-C-n-butyl-LABs for the SAR study of α-glucosidase inhibition. Tetrahedron, 2019, 75, 2866-2876.	1.0	4
30	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
31	Synthesis of 2′-aminouridine derivatives as an organocatalyst for Diels-Alder reaction. Nucleosides, Nucleotides and Nucleic Acids, 2020, 39, 365-383.	0.4	1
32	Synthesis of Drug Vaccine against Heroin Contaminated with Fentanyl and Their Biological Evaluation. Yuki Gosei Kagaku Kyokaishi/Journal of Synthetic Organic Chemistry, 2020, 78, 875-885.	0.0	0