

Piet Herdewijn

List of Publications by Year in descending order

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Version: 2024-02-01

640
papers

18,473
citations

22548

61
h-index

29333

108
g-index

771
all docs

771
docs citations

771
times ranked

11323
citing authors

#	ARTICLE	IF	CITATIONS
1	An Overview of Marketed Nucleoside and Nucleotide Analogs. <i>Current Protocols</i> , 2022, 2, e376.	1.3	11
2	In vivo assembly and expression of DNA containing non-canonical bases in the yeast <i>Saccharomyces cerevisiae</i> . <i>ChemBioChem</i> , 2022, , .	1.3	4
3	Reshaping an Acyclic Nucleoside Phosphonate into a Selective Anti-hepatitis B Virus Compound. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 9396-9417.	2.9	2
4	The Network of Replication, Transcription, and Reverse Transcription of a Synthetic Genetic Cassette. <i>Angewandte Chemie - International Edition</i> , 2021, 60, 4175-4182.	7.2	4
5	The Network of Replication, Transcription, and Reverse Transcription of a Synthetic Genetic Cassette. <i>Angewandte Chemie</i> , 2021, 133, 4221-4228.	1.6	1
6	Influence of 4-Substitution on the Activity of Gemcitabine and Its ProTide Against VZV and SARS-CoV-2. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 88-92.	1.3	16
7	Introduction of a cyano group at the 2-position of an (<i>R</i>)-3-hydroxy-2-(phosphonomethoxy)propyl (HPMP) derivative of thymine elicits selective anti-HBV activity. <i>RSC Medicinal Chemistry</i> , 2021, 12, 804-808.	1.7	1
8	Stable Hairpin Structures Formed by Xylose-Based Nucleic Acids. <i>ChemBioChem</i> , 2021, 22, 1638-1645.	1.3	4
9	Discovery of 3-phenyl- and 3-N-piperidinyl-isothiazolo[4,3-b]pyridines as highly potent inhibitors of cyclin G-associated kinase. <i>European Journal of Medicinal Chemistry</i> , 2021, 213, 113158.	2.6	10
10	Noncanonical DNA polymerization by aminoadenine-based siphoviruses. <i>Science</i> , 2021, 372, 520-524.	6.0	46
11	Functional Comparison of Laboratory-Evolved XNA Polymerases for Synthetic Biology. <i>ACS Synthetic Biology</i> , 2021, 10, 1429-1437.	1.9	16
12	Synthesis and in vitro antitumour activity of 4(R)-methyl-3-O-phosphonomethyl-L-threose nucleosides. <i>European Journal of Medicinal Chemistry</i> , 2021, 221, 113513.	2.6	3
13	Exploring the dNTP-binding site of HIV-1 reverse transcriptase for inhibitor design. <i>European Journal of Medicinal Chemistry</i> , 2021, 225, 113785.	2.6	3
14	Tenofovir-Amino Acid Conjugates Act as Polymerase Substrates—Implications for Avoiding Cellular Phosphorylation in the Discovery of Nucleotide Analogues. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 782-796.	2.9	2
15	Anno 2021: Which antivirals for the coming decade?. <i>Annual Reports in Medicinal Chemistry</i> , 2021, 57, 49-107.	0.5	4
16	Sliding of HIV-1 reverse transcriptase over DNA creates a transient P pocket—targeting P-pocket by fragment screening. <i>Nature Communications</i> , 2021, 12, 7127.	5.8	6
17	In Vivo Expression of Genetic Information from Phosphoramidate-DNA. <i>ChemBioChem</i> , 2020, 21, 272-278.	1.3	14
18	Orthogonal Genetic Systems. <i>ChemBioChem</i> , 2020, 21, 1408-1411.	1.3	25

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19	Structure-activity relationship study of the pyridine moiety of isothiazolo[4,3-b]pyridines as antiviral agents targeting cyclin G-associated kinase. <i>Bioorganic and Medicinal Chemistry</i> , 2020, 28, 115188.	1.4	14
20	Iron/Copper Co-Catalyzed Cross-Coupling Reaction for the Synthesis of 6-Substituted 7-Deazapurines and the Corresponding Nucleosides. <i>Journal of Organic Chemistry</i> , 2020, 85, 403-418.	1.7	14
21	Enzymatic Formation of an Artificial Base Pair Using a Modified Purine Nucleoside Triphosphate. <i>ACS Chemical Biology</i> , 2020, 15, 2872-2884.	1.6	21
22	Synthesis and Antiviral Evaluation of 3'-Hydroxymethyl-5'-Phosphonomethyl-2'-deoxyribose Nucleosides. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 4995-5002.	1.3	5
23	Synthesis and Antitumor Activity of C-7-Alkynylated and Arylated Pyrrolotriazine C-Ribonucleosides. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 1605-1610.	1.3	5
24	Scalable Synthesis, In Vitro cccDNA Reduction, and In Vivo Antihepatitis B Virus Activity of a Phosphonomethoxydeoxythreosyl Adenine Prodrug. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 13851-13860.	2.9	8
25	Effect of Molecular Crowding on DNA Polymerase Reactions along Unnatural DNA Templates. <i>Molecules</i> , 2020, 25, 4120.	1.7	5
26	Structural Studies of HNA Substrate Specificity in Mutants of an Archaeal DNA Polymerase Obtained by Directed Evolution. <i>Biomolecules</i> , 2020, 10, 1647.	1.8	7
27	Beneath the XNA world: Tools and targets to build novel biology. <i>Current Opinion in Systems Biology</i> , 2020, 24, 142-152.	1.3	5
28	The Kalimantacin Polyketide Antibiotics Inhibit Fatty Acid Biosynthesis in <i>Staphylococcus aureus</i> by Targeting the Enoyl-Acyl Carrier Protein Binding Site of FabI. <i>Angewandte Chemie</i> , 2020, 132, 10636-10643.	1.6	6
29	Structure-Activity Relationship Study of a Potent Thrombin Binding Aptamer Incorporating Hexitol Nucleotides. <i>Chemistry - A European Journal</i> , 2020, 26, 9589-9597.	1.7	17
30	New Metal-Free Route towards Imidazole-Substituted Uridine. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 4022-4025.	1.2	5
31	Anti-norovirus activity of C7-modified 4-amino-pyrrolo[2,1-f][1,2,4]triazine C-nucleosides. <i>European Journal of Medicinal Chemistry</i> , 2020, 195, 112198.	2.6	14
32	The Kalimantacin Polyketide Antibiotics Inhibit Fatty Acid Biosynthesis in <i>Staphylococcus aureus</i> by Targeting the Enoyl-Acyl Carrier Protein Binding Site of FabI. <i>Angewandte Chemie - International Edition</i> , 2020, 59, 10549-10556.	7.2	20
33	Chimeric siRNAs with chemically modified pentofuranose and hexopyranose nucleotides: altritol-nucleotide (ANA) containing GalNAc-siRNA conjugates: in vitro and in vivo RNAi activity and resistance to 5'-exonuclease. <i>Nucleic Acids Research</i> , 2020, 48, 4028-4040.	6.5	27
34	Amidate Prodrugs of O-2-Alkylated Pyrimidine Acyclic Nucleosides Display Potent Anti-Herpesvirus Activity. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 1410-1415.	1.3	7
35	Vitamin-guanosine monophosphate conjugates for in vitro transcription priming. <i>Chemical Communications</i> , 2020, 56, 2787-2790.	2.2	1
36	Synthesis of tetradialdose phosphonate nucleosides as mimics of l-nucleotides. <i>Tetrahedron</i> , 2019, 75, 130497.	1.0	1

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37	Rational design of an XNA ligase through docking of unbound nucleic acids to toroidal proteins. <i>Nucleic Acids Research</i> , 2019, 47, 7130-7142.	6.5	23
38	Synthesis of Poly(ADP-ribose) Monomer Containing 2'-O-Methyl-Ribofuranosyl Adenosine. <i>Current Protocols in Nucleic Acid Chemistry</i> , 2019, 78, e92.	0.5	1
39	Invading <i>Escherichia coli</i> Genetics with a Xenobiotic Nucleic Acid Carrying an Acyclic Phosphonate Backbone (ZNA). <i>Journal of the American Chemical Society</i> , 2019, 141, 10844-10851.	6.6	25
40	⁸ -Glycosylated 8-Azapurine and Methylated Purine Nucleobases: Synthesis and Study of Base Pairing Properties. <i>Journal of Organic Chemistry</i> , 2019, 84, 13394-13409.	1.7	2
41	Synthesis of a Threosyl-nucleoside Phosphonate. <i>European Journal of Organic Chemistry</i> , 2019, 2019, 6666-6672.	1.2	2
42	Full Pre-steady-State Kinetic Analysis of Single Nucleotide Incorporation by DNA Polymerases. <i>Current Protocols in Nucleic Acid Chemistry</i> , 2019, 78, e98.	0.5	1
43	Synthesis of 3-fluoro-4-amino-hexitol nucleosides with a pyrimidine nucleobase as building blocks for oligonucleotides. <i>Tetrahedron</i> , 2019, 75, 1107-1114.	1.0	2
44	Enzymatic Synthesis of Backbone-Modified Oligonucleotides Using T4 DNA Ligase. <i>Current Protocols in Chemical Biology</i> , 2019, 11, e62.	1.7	3
45	Synthesis and Structure-Activity Relationship Studies of Benzo[b][1,4]oxazin-3(4H)-one Analogues as Inhibitors of Mycobacterial Thymidylate Synthase...X. <i>ChemMedChem</i> , 2019, 14, 645-662.	1.6	9
46	What Is XNA?. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 11570-11572.	7.2	78
47	On the Enzymatic Formation of Metal Base Pairs with Thiolated and pKa-Perturbed Nucleotides. <i>ChemBioChem</i> , 2019, 20, 3032-3040.	1.3	15
48	Was ist XNA?. <i>Angewandte Chemie</i> , 2019, 131, 11694-11696.	1.6	10
49	Synthesis and Structure-Activity Relationships of 3,5-Disubstituted-pyrrolo[2,3-b]pyridines as Inhibitors of Adaptor-Associated Kinase 1 with Antiviral Activity. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 5810-5831.	2.9	44
50	Synthesis and Conformation of Pentopyranoside Nucleoside Phosphonates. <i>Journal of Organic Chemistry</i> , 2019, 84, 6589-6603.	1.7	4
51	Highly stable hexitol based XNA aptamers targeting the vascular endothelial growth factor. <i>Nucleic Acids Research</i> , 2019, 47, 4927-4939.	6.5	73
52	Xylo-C-nucleosides with a pyrrolo[2,1-f][1,2,4]triazin-4-amine heterocyclic base: Synthesis and antiproliferative properties. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2019, 29, 1450-1453.	1.0	6
53	Kinetic analysis of N-alkylaryl carboxamide hexitol nucleotides as substrates for evolved polymerases. <i>Nucleic Acids Research</i> , 2019, 47, 2160-2168.	6.5	10
54	Reprint of: Non Canonical Genetic Material. <i>Current Opinion in Biotechnology</i> , 2019, 60, 259-267.	3.3	10

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55	Non canonical genetic material. <i>Current Opinion in Biotechnology</i> , 2019, 57, 25-33.	3.3	30
56	Cyclin G-associated kinase (GAK) affinity and antiviral activity studies of a series of 3-C-substituted isothiazolo[4,3-b]pyridines. <i>European Journal of Medicinal Chemistry</i> , 2019, 163, 256-265.	2.6	10
57	Synthesis and Anti-HIV Activity of Guanine Modified Fluorinated Acyclic Nucleoside Phosphonate Derivatives. <i>Chemistry and Biodiversity</i> , 2019, 16, e1800532.	1.0	3
58	Towards the enzymatic formation of artificial metal base pairs with a carboxy-imidazole-modified nucleotide. <i>Journal of Inorganic Biochemistry</i> , 2019, 191, 154-163.	1.5	31
59	A Scaffold-Hopping Strategy toward the Identification of Inhibitors of Cyclin-G Associated Kinase. <i>ChemMedChem</i> , 2019, 14, 237-254.	1.6	1
60	Where cone snails and spiders meet: design of small cyclic sodium-channel inhibitors. <i>FASEB Journal</i> , 2019, 33, 3693-3703.	0.2	23
61	Synthesis of a C-Nucleoside Phosphonate by Base-Promoted Epimerization. <i>Organic Letters</i> , 2018, 20, 1203-1206.	2.4	3
62	Modulation of BACE1 Activity by Chemically Modified Aptamers. <i>ChemBioChem</i> , 2018, 19, 754-763.	1.3	23
63	Phosphonomethyl Oligonucleotides as Backbone-Modified Artificial Genetic Polymers. <i>Journal of the American Chemical Society</i> , 2018, 140, 6690-6699.	6.6	48
64	Amidate Prodrugs of Cyclic 9-[3-Hydroxy-2-(phosphonomethoxy)propyl]adenine with Potent Anti-Herpesvirus Activity. <i>ACS Medicinal Chemistry Letters</i> , 2018, 9, 381-385.	1.3	13
65	Synthesis and antiviral evaluation of cyclopentyl nucleoside phosphonates. <i>European Journal of Medicinal Chemistry</i> , 2018, 150, 616-625.	2.6	5
66	Emimycin and its nucleoside derivatives: Synthesis and antiviral activity. <i>European Journal of Medicinal Chemistry</i> , 2018, 144, 93-103.	2.6	6
67	Synthesis and Biological Evaluation of Pyrrolo[2,1-f][1,2,4]triazine Nucleosides with a Ribose, 2-Deoxyribose, and 3-Deoxyribose Sugar Moiety. <i>ChemMedChem</i> , 2018, 13, 97-104.	1.6	17
68	Synthesis of Protected Amino Hexitol Nucleosides as Building Blocks for Oligonucleotide Synthesis. <i>Journal of Organic Chemistry</i> , 2018, 83, 15155-15169.	1.7	8
69	Incorporation of a minimal nucleotide into DNA. <i>Tetrahedron Letters</i> , 2018, 59, 4241-4244.	0.7	7
70	Synthesis of a 2-Deoxy-5-C-Nucleoside Phosphonate Bearing 9-Deazaadenine as Base Moiety. <i>European Journal of Organic Chemistry</i> , 2018, 2018, 6657-6664.	1.2	1
71	A Single Amino Acid Substitution in Terminator DNA Polymerase Increases Incorporation Efficiency of Deoxyxylonucleotides. <i>ChemBioChem</i> , 2018, 19, 2410-2420.	1.3	4
72	Frontispiece: Chimeric XNA: An Unconventional Design for Orthogonal Informational Systems. <i>Chemistry - A European Journal</i> , 2018, 24, .	1.7	0

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73	XNA ligation using T4 DNA ligase in crowding conditions. <i>Chemical Communications</i> , 2018, 54, 6408-6411.	2.2	30
74	Discovery of a Potent, Orally Bioavailable PI4KIII β Inhibitor (UCB9608) Able To Significantly Prolong Allogeneic Organ Engraftment <i>in Vivo</i> . <i>Journal of Medicinal Chemistry</i> , 2018, 61, 6705-6723.	2.9	18
75	Optimization of Isothiazolo[4,3- <i>b</i>]pyridine-Based Inhibitors of Cyclin G Associated Kinase (GAK) with Broad-Spectrum Antiviral Activity. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 6178-6192.	2.9	36
76	Metabolic Recruitment and Directed Evolution of Nucleoside Triphosphate Uptake in <i>Escherichia coli</i> . <i>ACS Synthetic Biology</i> , 2018, 7, 1565-1572.	1.9	14
77	Mutant Variants of the Substrate-Binding Protein DppA from <i>Escherichia coli</i> Enhance Growth on Nonstandard β -Glutamyl Amide-Containing Peptides. <i>Applied and Environmental Microbiology</i> , 2018, 84, .	1.4	3
78	Phosphorus Pentachloride Promoted gem-Dichlorination of 2 α - and 3 α -Deoxynucleosides. <i>Molecules</i> , 2018, 23, 1457.	1.7	1
79	Methylated Nucleobases: Synthesis and Evaluation for Base Pairing <i>In Vitro</i> and <i>In Vivo</i> . <i>Chemistry - A European Journal</i> , 2018, 24, 12695-12707.	1.7	6
80	Chimeric XNA: An Unconventional Design for Orthogonal Informational Systems. <i>Chemistry - A European Journal</i> , 2018, 24, 12811-12819.	1.7	9
81	Peptidoglycan glycosyltransferase-ligand binding assay based on tryptophan fluorescence quenching. <i>Biochimie</i> , 2018, 152, 1-5.	1.3	5
82	PCR Amplification of Base-Modified DNA. <i>Current Protocols in Chemical Biology</i> , 2018, 10, 18-48.	1.7	3
83	Random-sequence genetic oligomer pools display an innate potential for ligation and recombination. <i>ELife</i> , 2018, 7, .	2.8	43
84	Oligonucleotide promoted peptide bond formation using a tRNA mimicking approach. <i>Chemical Communications</i> , 2017, 53, 5013-5016.	2.2	0
85	On the enzymatic incorporation of an imidazole nucleotide into DNA. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 4449-4455.	1.5	35
86	Synthesis and antiviral evaluation of base-modified deoxythreosyl nucleoside phosphonates. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 5513-5528.	1.5	4
87	Base-Modified Nucleic Acids as a Powerful Tool for Synthetic Biology and Biotechnology. <i>Chemistry - A European Journal</i> , 2017, 23, 9560-9576.	1.7	28
88	Discovery of a new <i>Mycobacterium tuberculosis</i> thymidylate synthase X inhibitor with a unique inhibition profile. <i>Biochemical Pharmacology</i> , 2017, 135, 69-78.	2.0	16
89	The 5-chlorouracil:7-deazaadenine base pair as an alternative to the dT:dA base pair. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 168-176.	1.5	20
90	Astemizole analogues with reduced hERG inhibition as potent antimalarial compounds. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 6332-6344.	1.4	17

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91	Enzymatic Incorporation of Modified Purine Nucleotides in DNA. <i>ChemBioChem</i> , 2017, 18, 2408-2415.	1.3	2
92	Synthesis of a 3-Fluoro-3-deoxytetrose Adenine Phosphonate. <i>Journal of Organic Chemistry</i> , 2017, 82, 9464-9478.	1.7	5
93	Molecular Dynamics of Double Stranded Xylo-Nucleic Acid. <i>Journal of Chemical Theory and Computation</i> , 2017, 13, 5028-5038.	2.3	9
94	Facile immobilization of DNA using an enzymatic his-tag mimic. <i>Chemical Communications</i> , 2017, 53, 13031-13034.	2.2	23
95	Expanding the Antiviral Spectrum of 3-Fluoro-2-(phosphonomethoxy)propyl Acyclic Nucleoside Phosphonates: Diamyl Aspartate Amidate Prodrugs. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 6220-6238.	2.9	22
96	Overcoming the membrane barrier: Recruitment of β -glutamyl transferase for intracellular release of metabolic cargo from peptide vectors. <i>Metabolic Engineering</i> , 2017, 39, 60-70.	3.6	5
97	Substrate-Dependence of Competitive Nucleotide Pyrophosphatase/Phosphodiesterase1 (NPP1) Inhibitors. <i>Frontiers in Pharmacology</i> , 2017, 8, 54.	1.6	36
98	Anticancer kinase inhibitors impair intracellular viral trafficking and exert broad-spectrum antiviral effects. <i>Journal of Clinical Investigation</i> , 2017, 127, 1338-1352.	3.9	188
99	Structural and Functional Elucidation of Peptide Ts11 Shows Evidence of a Novel Subfamily of Scorpion Venom Toxins. <i>Toxins</i> , 2016, 8, 288.	1.5	26
100	Incorporation of Amino Acids with Long-Chain Terminal Olefins into Proteins. <i>Molecules</i> , 2016, 21, 287.	1.7	10
101	Chemical Morphing of DNA Containing Four Noncanonical Bases. <i>Angewandte Chemie</i> , 2016, 128, 7641-7645.	1.6	11
102	Chemical Morphing of DNA Containing Four Noncanonical Bases. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 7515-7519.	7.2	40
103	Syntheses of 5-Nucleoside Monophosphate Derivatives with Unique Amino, Hemiaminal, and Hemithioaminal Functionalities: A New Class of 5-Peptidyl Nucleotides. <i>Chemistry - A European Journal</i> , 2016, 22, 8167-8180.	1.7	7
104	L-Aspartic and L-glutamic acid ester-based ProTides of anticancer nucleosides: Synthesis and antitumoral evaluation. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 2142-2146.	1.0	12
105	Lipophilic nalmefene prodrugs to achieve a one-month sustained release. <i>Journal of Controlled Release</i> , 2016, 232, 196-202.	4.8	10
106	Bifunctional aryloxyphosphoramidate prodrugs of 2-C-Me-uridine: synthesis and anti-HCV activity. <i>Organic and Biomolecular Chemistry</i> , 2016, 14, 8743-8757.	1.5	4
107	Amidate Prodrugs of Deoxythreosyl Nucleoside Phosphonates as Dual Inhibitors of HIV and HBV Replication. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 9513-9531.	2.9	26
108	Synthesis of a Nucleobase-Modified ProTide Library. <i>Organic Letters</i> , 2016, 18, 5816-5819.	2.4	9

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109	Evaluation of anhydrohexitol nucleic acid, cyclohexenyl nucleic acid and α -altritol nucleic acid-modified 2'-O-methyl RNA mixer antisense oligonucleotides for exon skipping in vitro. <i>Chemical Communications</i> , 2016, 52, 13467-13470.	2.2	39
110	Theoretical Analysis of a Self-Replicator With Reduced Template Inhibition Based on an Informational Leaving Group. <i>Journal of Molecular Evolution</i> , 2016, 82, 93-109.	0.8	2
111	Thiazolo[3,2-a]benzimidazol-3(2H)-one derivatives: Structure-activity relationships of selective nucleotide pyrophosphatase/phosphodiesterase1 (NPP1) inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 3157-3165.	1.4	19
112	Nanostructures from Synthetic Genetic Polymers. <i>ChemBioChem</i> , 2016, 17, 1107-1110.	1.3	57
113	Molecular simulation of cyclohexanyl nucleic acid (CNA) duplexes with CNA, DNA and RNA and CNA triloop and tetraloop hairpin structures. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 1778-1785.	1.4	2
114	Base pairing involving artificial bases in vitro and in vivo. <i>Chemical Science</i> , 2016, 7, 995-1010.	3.7	19
115	Synthesis and <i>In Vitro</i> Antiviral Activities of [(Dihydrofuran-2-yl)oxy]methylphosphonate Nucleosides with ϵ -Substituted Adenine as Base. <i>Chemistry and Biodiversity</i> , 2015, 12, 813-822.	1.0	5
116	Nucleosides with Transposed Base or 4'-Hydroxymethyl Moieties and Their Corresponding Oligonucleotides. <i>Chemical Reviews</i> , 2015, 115, 13484-13525.	23.0	21
117	Positive cooperativity between acceptor and donor sites of the peptidoglycan glycosyltransferase. <i>Biochemical Pharmacology</i> , 2015, 93, 141-150.	2.0	9
118	Isoguanine and 5-Methylisocytosine Bases, <i>In Vitro</i> and <i>In Vivo</i> . <i>Chemistry - A European Journal</i> , 2015, 21, 5009-5022.	1.7	33
119	Achiral, acyclic nucleic acids: synthesis and biophysical studies of a possible prebiotic polymer. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 9249-9260.	1.5	10
120	NMR-based conformational analysis of 2',6-disubstituted uridines and antiviral evaluation of new phosphoramidate prodrugs. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 5809-5815.	1.4	5
121	<i>In vitro</i> disposition profiling of heterocyclic compounds. <i>International Journal of Pharmaceutics</i> , 2015, 491, 78-90.	2.6	2
122	1',5'-Anhydro- α -D-ribohexitol Adenine Nucleic Acids (α -HNA-A): Synthesis and Chiral Selection Properties in the Mirror Image World. <i>Journal of Organic Chemistry</i> , 2015, 80, 5014-5022.	1.7	13
123	Selective Inhibitors of Cyclin G Associated Kinase (GAK) as Anti-Hepatitis C Agents. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 3393-3410.	2.9	54
124	Aspartic acid based nucleoside phosphoramidate prodrugs as potent inhibitors of hepatitis C virus replication. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 5158-5174.	1.5	23
125	Oligonucleotides containing a ribo-configured cyclohexanyl nucleoside: probing the role of sugar conformation in base pairing selectivity. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 10041-10049.	1.5	4
126	Synthesis and evaluation of C-5 modified 2'-deoxyuridine monophosphates as inhibitors of <i>M. tuberculosis</i> thymidylate synthase. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 7131-7137.	1.4	25

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127	Isothiazolo[4,3-b]pyridines as inhibitors of cyclin G associated kinase: synthesis, structure-activity relationship studies and antiviral activity. <i>MedChemComm</i> , 2015, 6, 1666-1672.	3.5	16
128	Xylonucleic acid: synthesis, structure, and orthogonal pairing properties. <i>Nucleic Acids Research</i> , 2015, 43, 7189-7200.	6.5	23
129	Catalysts from synthetic genetic polymers. <i>Nature</i> , 2015, 518, 427-430.	13.7	230
130	Nucleoside Phosphate-Conjugates Come of Age: Catalytic Transformation, Polymerase Recognition and Antiviral Properties. <i>Current Medicinal Chemistry</i> , 2015, 22, 3980-3990.	1.2	5
131	A Convenient Route for the Synthesis of 3-Deazaspongosine. <i>European Journal of Organic Chemistry</i> , 2014, 2014, 231-236.	1.2	5
132	Design and synthesis of nucleolipids as possible activated precursors for oligomer formation via intramolecular catalysis: stability study and supramolecular organization. <i>Journal of Systems Chemistry</i> , 2014, 5, 5.	1.7	11
133	Tailoring Peptide-Nucleotide Conjugates (PNCs) for Nucleotide Delivery in Bacterial Cells. <i>European Journal of Organic Chemistry</i> , 2014, 2014, 2322-2348.	1.2	4
134	Synthesis of an Apionucleoside Family and Discovery of a Prodrug with Anti-HIV Activity. <i>Journal of Organic Chemistry</i> , 2014, 79, 5097-5112.	1.7	27
135	Synthesis of new biocarrier-nucleotide systems for cellular delivery in bacterial auxotrophic strains. <i>Tetrahedron</i> , 2014, 70, 8843-8851.	1.0	1
136	Mutations in the chikungunya virus non-structural proteins cause resistance to favipiravir (T-705), a broad-spectrum antiviral. <i>Journal of Antimicrobial Chemotherapy</i> , 2014, 69, 2770-2784.	1.3	187
137	Discovery of a new subclass of $\hat{\pm}$ -conotoxins in the venom of <i>Conus australis</i> . <i>Toxicon</i> , 2014, 91, 145-154.	0.8	25
138	Discovery of Dual Death-Associated Protein Related Apoptosis Inducing Protein Kinase 1 and 2 Inhibitors by a Scaffold Hopping Approach. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 7624-7643.	2.9	38
139	Probing Ambiguous Base-Pairs by Genetic Transformation with XNA Templates. <i>ChemBioChem</i> , 2014, 15, 2255-2258.	1.3	18
140	Organophosphorus-catalyzed diaza-Wittig reaction: application to the synthesis of pyridazines. <i>Organic and Biomolecular Chemistry</i> , 2014, 12, 7159-7166.	1.5	28
141	Hydroxy fatty acids for the delivery of dideoxynucleosides as anti-HIV agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 817-820.	1.0	5
142	Phospho-carboxylic anhydride of a homologated nucleoside leads to primer degradation in the presence of a polymerase. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 2720-2723.	1.0	10
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