

Jef Rozenski

List of Publications by Year in descending order

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152
papers

3,788
citations

126708

33
h-index

161609

54
g-index

166
all docs

166
docs citations

166
times ranked

4483
citing authors

#	ARTICLE	IF	CITATIONS
1	Overview of inâ€capillary enzymatic reactions using capillary electrophoresis. <i>Electrophoresis</i> , 2022, 43, 57-73.	1.3	9
2	In vivo assembly and expression of DNA containing nonâ€canonical bases in the yeast <i>Saccharomyces cerevisiae</i> . <i>ChemBioChem</i> , 2022, . .	1.3	4
3	Ways to Improve Insights into Clindamycin Pharmacology and Pharmacokinetics Tailored to Practice. <i>Antibiotics</i> , 2022, 11, 701.	1.5	9
4	Synthesis and structure-activity studies of novel anhydrohexitol-based Leucyl-tRNA synthetase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2021, 211, 113021.	2.6	14
5	Noncanonical DNA polymerization by aminoadenine-based siphoviruses. <i>Science</i> , 2021, 372, 520-524.	6.0	46
6	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
7	Synthesis and Biological Evaluation of 1,3-Dideazapurine-Like 7-Amino-5-Hydroxymethyl-Benzimidazole Ribonucleoside Analogues as Aminoacyl-tRNA Synthetase Inhibitors. <i>Molecules</i> , 2020, 25, 4751.	1.7	2
8	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
9	Phenyltriazole-functionalized sulfamate inhibitors targeting tyrosyl- or isoleucyl-tRNA synthetase. <i>Bioorganic and Medicinal Chemistry</i> , 2020, 28, 115580.	1.4	6
10	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
11	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
12	Synthesis and Biological Evaluation of Lipophilic Nucleoside Analogues as Inhibitors of Aminoacyl-tRNA Synthetases. <i>Antibiotics</i> , 2019, 8, 180.	1.5	2
13	Synthesis and Structureâ€Activity Relationship Studies of Benzo[b][1,4]oxazinâ€(3 H)â€one Analogues as Inhibitors of Mycobacterial Thymidylate Synthaseâ€...X. <i>ChemMedChem</i> , 2019, 14, 645-662.	1.6	9
14	Acylated sulfonamide adenosines as potent inhibitors of the adenylate-forming enzyme superfamily. <i>European Journal of Medicinal Chemistry</i> , 2019, 174, 252-264.	2.6	10
15	Phosphonomethyl Oligonucleotides as Backbone-Modified Artificial Genetic Polymers. <i>Journal of the American Chemical Society</i> , 2018, 140, 6690-6699.	6.6	48
16	A Single Amino Acid Substitution in Terminator DNA Polymerase Increases Incorporation Efficiency of Deoxyxylonucleotides. <i>ChemBioChem</i> , 2018, 19, 2410-2420.	1.3	4
17	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
18	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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19	Aminopurine and aminoquinazoline scaffolds for development of potential dengue virus inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2017, 126, 101-109.	2.6	27
20	Chemical Morphing of DNA Containing Four Noncanonical Bases. <i>Angewandte Chemie</i> , 2016, 128, 7641-7645.	1.6	11
21	Chemical Morphing of DNA Containing Four Noncanonical Bases. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 7515-7519.	7.2	40
22	Synthetic strategy and antiviral evaluation of diamide containing heterocycles targeting dengue and yellow fever virus. <i>European Journal of Medicinal Chemistry</i> , 2016, 121, 158-168.	2.6	34
23	Characterization of insulin-degrading enzyme-mediated cleavage of A β 2 in distinct aggregation states. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2016, 1860, 1281-1290.	1.1	12
24	Base pairing involving artificial bases in vitro and in vivo. <i>Chemical Science</i> , 2016, 7, 995-1010.	3.7	19
25	Abl1 inhibitory contaminants leach from plastic tubes. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016, 31, 340-343.	2.5	3
26	Hybridisation Potential of 1',3'-Di-O-methylaltropyranoside Nucleic Acids. <i>Molecules</i> , 2015, 20, 4020-4041.	1.7	2
27	5 α -(N-aminoacyl)-sulfonamido-5 β -deoxyadenosine: Attempts for a stable alternative for aminoacyl-sulfamoyl adenosines as aaRS inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2015, 93, 227-236.	2.6	14
28	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
29	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
30	Base substituted 5 α -O-(N-isoleucyl)sulfamoyl nucleoside analogues as potential antibacterial agents. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 2875-2886.	1.4	18
31	Synthesis and evaluation of imidazole-4,5- and pyrazine-2,3-dicarboxamides targeting dengue and yellow fever virus. <i>European Journal of Medicinal Chemistry</i> , 2014, 87, 529-539.	2.6	57
32	In search of Flavivirus inhibitors part 2: Tritylated, diphenylmethylated and other alkylated nucleoside analogues. <i>European Journal of Medicinal Chemistry</i> , 2014, 76, 98-109.	2.6	25
33	Discovery of an acyclic nucleoside phosphonate that inhibits Mycobacterium Tuberculosis ThyX based on the binding mode of a 5-alkynyl substrate analogue. , 2014, , .		0
34	In search of flavivirus inhibitors: Evaluation of different tritylated nucleoside analogues. <i>European Journal of Medicinal Chemistry</i> , 2013, 65, 249-255.	2.6	28
35	The Antimicrobial Compound Xantholysin Defines a New Group of Pseudomonas Cyclic Lipopeptides. <i>PLoS ONE</i> , 2013, 8, e62946.	1.1	84
36	N-Alkylated Aminoacyl sulfamoyl adenosines as Potential Inhibitors of Aminoacylation Reactions and Microcin C Analogues Containing D-Amino Acids. <i>PLoS ONE</i> , 2013, 8, e79234.	1.1	9

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37	Molecular Plasticity Regulates Oligomerization and Cytotoxicity of the Multipetide-length Amyloid- β^2 Peptide Pool. <i>Journal of Biological Chemistry</i> , 2012, 287, 36732-36743.	1.6	37
38	Pretargeting of necrotic tumors with biotinylated hypericin using 123I-labeled avidin: evaluation of a two-step strategy. <i>Investigational New Drugs</i> , 2012, 30, 2132-2140.	1.2	14
39	Genetic and Functional Characterization of Cyclic Lipopeptide White-Line-Inducing Principle (WLIP) Production by Rice Rhizosphere Isolate <i>Pseudomonas putida</i> RW10S2. <i>Applied and Environmental Microbiology</i> , 2012, 78, 4826-4834.	1.4	67
40	Synthesis and Evaluation of 5-Substituted 2'-deoxyuridine Monophosphate Analogues As Inhibitors of Flavin-Dependent Thymidylate Synthase in <i>Mycobacterium tuberculosis</i> . <i>Journal of Medicinal Chemistry</i> , 2011, 54, 4847-4862.	2.9	68
41	Extended targeting potential and improved synthesis of Microcin C analogs as antibacterials. <i>Biorganic and Medicinal Chemistry</i> , 2011, 19, 5462-5467.	1.4	23
42	Promysalin, a Salicylate-Containing <i>Pseudomonas putida</i> Antibiotic, Promotes Surface Colonization and Selectively Targets Other <i>Pseudomonas</i> . <i>Chemistry and Biology</i> , 2011, 18, 1320-1330.	6.2	53
43	Analysis of nucleosides using the atmospheric-pressure solids analysis probe for ionization. <i>International Journal of Mass Spectrometry</i> , 2011, 304, 204-208.	0.7	12
44	A standardized and biocompatible preparation of aggregate-free amyloid beta peptide for biophysical and biological studies of Alzheimer's disease. <i>Protein Engineering, Design and Selection</i> , 2011, 24, 743-750.	1.0	97
45	Characterization of Peptide Chain Length and Constituency Requirements for YejABEF-Mediated Uptake of Microcin C Analogues. <i>Journal of Bacteriology</i> , 2011, 193, 3618-3623.	1.0	27
46	Synthesis and evaluation of 5-substituted-2'-deoxyuridine monophosphate analogues as inhibitors of flavin-dependent thymidylate synthase in <i>Mycobacterium tuberculosis</i> . , 2011, , .		0
47	Neurotoxicity of Alzheimer's disease $A\beta^2$ peptides is induced by small changes in the $A\beta^{242}$ to $A\beta^{240}$ ratio. <i>EMBO Journal</i> , 2010, 29, 3408-3420.	3.5	455
48	MccE Provides Resistance to Protein Synthesis Inhibitor Microcin C by Acetylating the Processed Form of the Antibiotic. <i>Journal of Biological Chemistry</i> , 2010, 285, 12662-12669.	1.6	35
49	Proof of concept for the reduction of classical swine fever infection in pigs by a novel viral polymerase inhibitor. <i>Journal of General Virology</i> , 2009, 90, 1335-1342.	1.3	24
50	Biological effects of hexitol and altritol-modified siRNAs targeting B-Raf. <i>European Journal of Pharmacology</i> , 2009, 606, 38-44.	1.7	40
51	Synthesis and Base Pairing Properties of 1,5-Anhydro-Hexitol Nucleic Acids (HNA). <i>Chemistry - A European Journal</i> , 2009, 15, 10121-10131.	1.7	30
52	Detection of RNA Hybridization by Pyrene-Labeled Probes. <i>ChemBioChem</i> , 2009, 10, 1175-1185.	1.3	32
53	Dendritic Nucleotides: Interaction with an Aliphatic Acid Monolayer. <i>Chemistry and Biodiversity</i> , 2008, 5, 1675-1682.	1.0	1
54	Characterization of the Posttranscriptional Modifications in <i>Legionella pneumophila</i> Small Subunit Ribosomal RNA. <i>Chemistry and Biodiversity</i> , 2008, 5, 2640-2653.	1.0	8

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55	Post-transcriptional modification mapping in the <i>Clostridium acetobutylicum</i> 16S rRNA by mass spectrometry and reverse transcriptase assays. <i>Nucleic Acids Research</i> , 2007, 35, 3494-3503.	6.5	34
56	2'-O-Hydroxyalkoxymethylribonucleosides and their Incorporation into Oligoribonucleotides. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2007, 26, 1509-1512.	0.4	4
57	Synthesis of Oligoribonucleotides Containing Pyrimidine 2'-O-[(Hydroxyalkoxy)methyl]ribonucleosides. <i>Collection of Czechoslovak Chemical Communications</i> , 2006, 71, 804-819.	1.0	6
58	Synthesis and Anti-BVDV Activity of Acridones As New Potential Antiviral Agents1. <i>Journal of Medicinal Chemistry</i> , 2006, 49, 2621-2627.	2.9	71
59	Substituted 5-benzyl-2-phenyl-5H-imidazo[4,5-c]pyridines: A new class of pestivirus inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006, 16, 5345-5349.	1.0	28
60	Characterization and sequence verification of thiolated deoxyoligonucleotides used for microarray construction. <i>Journal of the American Society for Mass Spectrometry</i> , 2006, 17, 1397-1400.	1.2	6
61	A Novel, Highly Selective Inhibitor of Pestivirus Replication That Targets the Viral RNA-Dependent RNA Polymerase. <i>Journal of Virology</i> , 2006, 80, 149-160.	1.5	78
62	Development of Synthetic Strategies for the Construction of Pyrido[4,3-d]pyrimidine Libraries – the Discovery of a New Class of PDE-4 Inhibitors. <i>European Journal of Organic Chemistry</i> , 2006, 2006, 4257-4269.	1.2	14
63	Synthetic dsDNA-Binding Peptides Using Natural Compounds as Model. <i>Helvetica Chimica Acta</i> , 2006, 89, 1194-1219.	1.0	6
64	Synthesis, in vitro cellular uptake and photo-induced antiproliferative effects of lipophilic hypericin acid derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2005, 13, 6347-6353.	1.4	13
65	Synthesis and Properties of Oligonucleotides Containing 2,4-Dihydroxycyclohexyl Nucleosides. <i>Helvetica Chimica Acta</i> , 2005, 88, 3210-3224.	1.0	3
66	Synthesis of RNA Containing O ² -D-Ribofuranosyl-(1 ^α ;2 ^α)-adenosine-5 ^α -phosphate and 1-Methyladenosine, Minor Components of tRNA. <i>Chemistry and Biodiversity</i> , 2005, 2, 1153-1163.	1.0	11
67	Synthesis and Properties of Aminopropyl Nucleic Acids. <i>ChemBioChem</i> , 2005, 6, 2298-2304.	1.3	25
68	Synthesis of Peptidoglycan Units with UDP at the Anomeric Position. <i>Collection of Czechoslovak Chemical Communications</i> , 2005, 70, 1615-1641.	1.0	16
69	Delivery of Antisense Oligonucleotides Using Cholesterol-Modified Sense Dendrimers and Cationic Lipids. <i>Bioconjugate Chemistry</i> , 2005, 16, 827-836.	1.8	24
70	ENZYMATIC RESOLUTION AND BASE PAIRING PROPERTIES OF D- AND L-CYCLOHEXYNYL NUCLEIC ACIDS (CeNA). <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2005, 24, 993-998.	0.4	3
71	Chemical incorporation of minor tRNA component O ² -D-ribofuranosyl-(1''-2')-adenosine-5''-phosphate into oligoribonucleotides. , 2005, , .		0
72	Synthesis of enantiomeric-pure cyclohexenyl nucleoside building blocks for oligonucleotide synthesis. <i>Tetrahedron</i> , 2004, 60, 2111-2123.	1.0	12

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73	Synthesis and Stability of Oligonucleotides Containing Acyclic Achiral Nucleoside Analogues with Two Base Moieties. <i>Organic Letters</i> , 2004, 6, 51-54.	2.4	28
74	The Small Subunit rRNA Modification Database. <i>Nucleic Acids Research</i> , 2004, 33, D135-D138.	6.5	61
75	Synthesis and Properties of O-D-ribofuranosyl-(1 α €³ α †'2 α €²)-guanosine-5 α €³- O-phosphate and Its Derivatives. <i>Helvetica Chimica Acta</i> , 2003, 86, 504-514.	1.0	9
76	New dsDNA-Binding Hybrid Molecules Combining an Unnatural Peptide and an Intercalating Moiety. <i>Helvetica Chimica Acta</i> , 2003, 86, 533-547.	1.0	5
77	Mycobacterium tuberculosis Thymidine Monophosphate Kinase Inhibitors: Biological Evaluation and Conformational Analysis of 2 α €²- and 3 α €²-Modified Thymidine Analogues. <i>European Journal of Organic Chemistry</i> , 2003, 2003, 2911-2918.	1.2	9
78	Straightforward Synthesis of Labeled and Unlabeled Pyrimidine d4Ns via 2 α €²,3 α €²-Diyne seco Analogues through Olefin Metathesis Reactions. <i>European Journal of Organic Chemistry</i> , 2003, 2003, 666-671.	1.2	16
79	2,5-Bis-(2-hydroxybenzoylamino)pentanoic acid, a salicylic acid-metabolite isolated from chicken: characterization and independent synthesis. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2003, 13, 335-337.	1.0	4
80	Synthesis and Conformational Properties of O- $\hat{1}$ ²-D-Ribofuranosyl-(1 α €³-2 α €²)-guanosine and (Adenosine)-5 α €³-phosphate. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2003, 22, 1109-1111.	0.4	4
81	Ribavirin Derivatives with a Hexitol Moiety: Synthesis and Antiviral Evaluation. <i>Antiviral Chemistry and Chemotherapy</i> , 2003, 14, 23-30.	0.3	3
82	Synthesis and Antiviral Activity of a Series of New Cyclohexenyl Nucleosides. <i>Antiviral Chemistry and Chemotherapy</i> , 2003, 14, 31-37.	0.3	8
83	Base Pairing Properties of D- and L-Cyclohexene Nucleic Acids (CeNA). <i>Oligonucleotides</i> , 2003, 13, 479-489.	2.7	12
84	Evaluation of Capillary HPLC/Mass Spectrometry as an Alternative Analysis Method for Gel Electrophoresis of Oligonucleotides. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2003, 22, 1513-1516.	0.4	3
85	Chemical Incorporation of 1-Methyladenosine, Minor tRNA Component, into Oligonucleotides. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2003, 22, 1113-1115.	0.4	1
86	Methylated Hexitol Nucleic Acids, Towards Congeners with Improved Antisense Potential. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2003, 22, 1227-1229.	0.4	2
87	Synthesis and Biological Evaluation of a Series of New Cyclohexenyl Nucleosides. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2003, 22, 845-847.	0.4	3
88	Cleavage of DNA without loss of genetic information by incorporation of a disaccharide nucleoside. <i>Nucleic Acids Research</i> , 2003, 31, 6758-6769.	6.5	4
89	Recognition of threosyl nucleotides by DNA and RNA polymerases. <i>Nucleic Acids Research</i> , 2003, 31, 6221-6226.	6.5	76
90	Synthesis and Properties of Phosphorylated 3 α €²-O- $\hat{1}$ ²-D-Ribofuranosyl-2 α €²-deoxythymidine. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2003, 22, 359-371.	0.4	7

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91	Chemical incorporation of 1-methyladenosine into oligonucleotides. <i>Nucleic Acids Research</i> , 2002, 30, 1124-1131.	6.5	32
92	Modification of the universally unmodified uridine-33 in a mitochondria-imported edited tRNA and the role of the anticodon arm structure on editing efficiency. <i>Rna</i> , 2002, 8, 752-761.	1.6	47
93	5â€-Deoxy Congeners of 9-(3-Amido-3-deoxy-Î²-d-xylofuranosyl)-N6-cyclopentyladenine:Â New Adenosine A1Receptor Antagonists and Inverse Agonists. <i>Journal of Medicinal Chemistry</i> , 2002, 45, 1845-1852.	2.9	13
94	Recognition of HNA and 1,5-anhydrohexitol nucleotides by DNA metabolizing enzymes. <i>BBA - Proteins and Proteomics</i> , 2002, 1597, 115-122.	2.1	10
95	Selection of New Sequence-Selective Unnatural Peptides Binding to Double-Stranded Deoxyribonucleic Acids (dsDNA) by Means of a Gel-Retardation Experiment for Library Analysis. <i>Helvetica Chimica Acta</i> , 2002, 85, 2258-2283.	1.0	14
96	SOS: A simple interactive program for ab initio oligonucleotide sequencing by mass spectrometry. <i>Journal of the American Society for Mass Spectrometry</i> , 2002, 13, 200-203.	1.2	94
97	Characterization and sequence confirmation of unnatural amino acid containing peptide libraries using electrospray ionization mass spectrometry. <i>Rapid Communications in Mass Spectrometry</i> , 2002, 16, 982-987.	0.7	3
98	New dsDNA binding unnatural oligopeptides with pyrimidine selectivity. <i>Bioorganic and Medicinal Chemistry</i> , 2002, 10, 3401-3413.	1.4	14
99	Protection of 1-methyladenosine and its chemical incorporation into oligonucleotides. , 2002, , .		0
100	INCREASED RNA AFFINITY OF HNA ANALOGUES BY INTRODUCING ALKOXY SUBSTITUENTS AT THE C-1 OR C-3 POSITION. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2001, 20, 781-784.	0.4	3
101	Hybridization between â€Six-Memberedâ€-Nucleic Acids:â€% RNA as a Universal Information System. <i>Organic Letters</i> , 2001, 3, 4129-4132.	2.4	19
102	Oligonucleotides Containing Disaccharide Nucleosides. <i>Helvetica Chimica Acta</i> , 2001, 84, 2387-2397.	1.0	22
103	Î±-Homo-DNA and RNA Form a Parallel Oriented Non-A, Non-B-Type Double Helical Structure. <i>Chemistry - A European Journal</i> , 2001, 7, 5183-5194.	1.7	19
104	Suzuki reactions on chloropyridazinones: an easy approach towards arylated 3(2 H)-pyridazinones. <i>Tetrahedron</i> , 2001, 57, 1323-1330.	1.0	55
105	Synthesis and Properties of O-Î²-D-Ribofuranosyl-(1â€³-2â€²)-Adenosine-5â€³-O-Phosphate and Its Derivatives. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2000, 19, 1847-1859.	0.4	14
106	Problems and Prospects in the Characterization of Posttranscriptional Modifications in Large RNAs. , 2000, , 531-551.		0
107	Synthesis of protected D-altritol nucleosides as building blocks for oligonucleotide synthesis. <i>Tetrahedron</i> , 1999, 55, 6527-6546.	1.0	35
108	Characterization of Oligonucleotide Sequence Isomers in Mixtures Using HPLC/MS. <i>Nucleosides & Nucleotides</i> , 1999, 18, 1539-1540.	0.5	3

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109	Determination of Nearest Neighbors in Nucleic Acids by Mass Spectrometry. <i>Analytical Chemistry</i> , 1999, 71, 1454-1459.	3.2	21
110	DNA-Binding Ligands from Peptide Libraries Containing Unnatural Amino Acids. <i>Chemistry - A European Journal</i> , 1998, 4, 425-433.	1.7	26
111	²⁹ Si NMR spectra of trimethylsilyl and tert-butyl dimethylsilyl derivatives of purines and pyrimidines. <i>Magnetic Resonance in Chemistry</i> , 1998, 36, 55-63.	1.1	6
112	Oligonucleotides Composed of 2'-Deoxy-5'-anhydro-d-mannitol Nucleosides with a Purine Base Moiety. <i>Journal of Organic Chemistry</i> , 1998, 63, 1574-1582.	1.7	22
113	Enantioselective Approach to the Synthesis of Cyclohexane Carbocyclic Nucleosides. <i>Journal of Organic Chemistry</i> , 1998, 63, 3051-3058.	1.7	31
114	Synthesis and hybridization properties of inverse oligonucleotides. <i>Nucleic Acids Research</i> , 1997, 25, 3034-3041.	6.5	7
115	An Efficient Synthesis and Physico-Chemical Properties OF 2'-O-d-Ribofuranosyl nucleosides, Minor tRNA Components. <i>Journal of Carbohydrate Chemistry</i> , 1997, 16, 75-92.	0.4	37
116	Synthesis and Antiviral Activity of the β -Analogues of 1,5-Anhydrohexitol Nucleosides (1,5-Anhydro-2,3-dideoxy-d-ribohexitol Nucleosides). <i>Journal of Organic Chemistry</i> , 1997, 62, 2442-2447.	1.7	42
117	Synthesis and Conformational Study of 3-Hydroxy-4-(Hydroxymethyl)-1-Cyclohexanyl Purines and Pyrimidines. <i>Journal of Organic Chemistry</i> , 1997, 62, 2861-2871.	1.7	66
118	N ⁶ -Cyclopentyl-3'-substituted-xylofuranosyladenosines: A New Class of Non-Xanthine Adenosine A ₁ Receptor Antagonists. <i>Journal of Medicinal Chemistry</i> , 1997, 40, 3765-3772.	2.9	25
119	Screening a Random Pentapeptide Library, Composed of 14 D-Amino Acids, against the COOH-terminal Sequence of Fructose-1,6-bisphosphate Aldolase from <i>Trypanosoma brucei</i> . <i>Journal of Biological Chemistry</i> , 1997, 272, 11378-11383.	1.6	4
120	Oligonucleotides with 3-hydroxy-N-acetylprolinol as sugar substitute. <i>Tetrahedron</i> , 1997, 53, 14957-14974.	1.0	13
121	Improved Synthesis of Anhydrohexitol Building Blocks for Oligonucleotide Synthesis. <i>Liebigs Annalen</i> , 1997, 1997, 1453-1461.	0.8	17
122	Oligonucleotide Analogues with 4-Hydroxy-N-Acetylprolinol as Sugar Substitute. <i>Chemistry - A European Journal</i> , 1997, 3, 1997-2010.	1.7	20
123	Fast-atom bombardment mass spectrometric study of SATE Foscarnet prodrugs and of a series of Foscarnet-AZT conjugates. <i>Rapid Communications in Mass Spectrometry</i> , 1997, 11, 1212-1218.	0.7	2
124	β -Amino acids derived from ornithine as building blocks for peptide synthesis. <i>Chemical Biology and Drug Design</i> , 1997, 49, 183-189.	1.2	0
125	Interpretation of Oligonucleotide Mass Spectra for Determination of Sequence Using Electrospray Ionization and Tandem Mass Spectrometry. <i>Analytical Chemistry</i> , 1996, 68, 1989-1999.	3.2	189
126	Synthesis of homo-N-nucleosides, a series of C1' branched-chain nucleosides. <i>Tetrahedron</i> , 1996, 52, 5563-5578.	1.0	38

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127	Incorporation of 5-hydroxytryptophan in oligopeptides. <i>Tetrahedron</i> , 1996, 52, 6965-6972.	1.0	7
128	Synthesis and antiviral activity of acyclic analogues of 1,5-anhydrohexitol nucleosides using Mitsunobu reaction. <i>Tetrahedron</i> , 1996, 52, 13655-13670.	1.0	28
129	Synthesis of 1,5-Anhydrohexitol Nucleosides as Mimics of AZT, D4T and DDC. <i>Nucleosides & Nucleotides</i> , 1996, 15, 325-335.	0.5	2
130	Amino acids derived from ornithine. <i>International Journal of Peptide Research and Therapeutics</i> , 1995, 2, 206-208.	0.1	2
131	5-Hydroxytryptophan as a building block in oligopeptides using Fmoc/tBu SPPS. <i>International Journal of Peptide Research and Therapeutics</i> , 1995, 2, 225-228.	0.1	0
132	Screening of a synthetic pentapeptide library composed of d-amino acids against fructose-1,6-biphosphate aldolase. <i>International Journal of Peptide Research and Therapeutics</i> , 1995, 2, 259-260.	0.1	3
133	The effect of addition of carbon powder to samples in liquid secondary ion mass spectrometry: Improved ionization of apolar compounds. <i>Rapid Communications in Mass Spectrometry</i> , 1995, 9, 1499-1501.	0.7	4
134	Identification of a peptide inhibitor against glycosomal phosphoglycerate kinase of <i>Trypanosoma brucei</i> by a synthetic peptide library approach. <i>Bioorganic and Medicinal Chemistry</i> , 1995, 3, 257-265.	1.4	23
135	Stereocontrolled synthesis of phosphonate derivatives of tetrahydro- and dihydro-2H-pyranil nucleosides: The selectivity of the Ferrier rearrangement. <i>Tetrahedron: Asymmetry</i> , 1995, 6, 973-984.	1.8	15
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