Roger Stromberg

List of Publications by Year in descending order

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218677 243625 2,323 112 26 44 citations g-index h-index papers 113 113 113 1910 docs citations times ranked citing authors all docs

#	Article	lF	CITATIONS
1	Influence of sequence variation on the RNA cleavage activity of Zn ²⁺ -dimethyl-dppz-PNA-based artificial enzymes. RSC Advances, 2022, 12, 5398-5406.	3.6	3
2	NanoSIMS Imaging Reveals the Impact of Ligand-ASO Conjugate Stability on ASO Subcellular Distribution. Pharmaceutics, 2022, 14, 463.	4.5	4
3	Innovative developments and emerging technologies in RNA therapeutics. RNA Biology, 2022, 19, 313-332.	3.1	19
4	2′- <i>O</i> -(<i>N</i> -(Aminoethyl)carbamoyl)methyl Modification Allows for Lower Phosphorothioate Content in Splice-Switching Oligonucleotides with Retained Activity. Nucleic Acid Therapeutics, 2022, , .	3.6	4
5	Synthesis and biological evaluation of modified laminin peptide (N2S2-KDP) with enhanced affinity for neuronal growth and targeted molecular imaging (SPECT). Bioorganic Chemistry, 2021, 107, 104516.	4.1	4
6	The Mechanism of Cleavage of RNA Phosphodiesters by a Gold Nanoparticle Nanozyme. Chemistry - A European Journal, 2021, 27, 8143-8148.	3.3	7
7	34S-SIL of PCSK9-Active Oligonucleotide as Tools for Accurate Quantification by Mass Spectrometry. Nucleic Acid Therapeutics, 2021, 31, 375-381.	3.6	0
8	Zn ²⁺ -Dependent peptide nucleic acid-based artificial ribonucleases with unprecedented efficiency and specificity. Chemical Communications, 2021, 57, 10911-10914.	4.1	7
9	New Alkyne and Amine Linkers for Versatile Multiple Conjugation of Oligonucleotides. ACS Omega, 2021, 6, 579-593.	3.5	8
10	A Study on Synthesis and Upscaling of 2′-O-AECM-5-methyl Pyrimidine Phosphoramidites for Oligonucleotide Synthesis. Molecules, 2021, 26, 6927.	3.8	1
11	Copperâ€Catalyzed Huisgen 1,3â€Dipolar Cycloaddition Tailored for Phosphorothioate Oligonucleotides. Current Protocols in Nucleic Acid Chemistry, 2020, 80, e102.	0.5	1
12	Attachment of Peptides to Oligonucleotides on Solid Support Using Copper(I)-Catalyzed Huisgen 1,3-Dipolar Cycloaddition. Methods in Molecular Biology, 2019, 2036, 165-171.	0.9	1
13	Amyloid- \hat{l}^2 Peptide Targeting Peptidomimetics for Prevention of Neurotoxicity. ACS Chemical Neuroscience, 2019, 10, 1462-1477.	3.5	7
14	Novel aroylated phenylenediamine compounds enhance antimicrobial defense and maintain airway epithelial barrier integrity. Scientific Reports, 2019, 9, 7114.	3.3	12
15	Efficient Conjugation to Phosphorothioate Oligonucleotides by Cu-Catalyzed Huisgen 1,3-Dipolar Cycloaddition. Bioconjugate Chemistry, 2019, 30, 1622-1628.	3.6	14
16	Further Probing of Cu2+-Dependent PNAzymes Acting as Artificial RNA Restriction Enzymes. Molecules, 2019, 24, 672.	3.8	9
17	A Versatile and Convenient Synthesis of ³⁴ S‣abeled Phosphorothioate Oligonucleotides. ChemBioChem, 2018, 19, 2114-2119.	2.6	7
18	Facile Access to Bromonucleosides Using Sodium Monobromoisocyanurate (SMBI). Current Protocols in Nucleic Acid Chemistry, 2017, 68, 1.39.1-1.39.9.	0.5	1

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19	Treatment with Entinostat Heals Experimental Cholera by Affecting Physical and Chemical Barrier Functions of Intestinal Epithelia. Antimicrobial Agents and Chemotherapy, 2017, 61, .	3.2	16
20	¹⁹ Fâ€NMR Spectroscopic Analysis of the Binding Modes in Tripleâ€Helical Peptide Nucleic Acid (PNA)/MicroRNA Complexes. Chemistry - A European Journal, 2017, 23, 7113-7124.	3.3	24
21	Facile functionalization of peptide nucleic acids (PNAs) for antisense and single nucleotide polymorphism detection. Organic and Biomolecular Chemistry, 2017, 15, 6710-6714.	2.8	6
22	Zinc Ion-Dependent Peptide Nucleic Acid-Based Artificial Enzyme that Cleaves RNAâ€"Bulge Size and Sequence Dependence. Molecules, 2017, 22, 1856.	3.8	14
23	Clamping of RNA with PNA enables targeting of microRNA. Organic and Biomolecular Chemistry, 2016, 14, 5210-5213.	2.8	6
24	Entinostat up-regulates the CAMP gene encoding LL-37 via activation of STAT3 and HIF- $1\hat{l}\pm$ transcription factors. Scientific Reports, 2016, 6, 33274.	3.3	38
25	Enabling Multiple Conjugation to Oligonucleotides Using "Click Cycles― Bioconjugate Chemistry, 2016, 27, 2620-2628.	3.6	10
26	Clickable trimethylguanosine cap analogs modified within the triphosphate bridge: synthesis, conjugation to RNA and susceptibility to degradation. RSC Advances, 2016, 6, 8317-8328.	3.6	9
27	Sequence-specific RNA cleavage by PNA conjugates of the metal-free artificial ribonuclease tris(2-aminobenzimidazole). Beilstein Journal of Organic Chemistry, 2015, 11, 493-498.	2.2	26
28	Synthesis of Triamino Acid Building Blocks with Different Lipophilicities. PLoS ONE, 2015, 10, e0124046.	2.5	2
29	Synthesis of fluorescent d-amino acids with 4-acetamidobiphenyl and 4-N,N-dimethylamino-1,8-naphthalimido containing side chains. Tetrahedron Letters, 2015, 56, 4780-4783.	1.4	8
30	Nuclease resistant oligonucleotides with cell penetrating properties. Chemical Communications, 2015, 51, 4044-4047.	4.1	18
31	Sequence-selective DNA recognition and enhanced cellular up-take by peptide–steroid conjugates. Chemical Communications, 2015, 51, 17552-17555.	4.1	8
32	Studies on Tris(2-aminobenzimidazole)-PNA Based Artificial Nucleases: A Comparison of Two Analytical Techniques. Bioconjugate Chemistry, 2015, 26, 2514-2519.	3.6	9
33	Synthesis of PNA Oligoether Conjugates. Molecules, 2014, 19, 3135-3148.	3.8	2
34	N2-tert-Butoxycarbonyl-N5-[N-(9-fluorenylmethyloxycarbonyl)-2-aminoethyl]-(S)-2,5-diaminopentanoic Acid. MolBank, 2014, 2014, M833.	0.5	1
35	Boosting innate immunity: Development and validation of a cell-based screening assay to identify LL-37 inducers. Innate Immunity, 2014, 20, 364-376.	2.4	28
36	Amyloid- \hat{l}^2 -Induced Action Potential Desynchronization and Degradation of Hippocampal Gamma Oscillations Is Prevented by Interference with Peptide Conformation Change and Aggregation. Journal of Neuroscience, 2014, 34, 11416-11425.	3.6	91

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37	Synthesis and evaluation of antineurotoxicity properties of an amyloid \hat{l}^2 peptide targeting ligand containing a triamino acid. Organic and Biomolecular Chemistry, 2014, 12, 6684-6693.	2.8	6
38	Synthesis and evaluation of stability of m3G-CAP analogues in serum-supplemented medium and cytosolic extract. Bioorganic and Medicinal Chemistry, 2013, 21, 7921-7928.	3.0	10
39	Synthesis and Stability of a 2′â€ <i>O</i> àâ€[<i>N</i> â€(Aminoethyl)carbamoyl]methyladenosineâ€Containing Dinucleotide. European Journal of Organic Chemistry, 2013, 2013, 7184-7192.	2.4	7
40	An Efficient and Facile Methodology for Bromination of Pyrimidine and Purine Nucleosides with Sodium Monobromoisocyanurate (SMBI). Molecules, 2013, 18, 12740-12750.	3.8	7
41	Lactose in Human Breast Milk an Inducer of Innate Immunity with Implications for a Role in Intestinal Homeostasis. PLoS ONE, 2013, 8, e53876.	2.5	76
42	Capping of oligonucleotides with "clickable―m3G-CAPs. RSC Advances, 2012, 2, 12949.	3.6	17
43	Diaminopropionic acid lipopeptides: Characterization studies of polyplexes aimed at pDNA delivery. Bioorganic and Medicinal Chemistry Letters, 2012, 22, 5635-5638.	2.2	1
44	Effects of Ligands on Unfolding of the Amyloid \hat{l}^2 -Peptide Central Helix: Mechanistic Insights from Molecular Dynamics Simulations. PLoS ONE, 2012, 7, e30510.	2.5	17
45	Stability of a 2′â€∢i>Oàâ€(Carbamoylmethyl)adenosineâ€Containing Dinucleotide. European Journal of Organic Chemistry, 2012, 2012, 539-543.	2.4	3
46	An activated triple bond linker enables â€ ⁻ clickâ€ ^{-™} attachment of peptides to oligonucleotides on solid support. Nucleic Acids Research, 2011, 39, 9047-9059.	14.5	34
47	Synthesis of estradiol backbone mimics via the Stille reaction using copper(II) oxide as co-reagent. Tetrahedron Letters, 2011, 52, 209-211.	1.4	14
48	Solid phase synthesis, radiolabeling and biological evaluation of a \sup 99m $ $ sup $ Tc$ -labeled $ $ 1 $ t$ 2 $ t$ 3 $ t$ 4 $ t$ 6 phase synthesis, radiolabeling and biological evaluation of a t 8 $ t$ 99m $ $ 1 $ t$ 99m $ $ 2 $ t$ 9 $ t$ 90m $ $ 2 $ t$ 9 $ t$ 90m $ $ 2 $ t$ 9 $ t$ 90m $ $ 2 $ t$ 9	3.4	11
49	Unfolding of the Amyloid \hat{l}^2 -Peptide Central Helix: Mechanistic Insights from Molecular Dynamics Simulations. PLoS ONE, 2011, 6, e17587.	2.5	26
50	Phenylbutyrate Counteracts Shigella Mediated Downregulation of Cathelicidin in Rabbit Lung and Intestinal Epithelia: A Potential Therapeutic Strategy. PLoS ONE, 2011, 6, e20637.	2.5	78
51	Cationic Peptides that Increase the Thermal Stabilities of 2′â€∢i>Oâ€MeRNA/RNA Duplexes but Do Not Affect DNA/DNA Melting. ChemBioChem, 2010, 11, 2606-2612.	2.6	15
52	PNAzymes That Are Artificial RNA Restriction Enzymes. Journal of the American Chemical Society, 2010, 132, 8984-8990.	13.7	61
53	A synthetic snRNA m3G-CAP enhances nuclear delivery of exogenous proteins and nucleic acids. Nucleic Acids Research, 2009, 37, 1925-1935.	14.5	29
54	Investigation on Condensing Agents for Phosphinate Ester Formation with Nucleoside 5â€2â€Hydroxyl Functions. European Journal of Organic Chemistry, 2008, 2008, 1705-1714.	2.4	12

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55	PNA based artificial nucleases displaying catalysis with turnover in the cleavage of a leukemia related RNA model. Organic and Biomolecular Chemistry, 2008, 6, 3837.	2.8	37
56	Analysis of the Stability and Flexibility of RNA Complexes Containing Bulge Loops of Different Sizes. Journal of Biomolecular Structure and Dynamics, 2008, 26, 163-173.	3.5	21
57	Solid Support Post-Conjugation of Amino Acids and a Phenanthroline Derivative to a Central Position in Peptide Nucleic Acids. Nucleosides, Nucleotides and Nucleic Acids, 2007, 26, 1485-1489.	1.1	9
58	RNA Cleavage by 2,9-Diamino-1,10-Phenanthroline PNA Conjugates. Nucleosides, Nucleotides and Nucleic Acids, 2007, 26, 1479-1483.	1.1	14
59	Oligoribonucleotide Analogues Containing a Mixed Backbone of Phosphodiester and Formacetal Internucleoside Linkages, Together with Vicinal 2′-O-Methyl Groups. ChemBioChem, 2007, 8, 537-545.	2.6	12
60	Biological Activity and Biotechnological Aspects of Peptide Nucleic Acid. Advances in Genetics, 2006, 56, 1-51.	1.8	97
61	Synthesis of 8-aminoadenosine 5′-(aminoalkyl phosphates), analogues of aminoacyl adenylates. Bioorganic and Medicinal Chemistry, 2006, 14, 2653-2659.	3.0	4
62	STUDIES IN OLIGONUCLEOTIDE-BASED ARTIFICIAL NUCLEASE SYSTEMS. INTRAMOLECULAR COPPER (II) COMPLEX FORMATION IN AN OLIGONUCLEOTIDE BIS-PHENANTHROLINE CONJUGATE. Nucleosides, Nucleotides and Nucleic Acids, 2005, 24, 901-905.	1.1	6
63	A SOLID SUPPORTED REAGENT FOR INTERNUCLEOSIDE H-PHOSPHONATE LINKAGE FORMATION. Nucleosides, Nucleotides and Nucleic Acids, 2005, 24, 897-899.	1.1	1
64	Compelling evidence for a stepwise mechanism of the alkaline cyclisation of uridine 3′-phosphate esters. Organic and Biomolecular Chemistry, 2004, 2, 2165-2167.	2.8	65
65	Acidity of Secondary Hydroxyls in ATP and Adenosine Analogues and the Question of a 2â€~,3â€~-Hydrogen Bond in Ribonucleosides. Journal of the American Chemical Society, 2004, 126, 14710-14711.	13.7	24
66	Synthesis of new OBAN's and further studies on positioning of the catalytic group. Organic and Biomolecular Chemistry, 2004, 2, 1901-1907.	2.8	43
67	A Method for Solid-Phase Synthesis of Oligonucleotide 5â€~-Peptide-Conjugates Using Acid-Labile α-Amino Protections. Journal of the American Chemical Society, 2004, 126, 14029-14035.	13.7	41
68	Characterization of an RNA bulge structure by Fourier transform infrared spectroscopy. Biochemical and Biophysical Research Communications, 2004, 324, 634-639.	2.1	9
69	Synthesis and Properties of RNA Analogues Having Amides as Interuridine Linkages at Selected Positions. Journal of the American Chemical Society, 2003, 125, 12125-12136.	13.7	62
70	Application of Nim-2,6-Dimethoxybenzoyl Histidine in Solid-Phase Peptide Synthesis. European Journal of Organic Chemistry, 2003, 2003, 2454-2461.	2.4	3
71	Stabilisation of RNA Bulges by Oligonucleotide Complements Containing an Adenosine Analogue. ChemBioChem, 2003, 4, 1194-1200.	2.6	8
72	Facile Determination of the Protecting Group Location of Nim-Protected Histidine Derivatives by 1Hâ^15N Heteronuclear Correlation NMR. Journal of Organic Chemistry, 2003, 68, 7521-7523.	3.2	8

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73	Oligonucleotide based artificial nuclease (OBAN) systems. Bulge size dependence and positioning of catalytic group in cleavage of RNA-bulges. Organic and Biomolecular Chemistry, 2003, 1, 1461-1465.	2.8	48
74	Side Reactions in the H-Phosphonate Approach to Oligonucleotide Synthesis: A Kinetic Investigation on Bisacylphosphite Formation and 5′-O-Acylation. Nucleosides, Nucleotides and Nucleic Acids, 2003, 22, 1-12.	1.1	8
75	Synthesis of 2′-Deuterio and 3′-Deuterio Cytidine 5′-Diphosphate. Nucleosides, Nucleotides and Nucleic Acids, 2003, 22, 1657-1659.	1.1	3
76	Comparison of Some Computational Methods for Geometry Optimisation of Phosphorus Acid Derivatives. Phosphorus, Sulfur and Silicon and the Related Elements, 2002, 177, 2711-2724.	1.6	4
77	Reactions of 3′-C-Halomethyl and 3′-C-Sulfonylmethyl Uridines with Phosphinic Acid Derivatives − Synthesis of Building Blocks for Oligonucleotides Containing 3′-C-Methylenephosphonate Linkages. European Journal of Organic Chemistry, 2002, 2002, 1509-1515.	2.4	5
78	Stability Studies of N-Acylimidazoles. European Journal of Organic Chemistry, 2002, 2002, 2633.	2.4	11
79	Synthesis of Nucleic Acid Fragments with 3′-Deoxy-3′-C-Methylenephosphonate Linkages ⬲ Oxidation Of Nucleoside 3′-Deoxy-3′-C-Methylenephosphinate Esters. European Journal of Organic Chemistry, 2002, 2002, 3140-3144.	2.4	3
80	A METHOD FOR SYNTHESIS OF AN ARTIFICIAL RIBONUCLEASE. Nucleosides, Nucleotides and Nucleic Acids, 2001, 20, 1385-1388.	1.1	14
81	EVALUATION OF SEVERAL ECONOMICAL COMPUTATIONAL METHODS FOR GEOMETRY OPTIMISATION OF PHOSPHORUS ACID DERIVATIVES. Nucleosides, Nucleotides and Nucleic Acids, 2001, 20, 1381-1384.	1.1	4
82	Specific metal-ion binding sites in a model of the P4-P6 triple-helical domain of a group I intron. Rna, 2001, 7, 1115-1125.	3.5	9
83	Stereoselectivity in the Synthesis of 3′-Deoxy-3′-C-(hydroxymethyl)uridines by Hydroboration and Conversion into a Building Block for Various 3′-Deoxy-3′-C-(methylene)uridine Analogues. European Journal of Organic Chemistry, 2001, 2001, 4305.	2.4	12
84	PREPARATION OF $3\hat{a}\in^2$ -C-BRANCHED URIDINE ANALOGUES, SUITABLE FOR CONVERSION INTO FUNCTIONALISED $3\hat{a}\in^2$ -C-METHYLENE DERIVATIVES. Nucleosides, Nucleotides and Nucleic Acids, 2001, 20, 1389-1392.	1.1	1
85	Mechanism of RNase T1: concerted triester-like phosphoryl transfer via a catalytic three-centered hydrogen bond. Chemistry and Biology, 2000, 7, 651-658.	6.0	28
86	The mechanism of the metal ion promoted cleavage of RNA phosphodiester bonds involves a general acid catalysis by the metal aquo ion on the departure of the leaving group. Journal of the Chemical Society Perkin Transactions II, 1999, , 1619-1626.	0.9	52
87	An engineered ribonuclease preferring phosphorothioate RNA. Nature Structural Biology, 1998, 5, 365-368.	9.7	11
88	Synthesis and Properties of Oligoribonucleotide Analogs Having Formacetal Internucleoside Linkages. Journal of Organic Chemistry, 1997, 62, 1846-1850.	3.2	23
89	Base Catalysis and Leaving Group Dependence in Intramolecular Alcoholysis of Uridine 3â€~-(Aryl) Tj ETQq1 1 0.78	4314 rgB ⁻ 13.7	T /Overlock
90	An approach towards the synthesis of oligomers containing a N-2-hydroxyethyl-aminomethylphosphonate backbone: A novel PNA analogue. Tetrahedron Letters, 1996, 37, 7857-7860.	1.4	42

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91	Solid support synthesis of all-Rp-oligo(ribonucleoside phosphorothioate)s. Nucleic Acids Research, 1996, 24, 3811-3820.	14.5	29
92	Synthesis and properties of 2'-O-methoxymethyl oligonucleotides. Collection of Czechoslovak Chemical Communications, 1996, 61, 283-286.	1.0	2
93	Chemical Synthesis of RNA-Fragment Analogues That Have Phosphorothioate Linkages of Identical Configuration. Nucleosides, Nucleotides and Nucleic Acids, 1995, 14, 879-881.	1.1	3
94	RNA-Synthesis Using the H-Phosphonate Approach and an Improved Protecting Group Strategy. Nucleosides, Nucleotides and Nucleic Acids, 1995, 14, 883-887.	1.1	2
95	Synthesis of RNA Fragments Using the H-Phosphonate Method and 2'-(2'-Chlorobenzoyl) Protection. Nucleosides, Nucleotides and Nucleic Acids, 1995, 14, 855-857.	1.1	3
96	Synthesis of Oligoarabinonucleotides Using H-Phosphonates. Nucleosides, Nucleotides and Nucleic Acids, 1995, 14, 851-853.	1.1	2
97	Hydrolytic Reactions of the Diastereomeric Phosphoromonothioate Analogs of Uridylyl(3',5')uridine: Kinetics and Mechanisms for Desulfurization, Phosphoester Hydrolysis, and Transesterification to the 2',5'-Isomers. Journal of Organic Chemistry, 1995, 60, 5620-5627.	3.2	59
98	Removal of f-butyldimethylsilyl protection in RNA-synthesis. Triethylamine trihydrofluoride (TEA, 3HF) is a more reliable alternative to tetrabutylammonium fluoride (TBAF). Nucleic Acids Research, 1994, 22, 2430-2431.	14.5	110
99	RNA-synthesis using H-phosphonates. Synchronizing 2'-OH and N-protection. Collection of Czechoslovak Chemical Communications, 1993, 58, 236-237.	1.0	9
100	Hydrolytic stability of nucleoside H-phosphonate and H-phosphonothioate diesters. Collection of Czechoslovak Chemical Communications, 1993, 58, 79-81.	1.0	0
101	2′-Amino-2′-deoxyguanosine is a cofactor for self-splicing in group I catalytic RNA. Biochemical and Biophysical Research Communications, 1992, 183, 842-847.	2.1	5
102	Stereospecific oxidation and oxidative coupling of H-phosphonate and H-phosphonothioate diesters. Tetrahedron Letters, 1992, 33, 3185-3188.	1.4	31
103	Intramolecular transesterification in thiophosphate-analogues of an RNA-dimer Tetrahedron Letters, 1991, 32, 3723-3726.	1.4	33
104	Synthesis of nucleoside methylphosphonates and nucleoside methylthiophosphonates via phosphinate intermediates. Collection of Czechoslovak Chemical Communications, 1990, 55, 145-148.	1.0	4
105	Studies on the t-butyldimethylsilyl group as 2'-O-protection in oligoribonucleotide synthesis via the H-phosphonate approach. Nucleic Acids Research, 1988, 16, 9285-9298.	14.5	81
106	Studies on Ribonucleoside Hydrogenphosphonates. Effect of a Vicinal Hydroxyl Function on the Stability of H-Phosphonate Diester Bond. Nucleosides & Nucleotides, 1988, 7, 321-337.	0.5	13
107	Studies on the Synthesis of Oligonucleotides <i>via</i> the Hydrogenphosphonate Approach. Nucleosides & Nucleotides, 1987, 6, 283-286.	0.5	10
108	Studies on the Oxidation of Nucleoside Hydrogenphosphonates. Nucleosides & Nucleotides, 1987, 6, 429-432.	0.5	9

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109	Activation of Nucleoside Hydrogenphosphonates by Use of Aryl Sulfonyl Chlorides. Nucleosides & Nucleotides, 1987, 6, 425-427.	0.5	2
110	Studies on the reaction of nucleoside phosphorodiesters with aryl sulfonyl chlorides. Tetrahedron Letters, 1986, 27, 2665-2666.	1.4	9
111	Nucleoside H-phosphonates. III. Chemical synthesis of oligodeoxyribonucleotides by the hydrogenphosphonate approach. Tetrahedron Letters, 1986, 27, 4051-4054.	1.4	179
112	Nucleoside H-phosphonates. IV. Automated solid phase synthesis of oligoribonucleotides by the hydrogenphosphonate approach. Tetrahedron Letters, 1986, 27, 4055-4058.	1.4	111