Gennaro Piccialli

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3024089/publications.pdf

Version: 2024-02-01

186265 315739 2,319 129 28 38 citations h-index g-index papers 136 136 136 2071 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Exploring a peptide nucleic acid-based antisense approach for CD5 targeting in chronic lymphocytic leukemia. PLoS ONE, 2022, 17, e0266090.	2.5	5
2	Nucleoside Analogs and Nucleoside Precursors as Drugs in the Fight against SARS-CoV-2 and Other Coronaviruses. Molecules, 2021, 26, 986.	3.8	60
3	Silver (I) N-Heterocyclic Carbene Complexes: A Winning and Broad Spectrum of Antimicrobial Properties. International Journal of Molecular Sciences, 2021, 22, 2497.	4.1	21
4	Transcriptomics and Metabolomics Integration Reveals Redox-Dependent Metabolic Rewiring in Breast Cancer Cells. Cancers, 2021, 13, 5058.	3.7	10
5	Synthesis, self-assembly-behavior and biomolecular recognition properties of thyminyl dipeptides. Arabian Journal of Chemistry, 2020, 13, 1966-1974.	4.9	18
6	From computational genomics to systems metabolomics for precision cancer medicine and drug discovery. Pharmacological Research, 2020, 151, 104479.	7.1	1
7	PNA-Based Graphene Oxide/Porous Silicon Hybrid Biosensor: Towards a Label-Free Optical Assay for Brugada Syndrome. Nanomaterials, 2020, 10, 2233.	4.1	10
8	Probing the DNA Reactivity and the Anticancer Properties of a Novel Tubercidin-Pt(II) Complex. Pharmaceutics, 2020, 12, 627.	4.5	6
9	π–π stacked DNA G-wire nanostructures formed by a short G-rich oligonucleotide containing a 3′–3′ inversion of polarity site. Organic Chemistry Frontiers, 2020, 7, 2187-2195.	4.5	8
10	Evaluation of an Analogue of the Marine Îμ-PLL Peptide as a Ligand of G-quadruplex DNA Structures. Marine Drugs, 2020, 18, 49.	4.6	24
11	Endogenous and artificial miRNAs explore a rich variety of conformations: a potential relationship between secondary structure and biological functionality. Scientific Reports, 2020, 10, 453.	3.3	7
12	New Linear Precursors of cIDPR Derivatives as Stable Analogs of cADPR: A Potent Second Messenger with Ca2+-Modulating Activity Isolated from Sea Urchin Eggs. Marine Drugs, 2019, 17, 476.	4.6	6
13	Porous Silicon-Based Aptasensors: The Next Generation of Label-Free Devices for Health Monitoring. Molecules, 2019, 24, 2216.	3.8	25
14	New G-Quadruplex-Forming Oligodeoxynucleotides Incorporating a Bifunctional Double-Ended Linker (DEL): Effects of DEL Size and ODNs Orientation on the Topology, Stability, and Molecularity of DEL-G-Quadruplexes. Molecules, 2019, 24, 654.	3.8	7
15	Peptide Nucleic Acid-Functionalized Adenoviral Vectors Targeting G-Quadruplexes in the P1 Promoter of Bcl-2 Proto-Oncogene: A New Tool for Gene Modulation in Anticancer Therapy. Bioconjugate Chemistry, 2019, 30, 572-582.	3.6	25
16	Anti-HIV activity of new higher order G-quadruplex aptamers obtained from tetra-end-linked oligonucleotides. Organic and Biomolecular Chemistry, 2018, 16, 2349-2355.	2.8	16
17	Synthesis and Biological Evaluation of a New Structural Simplified Analogue of cADPR, a Calcium-Mobilizing Secondary Messenger Firstly Isolated from Sea Urchin Eggs. Marine Drugs, 2018, 16, 89.	4.6	10
18	Design, Synthesis and Characterization of Novel Co-Polymers Decorated with Peptides for the Selective Nanoparticle Transport across the Cerebral Endothelium. Molecules, 2018, 23, 1655.	3.8	18

#	Article	IF	CITATIONS
19	Studies toward the Synthesis of Smenamide A, an Antiproliferative Metabolite from <i>Smenospongia aurea</i> : Total Synthesis of <i>ent</i> -Smenamide A and 16- <i>epi</i> -Smenamide A. ACS Omega, 2017, 2, 1477-1488.	3.5	19
20	Stabilization vs. destabilization of G-quadruplex superstructures: the role of the porphyrin derivative having spermine arms. Physical Chemistry Chemical Physics, 2017, 19, 17404-17410.	2.8	15
21	Synthesis and label free characterization of a bimolecular PNA homo quadruplex. Biochimica Et Biophysica Acta - General Subjects, 2017, 1861, 1222-1228.	2.4	8
22	Selfâ€Assembly of Gâ€Rich Oligonucleotides Incorporating a 3′–3′ Inversion of Polarity Site: A New Route Towards Gâ€Wire DNA Nanostructures. ChemistryOpen, 2017, 6, 599-605.	1.9	24
23	Peptide Nucleic Acids as miRNA Target Protectors for the Treatment of Cystic Fibrosis. Molecules, 2017, 22, 1144.	3.8	29
24	5-Amino-1-($2\hat{a}\in^2$, $3\hat{a}\in^2$ - <i>O</i> -isopropylidene- <scp>D</scp> -ribityl)-1 <i>H</i> -imidazole-4-carboxamide: a crystal structure with <i>Z</i> $\hat{a}\in^2$ = 4. Acta Crystallographica Section E: Crystallographic Communications, 2017, 73, 183-187.	0.5	1
25	Nanogravimetric and Optical Characterizations of Thrombin Interaction with a Self-Assembled Thiolated Aptamer. Journal of Sensors, 2016, 2016, 1-8.	1.1	8
26	Solid phase synthesis of a thrombin binding aptamer on macroporous silica for label free optical quantification of thrombin. RSC Advances, 2016, 6, 86762-86769.	3.6	39
27	Degradation of some representative polycyclic aromatic hydrocarbons by the water-soluble protein extracts from Zea mays L. cv PR32-B10. Chemosphere, 2016, 160, 258-265.	8.2	5
28	Screening Platform toward New Anti-HIV Aptamers Set on Molecular Docking and Fluorescence Quenching Techniques. Analytical Chemistry, 2016, 88, 2327-2334.	6.5	18
29	New synthetic AICAR derivatives with enhanced AMPK and ACC activation. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 748-753.	5.2	15
30	Synthesis and Evaluation of the Antiproliferative Properties of a Tethered Tubercidin–Platinum(II) Complex. European Journal of Organic Chemistry, 2015, 2015, 7550-7556.	2.4	6
31	Synthesis of 5â€Aminoimidazoleâ€4â€Carboxamide Riboside (AICAR) and Its Derivatives Using Inosine as Starting Material. Current Protocols in Nucleic Acid Chemistry, 2015, 63, 1.35.1-1.35.24.	0.5	3
32	Exploitation of a Very Small Peptide Nucleic Acid as a New Inhibitor of miR-509-3p Involved in the Regulation of Cystic Fibrosis Disease-Gene Expression. BioMed Research International, 2014, 2014, 1-10.	1.9	45
33	Outstanding effects on antithrombin activity of modified TBA diastereomers containing an optically pure acyclic nucleotide analogue. Organic and Biomolecular Chemistry, 2014, 12, 5235-5242.	2.8	27
34	Design, synthesis and biochemical investigation, by in vitro luciferase reporter system, of peptide nucleic acids as new inhibitors of miR-509-3p involved in the regulation of cystic fibrosis disease-gene expression. MedChemComm, 2014, 5, 68-71.	3.4	16
35	Aminosilane-modified mesoporous oxidized silicon for in situ oligonucleotides synthesis and detection. , $2014, , .$		O
36	Synthesis of mixed-sequence oligonucleotides on mesoporous silicon: chemical strategies and material stability. Nanoscale Research Letters, 2014, 9, 317.	5.7	9

#	Article	IF	CITATIONS
37	DNA-based nanostructures: The effect of the base sequence on octamer formation from d(XGGYGGT) tetramolecular G-quadruplexes. Biochimie, 2014, 99, 119-128.	2.6	20
38	Synthesis of 2,6â€Dialkyl(aryl)purine Nucleosides by Exploiting the Reactivity of Nebularine ⟨i>N⟨ i>1â€Oxide towards Grignard Reagents. European Journal of Organic Chemistry, 2013, 2013, 6948-6954.	2.4	7
39	PNA as a potential modulator of COL7A1 gene expression in dominant dystrophic epidermolysis bullosa: a physico-chemical study. Molecular BioSystems, 2013, 9, 3166.	2.9	9
40	Aminosilane functionalizations of mesoporous oxidized silicon for oligonucleotide synthesis and detection. Journal of the Royal Society Interface, 2013, 10, 20130160.	3.4	60
41	Synthesis of New Acadesine (AlCA-riboside) Analogues Having Acyclic d-Ribityl or 4-Hydroxybutyl Chains in Place of the Ribose. Molecules, 2013, 18, 9420-9431.	3.8	12
42	Investigating the Role of T ₇ and T ₁₂ Residues on the Biological Properties of Thrombin-Binding Aptamer: Enhancement of Anticoagulant Activity by a Single Nucleobase Modification. Journal of Medicinal Chemistry, 2012, 55, 10716-10728.	6.4	42
43	Synthesis and biological evaluation of unprecedented ring-expanded nucleosides (RENs) containing the imidazo [4,5-d][1,2,6] oxadiazepine ring system. Chemical Communications, 2012, 48, 9310.	4.1	33
44	New anti-HIV aptamers based on tetra-end-linked DNA G-quadruplexes: effect of the base sequence on anti-HIV activity. Chemical Communications, 2012, 48, 9516.	4.1	31
45	G-Quadruplex-Forming Oligonucleotide Conjugated to Magnetic Nanoparticles: Synthesis, Characterization, and Enzymatic Stability Assays. Bioconjugate Chemistry, 2012, 23, 382-391.	3.6	27
46	A Facile Synthesis of 5'-Fluoro-5'-deoxyacadesine (5'-F-AICAR): A Novel Non-phosphorylable AICAR Analogue. Molecules, 2012, 17, 13036-13044.	3.8	30
47	Targeting G-Quadruplex Structure in the Human c-Kit Promoter with Short PNA Sequences. Bioconjugate Chemistry, 2011, 22, 654-663.	3.6	45
48	Label-Free Probing of G-Quadruplex Formation by Surface-Enhanced Raman Scattering. Analytical Chemistry, 2011, 83, 6849-6855.	6.5	56
49	Solid-Phase Synthesis of a New Diphosphate 5-Aminoimidazole-4-carboxamide Riboside (AICAR) Derivative and Studies toward Cyclic AICAR Diphosphate Ribose. Molecules, 2011, 16, 8110-8118.	3.8	20
50	Solid-phase synthesis and pharmacological evaluation of novel nucleoside-tethered dinuclear platinum(II) complexes. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 5835-5838.	2.2	15
51	Probing the reactivity of nebularine N1-oxide. A novel approach to C-6 C-substituted purine nucleosides. Tetrahedron, 2011, 67, 6138-6144.	1.9	18
52	d(CGGTGGT) forms an octameric parallel G-quadruplex via stacking of unusual G(:C):G(:C):G(:C):G(:C) octads. Nucleic Acids Research, 2011, 39, 7848-7857.	14.5	42
53	Facile Solidâ€Phase Synthesis of AICAR 5′â€Monophosphate (ZMP) and Its 4â€∢i>Nà€Alkyl Derivatives. European Journal of Organic Chemistry, 2010, 2010, 1517-1524.	2.4	31
54	A solid-phase approach to the synthesis of N-1-alkyl analogues of cyclic inosine-diphosphate-ribose (cIDPR). Tetrahedron, 2010, 66, 1931-1936.	1.9	30

#	Article	IF	Citations
55	Direct Synthesis of Oligonucleotides on Nanostructured Silica Multilayers. Journal of Physical Chemistry C, 2010, 114, 2617-2621.	3.1	14
56	Tetra-end-linked oligonucleotides forming DNA G-quadruplexes: a new class of aptamers showing anti-HIV activity. Chemical Communications, 2010, 46, 8971.	4.1	39
57	Synthesis of quadruplexâ€forming tetraâ€endâ€linked oligonucleotides: Effects of the linker size on quadruplex topology and stability. Biopolymers, 2009, 91, 466-477.	2.4	31
58	Evidences for complex formation between l-dabPNA and aegPNA. Bioorganic and Medicinal Chemistry Letters, 2008, 18, 4757-4760.	2.2	7
59	Synthesis of 4-N-alkyl and ribose-modified AICAR analogues on solid support. Tetrahedron, 2008, 64, 6475-6481.	1.9	34
60	Oligonucleotides direct synthesis on porous silicon chip. Nucleic Acids Symposium Series, 2008, 52, 721-722.	0.3	1
61	Synthesis and Characterization of Tetra-End Linked Oligonucleotides Capable of Forming Monomolecular G-Quadruplexes. Nucleosides, Nucleotides and Nucleic Acids, 2007, 26, 1231-1236.	1.1	O
62	Optical Tweezers as a Probe for Oligodeoxyribonucleotide Structuration. Nucleosides, Nucleotides and Nucleic Acids, 2007, 26, 1295-1299.	1.1	0
63	Synthesis of A New Ribose Modified Analogue of Cyclic Inosine Diphosphate Ribose. Nucleosides, Nucleotides and Nucleic Acids, 2007, 26, 1321-1324.	1.1	2
64	Physico-chemical analysis of G-quadruplex containing bunch-oligonucleotides. International Journal of Biological Macromolecules, 2007, 40, 242-247.	7.5	4
65	Solid Phase Synthesis of Nucleobase and Ribose Modified Inosine Nucleoside Analogues. Nucleosides, Nucleotides and Nucleic Acids, 2007, 26, 1649-1652.	1.1	6
66	Synthesis of N-1 and ribose modified inosine analogues on solid support. Tetrahedron Letters, 2007, 48, 397-400.	1.4	34
67	Synthesis and Characterization of Monomolecular DNA G-Quadruplexes Formed by Tetra-End-Linked Oligonucleotides. Bioconjugate Chemistry, 2006, 17, 889-898.	3.6	28
68	Synthesis and characterization of DNA quadruplexes containing T-tetrads formed by bunch-oligonucleotides. Biopolymers, 2006, 81, 194-201.	2.4	22
69	A BUNCH-OLIGONUCLEOTIDE FORMING STABLE MONOMOLECULAR QUADRUPLEX CONTAINING A T-TETRAD. Nucleosides, Nucleotides and Nucleic Acids, 2005, 24, 443-446.	1.1	2
70	EFFECTS OF ACROLEIN ON THE QUADRUPLEX FORMING d(TTAGGG)4 TELOMERIC REPEAT SEQUENCE. Nucleosides, Nucleotides and Nucleic Acids, 2005, 24, 447-450.	1.1	0
71	Reactions of Pd(PPh3)4with 3â€~,5â€~-Di-O-acetylthymidine: Oxidative Addition of Pd(PPh3)4on Thymidine N3 and C4 Atoms. Organometallics, 2005, 24, 3401-3406.	2.3	12
72	Thermodynamics and Kinetics of PNAâ^'DNA Quadruplex-Forming Chimeras. Journal of the American Chemical Society, 2005, 127, 16215-16223.	13.7	44

#	Article	IF	Citations
73	UNUSUAL MONOMOLECULAR DNA QUADRUPLEX STRUCTURES USING BUNCH-OLIGONUCLEOTIDES. Nucleosides, Nucleotides and Nucleic Acids, 2005, 24, 739-741.	1.1	1
74	SYNTHESIS OF A NEW N-9 RIBITYL ANALOGUE OF CYCLIC INOSINE DIPHOSPHATE RIBOSE (cIDPR) AS A MIMIC OF CYCLIC ADP RIBOSE (cADPR). Nucleosides, Nucleotides and Nucleic Acids, 2005, 24, 735-738.	1.1	4
75	In-water reactivity of nucleosides and nucleotides: one-step preparation and biological evaluation of novel ferrocenyl-derivatives. Tetrahedron, 2004, 60, 6555-6563.	1.9	23
76	Targeting duplex DNA with DNA-PNA chimeras? Physico-chemical characterization of a triplex DNA-PNA/DNA/DNA. Biopolymers, 2004, 73, 434-442.	2.4	6
77	Synthesis of 3′â^'3′-Linked Pyrimidine Oligonucleotides Containing an Acridine Moiety for Alternate Strand Triple Helix Formation. European Journal of Organic Chemistry, 2004, 2004, 2331-2336.	2.4	6
78	Synthesis and characterization of a bunchy oligonucleotide forming a monomolecular parallel quadruplex structure in solution. Tetrahedron Letters, 2004, 45, 4869-4872.	1.4	29
79	Effect of Î ³ -hydroxypropano deoxyguanosine, the major acrolein-derived adduct, on monomolecular quadruplex structure of telomeric repeat d(TTAGGG)4. Bioorganic and Medicinal Chemistry Letters, 2004, 14, 5417-5421.	2.2	3
80	Effects of an 8-bromodeoxyguanosine incorporation on the parallel quadruplex structure $[d(TGGGT)]4$. Organic and Biomolecular Chemistry, 2004, 2, 313.	2.8	73
81	Excess electron transfer in G-quadruplex. Chemical Communications, 2004, , 1756-1757.	4.1	17
82	Synthesis and Structural Characterization of PNA-DNA Quadruplex-Forming Chimeras. European Journal of Organic Chemistry, 2003, 2003, 3364-3371.	2.4	8
83	Physico-chemical studies of a DNA triplex containing a new ferrocenemethyl-thymidine residue in the third strand. Biophysical Chemistry, 2003, 104, 259-270.	2.8	8
84	PNA-DNA Chimeras Forming Quadruplex Structures. Nucleosides, Nucleotides and Nucleic Acids, 2003, 22, 1681-1684.	1.1	7
85	Oligonucleotides Containing an Acridine Group Covalently Bonded to the Nucleotide Flanking the 3′-3′ Phosphodiester Junction for Alternate Strand Triple Helix Formation. Nucleosides, Nucleotides and Nucleic Acids, 2003, 22, 1069-1071.	1.1	3
86	New Solid Supports Linking Nucleoside Scaffolds. Nucleosides, Nucleotides and Nucleic Acids, 2003, 22, 695-697.	1.1	4
87	Synthesis and DNA Binding Properties of DNA-PNA Chimeras. Nucleosides, Nucleotides and Nucleic Acids, 2003, 22, 1089-1091.	1.1	4
88	Cyclic Uridine Diphosphate Glucose: A New Pyrimidine Analog of Cyclic ADP Ribose. Nucleosides, Nucleotides and Nucleic Acids, 2003, 22, 663-666.	1.1	3
89	Synthesis of a New N1-Pentyl Analogue of Cyclic Inosine Diphosphate Ribose (cIDPR) as a Stable Potential Mimic of Cyclic ADP Ribose (cADPR). European Journal of Organic Chemistry, 2002, 2002, 4234-4238.	2.4	15
90	Interaction of the ADP-ribosylating enzyme from the hyperthermophilic archaeonS. solfataricuswith DNA and ss-oligo deoxy ribonucleotides. Journal of Cellular Biochemistry, 2002, 85, 146-157.	2.6	6

#	Article	IF	CITATIONS
91	Solid phase synthesis of oligonucleotides tethered to oligo-glucose phosphate tails. Tetrahedron, 2002, 58, 6697-6704.	1.9	32
92	Synthesis of a novel N-1 carbocyclic, N-9 butyl analogue of cyclic ADP ribose (cADPR). Tetrahedron, 2002, 58, 363-368.	1.9	29
93	Fully automated synthesis of peptide-oligonucleotide conjugates. , 2002, , 784-785.		0
94	Physico-chemical studies on DNA triplexes containing an alternate third strand with a non-nucleotide linker. International Journal of Biological Macromolecules, 2001, 28, 387-394.	7.5	5
95	New nucleoside based solid supports. Synthesis of 5′,3′-derivatized thymidine analoguesElectronic supplementary information (ESI) available: experimental details. See http://www.rsc.org/suppdata/cc/b1/b107200p/. Chemical Communications, 2001, , 2598-2599.	4.1	16
96	Solid phase synthesis of DNA-3′-PNA chimeras by using Bhoc/Fmoc PNA monomers. Tetrahedron, 2001, 57, 9481-9486.	1.9	16
97	Oligonucleotides containing a lysine residue as 3′–3′ junction for alternate strand triple helix formation. Bioorganic and Medicinal Chemistry, 2001, 9, 2895-2900.	3.0	1
98	Solid-Phase Synthesis of Glyco-Oligonucleotide Conjugates. Synlett, 2001, 2001, 0745-0748.	1.8	13
99	A Facile Solid-Phase Strategy for the Synthesis of Oligonucleotide-Tetraphenylporphyrin Conjugates. European Journal of Organic Chemistry, 2000, 2000, 1013-1018.	2.4	9
100	Studies toward the Synthesis of Pinolidoxin, a Phytotoxic Nonenolide from the FungusAscochytapinodes. Determination of the Configuration at the C-7, C-8, and C-9 Chiral Centers and Stereoselective Synthesis of the C6â^'C18Fragment. Journal of Organic Chemistry, 2000, 65, 3432-3442.	3.2	27
101	Hydrolysis of oleuropein by recombinant \hat{l}^2 -glycosidase from hyperthermophilic archaeon Sulfolobus solfataricus immobilised on chitosan matrix. Journal of Biotechnology, 2000, 77, 275-286.	3.8	67
102	Solid phase glycosidation of oligonucleotides. Tetrahedron Letters, 1999, 40, 2607-2610.	1.4	31
103	Synthesis and characterization of new $3\hat{a}\in^2$ - $3\hat{a}\in^2$ linked oligodeoxyribonucleotides for alternate strand triple helix formation. Tetrahedron, 1999, 55, 9899-9914.	1.9	16
104	A new ferrocenemethyl-thymidine nucleoside: Synthesis, incorporation into oligonucleotides and optical spectroscopic studies on the resulting single strand, duplex and triplex structures. Tetrahedron, 1999, 55, 14435-14450.	1.9	37
105	A new solid-phase synthesis of oligonucleotides 3′-conjugated with peptides. Bioorganic and Medicinal Chemistry, 1999, 7, 395-400.	3.0	32
106	Thermodynamics of a 24-Mer Triple Helix Formation and Stability. Magyar Apróvad Közlemények, 1999, 56, 1177-1184.	1.4	8
107	Synthetic studies on the glycosylation of the base residues of inosine and uridine. Journal of the Chemical Society Perkin Transactions 1, 1999, , 3489-3493.	0.9	13
108	A Facile Solid-Phase Synthesis of Oligonucleotides Containing a 3′â^'3′ Phosphodiester Bond for Alternate Strand Triple-Helix Formation. European Journal of Organic Chemistry, 1998, 1998, 2119-2125.	2.4	7

#	Article	IF	CITATIONS
109	Use of controlled pore glass in solid phase oligosaccharide synthesis. Application to the semiautomated synthesis of a glyconucleotide conjugate. Tetrahedron Letters, 1998, 39, 1953-1956.	1.4	32
110	1-Substituted 2′-deoxyinosine analogues. Journal of the Chemical Society Perkin Transactions 1, 1997, , 2079-2082.	0.9	31
111	Identification of the Active Site Nucleophile in the Thermostable β-Glycosidase from the ArchaeonSulfolobus solfataricusExpressed inEscherichia coliâ€. Biochemistry, 1997, 36, 3068-3075.	2.5	28
112	Synthesis and Triple Helix Formation by Alternate Strand Recognition of Oligonucleotides Containing 3â€~-3â€~ Phosphodiester Bonds. Journal of Organic Chemistry, 1997, 62, 9024-9030.	3.2	20
113	Solid phase synthesis of oligosaccharides. Tetrahedron Letters, 1996, 37, 5007-5010.	1.4	51
114	Automated solid phase synthesis of cyclic oligonucleotides: a further improvement. Bioorganic and Medicinal Chemistry, 1995, 3, 1325-1329.	3.0	22
115	Synthesis of thymidine dimers containing a new internucleosidic amide linkage and their incorporation into oligodeoxyribonucleotides. Bioorganic and Medicinal Chemistry Letters, 1995, 5, 1647-1652.	2.2	13
116	Synthesis of [1-15N]-Labeled 2'-Deoxyinosine and 2'-Deoxyadenosine. Journal of Organic Chemistry, 1995, 60, 2251-2253.	3.2	27
117	Interaction of the High-Affinity Inhibitor Tetrahydro-Dump with the Allosteric Enzyme Deoxycytidylate Aminohydrolase. Archives of Biochemistry and Biophysics, 1994, 310, 49-53.	3.0	O
118	Reaction of 3′,5′-di-O-acetyl-2′-deoxyinosine with the chlorinating agent PPh3–CCl4: synthesis of the 6-chloroderivative and of a new base linked dimer, useful intermediate to15N-1-labelled 2′-deoxyinosine. Journal of the Chemical Society Perkin Transactions 1, 1994, , 923-925.	0.9	14
119	Facile preparation of cyclic oligoribonucleotides. Journal of the Chemical Society Perkin Transactions 1, 1993, , 747.	0.9	8
120	PEG-Supported Synthesis of Cyclic Oligodeoxyribonucleotides. Nucleosides & Nucleotides, 1993, 12, 21-30.	0.5	27
121	Automated Synthesis of Cyclic Oligodeoxyribonucleotides via Phosphoramidite Method. Nucleosides & Nucleotides, 1993, 12, 351-358.	0.5	12
122	Synthesis of 2′,3′-dideoxy-2′,3′-didehydronucleoside analogues as potential anti HIV agents. Bioorgan and Medicinal Chemistry Letters, 1992, 2, 315-318.	ic _{2.2}	8
123	Solid-Phase Synthesis of Oligodeoxyribonucleotide Analogues Containing 5, 6-Dihydroimidazo [1, 2-c] Pyrimidin-5-One as a Base Moiety. Nucleosides & Nucleotides, 1991, 10, 867-882.	0.5	1
124	A polymer-nucleotide linkage useful for the solid phase synthesis of cyclic oligodeoxyribonucleotides. Tetrahedron, 1989, 45, 4523-4536.	1.9	25
125	Synthesis of Bridged Pyrimidine Nucleosides and Triazo [4, 3-c] Pyrimidine Nucleoside Analogues. Nucleosides & Nucleotides, 1989, 8, 515-528.	0.5	2
126	Solid phase synthesis of 5'-phosphate labelled polynucleotides. Tetrahedron, 1988, 44, 215-220.	1.9	3

#	Article	IF	CITATIONS
127	New derivatives of pyrimidine nucleosides: Synthesis, physicoâ€chemical properties and biological activity. Journal of Heterocyclic Chemistry, 1988, 25, 1039-1042.	2.6	4
128	Synthesis of novel pyrimidine nucleoside analogues. Journal of Heterocyclic Chemistry, 1986, 23, 1401-1403.	2.6	18
129	Use of fast protein liquid chromatography for the purification of synthetic oligonucleotides. Journal of Chromatography A, 1985, 329, 406-414.	3.7	22