

# Herman O Sintim

## List of Publications by Year in descending order

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152  
papers

6,044  
citations

66343

42  
h-index

85541

71  
g-index

164  
all docs

164  
docs citations

164  
times ranked

7603  
citing authors

#	ARTICLE	IF	CITATIONS
1	Comparative Studies to Uncover Mechanisms of Action of <i>N</i> -(1,3,4-Oxadiazol-2-yl)benzamide Containing Antibacterial Agents. <i>ACS Infectious Diseases</i> , 2022, 8, 865-877.	3.8	2
2	Global proteomics of fibroblast cells treated with bacterial cyclic dinucleotides, c-di-GMP and c-di-AMP. <i>Journal of Oral Microbiology</i> , 2022, 14, 2003617.	2.7	5
3	Mechanistic Studies and <i>In Vivo</i> Efficacy of an Oxadiazole-Containing Antibiotic. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 6612-6630.	6.4	6
4	Membrane acting Povarov-Doebner derived compounds potently disperse preformed multidrug resistant Gram-positive bacterial biofilms. <i>European Journal of Medicinal Chemistry</i> , 2022, , 114550.	5.5	2
5	Identification of a <i>Mycobacterium tuberculosis</i> Cyclic Dinucleotide Phosphodiesterase Inhibitor. <i>ACS Infectious Diseases</i> , 2021, 7, 309-317.	3.8	8
6	Zwitterionic liquid crystalline polythiophene as an antibiofouling biomaterial. <i>Journal of Materials Chemistry B</i> , 2021, 9, 349-356.	5.8	5
7	SF <sup>5</sup> - and SCF <sup>3</sup> -substituted tetrahydroquinoline compounds as potent bactericidal agents against multidrug-resistant persister Gram-positive bacteria. <i>RSC Medicinal Chemistry</i> , 2021, 12, 1879-1893.	3.9	9
8	A STING-based fluorescent polarization assay for monitoring activities of cyclic dinucleotide metabolizing enzymes. <i>RSC Chemical Biology</i> , 2021, 2, 206-214.	4.1	3
9	<i>N</i> -(1,3,4-Oxadiazol-2-yl)Benzamides as Antibacterial Agents against <i>Neisseria gonorrhoeae</i> . <i>International Journal of Molecular Sciences</i> , 2021, 22, 2427.	4.1	12
10	Bacterial Cyclic Dinucleotides and the cGAS/cGAMP/STING Pathway: A Role in Periodontitis?. <i>Pathogens</i> , 2021, 10, 675.	2.8	7
11	Starving Bacteria of Iron: A Potential Strategy to Disperse Bacterial Biofilms. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 7272-7274.	6.4	4
12	3 <i>H</i> -Pyrazolo[4,3- <i>f</i> ]quinoline-Based Kinase Inhibitors Inhibit the Proliferation of Acute Myeloid Leukemia Cells <i>In Vivo</i> . <i>Journal of Medicinal Chemistry</i> , 2021, 64, 10981-10996.	6.4	10
13	c-di-GMP Induces COX-2 Expression in Macrophages in a STING-Independent Manner. <i>ACS Chemical Biology</i> , 2021, 16, 1663-1670.	3.4	4
14	Potent trifluoromethoxy, trifluoromethylsulfonyl, trifluoromethylthio and pentafluorosulfonyl containing (1,3,4-oxadiazol-2-yl)benzamides against drug-resistant Gram-positive bacteria. <i>RSC Medicinal Chemistry</i> , 2020, 11, 102-110.	3.9	19
15	Use of High-Throughput Tools for Telescoped Continuous Flow Synthesis of an Alkynyl naphthyridine Anticancer Agent, HSN608. <i>Organic Process Research and Development</i> , 2020, 24, 2240-2251.	2.7	21
16	Activation of Gingival Fibroblasts by Bacterial Cyclic Dinucleotides and Lipopolysaccharide. <i>Pathogens</i> , 2020, 9, 792.	2.8	7
17	Ultrapotent Inhibitor of <i>Clostridioides difficile</i> Growth, Which Suppresses Recurrence <i>In Vivo</i> . <i>Journal of Medicinal Chemistry</i> , 2020, 63, 11934-11944.	6.4	18
18	HSD1787, a Tetrahydro-3 <i>H</i> -Pyrazolo[4,3- <i>f</i> ]Quinoline Compound Synthesized via Povarov Reaction, Potently Inhibits Proliferation of Cancer Cell Lines at Nanomolar Concentrations. <i>ACS Omega</i> , 2020, 5, 23799-23807.	3.5	12

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19	Homologous Quorum Sensing Regulatory Circuit: A Dual-Input Genetic Controller for Modulating Quorum Sensing-Mediated Protein Expression in <i>E. coli</i> . <i>ACS Synthetic Biology</i> , 2020, 9, 2692-2702.	3.8	9
20	Lipoteichoic Acid Biosynthesis Inhibitors as Potent Inhibitors of <i>S. aureus</i> and <i>E. faecalis</i> Growth and Biofilm Formation. <i>Molecules</i> , 2020, 25, 2277.	3.8	28
21	Multiple ways to kill bacteria via inhibiting novel cell wall or membrane targets. <i>Future Medicinal Chemistry</i> , 2020, 12, 1253-1279.	2.3	26
22	Nicotinamideâ€Ponatinib Analogues as Potent Anti-CML and Anti-AML Compounds. <i>ACS Omega</i> , 2020, 5, 2690-2698.	3.5	8
23	Global Proteomic Analyses of STINGâ€Positive and â€Negative Macrophages Reveal STING and Nonâ€STING Differentially Regulated Cellular and Molecular Pathways. <i>Proteomics - Clinical Applications</i> , 2020, 14, e1900109.	1.6	7
24	Quorum Sensing Autoinducer-3 Finally Yields to Structural Elucidation. <i>ACS Central Science</i> , 2020, 6, 93-96.	11.3	11
25	Targeting Cyclic Dinucleotide Signaling with Small Molecules. , 2020, , 577-591.		2
26	One-Step Large-Scale Nanotexturing of Nonplanar PTFE Surfaces to Induce Bactericidal and Anti-inflammatory Properties. <i>ACS Applied Materials &amp; Interfaces</i> , 2020, 12, 26893-26904.	8.0	14
27	Zwitterionic Porous Conjugated Polymers as a Versatile Platform for Antibiofouling Implantable Bioelectronics. <i>ACS Applied Polymer Materials</i> , 2020, 2, 528-536.	4.4	17
28	Interrupting cyclic dinucleotide-cGASâ€STING axis with small molecules. <i>MedChemComm</i> , 2019, 10, 1999-2023.	3.4	19
29	Potently inhibiting cancer cell migration with novel 3H-pyrazolo[4,3-f]quinoline boronic acid ROCK inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2019, 180, 449-456.	5.5	19
30	Squarate desymmetrisationâ€ozonolysis as an approach to Î²-substituted-Î±-ketosuccinates and squalestatin synthesis. <i>Tetrahedron</i> , 2019, 75, 130747.	1.9	2
31	Inhibitors of Intracellular Gram-Positive Bacterial Growth Synthesized via Povarovâ€Doebner Reactions. <i>ACS Infectious Diseases</i> , 2019, 5, 1820-1830.	3.8	11
32	Amino alkynylisoquinoline and alkynylnaphthyridine compounds potently inhibit acute myeloid leukemia proliferation in mice. <i>EBioMedicine</i> , 2019, 40, 231-239.	6.1	11
33	Identification of nicotinamide aminonaphthyridine compounds as potent RET kinase inhibitors and antitumor activities against RET rearranged lung adenocarcinoma. <i>Bioorganic Chemistry</i> , 2019, 90, 103052.	4.1	13
34	Alkylation of lithiated dimethyl tartrate acetonide with unactivated alkyl halides and application to an asymmetric synthesis of the 2,8-dioxabicyclo[3.2.1]octane core of squalestatins/zaragozic acids. <i>Beilstein Journal of Organic Chemistry</i> , 2019, 15, 1194-1202.	2.2	2
35	Ligand-Based Stability Changes in Duplex DNA Measured with a Microscale Electrochemical Platform. <i>Biosensors</i> , 2019, 9, 54.	4.7	6
36	Proteomic analysis of bacterial response to a 4-hydroxybenzylidene indolinone compound, which re-sensitizes bacteria to traditional antibiotics. <i>Journal of Proteomics</i> , 2019, 202, 103368.	2.4	27

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37	Antibacterial Small Molecules That Potently Inhibit <i>Staphylococcus aureus</i> Lipoteichoic Acid Biosynthesis. <i>ChemMedChem</i> , 2019, 14, 1000-1004.	3.2	25
38	ENPP1, an Old Enzyme with New Functions, and Small Molecule Inhibitors—A STING in the Tale of ENPP1. <i>Molecules</i> , 2019, 24, 4192.	3.8	66
39	Regulation of gingival epithelial cytokine response by bacterial cyclic dinucleotides. <i>Journal of Oral Microbiology</i> , 2019, 11, 1538927.	2.7	18
40	Discriminating cyclic from linear nucleotides—CRISPR/Cas-related cyclic hexaadenosine monophosphate as a case study. <i>Analytical Biochemistry</i> , 2019, 567, 21-26.	2.4	3
41	3H-pyrazolo[4,3-f]quinoline haspin kinase inhibitors and anticancer properties. <i>Bioorganic Chemistry</i> , 2018, 78, 418-426.	4.1	35
42	On the ozonolysis of unsaturated tosylhydrazones as a direct approach to diazocarbonyl compounds. <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 2876-2884.	2.8	10
43	Alkynylnicotinamide-Based Compounds as ABL1 Inhibitors with Potent Activities against Drug-Resistant CML Harboring ABL1(T315I) Mutant Kinase. <i>ChemMedChem</i> , 2018, 13, 1172-1180.	3.2	12
44	Dual FLT3/TOPK inhibitor with activity against FLT3-ITD secondary mutations potently inhibits acute myeloid leukemia cell lines. <i>Future Medicinal Chemistry</i> , 2018, 10, 823-835.	2.3	17
45	Quorum-sensing molecule dihydroxy-2,3-pentanedione and its analogs as regulators of epithelial integrity. <i>Journal of Periodontal Research</i> , 2018, 53, 414-421.	2.7	7
46	Tetrahydro-3H-pyrazolo[4,3-a]phenanthridine-based CDK inhibitor. <i>Chemical Communications</i> , 2018, 54, 4521-4524.	4.1	11
47	Suramin potently inhibits cGAMP synthase, cGAS, in THP1 cells to modulate IFN- $\beta$ levels. <i>Future Medicinal Chemistry</i> , 2018, 10, 1301-1317.	2.3	78
48	Proteomic analysis of RAW macrophages treated with cGAMP or c-di-GMP reveals differentially activated cellular pathways. <i>RSC Advances</i> , 2018, 8, 36840-36851.	3.6	15
49	Berenil Binds Tightly to Parallel and Mixed Parallel/Antiparallel G-Quadruplex Motifs with Varied Thermodynamic Signatures. <i>ACS Omega</i> , 2018, 3, 11582-11591.	3.5	7
50	Miniaturized whole-cell bacterial bioreporter assay for identification of quorum sensing interfering compounds. <i>Journal of Microbiological Methods</i> , 2018, 154, 40-45.	1.6	6
51	Quorum sensing molecules regulate epithelial cytokine response and biofilm-related virulence of three <i>Prevotella</i> species. <i>Anaerobe</i> , 2018, 54, 128-135.	2.1	16
52	Evidence of link between quorum sensing and sugar metabolism in <i>Escherichia coli</i> revealed via cocrystal structures of LsrK and HPr. <i>Science Advances</i> , 2018, 4, eaar7063.	10.3	68
53	Alkene protection against acid using a bromide substituent: application in a total synthesis of ( $\alpha^{\omega}$ )-6,7-dideoxysqualenol H5. <i>Chemical Communications</i> , 2018, 54, 5354-5356.	4.1	6
54	N-(1,3,4-oxadiazol-2-yl)benzamide analogs, bacteriostatic agents against methicillin- and vancomycin-resistant bacteria. <i>European Journal of Medicinal Chemistry</i> , 2018, 155, 797-805.	5.5	34

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55	KRX-101, a Novel FLT3 Inhibitor, Potently Active Against Resistant FLT3-ITD/FLT3-TKD Mutant AML in Vitro and In Vivo. <i>Blood</i> , 2018, 132, 4049-4049.	1.4	0
56	Fluorescent analogs of cyclic and linear dinucleotides as phosphodiesterase and oligoribonuclease activity probes. <i>RSC Advances</i> , 2017, 7, 5421-5426.	3.6	11
57	Identification of New FLT3 Inhibitors That Potently Inhibit AML Cell Lines via an Azo Click-It/Staple-It Approach. <i>ACS Medicinal Chemistry Letters</i> , 2017, 8, 492-497.	2.8	16
58	Synthesis of (âˆ)—6,7-Dideoxysqualestatin H5 by Carbonyl Ylide Cycloadditionâ€“Rearrangement and Cross-electrophile Coupling. <i>Organic Letters</i> , 2017, 19, 3540-3543.	4.6	18
59	Aminoisoquinoline benzamides, FLT3 and Src-family kinase inhibitors, potently inhibit proliferation of acute myeloid leukemia cell lines. <i>Future Medicinal Chemistry</i> , 2017, 9, 1213-1225.	2.3	15
60	Hydroxybenzylidene-indolinones, c-di-AMP synthase inhibitors, have antibacterial and anti-biofilm activities and also re-sensitize resistant bacteria to methicillin and vancomycin. <i>RSC Advances</i> , 2017, 7, 8288-8294.	3.6	19
61	Inhibition of innate immune cytosolic surveillance by an <i>M. tuberculosis</i> phosphodiesterase. <i>Nature Chemical Biology</i> , 2017, 13, 210-217.	8.0	96
62	Dipeptidyl peptidase IV and quorum sensing signaling in biofilm-related virulence of <i>Prevotella aurantiaea</i> . <i>Anaerobe</i> , 2017, 48, 152-159.	2.1	9
63	Targeting c-di-GMP Signaling, Biofilm Formation, and Bacterial Motility with Small Molecules. <i>Methods in Molecular Biology</i> , 2017, 1657, 419-430.	0.9	28
64	Fluorescent 2-Aminopurine c-di-GMP and GpG Analogs as PDE Probes. <i>Methods in Molecular Biology</i> , 2017, 1657, 245-261.	0.9	1
65	Cyclic Dinucleotides in Oral Bacteria and in Oral Biofilms. <i>Frontiers in Cellular and Infection Microbiology</i> , 2017, 7, 273.	3.9	17
66	Supramolecular polymer formation by cyclic dinucleotides and intercalators affects dinucleotide enzymatic processing. <i>Future Science OA</i> , 2016, 2, FSO93.	1.9	8
67	Biofilms as â€œConnectorsâ€“for Oral and Systems Medicine: A New Opportunity for Biomarkers, Molecular Targets, and Bacterial Eradication. <i>OMICS A Journal of Integrative Biology</i> , 2016, 20, 3-11.	2.0	28
68	Insightful directed evolution of <i>Escherichia coli</i> quorum sensing promoter region of the <i>srACDBFG</i> operon: a tool for synthetic biology systems and protein expression. <i>Nucleic Acids Research</i> , 2016, 44, gkw981.	14.5	9
69	Autoinducer-2 analogs and electric fields - an antibiotic-free bacterial biofilm combination treatment. <i>Biomedical Microdevices</i> , 2016, 18, 95.	2.8	12
70	Construction and characterization of a multilayered gingival keratinocyte culture model: the TURK-U model. <i>Cytotechnology</i> , 2016, 68, 2345-2354.	1.6	6
71	Inhibition of cyclic diadenylate cyclase, DisA, by polyphenols. <i>Scientific Reports</i> , 2016, 6, 25445.	3.3	24
72	Computational understanding and experimental characterization of twice-as-smart quadruplex ligands as chemical sensors of bacterial nucleotide second messengers. <i>Scientific Reports</i> , 2016, 6, 33888.	3.3	11

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73	Cyclic dinucleotide (c-di-GMP, c-di-AMP, and cGAMP) signalings have come of age to be inhibited by small molecules. <i>Chemical Communications</i> , 2016, 52, 9327-9342.	4.1	78
74	Inhibition of <i>P. aeruginosa</i> c-di-GMP phosphodiesterase RocR and swarming motility by a benzoisothiazolinone derivative. <i>Chemical Science</i> , 2016, 7, 6238-6244.	7.4	39
75	Alkyne-substituted diminazene as G-quadruplex binders with anticancer activities. <i>European Journal of Medicinal Chemistry</i> , 2016, 118, 266-275.	5.5	23
76	Potent inhibition of cyclic diadenylate monophosphate cyclase by the antiparasitic drug, suramin. <i>Chemical Communications</i> , 2016, 52, 3754-3757.	4.1	19
77	Cyclic dinucleotide detection with riboswitchâ€“G-quadruplex hybrid. <i>Molecular BioSystems</i> , 2016, 12, 773-777.	2.9	6
78	Fluorescence, Phosphorescence, and Chemiluminescence. <i>Analytical Chemistry</i> , 2016, 88, 170-202.	6.5	95
79	Structureâ€“activity relationship studies of c-di-AMP synthase inhibitor, bromophenol-thiohydantoin. <i>Tetrahedron</i> , 2016, 72, 3554-3558.	1.9	7
80	Membrane Affinity of Platensimycin and Its Dialkylamine Analogs. <i>International Journal of Molecular Sciences</i> , 2015, 16, 17909-17932.	4.1	6
81	Alkaloid Synthesis via Carbenoid Intermediates. <i>Current Organic Chemistry</i> , 2015, 20, 82-101.	1.6	9
82	Bacterial Secretions of Nonpathogenic <i>Escherichia coli</i> Elicit Inflammatory Pathways: a Closer Investigation of Interkingdom Signaling. <i>MBio</i> , 2015, 6, e00025.	4.1	67
83	3-Aminooxazolidinone AHL analogs as hydrolytically-stable quorum sensing agonists in Gram-negative bacteria. <i>MedChemComm</i> , 2015, 6, 1086-1092.	3.4	9
84	Biofilm formation mechanisms and targets for developing antibiofilm agents. <i>Future Medicinal Chemistry</i> , 2015, 7, 493-512.	2.3	492
85	Agents that inhibit bacterial biofilm formation. <i>Future Medicinal Chemistry</i> , 2015, 7, 647-671.	2.3	226
86	Unraveling Curcumin Degradation. <i>Journal of Biological Chemistry</i> , 2015, 290, 4817-4828.	3.4	129
87	Oligoribonuclease is the primary degradative enzyme for pCpG in <i>Pseudomonas aeruginosa</i> that is required for cyclic-di-GMP turnover. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, E5048-57.	7.1	117
88	Geminal dihalogen isosteric replacement in hydrated AI-2 affords potent quorum sensing modulators. <i>Chemical Communications</i> , 2015, 51, 2617-2620.	4.1	9
89	Rapid nucleic acid melting analyses using a microfabricated electrochemical platform. <i>Analytica Chimica Acta</i> , 2015, 853, 265-270.	5.4	12
90	Synthesis of a Biotinylated Photocleavable Nucleotide Monophosphate for the Preparation of Natively Folded RNAs. <i>Methods in Enzymology</i> , 2014, 549, 115-131.	1.0	0

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91	Isothermal amplified detection of DNA and RNA. <i>Molecular BioSystems</i> , 2014, 10, 970.	2.9	354
92	Evolved Quorum Sensing Regulator, LsrR, for Altered Switching Functions. <i>ACS Synthetic Biology</i> , 2014, 3, 210-219.	3.8	28
93	Molecular Insights into How Ligands Activate or Inactivate LasR. <i>Chemistry and Biology</i> , 2014, 21, 1261-1263.	6.0	3
94	Diminazene or berenil, a classic duplex minor groove binder, binds to G-quadruplexes with low nanomolar dissociation constants and the amidine groups are also critical for G-quadruplex binding. <i>Molecular BioSystems</i> , 2014, 10, 2724-2734.	2.9	35
95	E88, a new cyclic-di-GMP class I riboswitch aptamer from <i>Clostridium tetani</i> , has a similar fold to the prototypical class I riboswitch, Vc2, but differentially binds to c-di-GMP analogs. <i>Molecular BioSystems</i> , 2014, 10, 384-390.	2.9	10
96	Identification of bromophenol thiohydantoin as an inhibitor of DisA, a c-di-AMP synthase, from a 1000 compound library, using the coralyne assay. <i>Chemical Communications</i> , 2014, 50, 11234-11237.	4.1	30
97	Convenient detection of HPV virus in a clinical sample using concurrent rolling circle and junction probe amplifications. <i>Chemical Communications</i> , 2014, 50, 7147-7149.	4.1	5
98	Unexpected Complex Formation between Coralyne and Cyclic Diadenosine Monophosphate Providing a Simple Fluorescent Turn-on Assay to Detect This Bacterial Second Messenger. <i>Analytical Chemistry</i> , 2014, 86, 2412-2420.	6.5	32
99	A cyclic dinucleotide containing 2-aminopurine is a general fluorescent sensor for c-di-GMP and cGAMP. <i>Molecular BioSystems</i> , 2014, 10, 1568-1575.	2.9	18
100	STING Ligand c-di-GMP Improves Cancer Vaccination against Metastatic Breast Cancer. <i>Cancer Immunology Research</i> , 2014, 2, 901-910.	3.4	187
101	Crystal Structures of the LsrR Proteins Complexed with Phospho-AI-2 and Two Signal-Interrupting Analogues Reveal Distinct Mechanisms for Ligand Recognition. <i>Journal of the American Chemical Society</i> , 2013, 135, 15526-15535.	13.7	21
102	Probe design rules and effective enzymes for endonuclease-based detection of nucleic acids. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 6181-6185.	3.0	4
103	Octameric G8 c-di-GMP is an efficient peroxidase and this suggests that an open G-tetrad site can effectively enhance hemin peroxidation reactions. <i>RSC Advances</i> , 2013, 3, 6305.	3.6	12
104	Nucleotide, c-di-GMP, c-di-AMP, cGMP, cAMP, (p)ppGpp signaling in bacteria and implications in pathogenesis. <i>Chemical Society Reviews</i> , 2013, 42, 305-341.	38.1	315
105	Selective binding of 2-F-c-di-GMP to Ct-E88 and Cb-E43, new class I riboswitches from <i>Clostridium tetani</i> and <i>Clostridium botulinum</i> respectively. <i>Molecular BioSystems</i> , 2013, 9, 1535.	2.9	9
106	Potent suppression of c-di-GMP synthesis via I-site allosteric inhibition of diguanylate cyclases with 2-F-c-di-GMP. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 4396-4404.	3.0	45
107	N <sup>o</sup> Tethered Carbenoid Cyclopropanation Facilitates the Synthesis of a Functionalized Cyclopropyl-Fused Pyrrolidine. <i>Journal of Organic Chemistry</i> , 2013, 78, 6131-6142.	3.2	13
108	Temperature-controlled electrochemical microwell platform for studying biomolecular interactions. , 2013, , .		0



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109	Nucleic acid detection using G-quadruplex amplification methodologies. <i>Methods</i> , 2013, 64, 185-198.	3.8	43
110	AI-2 analogs and antibiotics: a synergistic approach to reduce bacterial biofilms. <i>Applied Microbiology and Biotechnology</i> , 2013, 97, 2627-2638.	3.6	87
111	Small Molecule Inhibitors of AI-2 Signaling in Bacteria: State-of-the-Art and Future Perspectives for Anti-Quorum Sensing Agents. <i>International Journal of Molecular Sciences</i> , 2013, 14, 17694-17728.	4.1	60
112	“Rolling out” High-Molecular-Weight Proteins, which Contain Repeating Polypeptide Motif, by Using Rolling Circle Amplification. <i>ChemBioChem</i> , 2013, 14, 1929-1930.	2.6	0
113	Detection of Single-Stranded Nucleic Acids via Colorimetric Means, Using G-Quadruplex Probes. <i>Methods in Molecular Biology</i> , 2013, 1039, 153-159.	0.9	2
114	A Pro-Drug Approach for Selective Modulation of AI-2-Mediated Bacterial Cell-to-Cell Communication. <i>Sensors</i> , 2012, 12, 3762-3772.	3.8	20
115	A rapid assay for affinity and kinetics of molecular interactions with nucleic acids. <i>Nucleic Acids Research</i> , 2012, 40, e48-e48.	14.5	30
116	Nanomolar fluorescent detection of c-di-GMP using a modular aptamer strategy. <i>Chemical Communications</i> , 2012, 48, 9059.	4.1	68
117	Signal-on electrochemical Y or junction probe detection of nucleic acid. <i>Chemical Communications</i> , 2012, 48, 7580.	4.1	24
118	A C-di-GMP “proflavine” hemin supramolecular complex has peroxidase activity” implication for a simple colorimetric detection. <i>Molecular BioSystems</i> , 2012, 8, 726.	2.9	30
119	Investigating the interactions between cations, peroxidation substrates and G-quadruplex topology in DNAzyme peroxidation reactions using statistical testing. <i>Analytica Chimica Acta</i> , 2012, 747, 1-6.	5.4	28
120	Differential binding of 2- <sup>2</sup> -biotinylated analogs of c-di-GMP with c-di-GMP riboswitches and binding proteins. <i>Molecular BioSystems</i> , 2012, 8, 772-778.	2.9	21
121	Inhibitors of fatty acid synthesis in prokaryotes and eukaryotes as anti-infective, anticancer and anti-obesity drugs. <i>Future Medicinal Chemistry</i> , 2012, 4, 1113-1151.	2.3	18
122	Endo-S-c-di-GMP Analogues-Polymorphism and Binding Studies with Class I Riboswitch. <i>Molecules</i> , 2012, 17, 13376-13389.	3.8	16
123	Molecular Fluorescence, Phosphorescence, and Chemiluminescence Spectrometry. <i>Analytical Chemistry</i> , 2012, 84, 597-625.	6.5	83
124	Altering the Communication Networks of Multispecies Microbial Systems Using a Diverse Toolbox of AI-2 Analogues. <i>ACS Chemical Biology</i> , 2012, 7, 1023-1030.	3.4	45
125	Diamidinium and iminium aromatics as new aggregators of the bacterial signaling molecule, c-di-GMP. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2012, 22, 881-885.	2.2	19
126	Isothermal detection of RNA with restriction endonucleases. <i>Chemical Communications</i> , 2011, 47, 200-202.	4.1	21



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127	c-di-GMP can form remarkably stable G-quadruplexes at physiological conditions in the presence of some planar intercalators. <i>Chemical Communications</i> , 2011, 47, 4766.	4.1	49
128	Effects on Membrane Lateral Pressure Suggest Permeation Mechanisms for Bacterial Quorum Signaling Molecules. <i>Biochemistry</i> , 2011, 50, 6983-6993.	2.5	41
129	Thiazole Orange-Induced c-di-GMP Quadruplex Formation Facilitates a Simple Fluorescent Detection of This Ubiquitous Biofilm Regulating Molecule. <i>Journal of the American Chemical Society</i> , 2011, 133, 4856-4864.	13.7	74
130	Conservative Change to the Phosphate Moiety of Cyclic Diguanylic Monophosphate Remarkably Affects Its Polymorphism and Ability To Bind DGC, PDE, and PilZ Proteins. <i>Journal of the American Chemical Society</i> , 2011, 133, 9320-9330.	13.7	50
131	DNA-Based Peroxidation Catalyst—What Is the Exact Role of Topology on Catalysis and Is There a Special Binding Site for Catalysis?. <i>Chemistry - A European Journal</i> , 2011, 17, 5691-5698.	3.3	80
132	Dialkylamino-2,4-dihydroxybenzoic Acids as Easily Synthesized Analogues of Platensimycin and Platencin with Comparable Antibacterial Properties. <i>Chemistry - A European Journal</i> , 2011, 17, 3352-3357.	3.3	31
133	RNAs synthesized using photocleavable biotinylated nucleotides have dramatically improved catalytic efficiency. <i>Nucleic Acids Research</i> , 2011, 39, 8559-8571.	14.5	11
134	Differential radial capillary action of ligand assay for high-throughput detection of protein-metabolite interactions. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, 15528-15533.	7.1	177
135	Evidence of magnetic isotope effects during thermochemical sulfate reduction. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, 17635-17638.	7.1	85
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