

Regan M Leblanc

List of Publications by Year in descending order

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Version: 2024-02-01

25
papers

597
citations

687220

13
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26
all docs

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docs citations

26
times ranked

669
citing authors

#	ARTICLE	IF	CITATIONS
1	Structural insights of the conserved "priming loop" of hepatitis B virus pre-genomic RNA. <i>Journal of Biomolecular Structure and Dynamics</i> , 2022, 40, 9761-9773.	2.0	14
2	Extraction and Purification of (E)-Resveratrol from the Bark of Conifer Species. <i>Processes</i> , 2022, 10, 647.	1.3	4
3	Cross-correlated Relaxation Rates Provide Facile Exchange Signature in Selectively Labeled RNA. <i>Journal of Magnetic Resonance</i> , 2022, , 107245.	1.2	1
4	Strategies for Modeling Ligand Docking to Natural and Engineered RNA Structures. <i>Biophysical Journal</i> , 2021, 120, 315a.	0.2	0
5	Isotope-Labeled RNA Building Blocks for NMR Structure and Dynamics Studies. <i>Molecules</i> , 2021, 26, 5581.	1.7	8
6	A drug discovery toolbox for Nuclear Magnetic Resonance (NMR) characterization of ligands and their targets. <i>Drug Discovery Today: Technologies</i> , 2020, 37, 51-60.	4.0	6
7	Deleterious effects of carbon-carbon dipolar coupling on RNA NMR dynamics. <i>Journal of Biomolecular NMR</i> , 2020, 74, 321-331.	1.6	8
8	NMR probing of invisible excited states using selectively labeled RNAs. <i>Journal of Biomolecular NMR</i> , 2018, 71, 165-172.	1.6	14
9	Combining asymmetric ¹³ C-labeling and isotopic filter/edit NOESY: a novel strategy for rapid and logical RNA resonance assignment. <i>Nucleic Acids Research</i> , 2017, 45, e146-e146.	6.5	15
10	A magnesium-induced triplex pre-organizes the SAM-II riboswitch. <i>PLoS Computational Biology</i> , 2017, 13, e1005406.	1.5	24
11	SAM-II Riboswitch Samples at least Two Conformations in Solution in the Absence of Ligand: Implications for Recognition. <i>Angewandte Chemie</i> , 2016, 128, 2774-2777.	1.6	6
12	SAM-II Riboswitch Samples at least Two Conformations in Solution in the Absence of Ligand: Implications for Recognition. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 2724-2727.	7.2	39
13	Chemo-enzymatic synthesis of site-specific isotopically labeled nucleotides for use in NMR resonance assignment, dynamics and structural characterizations. <i>Nucleic Acids Research</i> , 2016, 44, e52-e52.	6.5	44
14	Chemo-enzymatic labeling for rapid assignment of RNA molecules. <i>Methods</i> , 2016, 103, 11-17.	1.9	13
15	Stable Isotope-Labeled RNA Phosphoramidites to Facilitate Dynamics by NMR. <i>Methods in Enzymology</i> , 2015, 565, 461-494.	0.4	18
16	Chemo-Enzymatic Synthesis of Selectively ¹³ C/ ¹⁵ N-Labeled RNA for NMR Structural and Dynamics Studies. <i>Methods in Enzymology</i> , 2014, 549, 133-162.	0.4	30
17	Region-Selective Chemical-Enzymatic Synthesis of Pyrimidine Nucleotides Facilitates RNA Structure and Dynamics Studies. <i>ChemBioChem</i> , 2014, 15, 1573-1577.	1.3	45
18	Design, Synthesis and Biological Testing of Cyclohexenone Derivatives of Combretastatin-A4. <i>Letters in Drug Design and Discovery</i> , 2007, 4, 144-148.	0.4	20

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19	Design, synthesis, and biological testing of pyrazoline derivatives of combretastatin-A4. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007, 17, 5897-5901.	1.0	131
20	Synthesis and Cytotoxic Properties of Chalcones: An Interactive and Investigative Undergraduate Laboratory Project at the Interface of Chemistry and Biology. <i>Journal of Chemical Education</i> , 2006, 83, 934.	1.1	11
21	Synthesis and cytotoxicity of epoxide and pyrazole analogs of the combretastatins. <i>Bioorganic and Medicinal Chemistry</i> , 2005, 13, 6025-6034.	1.4	79
22	SYNTHESIS AND CYTOTOXIC PROPERTIES OF NITRO- AND AMINOCHALCONES. <i>Medicinal Chemistry Research</i> , 2005, 14, 19-25.	1.1	33
23	SYNTHESIS AND BIOLOGICAL EVALUATION OF CIS-COMBRETASTATIN ANALOGS AND THEIR NOVEL 1,2,3-TRIAZOLE DERIVATIVES. <i>Heterocyclic Communications</i> , 2005, 11, .	0.6	30
24	REACTION OF CHALCONES WITH BASIC HYDROGEN PEROXIDE: A STRUCTURE AND REACTIVITY STUDY. <i>Heterocyclic Communications</i> , 2005, 11, .	0.6	4
25	AN EFFICIENT METHOD FOR THE SYNTHESIS OF SUBSTITUTED 4-ACETOXYNAPHTHALENE-2-CARBOXYLATE ESTERS, ETHYL 4-ACETOXYBENZOFURAN-6-CARBOXYLATE, AND ETHYL 4-ACETOXYBENZOTHIOPHENE-6-CARBOXYLATE. <i>Heterocyclic Communications</i> , 2003, 9, .	0.6	0