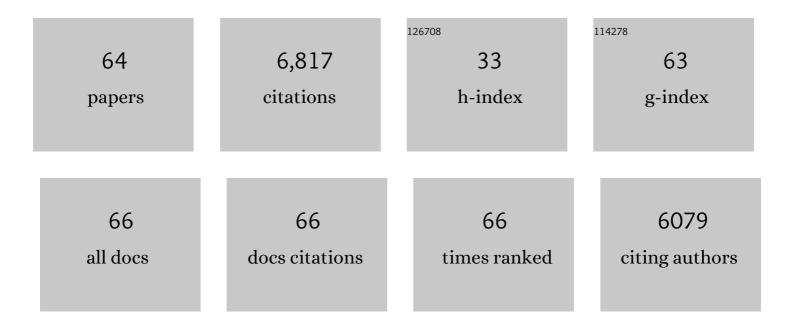
Louis-Charles Campeau

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Kilogram-Scale Synthesis of 2′- <i>C</i> -Methyl- <i>arabino</i> -Uridine from Uridine via Dynamic Selective Dipivaloylation. Organic Process Research and Development, 2022, 26, 698-709.	1.3	7
2	Polycationic Rh–JosiPhos Polymers Supported on Phosphotungstic Acid/Al ₂ O ₃ by Multiple Electrostatic Attractions. ACS Catalysis, 2022, 12, 2034-2044.	5.5	2
3	A kinase-cGAS cascade to synthesize a therapeutic STING activator. Nature, 2022, 603, 439-444.	13.7	58
4	Driving Aspirational Process Mass Intensity Using Simple Structure-Based Prediction. Organic Process Research and Development, 2022, 26, 1405-1410.	1.3	8
5	Synthesis of α-Aryl Secondary Amides via Nickel-Catalyzed Reductive Coupling of Redox-Active Esters. Organic Letters, 2022, 24, 3173-3178.	2.4	8
6	Practical and concise synthesis of nucleoside analogs. Nature Protocols, 2022, 17, 2008-2024.	5.5	5
7	Efficient synthesis of antiviral agent uprifosbuvir enabled by new synthetic methods. Chemical Science, 2021, 12, 9031-9036.	3.7	14
8	Diversity-oriented synthesis of glycomimetics. Communications Chemistry, 2021, 4, .	2.0	17
9	Invention of MK-8262, a Cholesteryl Ester Transfer Protein (CETP) Inhibitor Backup to Anacetrapib with Best-in-Class Properties. Journal of Medicinal Chemistry, 2021, 64, 13215-13258.	2.9	10
10	Preface: Modern Heterocycle Synthesis and Functionalization. Synlett, 2021, 32, 140-141.	1.0	0
11	Bringing amines back into aziridination. Nature Chemistry, 2021, 13, 1027-1028.	6.6	2
12	Development of a Green and Sustainable Manufacturing Process for Gefapixant Citrate (MK-7264) Part 1: Introduction and Process Overview. Organic Process Research and Development, 2020, 24, 2445-2452.	1.3	25
13	Harder, better, faster. Nature Chemistry, 2020, 12, 661-664.	6.6	25
14	A short de novo synthesis of nucleoside analogs. Science, 2020, 369, 725-730.	6.0	61
15	Analysis of Benzenoid Substitution Patterns in Small Molecule Active Pharmaceutical Ingredients. Journal of Medicinal Chemistry, 2020, 63, 13389-13396.	2.9	51
16	Cross-Coupling and Related Reactions: Connecting Past Success to the Development of New Reactions for the Future. Organometallics, 2019, 38, 3-35.	1.1	267
17	The Roles of Organometallic Chemistry in Pharmaceutical Research and Development. Organometallics, 2019, 38, 1-2.	1.1	6
18	A rational pre-catalyst design for bis-phosphine mono-oxide palladium catalyzed reactions. Chemical Science, 2017, 8, 2841-2851.	3.7	24

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19	Design and Synthesis of Novel, Selective GPR40 AgoPAMs. ACS Medicinal Chemistry Letters, 2017, 8, 221-226.	1.3	26
20	Discovery of phenyl acetamides as potent and selective GPR119 agonists. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 1124-1128.	1.0	10
21	A multifunctional catalyst that stereoselectively assembles prodrugs. Science, 2017, 356, 426-430.	6.0	116
22	Selective functionalization of complex heterocycles via an automated strong base screening platform. Reaction Chemistry and Engineering, 2017, 2, 446-450.	1.9	25
23	Discovery and optimization of 2-pyridinone aminal integrase strand transfer inhibitors for the treatment of HIV. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 2038-2046.	1.0	10
24	A Modular Synthesis of 2-Alkyl- and 2-Arylchromans via a Three-Step Sequence. Synthesis, 2017, 49, 657-666.	1.2	21
25	A Robust Kilo-Scale Synthesis of Doravirine. Organic Process Research and Development, 2016, 20, 1476-1481.	1.3	50
26	The Discovery of Quinoxaline-Based Metathesis Catalysts from Synthesis of Grazoprevir (MK-5172). Organic Letters, 2016, 18, 1952-1955.	2.4	34
27	Acridinium-Based Photocatalysts: A Sustainable Option in Photoredox Catalysis. Journal of Organic Chemistry, 2016, 81, 7244-7249.	1.7	259
28	Nickel-Catalyzed Asymmetric Alkene Hydrogenation of α,β-Unsaturated Esters: High-Throughput Experimentation-Enabled Reaction Discovery, Optimization, and Mechanistic Elucidation. Journal of the American Chemical Society, 2016, 138, 3562-3569.	6.6	165
29	Cobalt-Catalyzed Enantioselective Hydrogenation of Minimally Functionalized Alkenes: Isotopic Labeling Provides Insight into the Origin of Stereoselectivity and Alkene Insertion Preferences. Journal of the American Chemical Society, 2016, 138, 3314-3324.	6.6	179
30	Chemistry informer libraries: a chemoinformatics enabled approach to evaluate and advance synthetic methods. Chemical Science, 2016, 7, 2604-2613.	3.7	158
31	Enantioselective Synthesis of α-Methyl-β-cyclopropyldihydrocinnamates. Journal of Organic Chemistry, 2016, 81, 824-830.	1.7	38
32	Discovery and mechanistic study of a photocatalytic indoline dehydrogenation for the synthesis of elbasvir. Chemical Science, 2016, 7, 2066-2073.	3.7	103
33	Development of a novel class of potent and selective FIXa inhibitors. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 4945-4949.	1.0	16
34	P ₂ Et Phosphazene: A Mild, Functional Group Tolerant Base for Soluble, Room Temperature Pd-Catalyzed C–N, C–O, and C–C Cross-Coupling Reactions. Organic Letters, 2015, 17, 3370-3373.	2.4	52
35	Design of Potent and Orally Active GPR119 Agonists for the Treatment of Type II Diabetes. ACS Medicinal Chemistry Letters, 2015, 6, 936-941.	1.3	13
36	Enantioselective Synthesis of Hemiaminals via Pd-Catalyzed C–N Coupling with Chiral Bisphosphine Mono-oxides. Journal of the American Chemical Society, 2015, 137, 13728-13731.	6.6	88

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37	Development of a novel tricyclic class of potent and selective FIXa inhibitors. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 5437-5443.	1.0	16
38	Development of a Direct Photocatalytic C–H Fluorination for the Preparative Synthesis of Odanacatib. Organic Letters, 2015, 17, 5200-5203.	2.4	147
39	Nanomole-scale high-throughput chemistry for the synthesis of complex molecules. Science, 2015, 347, 49-53.	6.0	454
40	Development of a Palladium-Catalyzed α-Arylation of Cyclopropyl Nitriles. Organic Letters, 2014, 16, 6314-6317.	2.4	30
41	Discovery of MK-1439, an orally bioavailable non-nucleoside reverse transcriptase inhibitor potent against a wide range of resistant mutant HIV viruses. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 917-922.	1.0	81
42	<i>In Vitro</i> Characterization of MK-1439, a Novel HIV-1 Nonnucleoside Reverse Transcriptase Inhibitor. Antimicrobial Agents and Chemotherapy, 2014, 58, 1652-1663.	1.4	100
43	Modeling a Crowdsourced Definition of Molecular Complexity. Journal of Chemical Information and Modeling, 2014, 54, 1604-1616.	2.5	48
44	Convergent Kilogram-Scale Synthesis of Dual Orexin Receptor Antagonist. Organic Process Research and Development, 2013, 17, 61-68.	1.3	69
45	Response to the Comments by Rautenstrauch et al. on our Article, "Convergent Kilo-Scale Synthesis of a Potent Renin Inhibitor for the Treatment of Hypertension― Organic Process Research and Development, 2012, 16, 1187-1187.	1.3	1
46	Convergent, Fit-For-Purpose, Kilogram-Scale Synthesis of a 5-Lipoxygenase Inhibitor. Organic Process Research and Development, 2012, 16, 214-219.	1.3	20
47	Renin inhibitors for the treatment of hypertension: Design and optimization of a novel series of spirocyclic piperidines. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 7399-7404.	1.0	16
48	Convergent Kilo-Scale Synthesis of a Potent Renin Inhibitor for the Treatment of Hypertension. Organic Process Research and Development, 2011, 15, 1138-1148.	1.3	31
49	Efficient preparation of 3â€substituted quinazolinediones directly from anthranilic acids and isocyanates. Journal of Heterocyclic Chemistry, 2011, 48, 473-478.	1.4	11
50	Renin inhibitors for the treatment of hypertension: Design and optimization of a novel series of pyridone-substituted piperidines. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 3970-3975.	1.0	16
51	Chemoselective Staudinger Strategy in the Practical, Fit for Purpose, Gram-Scale Synthesis of an HCV RNA Polymerase Inhibitor. Synlett, 2011, 2011, 57-60.	1.0	8
52	Convergent Approach to Nonsymmetrical 2,5-Diester Pyrroles. Synlett, 2010, 2010, 3086-3088.	1.0	1
53	Mechanistic Analysis of Azine <i>N</i> -Oxide Direct Arylation: Evidence for a Critical Role of Acetate in the Pd(OAc) ₂ Precatalyst. Journal of Organic Chemistry, 2010, 75, 8180-8189.	1.7	203
54	Catalyst and base controlled site-selective sp2 and sp3 direct arylation of azine N-oxides. Tetrahedron, 2009. 65. 3155-3164.	1.0	109

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55	Palladium-Catalyzed Direct Arylation of Azine and Azole <i>N</i> -Oxides: Reaction Development, Scope and Applications in Synthesis. Journal of the American Chemical Society, 2009, 131, 3291-3306.	6.6	392
56	Palladium-Catalyzed Direct Arylation of Nitro-Substituted Aromatics with Aryl Halides. Organic Letters, 2008, 10, 4533-4536.	2.4	138
57	C2, C5, and C4 Azole <i>N</i> -Oxide Direct Arylation Including Room-Temperature Reactions. Journal of the American Chemical Society, 2008, 130, 3276-3277.	6.6	264
58	Site-Selective sp ² and Benzylic sp ³ Palladium-Catalyzed Direct Arylation. Journal of the American Chemical Society, 2008, 130, 3266-3267.	6.6	337
59	Applications of and alternatives to ï€-electron-deficient azine organometallics in metal catalyzed cross-coupling reactions. Chemical Society Reviews, 2007, 36, 1058-1068.	18.7	220
60	Palladium-catalyzed direct arylation of simple arenes in synthesis of biaryl molecules. Chemical Communications, 2006, , 1253.	2.2	518
61	Catalytic Direct Arylation with Aryl Chlorides, Bromides, and Iodides:Â Intramolecular Studies Leading to New Intermolecular Reactions. Journal of the American Chemical Society, 2006, 128, 581-590.	6.6	644
62	A Solution to the 2-Pyridyl Organometallic Cross-Coupling Problem:  Regioselective Catalytic Direct Arylation of Pyridine N-Oxides. Journal of the American Chemical Society, 2005, 127, 18020-18021.	6.6	508
63	High-Yielding Intramolecular Direct Arylation Reactions with Aryl Chlorides. Organic Letters, 2005, 7, 1857-1860.	2.4	217
64	Biaryl Synthesis via Direct Arylation:Â Establishment of an Efficient Catalyst for Intramolecular Processes. Journal of the American Chemical Society, 2004, 126, 9186-9187.	6.6	259