

Jean-Nicolas Desrosiers

List of Publications by Year in descending order

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#	ARTICLE	IF	CITATIONS
1	Construction of Quaternary Stereocenters by Nickel-Catalyzed Heck Cyclization Reactions. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 11921-11924.	7.2	58
2	Enantioselective Nickel-Catalyzed Mizoroki-Heck Cyclizations To Generate Quaternary Stereocenters. <i>Organic Letters</i> , 2017, 19, 3338-3341.	2.4	54
3	Sequential C-H Arylation and Enantioselective Hydrogenation Enables Ideal Asymmetric Entry to the Indenopiperidine Core of an 11 β -HSD-1 Inhibitor. <i>Journal of the American Chemical Society</i> , 2016, 138, 15473-15481.	6.6	48
4	Enantioselective Synthesis of \pm -(Hetero)aryl Piperidines through Asymmetric Hydrogenation of Pyridinium Salts and Its Mechanistic Insights. <i>Organic Letters</i> , 2018, 20, 1333-1337.	2.4	48
5	Synthesis of Enantioenriched 2-Alkyl Piperidine Derivatives through Asymmetric Reduction of Pyridinium Salts. <i>Organic Letters</i> , 2016, 18, 4920-4923.	2.4	46
6	Copper-catalyzed asymmetric hydrogenation of 2-substituted ketones via dynamic kinetic resolution. <i>Chemical Science</i> , 2018, 9, 4505-4510.	3.7	46
7	A Mild Dihydrobenzoxaphosphole Oxazoline/Iridium Catalytic System for Asymmetric Hydrogenation of Unfunctionalized Dialins. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 14428-14432.	7.2	41
8	Application of Biocatalytic Reductive Amination for the Synthesis of a Key Intermediate to a CDK 2/4/6 Inhibitor. <i>Organic Process Research and Development</i> , 2022, 26, 879-890.	1.3	20
9	A Scalable and Regioselective Synthesis of 2-Difluoromethyl Pyridines from Commodity Chemicals. <i>Organic Letters</i> , 2014, 16, 1724-1727.	2.4	18
10	Nickel-catalyzed C-3 direct arylation of pyridinium ions for the synthesis of 1-azafluorenes. <i>Chemical Science</i> , 2016, 7, 5581-5586.	3.7	18
11	Construction of Quaternary Stereocenters by Nickel-Catalyzed Heck Cyclization Reactions. <i>Angewandte Chemie</i> , 2016, 128, 12100-12103.	1.6	18
12	BABIPhos Family of Biaryl Dihydrobenzoxaphosphole Ligands for Asymmetric Hydrogenation. <i>Organic Letters</i> , 2018, 20, 1725-1729.	2.4	18
13	Continuous Process for Preparing the Difluoromethylating Reagent [(DMPU) ₂ Zn(CF ₂ H) ₂] and Improved Synthesis of the ICHF ₂ Precursor. <i>Organic Process Research and Development</i> , 2020, 24, 1077-1083.	1.3	11
14	A radical chlorodifluoromethylation protocol for late-stage difluoromethylation and its application to an oncology candidate. <i>Cell Reports Physical Science</i> , 2021, 2, 100394.	2.8	10
15	Early Development Scale-Up of a Structurally-Challenging 5-Lipoxygenase Activating Protein (FLAP) Inhibitor. <i>Organic Process Research and Development</i> , 2017, 21, 1427-1434.	1.3	9
16	Development of a Practical Sequence for Difluoromethylation of 2-Bromopyridines via Copper-Mediated Reductive Coupling and Decarboxylation. <i>Asian Journal of Organic Chemistry</i> , 2015, 4, 1262-1264.	1.3	6
17	Large-Scale Enantioselective Reduction of 2,3-Disubstituted Indenopyridine Enables a Practical Manufacturing Process for an 11 β -HSD-1 Inhibitor. <i>Organic Process Research and Development</i> , 0, , .	1.3	5