

Jeffery Richardson

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4526273/publications.pdf>

Version: 2024-02-01

37
papers

1,417
citations

394421

19
h-index

361022

35
g-index

41
all docs

41
docs citations

41
times ranked

1314
citing authors

#	ARTICLE	IF	CITATIONS
1	Synthesis and Preclinical Characterization of LY3154885, a Human Dopamine D1 Receptor Positive Allosteric Modulator with an Improved Nonclinical Drug-Drug Interaction Risk Profile. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 3786-3797.	6.4	3
2	Gold(I)-Catalyzed Synthesis of 3-Sulphenyl Pyrroles and Indoles by a Regioselective Annulation of Alkynyl Thioethers. <i>ACS Catalysis</i> , 2021, 11, 6357-6362.	11.2	27
3	Quinolin-8-yl Formate: A New Option for Small-Scale Carbonylation Reactions in Microwave Reactors. <i>Synlett</i> , 2020, 31, 1608-1612.	1.8	1
4	An Efficient Palladium-Catalyzed α -Arylation of Acetone Below its Boiling Point. <i>Synlett</i> , 2020, 31, 1532-1536.	1.8	4
5	Efficient Method for the Synthesis of Amino-1,3-Oxazines from Thioureas. <i>Organic Process Research and Development</i> , 2020, 24, 2853-2863.	2.7	3
6	Development of a Scalable Synthesis of Mevidalen (LY3154207), an Orally Available Positive Allosteric Modulator of the Human Dopamine D1 Receptor. <i>Organic Process Research and Development</i> , 2020, 24, 2549-2564.	2.7	10
7	Rapid Access to Azabicyclo[3.3.1]nonanes by a Tandem Diverted Tsuji-Trost Process. <i>Chemistry - A European Journal</i> , 2020, 26, 14330-14334.	3.3	6
8	Unlocking the potential of late-stage functionalisation: an accurate and fully automated method for the rapid characterisation of multiple regioisomeric products. <i>Reaction Chemistry and Engineering</i> , 2020, 5, 779-792.	3.7	4
9	An Efficient Palladium-Catalysed Aminocarbonylation of Benzyl Chlorides. <i>Synlett</i> , 2020, 31, 369-372.	1.8	7
10	Synthesis and Pharmacological Characterization of 2-(2,6-Dichlorophenyl)-1-(1 <i>S</i> ,3 <i>R</i>)-5-(3-hydroxy-3-methylbutyl)-3-(hydroxymethyl)-1-methyl-3,4-dihydroisoquinolin-2(1 <i>H</i>) (LY3154207), a Potent, Subtype Selective, and Orally Available Positive Allosteric Modulator of the Human Dopamine D1 Receptor. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 8711-8732.	6.4	49
11	Rhodium-Catalyzed Aldehyde Arylation via Formate-Mediated Transfer Hydrogenation: Beyond Metallic Reductants in Grignard/Nozaki-Kishi-Type Addition. <i>Journal of the American Chemical Society</i> , 2019, 141, 1828-1832.	13.7	50
12	Cyclometalated Iridium-PhanePhos Complexes Are Active Catalysts in Enantioselective Allene-Fluoral Reductive Coupling and Related Alcohol-Mediated Carbonyl Additions That Form Acyclic Quaternary Carbon Stereocenters. <i>Journal of the American Chemical Society</i> , 2019, 141, 2087-2096.	13.7	41
13	Improved Substrate Scope in the Potassium Hexacyanoferrate(II)-Based Cyanation for the Synthesis of Benzonitriles and Their Heterocyclic Analogues. <i>Journal of Organic Chemistry</i> , 2018, 83, 4922-4931.	3.2	9
14	Short, Gram-Scale Syntheses of $\hat{1}^2$ - and $\hat{1}^3$ -Lycorane Using Two Distinct Photochemical Approaches. <i>Organic Letters</i> , 2018, 20, 1272-1274.	4.6	23
15	Model Guided Development of a Simple Catalytic Method for the Synthesis of Unsymmetrical Stilbenes by Sequential Heck Reactions of Aryl Bromides with Ethylene. <i>Advanced Synthesis and Catalysis</i> , 2018, 360, 2678-2690.	4.3	9
16	Multi-Objective Computer-Aided Solvent Design for Selectivity and Rate in Reactions. <i>Computer Aided Chemical Engineering</i> , 2018, , 2437-2442.	0.5	4
17	A Method for Identifying and Developing Functional Group Tolerant Catalytic Reactions: Application to the Buchwald-Hartwig Amination. <i>Journal of Organic Chemistry</i> , 2017, 82, 3741-3750.	3.2	28
18	Kinetic Resolution of 2-Substituted Indolines by α -Sulfonylation using an Atropisomeric 4-DMAP-oxide Organocatalyst. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 5760-5764.	13.8	48

#	ARTICLE	IF	CITATIONS
19	Cation exchange media provide a multi-stage, orthogonal catch and release method for MIDA boronates bearing basic centers. <i>Tetrahedron Letters</i> , 2017, 58, 2578-2582.	1.4	1
20	Kinetic Resolution of 2-Substituted Indolines by N -Sulfonylation using an Atropisomeric 4-DMAP Oxide Organocatalyst. <i>Angewandte Chemie</i> , 2017, 129, 5854-5858.	2.0	12
21	Two-Step Cyanomethylation Protocol: Convenient Access to Functionalized Aryl- and Heteroarylacetonitriles. <i>Organic Letters</i> , 2015, 17, 476-479.	4.6	40
22	Enantioselective formal total synthesis of (β)-trachyspic acid. <i>Tetrahedron Letters</i> , 2009, 50, 1566-1567.	1.4	10
23	Total synthesis and determination of the absolute stereochemistry of the squalene synthase inhibitors CJ-13,981 and CJ-13,982. <i>Tetrahedron Letters</i> , 2009, 50, 3388-3390.	1.4	9
24	Total Synthesis of Aigialomycin D using a One-Pot Ketene Generation-Trapping-Aromatization Sequence. <i>Organic Letters</i> , 2009, 11, 4910-4913.	4.6	62
25	Total Synthesis of Citrafungin A. <i>Journal of Organic Chemistry</i> , 2008, 73, 9692-9697.	3.2	21
26	Palladium-Catalyzed Chemo- and Enantioselective Oxidation of Allylic Esters and Carbonates. <i>Journal of the American Chemical Society</i> , 2006, 128, 2540-2541.	13.7	44
27	Reactivity and Selectivity in the Wittig Reaction: A Computational Study. <i>Journal of the American Chemical Society</i> , 2006, 128, 2394-2409.	13.7	164
28	A practical synthesis of a [2.2.1] bicyclic chiral sulfide for asymmetric transformations. <i>Tetrahedron</i> , 2006, 62, 11297-11303.	1.9	30
29	On the Origin of High Selectivity in the Wittig Reaction of Stabilized Ylides: Importance of Dipole-Dipole Interactions. <i>Journal of the American Chemical Society</i> , 2005, 127, 13468-13469.	13.7	70
30	Asymmetric Catalysis Special Feature Part I: Effect of sulfide structure on enantioselectivity in catalytic asymmetric epoxidation of aldehydes: Mechanistic insights and implications. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2004, 101, 5467-5471.	7.1	32
31	The Complexity of Catalysis: Origins of Enantio- and Diastereocontrol in Sulfur Ylide Mediated Epoxidation Reactions. <i>ChemInform</i> , 2004, 35, no.	0.0	0
32	The complexity of catalysis: origins of enantio- and diastereocontrol in sulfur ylide mediated epoxidation reactions. <i>Chemical Communications</i> , 2003, , 2644.	4.1	125
33	Title is missing!. <i>Angewandte Chemie</i> , 2003, 115, 3396-3400.	2.0	34
34	Sulfur-Ylide-Mediated Synthesis of Functionalized and Trisubstituted Epoxides with High Enantioselectivity; Application to the Synthesis of CDP-840.. <i>ChemInform</i> , 2003, 34, no.	0.0	0
35	Sulfur-Ylide-Mediated Synthesis of Functionalized and Trisubstituted Epoxides with High Enantioselectivity; Application to the Synthesis of CDP-840. <i>Angewandte Chemie - International Edition</i> , 2003, 42, 3274-3278.	13.8	122
36	A New Protocol for the In Situ Generation of Aromatic, Heteroaromatic, and Unsaturated Diazo Compounds and Its Application in Catalytic and Asymmetric Epoxidation of Carbonyl Compounds. Extensive Studies To Map Out Scope and Limitations, and Rationalization of Diastereo- and Enantioselectivities. <i>Journal of the American Chemical Society</i> , 2003, 125, 10926-10940.	13.7	179

#	ARTICLE	IF	CITATIONS
37	Unraveling the Mechanism of Epoxide Formation from Sulfur Ylides and Aldehydes. Journal of the American Chemical Society, 2002, 124, 5747-5756.	13.7	136