

Craig Jamieson

List of Publications by Year in descending order

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60
papers

1,740
citations

218381
26
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301761
39
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67
all docs

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docs citations

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times ranked

2115
citing authors

#	ARTICLE	IF	CITATIONS
1	Amide Bond Formation via the Rearrangement of Nitrile Imines Derived from <i>N</i> -2-Nitrophenyl Hydrazonyl Bromides. <i>Organic Letters</i> , 2022, 24, 334-338.	2.4	5
2	Structure-Based Design of a Novel Class of Autotaxin Inhibitors Based on Endogenous Allosteric Modulators. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 6338-6351.	2.9	6
3	Cancer-associated fibroblasts require proline synthesis by PYCR1 for the deposition of pro-tumorigenic extracellular matrix. <i>Nature Metabolism</i> , 2022, 4, 693-710.	5.1	49
4	A type IV Autotaxin inhibitor ameliorates acute liver injury and nonalcoholic steatohepatitis. <i>EMBO Molecular Medicine</i> , 2022, 14, .	3.3	7
5	Design, synthesis and evaluation of E2K derived stapled peptides. <i>Peptide Science</i> , 2021, 113, e24158.	1.0	1
6	Recent Advances in the Generation of Nitrilium Betaine 1,3-Dipoles. <i>Synthesis</i> , 2021, 53, 2395-2407.	1.2	14
7	Ruthenium-Catalyzed Ester Reductions Applied to Pharmaceutical Intermediates. <i>Organic Process Research and Development</i> , 2020, 24, 2745-2751.	1.3	16
8	UV-Induced 1,3,4-Oxadiazole Formation from 5-Substituted Tetrazoles and Carboxylic Acids in Flow. <i>Chemistry - A European Journal</i> , 2020, 26, 14866-14870.	1.7	12
9	One-Pot Suzuki-Hydrogenolysis Protocol for the Modular Synthesis of 2,5-Diaryltetrazoles. <i>Journal of Organic Chemistry</i> , 2020, 85, 7413-7423.	1.7	7
10	Design and Development of a Macrocyclic Series Targeting Phosphoinositide 3-Kinase $\hat{\Gamma}$. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 1386-1391.	1.3	2
11	The Nitrile Imine 1,3-Dipole. , 2020, , .		22
12	Transition-Metal-Free Coupling of 1,3-Dipoles and Boronic Acids as a Sustainable Approach to C-C Bond Formation. <i>Chemistry - A European Journal</i> , 2020, 26, 10591-10597.	1.7	12
13	The Reactivity of Nitrile Imines. , 2020, , 37-97.		2
14	Applications of Nitrile Imine Derivatives. , 2020, , 99-150.		0
15	A fragment-like approach to PYCR1 inhibition. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2019, 29, 2626-2631.	1.0	28
16	A reappraisal of the Ni-[(Benzylpropyl)amino]benzophenone complex in the synthesis of $\hat{\Gamma}$, $\hat{\Gamma}$ -disubstituted amino acid derivatives. <i>Tetrahedron</i> , 2019, 75, 130485.	1.0	3
17	Photocatalytic <i>E</i> $\hat{\Gamma}$ <i>Z</i> Isomerization of $\hat{\Gamma}$ -Ionyl Derivatives. <i>Organic Letters</i> , 2019, 21, 9677-9680.	2.4	33
18	Metal-free C-C bond formation via coupling of nitrile imines and boronic acids. <i>Chemical Science</i> , 2019, 10, 10412-10416.	3.7	19

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19	Cyrene as a bio-based solvent for HATU mediated amide coupling. <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 2851-2854.	1.5	59
20	Surface Functionalization with Carboxylic Acids by Photochemical Microcontact Printing and Tetrazole Chemistry. <i>Langmuir</i> , 2018, 34, 2132-2138.	1.6	20
21	Practical synthesis of pharmaceutically relevant molecules enriched in sp ³ character. <i>Chemical Communications</i> , 2018, 54, 46-49.	2.2	18
22	Scalable total synthesis and comprehensive structure-activity relationship studies of the phytotoxin coronatine. <i>Nature Communications</i> , 2018, 9, 1105.	5.8	16
23	Cyrene as a Bio-Based Solvent for the Suzuki-Miyaura Cross-Coupling. <i>Synlett</i> , 2018, 29, 650-654.	1.0	53
24	Dimethylisorbide (DMI) as a Bio-Derived Solvent for Pd-Catalyzed Cross-Coupling Reactions. <i>Synlett</i> , 2018, 29, 2293-2297.	1.0	21
25	Atom Efficient Synthesis of Selectively Difluorinated Carbocycles through a Gold(I)-Catalyzed Cyclization. <i>Journal of Organic Chemistry</i> , 2018, 83, 8888-8905.	1.7	5
26	Emergence of Small-Molecule Non-RGD-Mimetic Inhibitors for RGD Integrins. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 3241-3251.	2.9	40
27	Rational Design of Autotaxin Inhibitors by Structural Evolution of Endogenous Modulators. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 2006-2017.	2.9	34
28	Amidation of unactivated ester derivatives mediated by trifluoroethanol. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 3507-3518.	1.5	31
29	Structure-Activity Relationships of Small Molecule Autotaxin Inhibitors with a Discrete Binding Mode. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 722-748.	2.9	29
30	A Multicomponent Route to Functionalized Amides and Oxazolidinones. <i>Organic Letters</i> , 2017, 19, 6736-6739.	2.4	7
31	One-Pot, Three-Step Synthesis of Cyclopropylboronic Acid Pinacol Esters from Synthetically Tractable Propargylic Silyl Ethers. <i>Organic Letters</i> , 2017, 19, 3891-3894.	2.4	22
32	Investigation of a Bicyclo[1.1.1]pentane as a Phenyl Replacement within an LpPLA ₂ Inhibitor. <i>ACS Medicinal Chemistry Letters</i> , 2017, 8, 43-48.	1.3	156
33	Scope and limitations of a DMF bio-alternative within Sonogashira cross-coupling and Cacchi-type annulation. <i>Beilstein Journal of Organic Chemistry</i> , 2016, 12, 2005-2011.	1.3	82
34	Synthetic Approaches to Coronafacic Acid, Coronamic Acid, and Coronatine. <i>Synthesis</i> , 2016, 48, 3429-3448.	1.2	12
35	Evolution of a Novel, Orally Bioavailable Series of PI3K γ Inhibitors from an Inhaled Lead for the Treatment of Respiratory Disease. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 7239-7251.	2.9	22
36	Structurally Diverse Mitochondrial Branched Chain Aminotransferase (BCATm) Leads with Varying Binding Modes Identified by Fragment Screening. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 2452-2467.	2.9	23

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37	Development of Autotaxin Inhibitors: An Overview of the Patent and Primary Literature. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 5604-5621.	2.9	59
38	Palladium-Catalyzed Synthesis of Aryl Amides through Silanoate-Mediated Hydrolysis of Nitriles. <i>Synlett</i> , 2015, 27, 88-92.	1.0	5
39	Rational Design of a Novel AMPA Receptor Modulator through a Hybridization Approach. <i>ACS Medicinal Chemistry Letters</i> , 2015, 6, 392-396.	1.3	10
40	Identification of a novel class of autotaxin inhibitors through cross-screening. <i>MedChemComm</i> , 2015, 6, 1149-1155.	3.5	7
41	The Discovery of in Vivo Active Mitochondrial Branched-Chain Aminotransferase (BCATm) Inhibitors by Hybridizing Fragment and HTS Hits. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 7140-7163.	2.9	29
42	Catalytic amidation of unactivated ester derivatives mediated by trifluoroethanol. <i>Chemical Communications</i> , 2015, 51, 9495-9498.	2.2	51
43	Regiospecific Synthesis of N2-Aryl 1,2,3-Triazoles from 2,5-Disubstituted Tetrazoles via Photochemically Generated Nitrile Imine Intermediates. <i>Synlett</i> , 2014, 25, 2480-2484.	1.0	12
44	Amidation of Esters with Amino Alcohols Using Organobase Catalysis. <i>Journal of Organic Chemistry</i> , 2014, 79, 9347-9354.	1.7	27
45	Molar Efficiency: A Useful Metric To Gauge Relative Reaction Efficiency in Discovery Medicinal Chemistry. <i>ACS Sustainable Chemistry and Engineering</i> , 2014, 2, 523-532.	3.2	29
46	Development of a solvent selection guide for aldehyde-based direct reductive amination processes. <i>Green Chemistry</i> , 2013, 15, 1159.	4.6	59
47	Development of a Sustainable Catalytic Ester Amidation Process. <i>ACS Sustainable Chemistry and Engineering</i> , 2013, 1, 1339-1344.	3.2	37
48	Evaluation of alternative solvents in common amide coupling reactions: replacement of dichloromethane and N,N-dimethylformamide. <i>Green Chemistry</i> , 2013, 15, 596.	4.6	118
49	Functional analysis of a novel positive allosteric modulator of AMPA receptors derived from a structure-based drug design strategy. <i>Neuropharmacology</i> , 2013, 64, 45-52.	2.0	31
50	Organobase-Catalyzed Amidation of Esters with Amino Alcohols. <i>Organic Letters</i> , 2013, 15, 2506-2509.	2.4	49
51	Replacement of dichloromethane within chromatographic purification: a guide to alternative solvents. <i>Green Chemistry</i> , 2012, 14, 3016.	4.6	41
52	Structure based evolution of a novel series of positive modulators of the AMPA receptor. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 805-811.	1.0	30
53	Dioxo-triazines as a novel series of cathepsin K inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 1488-1490.	1.0	21
54	A novel series of positive modulators of the AMPA receptor: Discovery and structure based hit-to-lead studies. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 5753-5756.	1.0	36

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55	A novel series of positive modulators of the AMPA receptor: Structure-based lead optimization. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 6072-6075.	1.0	27
56	Positive Allosteric Modulators of the $\hat{\pm}$ -Amino-3-hydroxy-5-methyl-4-isoxazolepropionic Acid (AMPA) Receptor. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 7271-7279.	2.9	37
57	Synthesis of the originally proposed structures of elatenyne and an enyne from <i>Laurencia majuscula</i> . <i>Organic and Biomolecular Chemistry</i> , 2009, 7, 238-252.	1.5	38
58	The Changing Faces of Halogenated Marine Natural Products: Total Synthesis of the Reported Structures of Elatenyne and an Enyne from <i>Laurencia majuscula</i> . <i>Angewandte Chemie - International Edition</i> , 2006, 45, 7199-7202.	7.2	59
59	Recent advances in positive allosteric modulators of the AMPA receptor. <i>Current Opinion in Drug Discovery & Development</i> , 2006, 9, 571-9.	1.9	18
60	Cathepsin K inhibitors, 2000 – 2004. <i>Expert Opinion on Therapeutic Patents</i> , 2005, 15, 33-48.	2.4	18