

Gui Lu

List of Publications by Year in descending order

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56
papers

1,571
citations

236925

25
h-index

330143

37
g-index

56
all docs

56
docs citations

56
times ranked

1867
citing authors

#	ARTICLE	IF	CITATIONS
1	Catalytic Decarboxylative Fluorosulfonylation Enabled by Energy-Transfer-Mediated Photocatalysis. <i>Organic Letters</i> , 2022, 24, 2474-2478.	4.6	36
2	Photoredox-catalyzed aminofluorosulfonylation of unactivated olefins. <i>Chemical Science</i> , 2021, 12, 9359-9365.	7.4	45
3	Recent progress in the synthesis of sulfonyl fluorides for SuFEx click chemistry. <i>Chinese Chemical Letters</i> , 2021, 32, 2736-2750.	9.0	41
4	Organocatalytic enantioselective S _N 1-type dehydrative nucleophilic substitution: access to bis(indolyl)methanes bearing quaternary carbon stereocenters. <i>Chemical Science</i> , 2021, 13, 170-177.	7.4	28
5	Visible-light-promoted radical cross-coupling of <i>para</i> -quinone methides with <i>N</i> -substituted anilines: an efficient approach to 2,2-diarylethylamines. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 860-864.	2.8	20
6	A Novel Aurora Kinase Inhibitor Attenuates Leukemic Cell Proliferation Induced by Mesenchymal Stem Cells. <i>Molecular Therapy - Oncolytics</i> , 2020, 18, 491-503.	4.4	5
7	Transition metal-free synthesis of α -aryl ketones <i>via</i> oxyallyl cation capture with arylboronic acids. <i>Organic Chemistry Frontiers</i> , 2020, 7, 2480-2485.	4.5	9
8	Organocatalytic synthesis of chiral CF ₃ -containing oxazolidines and 1,2-amino alcohols: asymmetric oxa-1,3-dipolar cycloaddition of trifluoroethylamine-derived azomethine ylides. <i>Organic Chemistry Frontiers</i> , 2020, 7, 3452-3458.	4.5	13
9	Enantioselective Dehydrative α -Arylation of α -Indolyl Propargylic Alcohols with Phenols: Access to Chiral Tetrasubstituted Allenes and Naphthopyrans. <i>Organic Letters</i> , 2020, 22, 6873-6878.	4.6	39
10	Catalytic Asymmetric Synthesis of Vicinal Tetrasubstituted Diamines via Umpolung Cross-Mannich Reaction of Cyclic Ketimines. <i>Organic Letters</i> , 2020, 22, 5014-5019.	4.6	30
11	Enantioselective Reformatsky Reaction of Ketones Catalyzed by Chiral Indolinylmethanol. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 4805-4811.	2.4	4
12	Copper-free Sandmeyer-type Reaction for the Synthesis of Sulfonyl Fluorides. <i>Organic Letters</i> , 2020, 22, 3072-3078.	4.6	78
13	Metal-Free Aerobic Sulfonyllactonization of Unsaturated Carboxylic Acids with Thiols Using Air as Sole Oxidant. <i>ChemistrySelect</i> , 2020, 5, 7382-7386.	1.5	3
14	Aurora kinase inhibitor restrains STAT5-activated leukemic cell proliferation by inducing mitochondrial impairment. <i>Journal of Cellular Physiology</i> , 2020, 235, 8358-8370.	4.1	15
15	Construction of Sulfonyl Phthalides via Copper-Catalyzed Oxysulfonylation of 2-Vinylbenzoic Acids with Sodium Sulfinates. <i>Journal of Organic Chemistry</i> , 2019, 84, 13465-13472.	3.2	29
16	An efficient approach to access 1,1,2-triarylethanes enabled by the organo-photoredox-catalyzed decarboxylative addition reaction. <i>Organic Chemistry Frontiers</i> , 2019, 6, 1955-1960.	4.5	25
17	Asymmetric Synthesis of Vicinally Bis(trifluoromethyl)-Substituted 3,3- α -Pyrrolidinyl Spirooxindoles via Organocatalytic 1,3-Dipolar Cycloaddition Reactions. <i>Synthesis</i> , 2019, 51, 1969-1979.	2.3	17
18	Enantioselective Synthesis of Triarylmethanes <i>via</i> Organocatalytic 1,6-Addition of Arylboronic Acids to <i>para</i> -Quinone Methides. <i>Advanced Synthesis and Catalysis</i> , 2019, 361, 1241-1246.	4.3	43

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19	Manganese(III)-Mediated and -Catalyzed Decarboxylative Hydroxysulfonylation of Arylpropionic Acids with Sodium Sulfinates in Water. <i>Advanced Synthesis and Catalysis</i> , 2018, 360, 1611-1616.	4.3	52
20	Organocatalytic Michael/cyclization cascade reactions of 3-isothiocyanato oxindoles with 3-trifluoroethylidene oxindoles: an approach for the synthesis of 3-trifluoromethyl substituted 3,2-pyrrolidinyl-bispirooxindoles. <i>Organic Chemistry Frontiers</i> , 2018, 5, 1375-1380.	4.5	31
21	Mannich Reaction of Indole with Cyclic Imines in Water. <i>Tetrahedron Letters</i> , 2018, 59, 457-461.	1.4	21
22	Synthesis and biological evaluation of aurora kinases inhibitors based on N-trisubstituted pyrimidine scaffold. <i>European Journal of Medicinal Chemistry</i> , 2018, 145, 805-812.	5.5	20
23	Copper-catalyzed <i>peri</i> -selective direct sulfenylation of 1-naphthylamines with disulfides. <i>Organic Chemistry Frontiers</i> , 2018, 5, 982-989.	4.5	34
24	Copper-catalyzed aerobic decarboxylative coupling between cyclic α -amino acids and diverse C-H nucleophiles with low catalyst loading. <i>RSC Advances</i> , 2018, 8, 16202-16206.	3.6	11
25	Identification of a novel compound targeting the nuclear export of influenza A virus nucleoprotein. <i>Journal of Cellular and Molecular Medicine</i> , 2018, 22, 1826-1839.	3.6	10
26	Visible-Light-Mediated Decarboxylative Benzoylation of Imines with Arylacetic Acids. <i>Journal of Organic Chemistry</i> , 2018, 83, 12559-12567.	3.2	25
27	Highly efficient construction of chiral dispirocyclic oxindole/thiobutyrolactam/chromanone complexes through Michael/cyclization cascade reactions with a rosin-based squaramide catalyst. <i>Tetrahedron</i> , 2018, 74, 3734-3741.	1.9	25
28	Nickel-catalyzed direct C-H bond sulfenylation of acylhydrazines. <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 6047-6056.	2.8	27
29	Structure-based drug design: Synthesis and biological evaluation of quinazolin-4-amine derivatives as selective Aurora A kinase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2018, 157, 1361-1375.	5.5	23
30	Copper-Catalyzed Remote C-H Functionalizations of Naphthylamides through a Coordinating Activation Strategy and Single-Electron-Transfer (SET) Mechanism. <i>ACS Catalysis</i> , 2017, 7, 2661-2667.	11.2	122
31	Asymmetric amination of 2-substituted indolin-3-ones catalyzed by natural cinchona alkaloids. <i>Organic Chemistry Frontiers</i> , 2017, 4, 1400-1406.	4.5	22
32	A rapid and sensitive UHPLC-MS/MS method for quantification of 83b1 in plasma and its application to bioavailability study in rats. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2017, 134, 71-76.	2.8	1
33	Asymmetric synthesis of trifluoromethyl-substituted 3,3-pyrrolidinyl-dispirooxindoles through organocatalytic 1,3-dipolar cycloaddition reactions. <i>Organic Chemistry Frontiers</i> , 2017, 4, 472-482.	4.5	68
34	Highly Efficient Construction of CF ₃ -Containing 3,3-Pyrrolidinyl-dispirooxindoles via Base-Catalyzed Diastereoselective [3+2] Annulation. <i>Heterocycles</i> , 2017, 94, 879.	0.7	9
35	Synthesis of Pelorol and Its Analogs and Their Inhibitory Effects on Phosphatidylinositol 3-Kinase. <i>Marine Drugs</i> , 2016, 14, 118.	4.6	7
36	Copper-catalyzed C5-regioselective C-H sulfonylation of 8-aminoquinoline amides with aryl sulfonyl chlorides. <i>Tetrahedron Letters</i> , 2016, 57, 2121-2124.	1.4	47

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37	Determination of a novel Aurora-A (AurA) kinase AKI603 by UPLC-MS/MS and its application to a bioavailability study in rat. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2016, 125, 303-309.	2.8	0
38	Cu(II)-catalyzed cross-dehydrogenative coupling reaction of N ² -acyl arylhydrazines and phosphites. <i>RSC Advances</i> , 2016, 6, 84587-84591.	3.6	12
39	Design, Synthesis, and Biological Evaluation of Substituted Pyrimidines as Potential Phosphatidylinositol 3-Kinase (PI3K) Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 7268-7274.	6.4	35
40	Aurora A Kinase Inhibitor AKI603 Induces Cellular Senescence in Chronic Myeloid Leukemia Cells Harboring T315I Mutation. <i>Scientific Reports</i> , 2016, 6, 35533.	3.3	29
41	Discovery of 2-(2-aminopyrimidin-5-yl)-4-morpholino- N -(pyridin-3-yl)quinazolin-7-amines as novel PI3K/mTOR inhibitors and anticancer agents. <i>European Journal of Medicinal Chemistry</i> , 2016, 108, 644-654.	5.5	28
42	Organocatalytic Diels-Alder Reaction of 2-Vinylindoles with Methyleneindolinones: An Efficient Approach to Functionalized Carbazolespirooxindoles. <i>Advanced Synthesis and Catalysis</i> , 2015, 357, 993-1003.	4.3	53
43	Stereoselective synthesis of epoxyisoprostanes: an organocatalytic and <i>pot-economy</i> approach. <i>Chemical Communications</i> , 2015, 51, 10170-10173.	4.1	17
44	A novel compound against oncogenic Aurora kinase A overcomes imatinib resistance in chronic myeloid leukemia cells. <i>International Journal of Oncology</i> , 2015, 46, 2488-2496.	3.3	17
45	Copper(II)-catalyzed coupling reaction: an efficient and regioselective approach to N ² ,N ² -diaryl acylhydrazines. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 2055-2063.	2.8	27
46	Synthesis of Terminal Vinylphosphonates Via Dbu-Promoted Tandem Phospha-Michael/Elimination Reactions. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 2014, 189, 1858-1866.	1.6	9
47	Design, synthesis and bioevaluation of N-trisubstituted pyrimidine derivatives as potent aurora A kinase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2014, 78, 65-71.	5.5	39
48	A Novel Small-Molecule Aurora Kinase Inhibitor Attenuates Breast Tumor-Initiating Cells and Overcomes Drug Resistance. <i>Molecular Cancer Therapeutics</i> , 2014, 13, 1991-2003.	4.1	51
49	Palladium-catalyzed reductive homocoupling of N ² -tosyl arylhydrazines. <i>Organic and Biomolecular Chemistry</i> , 2013, 11, 8014.	2.8	24
50	Chemomics and drug innovation. <i>Science China Chemistry</i> , 2013, 56, 71-85.	8.2	12
51	Palladium-catalyzed Suzuki cross-coupling of N ² -tosyl arylhydrazines. <i>Chemical Communications</i> , 2013, 49, 5268.	4.1	46
52	Asymmetric Domino Nitro-Michael/Horner-Wadsworth-Emmons Reaction for Disubstituted Cyclohexenecarboxylate Annulation: Efficient Synthesis of Dipeptidyl Peptidase IV Inhibitor ABT-341 and Influenza Neuraminidase Inhibitor. <i>Advanced Synthesis and Catalysis</i> , 2012, 354, 1961-1970.	4.3	13
53	Indolinylmethanol catalyzed enantioselective Reformatsky reaction with ketones. <i>Tetrahedron: Asymmetry</i> , 2010, 21, 2816-2824.	1.8	24
54	Zinc-Salen-Catalyzed Asymmetric Alkynylation of Alkyl Acylsilanes. <i>Advanced Synthesis and Catalysis</i> , 2009, 351, 1955-1960.	4.3	37

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55	Highly Efficient Asymmetric Michael Reaction of Aldehydes to Nitroalkenes with Diphenylperhydroindolinol Silyl Ethers as Organocatalysts. <i>Advanced Synthesis and Catalysis</i> , 2009, 351, 2449-2459.	4.3	58
56	Zinc-Salen-Catalyzed Asymmetric Alkynylation of Alkyl Acylsilanes. <i>Advanced Synthesis and Catalysis</i> , 2009, 351, 2541-2541.	4.3	5