

Zhi Zhou

List of Publications by Year in descending order

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57
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#	ARTICLE	IF	CITATIONS
1	Enantioselective synthesis of indenopyrazolopyrazolones enabled by dual directing groups-assisted and rhodium(III)-catalyzed tandem C-H alkenylation/[3+2] stepwise cycloaddition. <i>Chinese Chemical Letters</i> , 2022, 33, 842-846.	9.0	19
2	Rh(III)-Catalysed Switchable and Chemoselective Synthesis of Difluorinated Pyrazolo[1,2-a]indazolone and Indole Frameworks. <i>Asian Journal of Organic Chemistry</i> , 2022, 11, .	2.7	5
3	Hexafluoroisopropanol (HFIP)-prompted rearrangement of N-phenoxy-sulfonamides for the direct assembly of ortho-sulfonamide phenols: A combined experimental and computational study. <i>Tetrahedron Letters</i> , 2022, 89, 153601.	1.4	0
4	Rh(III)-Catalysed cascade C-H imidization/cyclization of N-methoxybenzamidates with isoxazolones for the assembly of dihydroquinazolin-4(1H)-one derivatives. <i>Organic Chemistry Frontiers</i> , 2022, 9, 1904-1910.	4.5	4
5	Specific assembly of dihydrobenzofuran frameworks via Rh(III)-catalysed C-H coupling of N-phenoxyacetamides with 2-alkenylphenols. <i>New Journal of Chemistry</i> , 2022, 46, 5705-5711.	2.8	3
6	Direct Assembly of Phthalides via Calcium(II)-Catalyzed Cascade ortho-C-Alkenylation/Hydroacyloxylation of 3-Aminobenzoic Acids with Alkynes in Hexafluoroisopropanol. <i>Organic Letters</i> , 2022, 24, 1575-1580.	4.6	8
7	Site-selective rhodium carbene transfer of 2-hydroxy-1-nitrostyrenes with diazo compounds En route to 2-alkylated benzofurans. <i>Organic Chemistry Frontiers</i> , 2022, 9, 3268-3273.	4.5	4
8	A novel 3-acyl isoquinolin-1(2H)-one induces G2 phase arrest, apoptosis and GSDME-dependent pyroptosis in breast cancer. <i>PLoS ONE</i> , 2022, 17, e0268060.	2.5	3
9	Difluoromethylene Alkyne-Enabled Diverse C-H Functionalization and Application to the on-DNA Synthesis of Difluorinated Isocoumarins. <i>Angewandte Chemie - International Edition</i> , 2021, 60, 1959-1966.	13.8	55
10	Difluoromethylene Alkyne-Enabled Diverse C-H Functionalization and Application to the on-DNA Synthesis of Difluorinated Isocoumarins. <i>Angewandte Chemie</i> , 2021, 133, 1987-1994.	2.0	8
11	Synthesis of 2-aminobenzofurans via base-mediated [3 + 2] annulation of N-phenoxy amides with gem-difluoroalkenes. <i>Organic Chemistry Frontiers</i> , 2021, 8, 4452-4458.	4.5	15
12	Chiral Allylic Amine Synthesis Enabled by the Enantioselective Cp* ^X Rh(III)-Catalyzed Carboaminations of 1,3-Dienes. <i>ACS Catalysis</i> , 2021, 11, 2279-2287.	11.2	33
13	Rh(III)-Catalyzed Redox-Neutral C-H Activation/[3 + 2] Annulation of N-Phenoxy Amides with Propargylic Monofluoroalkynes. <i>Organic Letters</i> , 2021, 23, 2285-2291.	4.6	10
14	Rh(III)-Catalyzed C-H Activation/[3 + 2] Annulation of N-Phenoxyacetamides via Carboxygenation of 1,3-Dienes. <i>Organic Letters</i> , 2021, 23, 3844-3849.	4.6	16
15	Chemodivergent assembly of ortho-functionalized phenols with tunable selectivity via rhodium(III)-catalyzed and solvent-controlled C-H activation. <i>Communications Chemistry</i> , 2021, 4, .	4.5	10
16	Rh(III)-Catalyzed Chemoselective C-H Alkenylation and [5 + 1] Annulation with gem-Difluoromethylene Enabled by the Distinctive Fluorine Effect. <i>Journal of Organic Chemistry</i> , 2021, 86, 9711-9722.	3.2	9
17	Rhodium(III)-Catalyzed Cascade C-H Coupling/Terminus Michael Addition of N-Phenoxy Amides with 1,6-Diynes. <i>ChemistrySelect</i> , 2021, 6, 6574-6578.	1.5	5
18	Mechanistic Insights into the Dual Directing Group-Mediated C-H Functionalization/Annulation via a Hydroxyl Group-Assisted M ^{III} -M ^V -M ^{III} Pathway. <i>ACS Omega</i> , 2021, 6, 17642-17650.	3.5	5

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19	TFA-Prompted/Rh(III)-Catalysed Chemoselective C3-or C2-H Functionalization of Indoles with Methylenecyclopropanes. <i>European Journal of Organic Chemistry</i> , 2021, 2021, 5507.	2.4	5
20	Identification of the anti-fungal drug fenticonazole nitrate as a novel PPAR δ -modulating ligand with good therapeutic index: Structure-based screening and biological validation. <i>Pharmacological Research</i> , 2021, 173, 105860.	7.1	7
21	Synthesis of Difluorinated Dihydrobenzo[<i>de</i>]chromenes via Rh(III)-Catalysed C-H Couplings of 1-Naphthols with Gem-Difluoromethylene Alkynes. <i>Advanced Synthesis and Catalysis</i> , 2021, 363, 1352-1357.	4.3	13
22	Rh-Catalyzed and synergistic dual directing group-enabled redox-neutral [3+3] annulation of N-phenoxyacetamides with β -allenols. <i>Chemical Communications</i> , 2021, 57, 9284-9287.	4.1	12
23	Chemo-, Regio-, and Stereoselective Assembly of Polysubstituted Furan-2(5H)-ones Enabled by Rh(III)-Catalyzed Domino C-H Alkenylation/Directing Group Migration/Lactonization: A Combined Experimental and Computational Study. <i>ACS Catalysis</i> , 2021, 11, 13921-13934.	11.2	20
24	Gem-Difluorocyclopropenes as Versatile β -Monofluorinated Three-sp ² Carbon Sources for Cp*Rh(III)-Catalyzed [4 + 3] Annulation: Experimental Development and Mechanistic Insight. <i>ACS Catalysis</i> , 2021, 11, 14694-14701.	11.2	27
25	Rh(III)-Catalyzed Redox-Neutral [4+2] Annulation for Direct Assembly of 3-Acyl Isoquinolin-1(2H)-ones as Potent Antitumor Agents. <i>ChemPlusChem</i> , 2020, 85, 405-410.	2.8	16
26	Rhodium(III)-Catalyzed Enantio- and Diastereoselective C-H Cyclopropylation of N-Phenoxy-sulfonamides: Combined Experimental and Computational Studies. <i>Angewandte Chemie - International Edition</i> , 2020, 59, 2890-2896.	13.8	80
27	Rhodium(III)-Catalyzed Enantio- and Diastereoselective C-H Cyclopropylation of N-Phenoxy-sulfonamides: Combined Experimental and Computational Studies. <i>Angewandte Chemie</i> , 2020, 132, 2912-2918.	2.0	19
28	Lossen Rearrangement vs C-N Reductive Elimination Enabled by Rh(III)-Catalyzed C-H Activation/Selective Lactone Ring-Opening: Chemodivergent Synthesis of Quinolinones and Dihydroisoquinolinones. <i>Organic Letters</i> , 2020, 22, 9677-9682.	4.6	18
29	Ru-Catalyzed and acidity-controlled tunable [5+1]/[5+2] annulation for building ring-fused quinazolines and 1,3-benzodiazepines. <i>Chemical Communications</i> , 2020, 56, 11315-11318.	4.1	14
30	Synthesis of Indenopyrazole Frameworks via Cascade C-H Functionalization/[3 + 2] Dipolar Cycloaddition/Aromatization Rearrangement Reactions. <i>Organic Letters</i> , 2020, 22, 7152-7157.	4.6	29
31	Identification and structural insight of an effective PPAR δ modulator with improved therapeutic index for anti-diabetic drug discovery. <i>Chemical Science</i> , 2020, 11, 2260-2268.	7.4	15
32	Experimental and Computational Studies on Cp*CyRh(III)/KOPiv-Catalyzed Intramolecular Dehydrogenative Cross-Couplings for Building Eight-Membered Sultam/Lactam Frameworks. <i>Organic Letters</i> , 2020, 22, 5473-5478.	4.6	14
33	Cascade Reductive Rearrangement for the Stereoselective Synthesis of Multifunctional Piperidinones: A Combined Experimental and Computational Study. <i>ChemistrySelect</i> , 2020, 5, 2332-2336.	1.5	0
34	Synergistic Dual Directing Groups-Enabled Diastereoselective C-H Cyclopropylation via Rh(III)-Catalyzed Couplings with Cyclopropenyl Alcohols. <i>Organic Letters</i> , 2020, 22, 1295-1300.	4.6	16
35	Cobalt(III)-Catalyzed and Dimethyl Sulfoxide-Involving Cross-Coupling of Ketones and Amides for Direct Synthesis of β -Amino Ketones. <i>Advanced Synthesis and Catalysis</i> , 2019, 361, 4278-4285.	4.3	17
36	Rh(III)-Catalyzed C-H Activation/Cycloisomerization of N-Phenoxyacetamides with Enynones for One-Pot Assembly of Furylated 2-Alkenylphenols. <i>Journal of Organic Chemistry</i> , 2019, 84, 15557-15566.	3.2	9

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37	Cobalt-Catalyzed Allylation of Amides with Styrenes Using DMSO as Both the Solvent and the β -Methylene Source. <i>Organic Letters</i> , 2019, 21, 7248-7253.	4.6	12
38	Chiral, <i>N</i> -phosphoryl sulfonamide Brønsted acids with an intramolecular hydrogen bond interaction that modulates organocatalysis. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 8690-8694.	2.8	13
39	Chemodivergent Couplings of <i>N</i> -Arylureas and Methyleneoxetanones via Rh(III)-Catalyzed and Solvent-Controlled $C\alpha$ -H Activation. <i>Organic Letters</i> , 2019, 21, 4143-4147.	4.6	27
40	Metal-Free [3,3]-Sigmatropic Rearrangement/[3+2] Annulation Cascade of <i>N</i> -Phenoxy Amides with Terminal Alkynes for the Diastereoselective Synthesis of trans-Dihydrobenzofurans. <i>Advanced Synthesis and Catalysis</i> , 2019, 361, 3980-3985.	4.3	7
41	Stereoselective β -F Elimination Enabled Redox-Neutral [4 + 1] Annulation via Rh(III)-Catalyzed $C\alpha$ -H Activation: Access to Z-Monofluoroalkenyl Dihydrobenzo[d]isoxazole Framework. <i>Organic Letters</i> , 2019, 21, 5229-5233.	4.6	36
42	Rhodium-catalyzed chemoselective $C\alpha$ -H functionalization of benzamides with methyleneoxetanones controlled by the solvent. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 6114-6118.	2.8	20
43	Cobalt(III)-Catalyzed, DMSO-Involved, and TFA-Controlled Regioselective $C\alpha$ -H Functionalization of Anilines with Alkynes for Specific Assembly of 3-Arylquinolines. <i>Advanced Synthesis and Catalysis</i> , 2019, 361, 3002-3007.	4.3	26
44	Redox-Neutral [4 + 2] Annulation of <i>N</i> -Methoxybenzamides with Alkynes Enabled by an Osmium(II)/HOAc Catalytic System. <i>Organic Letters</i> , 2019, 21, 9904-9908.	4.6	25
45	Hydroxyl Group-Prompted and Iridium(III)-Catalyzed Regioselective $C\alpha$ -H Annulation of <i>N</i> -phenoxyacetamides with Propargyl Alcohols. <i>Advanced Synthesis and Catalysis</i> , 2018, 360, 2470-2475.	4.3	48
46	One-pot regioselective synthesis of 2,4-disubstituted quinolines via copper-catalyzed cascade annulation. <i>Organic Chemistry Frontiers</i> , 2018, 5, 1713-1718.	4.5	20
47	Catalyst-Controlled [3 + 2] and [4 + 2] Annulations of Oximes with Propargyl Alcohols: Divergent Access to Indenamines and Isoquinolines. <i>Organic Letters</i> , 2018, 20, 182-185.	4.6	60
48	Rh(III)-Catalyzed Oxidative [5 + 2] Annulation Using Two Transient Assisting Groups: Stereospecific Assembly of 3-Alkenylated Benzoxepine Framework. <i>Organic Letters</i> , 2018, 20, 6812-6816.	4.6	29
49	Rh(III)-Catalyzed and Solvent-Controlled Chemoselective Synthesis of Chalcone and Benzofuran Frameworks via Synergistic Dual Directing Groups Enabled Regioselective $C\alpha$ -H Functionalization: A Combined Experimental and Computational Study. <i>ACS Catalysis</i> , 2018, 8, 9508-9519.	11.2	77
50	2-Chromene-3-carboxylic Acid Synthesis via Solvent-Controlled and Rhodium(III)-Catalyzed Redox-Neutral $C\alpha$ -H Activation/[3 + 3] Annulation Cascade. <i>Organic Letters</i> , 2018, 20, 3892-3896.	4.6	37
51	Regiocontrolled Coupling of Aromatic and Vinylic Amides with β -Allenols To Form β -Lactams via Rhodium(III)-Catalyzed $C\alpha$ -H Activation. <i>Organic Letters</i> , 2016, 18, 5668-5671.	4.6	85
52	Rhodium(III)-Catalyzed Redox-Neutral $C\alpha$ -H Annulation of Arylnitrones and Alkynes for the Synthesis of Indole Derivatives. <i>Advanced Synthesis and Catalysis</i> , 2015, 357, 2944-2950.	4.3	52
53	Cascade Synthesis of 3-Alkylidene Dihydrobenzofuran Derivatives via Rhodium(III)-Catalyzed Redox-Neutral $C\alpha$ -H Functionalization/Cyclization. <i>Organic Letters</i> , 2015, 17, 5874-5877.	4.6	64
54	Synthesis of benzofurans via ruthenium-catalyzed redox-neutral $C\alpha$ -H functionalization and reaction with alkynes under mild conditions. <i>Organic Chemistry Frontiers</i> , 2014, 1, 1161-1165.	4.5	60

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55	Rhodium(III)-Catalyzed Redox-Neutral Coupling of <i>N</i> -Phenoxyacetamides and Alkynes with Tunable Selectivity. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 6033-6037.	13.8	293
56	Rhodium(III)-Catalyzed C-H Olefination for the Synthesis of <i>ortho</i> -Alkenyl Phenols Using an Oxidizing Directing Group. <i>Organic Letters</i> , 2013, 15, 3366-3369.	4.6	152
57	One-pot self-assembly of three-dimensional graphene macroassemblies with porous core and layered shell. <i>Journal of Materials Chemistry</i> , 2011, 21, 12352.	6.7	64