

Wen-Hao Hu

List of Publications by Year in descending order

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papers

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#	ARTICLE	IF	CITATIONS
1	An asymmetric catalytic multi-component reaction enabled the green synthesis of isoserine derivatives and semi-synthesis of paclitaxel. <i>Green Synthesis and Catalysis</i> , 2023, 4, 58-63.	3.7	6
2	A diastereoselective three-component reaction for the assembly of succinimide and isatin hybrid molecules with potent anticancer activities. <i>Molecular Diversity</i> , 2023, 27, 837-843.	2.1	1
3	Synthesis of dihydrofuran-3-one and 9,10-phenanthrenequinone hybrid molecules and biological evaluation against colon cancer cells as selective Akt kinase inhibitors. <i>Molecular Diversity</i> , 2023, 27, 845-855.	2.1	2
4	Radical Cascade Multicomponent Minisci Reactions with Diazo Compounds. <i>ACS Catalysis</i> , 2022, 12, 1357-1363.	5.5	34
5	A Rh(II)-catalyzed highly stereoselective [3+2] annulation of vinyl diazoacetates with indole-2-carbaldehyde for the synthesis of indolyl dihydrofurans. <i>Molecular Diversity</i> , 2022, 26, 3379-3386.	2.1	1
6	Enantioselective Propargylation of Oxonium Ylide with $\hat{\pm}$ -Propargylic-3-Indolymethanol: Access to Chiral Propargylic Indoles. <i>Organic Letters</i> , 2022, 24, 1027-1032.	2.4	4
7	An asymmetric three-component reaction of a diazo compound with an alcohol and a seven-membered imine. <i>Organic Chemistry Frontiers</i> , 2022, 9, 2102-2108.	2.3	5
8	Functionalization of DNA-Tagged Alkenes with Diazo Compounds via Photocatalysis. <i>Organic Letters</i> , 2022, 24, 2208-2213.	2.4	28
9	Recent advances in gold-complex and chiral organocatalyst cooperative catalysis for asymmetric alkyne functionalization. <i>Chinese Chemical Letters</i> , 2022, 33, 4969-4979.	4.8	26
10	Diastereoselective aldol-type interception of phenolic oxonium ylides for the direct assembly of 2,2-disubstituted dihydrobenzofurans. <i>Organic and Biomolecular Chemistry</i> , 2022, 20, 4635-4639.	1.5	4
11	Discovery of Novel Benzo[4,5]imidazo[1,2- <i>a</i>]pyrazin-1-amine-3-amide-one Derivatives as Anticancer Human A _{2A} Adenosine Receptor Antagonists. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 8933-8947.	2.9	8
12	Chiral rhodium(II)-catalyzed asymmetric aldol-type interception of an oxonium ylide to assemble chiral 2,3-dihydropyrans. <i>Science China Chemistry</i> , 2022, 65, 1607-1614.	4.2	7
13	Photoredox-Catalyzed Carbonyl Alkylative Amination with Diazo Compounds: A Three-Component Reaction for the Construction of $\hat{\pm}$ -Amino Acid Derivatives. <i>Organic Letters</i> , 2022, 24, 4908-4913.	2.4	12
14	gem-Difunctionalization of $\hat{\pm}$ -diazoarylketones with diaryldiselenides and N-halosuccinimides: facile synthesis of $\hat{\pm}$ -halo- $\hat{\pm}$ -arylseleno ketones. <i>Molecular Diversity</i> , 2021, 25, 2459-2466.	2.1	3
15	Ruthenium(II)-catalyzed facile synthesis of 3-(phenylamino)-1H-indole-2-carboxylates from anilines and diazo pyruvates promoted by FeCl ₃ . <i>Tetrahedron</i> , 2021, 77, 131399.	1.0	2
16	Asymmetric Allylation by Chiral Organocatalyst-Promoted Formal Hetero-Ene Reactions of Alkylgold Intermediates. <i>Angewandte Chemie</i> , 2021, 133, 2020-2027.	1.6	4
17	Asymmetric Allylation by Chiral Organocatalyst-Promoted Formal Hetero-Ene Reactions of Alkylgold Intermediates. <i>Angewandte Chemie - International Edition</i> , 2021, 60, 1992-1999.	7.2	33
18	Design, synthesis and biological evaluation of novel scaffold benzo[4,5]imidazo[1,2- <i>a</i>]pyrazin-1-amine: Towards adenosine A _{2A} receptor (A _{2A} AR) antagonist. <i>European Journal of Medicinal Chemistry</i> , 2021, 210, 113040.	2.6	12

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19	A novel STAT3 inhibitor W2014-S regresses human non-small cell lung cancer xenografts and sensitizes EGFR-TKI acquired resistance. <i>Theranostics</i> , 2021, 11, 824-840.	4.6	50
20	Facile synthesis of 1,4-oxazines by ruthenium-catalyzed tandem N-H insertion/cyclization of β -arylamino ketones and diazo pyruvates. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 1769-1772.	1.5	5
21	Highly diastereoselective synthesis of vicinal diamines <i>via</i> a Rh-catalyzed three-component reaction of diazo compounds with diarylmethanimines and ketimines. <i>Organic Chemistry Frontiers</i> , 2021, 8, 2997-3003.	2.3	4
22	Gold(I)-catalyzed redox transformation of α -nitroalkynes with indoles for the synthesis of 2,3-biindole derivatives. <i>Organic Chemistry Frontiers</i> , 2021, 8, 1808-1816.	2.3	16
23	Catalyst-free <i>gem</i> -chlorosulfurization of difluoromethyl-substituted diazo compounds with disulfide and PhICl_2 . <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 8030-8034.	1.5	3
24	Gold(I)-catalyzed intramolecular cyclization/intermolecular cycloaddition cascade as a fast track to polycarbocycles and mechanistic insights. <i>Nature Communications</i> , 2021, 12, 1182.	5.8	43
25	Gold-catalyzed ketene dual functionalization and mechanistic insights: divergent synthesis of indenes and benzo[d]oxepines. <i>Science China Chemistry</i> , 2021, 64, 778-787.	4.2	23
26	Enantioselective Intermolecular Mannich-Type Interception of Phenolic Oxonium Ylide for the Direct Assembly of Chiral 2,2-Disubstituted Dihydrobenzofurans. <i>ACS Catalysis</i> , 2021, 11, 6750-6756.	5.5	21
27	Asymmetric Three-Component Propargyloxylation for Direct Assembly of Polyfunctionalized Chiral Succinate Derivatives. <i>CCS Chemistry</i> , 2021, 3, 1903-1912.	4.6	15
28	Gold-Catalyzed Carbocyclization/C=N Bond Formation Cascade of Alkyne-Tethered Diazo Compounds with Benzo[<i>c</i>]isoxazoles for the Assembly of 4-aminonaphthalenones and Indenes. <i>Advanced Synthesis and Catalysis</i> , 2021, 363, 4018-4023.	2.1	13
29	Enantioselective Oxidative Multi-Functionalization of Terminal Alkynes with Nitrones and Alcohols for Expedient Assembly of Chiral β -Alkoxy- β -amino-ketones. <i>Journal of the American Chemical Society</i> , 2021, 143, 14703-14711.	6.6	44
30	Structure-based discovery of potent and selective small-molecule inhibitors targeting signal transducer and activator of transcription 3 (STAT3). <i>European Journal of Medicinal Chemistry</i> , 2021, 221, 113525.	2.6	6
31	Enantioselective assembly of 3,3-disubstituted succinimides <i>via</i> three-component reaction of vinyl diazosuccinimides with alcohols and imines. <i>Chemical Communications</i> , 2021, 57, 8043-8046.	2.2	12
32	Enantioselective formal carbene insertion into C=N bond of amina as a concise track to chiral β -amino- β , β -amino acids and synthetic applications. <i>Green Synthesis and Catalysis</i> , 2021, 2, 337-344.	3.7	29
33	An asymmetric oxidative cyclization/Mannich-type addition cascade reaction for direct access to chiral pyrrolidin-3-ones. <i>Chemical Communications</i> , 2021, 57, 12171-12174.	2.2	7
34	Ternary Catalysis Enabled Three-Component Asymmetric Allylic Alkylation as a Concise Track to Chiral β , β -Disubstituted Ketones. <i>Journal of the American Chemical Society</i> , 2021, 143, 20818-20827.	6.6	60
35	Iron-catalyzed [3 + 2]-cycloaddition of <i>in situ</i> generated α -ylides with alkynes or olefins: access to multi-substituted/polycyclic pyrrole derivatives. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 409-414.	1.5	15
36	Revisiting signal transducer and activator of transcription 3 (STAT3) as an anticancer target and its inhibitor discovery: Where are we and where should we go?. <i>European Journal of Medicinal Chemistry</i> , 2020, 187, 111922.	2.6	56

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37	A gold(κ^2)-catalysed chemoselective three-component reaction between phenols, α -diazocarbonyl compounds and allenamides. <i>Chemical Communications</i> , 2020, 56, 1649-1652.	2.2	10
38	Blue Light-Promoted Formal [4+1]-Annulation of Diazoacetates with <i>o</i> -Aminoacetophenones: Synthesis of Polysubstituted Indolines and Computational Study. <i>Journal of Organic Chemistry</i> , 2020, 85, 13920-13928.	1.7	21
39	A Rh(II)/phosphoric acid co-catalyzed three-component reaction of diazo-ketones with alcohols and azonaphthalenes: access to indole derivatives via a formal [3 + 2]-cycloaddition. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 9805-9809.	1.5	7
40	Synthesis and biological evaluation of substituted pyrrolidines and pyrroles as potential anticancer agents. <i>Archiv Der Pharmazie</i> , 2020, 353, e2000136.	2.1	10
41	Diastereoselective Trapping of Transient Carboxylic Oxonium Ylides with α,β -Unsaturated α -Acyl Imidazoles. <i>Advanced Synthesis and Catalysis</i> , 2020, 362, 4662-4667.	2.1	6
42	Desaturation via Redox-Neutral Hydrogen Transfer Process: Synthesis of 2-Allyl Anilines, Mechanism and Applications. <i>IScience</i> , 2020, 23, 101168.	1.9	1
43	Discovery of Novel Antibiotics as Covalent Inhibitors of Fatty Acid Synthesis. <i>ACS Chemical Biology</i> , 2020, 15, 1826-1834.	1.6	10
44	Synthesis and Anticancer Activity of Novel Actinonin Derivatives as HsPDF Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 6959-6978.	2.9	18
45	A Cleavage-Modification-Reassembly Process Catalyzed by Rhodium and Brønsted Acid for the Synthesis of Multi-Substituted Anilines. <i>Advanced Synthesis and Catalysis</i> , 2020, 362, 1961-1965.	2.1	8
46	Enantioselective three-component aminomethylation of α -diazo ketones with alcohols and 1,3,5-triazines. <i>Nature Communications</i> , 2020, 11, 1511.	5.8	62
47	Enantioselective Synthesis of Fluoroalkyl-Substituted <i>syn</i> -Diamines by the Asymmetric <i>gem</i> -Difunctionalization of 2,2,2-Trifluorodiaoethane. <i>ACS Catalysis</i> , 2020, 10, 4559-4565.	5.5	43
48	Ruthenium-Catalyzed Diastereoselective Synthesis of Fully Substituted Pyrrolidines from Anilines and Diazo Pyruvates. <i>Organic Letters</i> , 2020, 22, 3094-3098.	2.4	8
49	Rh(II)/Ag(I)-Cocatalyzed Three-Component Reaction <i>via</i> S_N1/S_N1 -Type Trapping of Oxonium Ylide with the Nicholas Intermediate. <i>Journal of Organic Chemistry</i> , 2020, 85, 9850-9862.	1.7	11
50	Rhodium-Catalyzed Sequential Cycloisomerization/Aldol Addition of Cyclopropene Carboxylic Acids with Isatins. <i>Organic Letters</i> , 2020, 22, 5600-5604.	2.4	12
51	Rhodium catalyzed direct C3-ethoxycarbonylmethylation of imidazo[1,2-a]pyridines with ethyl diazoacetate. <i>Tetrahedron</i> , 2020, 76, 130998.	1.0	6
52	Brønsted Acid Catalyzed Enantioselective Assembly of Spirochroman-3,3-oxindoles. <i>Organic Letters</i> , 2020, 22, 2925-2930.	2.4	27
53	A Rh-catalyzed three-component reaction for the diastereoselective synthesis of pyrazolone derivatives with contiguous quaternary stereocenters. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 3466-3470.	1.5	8
54	Rh-Catalyzed nitrene alkyne metathesis/formal C-N bond insertion cascade: synthesis of 3-iminoindolines. <i>Organic Chemistry Frontiers</i> , 2020, 7, 1327-1333.	2.3	15

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55	Copper-catalyzed formal [1 + 2 + 2]-annulation of alkyne-tethered diazoacetates and pyridines: access to polycyclic indolizines. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 1926-1932.	1.5	15
56	Highly Enantioselective Trapping of Carboxylic Oxonium Ylides with Imines for Direct Assembly of Enantioenriched 1 ³ -Butenolides. <i>CCS Chemistry</i> , 2020, 2, 432-439.	4.6	32
57	Synergistic Activation Strategy to Achieve Rh ₂ (II)-Catalyzed Asymmetric Cycloisomerization of 1, <i>n</i> -Enynes. <i>Chinese Journal of Organic Chemistry</i> , 2020, 40, 4370.	0.6	5
58	Synthesis of spiro[2,3-dihydrofuran-3,3-oxindole] derivatives via a multi-component cascade reaction of $\hat{1}\pm$ -diazo esters, water, isatins and malononitrile/ethyl cyanoacetate. <i>Green Chemistry</i> , 2019, 21, 4936-4940.	4.6	28
59	An Isoform-Selective PTP1B Inhibitor Derived from Nitrogen-Atom Augmentation of Radicol. <i>Biochemistry</i> , 2019, 58, 3225-3231.	1.2	9
60	A highly diastereoselective [5+1] annulation to 2,2,3-trisubstituted tetrahydroquinoxalines via intramolecular Mannich-type trapping of ammonium ylides. <i>Chemical Communications</i> , 2019, 55, 9809-9812.	2.2	13
61	Asymmetric Multicomponent Reactions for Efficient Construction of Homopropargyl Amine Carboxylic Esters. <i>Organic Letters</i> , 2019, 21, 5737-5741.	2.4	35
62	Discovery of Novel Isothiazole, 1,2,3-Thiadiazole, and Thiazole-Based Cinnamamides as Fungicidal Candidates. <i>Journal of Agricultural and Food Chemistry</i> , 2019, 67, 12357-12365.	2.4	35
63	Zinc-Catalyzed Alkyne-Carbonyl Metathesis of Ynamides with Isatins: Stereoselective Access to Fully Substituted Alkenes. <i>Journal of Organic Chemistry</i> , 2019, 84, 15331-15342.	1.7	24
64	Gold(I)-Catalyzed Aromatization: Expedient Synthesis of Polyfunctionalized Naphthalenes. <i>IScience</i> , 2019, 21, 499-508.	1.9	19
65	Divergent Construction of Macrocyclic Alkynes via Catalytic Metal Carbene C(sp ²)-H Insertion and the Buchner Reaction. <i>ACS Catalysis</i> , 2019, 9, 10773-10779.	5.5	20
66	Catalytic asymmetric synthesis of 2,5-dihydrofurans using synergistic bifunctional Ag catalysis. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 8737-8744.	1.5	13
67	Selective Vinylogous Reactivity of Carbene Intermediate in Gold-Catalyzed Alkyne Carbocyclization: Synthesis of Indenols. <i>ACS Catalysis</i> , 2019, 9, 2440-2447.	5.5	40
68	Synthesis and biological evaluation of novel potent FFA1 agonists containing 2,3-dihydrobenzo[b][1,4]dioxine. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2019, 29, 848-852.	1.0	3
69	Cu(I)-Catalyzed Three-Component Reaction of $\hat{1}\pm$ -Diazo Amide with Terminal Alkyne and Isatin Ketimine via Electrophilic Trapping of Active Alkynoate-Copper Intermediate. <i>Organic Letters</i> , 2019, 21, 4571-4574.	2.4	17
70	Rhodium-Catalyzed Formal C=O Insertion in Carbene/Alkyne Metathesis Reactions: Synthesis of 3-Substituted 3-H-Indol-3-ols. <i>Organic Letters</i> , 2019, 21, 4322-4326.	2.4	13
71	Copper-catalyzed [4+1]-annulation of 2-alkenylindoles with diazoacetates: a facile access to dihydrocyclopenta[b]indoles. <i>Chemical Communications</i> , 2019, 55, 6393-6396.	2.2	22
72	Gold-catalyzed dual annulation of azide-tethered alkynes with nitriles: expeditious synthesis of oxazolo[4,5-c]quinolines. <i>Organic Chemistry Frontiers</i> , 2019, 6, 2404-2409.	2.3	25

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73	Trapping of Zwitterionic Intermediates by Isatins and Imines: Synthesis of Benzoxazines Bearing a C4-Quaternary Stereocenter. <i>Organic Letters</i> , 2019, 21, 4014-4018.	2.4	16
74	Gold-Catalyzed Dual Annulation of Homopropargyl Alcohols with Nitrones: Synthesis of Tetrahydropyrano[4,3- <i>b</i>]indole Scaffolds. <i>Advanced Synthesis and Catalysis</i> , 2019, 361, 3569-3574.	2.1	10
75	Metal-Dependent Umpolung Reactivity of Carbenes Derived from Cyclopropenes. <i>Science</i> , 2019, 14, 292-300.	1.9	28
76	Rhodium-Catalyzed Nitrene/Alkyne Metathesis: An Enantioselective Process for the Synthesis of <i>N</i> -Heterocycles. <i>Organic Letters</i> , 2019, 21, 3328-3331.	2.4	19
77	Catalyst-Free <i>gem</i> -Difunctionalization of Fluoroalkyl-Substituted Diazo Compound with Diselenide or Disulfide and NFSI. <i>Organic Letters</i> , 2019, 21, 2101-2105.	2.4	36
78	Gold-Catalyzed 1,2-Acyloxy Migration/Coupling Cascade of Propargyl Diazoacetates: Synthesis of Isomycin Derivatives. <i>Organic Letters</i> , 2019, 21, 1813-1817.	2.4	19
79	Optimization of P2Y ₁₂ Antagonist Ethyl 6-(4-((Benzylsulfonyl)carbamoyl)piperidin-1-yl)-5-cyano-2-methylnicotinate (AZD1283) Led to the Discovery of an Oral Antiplatelet Agent with Improved Druglike Properties. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 3088-3106.	2.9	22
80	Rh(I)/Sc(OTf) ₃ -co-catalyzed Michael addition of ammonium ylide to (E)-1,4-enediones: synthesis of functionalized 1,4-diketones. <i>Molecular Diversity</i> , 2019, 23, 997-1010.	2.1	7
81	A rhodium-catalysed three-component reaction to access C1-substituted tetrahydroisoquinolines. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 9844-9848.	1.5	8
82	A gold-catalysed three-component reaction <i>via</i> trapping oxonium ylides with allenamides. <i>Chemical Communications</i> , 2019, 55, 12675-12678.	2.2	11
83	A sustainable catalytic enantioselective synthesis of norstatine derivatives. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 9792-9798.	1.5	4
84	Privilege-Structure-Oriented Three-Component Asymmetric Aminomethylation: Assembly of Chiral 3-Aminomethyl Indolones. <i>Organic Letters</i> , 2019, 21, 9878-9883.	2.4	23
85	Gold-Catalyzed Oxidative Cyclization/Aldol Addition of Homopropargyl Alcohols with Isatins. <i>Organic Letters</i> , 2019, 21, 369-372.	2.4	37
86	Rhodium(II)-Catalyzed Formal [4+1] Cycloaddition of Pyridotriazoles and Propargyl Alcohols: Synthesis of 2,5-Dihydrofurans. <i>Advanced Synthesis and Catalysis</i> , 2019, 361, 1265-1270.	2.1	22
87	Asymmetric Counter-Anion-Directed Aminomethylation: Synthesis of Chiral β -Amino Acids via Trapping of an Enol Intermediate. <i>Journal of the American Chemical Society</i> , 2019, 141, 1473-1478.	6.6	116
88	Synthesis of Paclitaxel Side Chain via Multi-Component Reaction and Its Application to the Synthesis of Paclitaxel Analogues. <i>Chinese Journal of Organic Chemistry</i> , 2019, 39, 377.	0.6	1
89	Formal Carbene Insertion into C=O or C=N Bond: An Efficient Strategy for the Synthesis of α -Substituted β -Hydroxy Chromene Derivatives from Chromene Acetals or Hemiaminal Ethers. <i>Advanced Synthesis and Catalysis</i> , 2018, 360, 2446-2452.	2.1	17
90	A convenient one-pot approach to Paclitaxel (Taxol) side chain via 1,3-dipolar cycloaddition of carbonyl ylides and <i>N</i> -benzoylbenzyl imines. <i>Tetrahedron Letters</i> , 2018, 59, 2141-2144.	0.7	6

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91	Enantioselective Trapping of Oxonium Ylides by 3-Hydroxyisoindolinones via a Formal S _N 1 Pathway for Construction of Contiguous Quaternary Stereocenters. <i>Organic Letters</i> , 2018, 20, 983-986.	2.4	54
92	Synthesis of β -Sulfur-Substituted Ketones via Rh(II)/Sc(III) a Cocatalyzed Three-Component Reaction of Diazo Compounds with Thiophenols and Enones. <i>Journal of Organic Chemistry</i> , 2018, 83, 4786-4791.	1.7	15
93	Intramolecular cycloaddition/rearrangement cascade from gold(III)-catalysed reactions of propargyl aryldiazoesters with cinnamyl imines. <i>Chemical Communications</i> , 2018, 54, 12828-12831.	2.2	7
94	Improved Synthesis of Yt-14, A Potent Antibiotic to Multidrug-Resistant Strains. <i>Journal of Chemical Research</i> , 2018, 42, 354-360.	0.6	0
95	Diastereoselective synthesis of isochromans via the Cu(II)-catalysed intramolecular Michael-type trapping of oxonium ylides. <i>Chemical Communications</i> , 2018, 54, 12650-12653.	2.2	17
96	Enantioselective Oxidative Cyclization/Mannich Addition Enabled by Gold(I)/Chiral Phosphoric Acid Cooperative Catalysis. <i>Angewandte Chemie</i> , 2018, 130, 17446-17450.	1.6	16
97	Enantioselective Oxidative Cyclization/Mannich Addition Enabled by Gold(I)/Chiral Phosphoric Acid Cooperative Catalysis. <i>Angewandte Chemie - International Edition</i> , 2018, 57, 17200-17204.	7.2	86
98	Design, Synthesis and Biological Evaluation of Isothiazole Based 1,2,4-triazole Derivatives. <i>Chinese Journal of Chemistry</i> , 2018, 36, 731-736.	2.6	11
99	Cu(I)-Catalyzed Three-Component Reaction of Diazo Compound with Terminal Alkyne and Nitrosobenzene for the Synthesis of Trifluoromethyl Dihydroisoxazoles. <i>Organic Letters</i> , 2018, 20, 4843-4847.	2.4	35
100	Formal carbene insertion into C O double bond: A facile approach to the synthesis of 2H-chromenes. <i>Tetrahedron</i> , 2018, 74, 4551-4557.	1.0	7
101	Rh(II)/Chiral Phosphoric Acid-Cocatalyzed Enantioselective Synthesis of Spirooxindole-Fused Thiaindanes. <i>Organic Letters</i> , 2018, 20, 4531-4535.	2.4	42
102	Gold(I)-Catalyzed and H ₂ O-Mediated Carbene Cascade Reaction of Propargyl Diazoacetates: Furan Synthesis and Mechanistic Insights. <i>Organic Letters</i> , 2018, 20, 5332-5335.	2.4	25
103	The First Kilogram Synthesis of Beclabuvir, an HCV NS5B Polymerase Inhibitor. <i>Organic Process Research and Development</i> , 2018, 22, 1393-1408.	1.3	37
104	Protein Arginine Methyltransferase 5 (PRMT5) as an Anticancer Target and Its Inhibitor Discovery. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 9429-9441.	2.9	75
105	Efficient and Facile Synthesis of Chiral Sulfonamides via Asymmetric Multicomponent Reactions. <i>Acta Chimica Sinica</i> , 2018, 76, 895.	0.5	6
106	An efficient stereoselective synthesis of six stereoisomers of 3, 4-diaminocyclohexane carboxamide as key intermediates for the synthesis of factor Xa inhibitors. <i>Tetrahedron</i> , 2017, 73, 1381-1388.	1.0	9
107	A Diastereoselective Multicomponent Reaction for Construction of Alkynylamide-Substituted β,β' -Diamino Acid Derivatives To Hunt Hits. <i>Journal of Organic Chemistry</i> , 2017, 82, 2862-2869.	1.7	12
108	A Rh(II)-catalyzed multicomponent reaction by trapping an β -amino enol intermediate in a traditional two-component reaction pathway. <i>Science Advances</i> , 2017, 3, e1602467.	4.7	42

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109	Enantioselective Formal [3 + 1 + 1] Cycloaddition Reaction by Ru(II)/Iminium Cocatalysis for Construction of Multisubstituted Pyrrolidines. <i>Organic Letters</i> , 2017, 19, 1290-1293.	2.4	14
110	Deactivating Influence of 3-O-Glycosyl Substituent on Anomeric Reactivity of Thiomannoside Observed in Oligomannoside Synthesis. <i>Journal of Organic Chemistry</i> , 2017, 82, 2599-2621.	1.7	9
111	Iron catalyzed efficient synthesis of poly-functional primary amines via the direct use of ammonia. <i>Chemical Communications</i> , 2017, 53, 2854-2857.	2.2	16
112	Trapping of Transient Zwitterionic Intermediates by N-Acylpyridinium Salts: A Palladium-Catalyzed Diastereoselective Three-Component Reaction. <i>Journal of Organic Chemistry</i> , 2017, 82, 5952-5958.	1.7	13
113	Enantioselective Multicomponent Reaction for Rapid Construction of 1,2,5-Triol Derivatives with Vicinal Chiral Centers. <i>Journal of Organic Chemistry</i> , 2017, 82, 5212-5221.	1.7	13
114	Synthesis and biological evaluation of 3-amino-3-hydroxymethyloxindoles as potential anti-cancer agents. <i>RSC Advances</i> , 2017, 7, 23265-23271.	1.7	10
115	A DFT calculation-inspired Rh-catalyzed reaction via suppression of β -H shift in β -alkyldiazoacetates. <i>Chemical Science</i> , 2017, 8, 4312-4317.	3.7	28
116	Synthesis and biological activity evaluation of dolastatin 10 analogues with N-terminal modifications. <i>Tetrahedron</i> , 2017, 73, 2255-2266.	1.0	16
117	Discovery of core-structurally novel PTP1B inhibitors with specific selectivity containing oxindole-fused spirotetrahydrofurochroman by one-pot reaction. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 1105-1108.	1.0	12
118	Discovery of Bisindole as a Novel Scaffold for Protein Tyrosine Phosphatase 1B Inhibitors. <i>Archiv Der Pharmazie</i> , 2017, 350, e1600173.	2.1	2
119	Asymmetric Multicomponent Reactions Based on Trapping of Active Intermediates. <i>Chemical Record</i> , 2017, 17, 739-753.	2.9	118
120	Diastereoselective Intramolecular Aldol-Type Trapping of Zwitterionic Intermediates by Ketones for the Synthesis of Spiro[chroman-4,3-oxindole] Derivatives. <i>Advanced Synthesis and Catalysis</i> , 2017, 359, 58-63.	2.1	23
121	Structure-based design and synthesis of imidazo[1,2-a]pyridine derivatives as novel and potent Nek2 inhibitors with <i>in vitro</i> and <i>in vivo</i> antitumor activities. <i>European Journal of Medicinal Chemistry</i> , 2017, 126, 1083-1106.	2.6	41
122	Regio- and Diastereoselective Three-Component Reactions via Trapping of Ammonium Ylides with N-Alkylquinolinium Salts: Synthesis of Multisubstituted Tetra- and Dihydroquinoline Derivatives. <i>Organic Letters</i> , 2017, 19, 3783-3786.	2.4	44
123	Enantioselective trapping of oxonium ylide intermediates by N-benzhydryl- β -imino ester: Synthesis of β -tetrasubstituted β -amino acids. <i>Chinese Chemical Letters</i> , 2017, 28, 213-217.	4.8	12
124	Targeting NEK2 attenuates glioblastoma growth and radioresistance by destabilizing histone methyltransferase EZH2. <i>Journal of Clinical Investigation</i> , 2017, 127, 3075-3089.	3.9	86
125	Recent Advances in Asymmetric Metal-Catalyzed Carbene Transfer from Diazo Compounds Toward Molecular Complexity. <i>Advances in Organometallic Chemistry</i> , 2016, 66, 33-91.	0.5	33
126	A transformation of cyclopropyl carbene: a highly enantioselective three-component reaction via trapping oxonium ylide by imine. <i>Tetrahedron</i> , 2016, 72, 2929-2934.	1.0	9

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127	Enantioselective oxidative functionalization of the C _{sp3} -H bond adjacent to a nitrogen atom for rapid access to β -hydroxy- α -amino acid derivatives. <i>Chemical Communications</i> , 2016, 52, 11831-11833.	2.2	18
128	A Rh-catalyzed three-component reaction of 3-diazooxindoles with N,N-disubstituted anilines and glyoxylates for the synthesis of 3-aryl-3-substituted oxindoles. <i>Organic and Biomolecular Chemistry</i> , 2016, 14, 10157-10160.	1.5	8
129	Pd(II)/ β -ICD-Cocatalyzed Asymmetric Route to Oxindole Scaffold via Cascade Reaction of Diazoacetamides and MBH-Carbonates. <i>Journal of Organic Chemistry</i> , 2016, 81, 8537-8543.	1.7	15
130	Double C-H Functionalization of Indoles via Three-Component Reactions/CuCl ₂ -Catalyzed Aerobic Dehydrogenative Coupling for the Synthesis of Polyfunctional Cyclopenta[<i>b</i>]indoles. <i>ACS Catalysis</i> , 2016, 6, 6146-6150.	5.5	43
131	Enantioselective Synthesis of β -Mercapto- α -amino Esters via Rh(II)/Chiral Phosphoric Acid-Cocatalyzed Three-Component Reaction of Diazo Compounds, Thiols, and Imines. <i>Organic Letters</i> , 2016, 18, 6086-6089.	2.4	33
132	New peptide deformylase inhibitors design, synthesis and pharmacokinetic assessment. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 3714-3718.	1.0	8
133	Catalyst-Free Halogenation of β -Diazocarbonyl Compounds with <i>N</i> -Halosuccinimides: Synthesis of 3-Halooxindoles or Vinyl Halides. <i>Organic Letters</i> , 2016, 18, 3134-3137.	2.4	37
134	Diastereoselective Three-Component Cascade Reaction to Construct Oxindole-Fused Spirotetrahydrofurochroman Scaffolds for Drug Discovery. <i>European Journal of Organic Chemistry</i> , 2016, 2016, 2671-2680.	1.2	23
135	An enantioselective three-component reaction of diazoacetates with indoles and enals by iridium/iminium co-catalysis. <i>Chemical Communications</i> , 2016, 52, 2736-2739.	2.2	42
136	Rh ₂ (OAc) ₄ and InCl ₃ co-catalyzed diastereoselective trapping of carbamate ammonium ylides with aldehydes for the synthesis of β -hydroxy- α -amino acid derivatives. <i>Tetrahedron</i> , 2016, 72, 579-583.	1.0	10
137	Preclinical activity of MBM-5 in gastrointestinal cancer by inhibiting NEK2 kinase activity. <i>Oncotarget</i> , 2016, 7, 79327-79341.	0.8	11
138	One-pot Enantioselective Multi-component Cascade Reactions for Synthesis of Chiral Functionalized Hydro-epoxyisochromenes: A Rapid Access to Molecular Complexity. <i>Acta Chimica Sinica</i> , 2016, 74, 54.	0.5	15
139	Small molecules enhance functional O-mannosylation of Alpha-dystroglycan. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 7661-7670.	1.4	15
140	Divergent Synthesis of Multisubstituted Tetrahydrofurans and Pyrrolidines via Intramolecular Aldol-type Trapping of Onium Ylide Intermediates. <i>Chemistry - A European Journal</i> , 2015, 21, 19202-19207.	1.7	42
141	Metal-Catalyzed Cross-Coupling of Terminal Alkynes with Different Carbene Precursors. <i>Current Organic Chemistry</i> , 2015, 20, 41-60.	0.9	20
142	Alkaloid Synthesis via Carbenoid Intermediates. <i>Current Organic Chemistry</i> , 2015, 20, 82-101.	0.9	9
143	Bond cleavage, fragment modification and reassembly in enantioselective three-component reactions. <i>Nature Communications</i> , 2015, 6, 5801.	5.8	86
144	Design, synthesis and biological evaluation of LpxC inhibitors with novel hydrophilic terminus. <i>Chinese Chemical Letters</i> , 2015, 26, 763-767.	4.8	4

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145	Synthesis of 3-(hydroxymethyl)-3-indol-3-ylloxindoles via Rh(II)-catalyzed three-component reaction of 3-diazoindoles, indoles and formalin. <i>Tetrahedron</i> , 2015, 71, 3597-3602.	1.0	18
146	Divergent synthesis of chiral heterocycles via sequencing of enantioselective three-component reactions and one-pot subsequent cyclization reactions. <i>Chemical Communications</i> , 2015, 51, 10612-10615.	2.2	37
147	Pd(II)-catalyzed formal [4+1] cycloaddition reactions of diazoacetates and aryl propargyl alcohols to form 2,5-dihydrofurans. <i>Chemical Communications</i> , 2015, 51, 15204-15207.	2.2	55
148	Catalytic Asymmetric Four-Component Reaction for the Rapid Construction of 3,3-Disubstituted 3-Indol-3-ylloxindoles. <i>Organic Letters</i> , 2015, 17, 4336-4339.	2.4	56
149	Trapping of Carboxylic Oxonium Ylides with N-Boc Imines for the Facile Synthesis of β^2 -Amino Alcohol Derivatives. <i>Synlett</i> , 2014, 25, 1745-1750.	1.0	10
150	Rh ₂ (OAc) ₄ and Chiral Phosphoric Acid Cocatalyzed Highly Diastereo- and Enantioselective Four-Component Reactions: Facile Synthesis of Chiral β^2 -Diamino Acid Derivatives. <i>Synthesis</i> , 2014, 46, 1348-1354.	1.2	10
151	Step-economy etherification of acylated alcohols. <i>Tetrahedron Letters</i> , 2014, 55, 6836-6838.	0.7	3
152	An Ylide Transformation of Rhodium(I) Carbene: Enantioselective Three-Component Reaction through Trapping of Rhodium(I)-Associated Ammonium Ylides by β^2 -Nitroacrylates. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 13136-13139.	7.2	90
153	Facile Synthesis of 3-Aryloxindoles via Brønsted Acid Catalyzed Friedel-Crafts Alkylation of Electron-Rich Arenes with 3-Diazoindoles. <i>Organic Letters</i> , 2014, 16, 2934-2937.	2.4	80
154	Design, Synthesis, and Structure-Activity Relationship Studies of Novel Fused Heterocycles-Linked Triazoles with Good Activity and Water Solubility. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 3687-3706.	2.9	100
155	Ruthenium(II)/Chiral Brønsted Acid Co-catalyzed Enantioselective Four-Component Reaction/Cascade Aza-Michael Addition for Efficient Construction of 1,3,4-Tetrasubstituted Tetrahydroisoquinolines. <i>Chemistry - A European Journal</i> , 2014, 20, 1505-1509.	1.7	43
156	Recent advances in metal carbenoid mediated nitrogen-containing zwitterionic intermediate trapping process. <i>Tetrahedron Letters</i> , 2014, 55, 777-783.	0.7	52
157	A Facile Access to Polyfunctional Oxygen-containing Heterocycles via Intramolecularly Formed Protic Oxonium Ylide Trapping Processes. <i>Chemistry - an Asian Journal</i> , 2014, 9, 117-120.	1.7	12
158	Highly diastereoselective intermolecular 1,3-dipolar cycloaddition reactions of carbonyl ylides with aldimines to afford sterically disfavored cis-oxazolidines. <i>Organic Chemistry Frontiers</i> , 2014, 1, 181.	2.3	15
159	Interception of benzyne with thioethers: a facile access to sulfur ylides under mild conditions. <i>RSC Advances</i> , 2014, 4, 7623-7626.	1.7	37
160	Efficient synthesis of chiral cyclic acetals by metal and Brønsted acid co-catalyzed enantioselective four-component cascade reactions. <i>Chemical Communications</i> , 2014, 50, 2196-2198.	2.2	27
161	Highly diastereoselective synthesis of 3-hydroxy-2,2,3-trisubstituted indolines via intramolecular trapping of ammonium ylides with ketones. <i>Chemical Communications</i> , 2014, 50, 951-953.	2.2	41
162	Rh(II)/Brønsted Acid Cocatalyzed Intramolecular Trapping of Ammonium Ylides with Enones: Diastereoselective Synthesis of 2,2,3-Trisubstituted Indolines. <i>Journal of Organic Chemistry</i> , 2014, 79, 8440-8446.	1.7	34

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163	Catalytic Asymmetric Functionalization of Aromatic C–H Bonds by Electrophilic Trapping of Metal-Carbene-Induced Zwitterionic Intermediates. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 13098-13101.	7.2	146
164	Synthesis, antibacterial activity, and biological evaluation of formyl hydroxyamino derivatives as novel potent peptide deformylase inhibitors against drug-resistant bacteria. <i>European Journal of Medicinal Chemistry</i> , 2014, 86, 133-152.	2.6	11
165	Therapeutic Melting Pot of Never in Mitosis Gene A Related Kinase 2 (Nek2): A Perspective on Nek2 as an Oncology Target and Recent Advancements in Nek2 Small Molecule Inhibition. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 5835-5844.	2.9	21
166	Three-component reactions based on trapping ammonium ylides with N-sulfonyl aldimines via cooperative catalysis of squaramides and Rh ₂ (OAc) ₄ . <i>Tetrahedron</i> , 2014, 70, 1471-1477.	1.0	10
167	Stereoselective Synthesis of a Sulfated Tetrasaccharide Corresponding to a Rare Sequence in the Galactofucan Isolated from <i>Sargassum polycystum</i> . <i>Journal of Organic Chemistry</i> , 2014, 79, 4718-4726.	1.7	29
168	Regio- and Diastereoselective Construction of β -Hydroxy- α -amino Ester Derivatives via 1,4-Conjugate Addition of β,β -Unsaturated α -Sulfonylimines. <i>Journal of Organic Chemistry</i> , 2014, 79, 4142-4147.	1.7	17
169	Synthesis of 3-Amino-3-hydroxymethyloxindoles and 3-Hydroxy-3-hydroxymethyloxindoles by Rh ₂ (OAc) ₄ -Catalyzed Three-Component Reactions of 3-Diazoindoles with Formaldehyde and Anilines or Water. <i>Journal of Organic Chemistry</i> , 2014, 79, 3908-3916.	1.7	27
170	Asymmetric N-H Insertion Reaction of α -Diazoesters and Carbamates Co-catalyzed by Dirhodium Acetate, Sulfonic Acid and Chiral Sulfonamide Urea. <i>Chinese Journal of Organic Chemistry</i> , 2014, 34, 107.	0.6	8
171	Diazo Compounds-Involved Catalytic Asymmetric Multicomponent Reactions. <i>Chinese Journal of Organic Chemistry</i> , 2014, 34, 1268.	0.6	20
172	Enantioselective trapping of phosphoramidate ammonium ylides with imino esters for synthesis of 2,3-diaminosuccinic acid derivatives. <i>Chemical Communications</i> , 2013, 49, 4238.	2.2	52
173	Recent Advances in the Use of Chiral Brønsted Acids as Cooperative Catalysts in Cascade and Multicomponent Reactions. <i>Asian Journal of Organic Chemistry</i> , 2013, 2, 824-836.	1.3	65
174	Efficient synthesis of β -aryl serine derivatives via three-component reactions of aryldiazoacetates, anilines and formaldehyde. <i>Tetrahedron</i> , 2013, 69, 11203-11208.	1.0	16
175	A stereoselective synthesis of fully substituted tetrahydrofurans through 1,3-dipolar cycloaddition with cinnamaldehydes: an easy access to chroman derivatives. <i>RSC Advances</i> , 2013, 3, 20065.	1.7	7
176	Highly Diastereoselective Multicomponent Cascade Reactions: Efficient Synthesis of Functionalized β -Indanols. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 1539-1542.	7.2	41
177	CuSO ₄ -catalyzed three-component reaction of β -diazo ester, water and isatin: an efficient approach to oxindole derivatives. <i>Green Chemistry</i> , 2013, 15, 620.	4.6	35
178	Efficient synthesis of dihydropyrido[4,3-d]pyrimidines by microwave-promoted three-component aza-Diels-Alder reaction. <i>Tetrahedron Letters</i> , 2013, 54, 267-271.	0.7	16
179	Rhodium-Catalyzed Chemo- and Regioselective Cross-Dimerization of Two Terminal Alkynes. <i>Organic Letters</i> , 2013, 15, 840-843.	2.4	63
180	Vinylogous Reactivity of Enol Diazoacetates with Donor-Acceptor Substituted Hydrazones. Synthesis of Substituted Pyrazole Derivatives. <i>Journal of Organic Chemistry</i> , 2013, 78, 1583-1588.	1.7	46

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181	A highly enantioselective four-component reaction for the efficient construction of chiral β^2 -hydroxy- β^1 -amino acid derivatives. <i>Chemical Communications</i> , 2013, 49, 2700.	2.2	39
182	Bicyclic Pyrazolidinone Derivatives from Diastereoselective Catalytic [3 + 3]-Cycloaddition Reactions of Enoldiazoacetates with Azomethine Imines. <i>Organic Letters</i> , 2013, 15, 1564-1567.	2.4	88
183	Novel Multicomponent Reactions via Trapping of Protic Onium Ylides with Electrophiles. <i>Accounts of Chemical Research</i> , 2013, 46, 2427-2440.	7.6	552
184	Highly Efficient Synthesis of Mixed 3,3'-Bisindoles via Rh(II)-Catalyzed Three-Component Reaction of 3-Diazoindoles with Indoles and Ethyl Glyoxylate. <i>Organic Letters</i> , 2013, 15, 3578-3581.	2.4	53
185	Diversity-Oriented Three-Component Reactions of Diazo Compounds with Anilines and 4-Oxoenoates. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 9289-9292.	7.2	71
186	Iron Porphyrin-Catalyzed Three-Component Reaction of Ethyl Diazoacetate with Aliphatic Amines and β^2,β^3 -Unsaturated β^1 -Keto Esters. <i>Organic Letters</i> , 2013, 15, 6140-6143.	2.4	49
187	Highly Enantioselective Three-Component Reactions of tert-Butyl Diazoacetate with Arylamines and Imines: An Efficient Synthesis of β^1,β^2 -Bis(arylamino) Acid Derivatives. <i>Synthesis</i> , 2013, 45, 452-458.	1.2	11
188	Synthesis of Novel and Complex Tetrahydroquinazolinone and Dihydro- $\hat{\text{A}}$ pyrido[2,3-d]pyrimidine Derivatives via a One-Pot [4+2]-Cycloaddition Strategy. <i>Synlett</i> , 2013, 24, 471-474.	1.0	5
189	Enantioselective Palladium(II) Phosphate Catalyzed Three-Component Reactions of Pyrrole, Diazoesters, and Imines. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 13356-13360.	7.2	152
190	Rhodium(ii) catalyzed diastereoselective reactions of diazoacetamides with isatins: an efficient approach to 3-hydroxy-3,3'-bioxindoles. <i>Organic and Biomolecular Chemistry</i> , 2012, 10, 8808.	1.5	27
191	Highly enantioselective trapping of zwitterionic intermediates by imines. <i>Nature Chemistry</i> , 2012, 4, 733-738.	6.6	274
192	Efficient synthesis of oxazoles by dirhodium(ii)-catalyzed reactions of styryl diazoacetate with oximes. <i>Chemical Communications</i> , 2012, 48, 11522.	2.2	33
193	A highly diastereoselective three-component tandem 1,4-conjugated addition-cyclization reaction to multisubstituted pyrrolidines. <i>Organic and Biomolecular Chemistry</i> , 2012, 10, 2133.	1.5	27
194	Rhodium(II)- and Copper(II)-Catalyzed Reactions of Enol Diazoacetates with Nitrones: Metal Carbene versus Lewis Acid Directed Pathways. <i>Angewandte Chemie - International Edition</i> , 2012, 51, 5900-5903.	7.2	69
195	Practical and Scalable Synthesis of Ethyl (R)-Piperidine-3-acetate. <i>Synthetic Communications</i> , 2012, 42, 1137-1145.	1.1	4
196	InBr ₃ catalyzed three-component reactions of an aryldiazoacetate, an alcohol, and a carbonyl compound. <i>Tetrahedron Letters</i> , 2012, 53, 182-185.	0.7	9
197	A Novel Method for Synthesizing N-Alkoxycarbonyl Aryl β^1 -Amino Esters and Their Applications in Enantioselective Transformations. <i>Advanced Synthesis and Catalysis</i> , 2012, 354, 301-307.	2.1	57
198	Asymmetric C-H Functionalization of Indoles via Enantioselective Protonation. <i>Acta Chimica Sinica</i> , 2012, 70, 2484.	0.5	43

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199	Diastereoselective three-component reactions of aryldiazoacetates with alcohols/water and alkynals: application to substituted enelactones. <i>Organic and Biomolecular Chemistry</i> , 2011, 9, 3839.	1.5	17
200	Cooperative catalysis in highly enantioselective Mannich-type three-component reaction of a diazoacetophenone with an alcohol and an imine. <i>Chemical Communications</i> , 2011, 47, 797-799.	2.2	65
201	Diastereoselectivity Switch in Cooperatively Catalyzed Three-Component Reactions of an Aryldiazoacetate, an Alcohol, and a β,β -Unsaturated α -Keto Ester. <i>Journal of Organic Chemistry</i> , 2011, 76, 5821-5824.	1.7	25
202	Diastereoselectively Switchable Enantioselective Trapping of Carbamate Ammonium Ylides with Imines. <i>Journal of the American Chemical Society</i> , 2011, 133, 8428-8431.	6.6	215
203	Dual Catalysis in Highly Enantioselective Multicomponent Reaction with Water: An Efficient Approach to Chiral β -Amino α -Hydroxy Acid Derivatives. <i>ChemCatChem</i> , 2011, 3, 653-656.	1.8	31
204	Green chemistry approaches to the regioselective synthesis of spiro heterobicyclic rings using iodine as a new and efficient catalyst under solvent-free conditions. <i>Molecular Diversity</i> , 2011, 15, 257-261.	2.1	27
205	Highly Regioselective, Three-Component Reactions of Diazoacetates with Anilines and β,β -Unsaturated α -Keto Esters: 1,2-Addition versus 1,4-Addition. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 1113-1124.	1.2	25
206	Highly Enantioselective Catalytic Synthesis of Functionalized Chiral Diazoacetoacetates. <i>Angewandte Chemie - International Edition</i> , 2011, 50, 6392-6395.	7.2	55
207	Divergent Outcomes of Carbene Transfer Reactions from Dirhodium- and Copper-Based Catalysts Separately or in Combination. <i>Angewandte Chemie - International Edition</i> , 2011, 50, 11152-11155.	7.2	61
208	Design, synthesis and antibacterial activity of 3-methylenepyrrolidine formyl hydroxyamino derivatives as novel peptide deformylase inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 1060-1063.	1.0	18
209	3-Diazo-N-[(2S)-1-hydroxypropan-2-yl]-2-oxopropanamide. <i>Acta Crystallographica Section E: Structure Reports Online</i> , 2011, 67, o1192-o1192.	0.2	0
210	Trapping of Oxonium Ylides with Michael Acceptors: Highly Diastereoselective Three-Component Reactions of Diazo Compounds with Alcohols and Benzylidene Meldrum's Acids/4-Oxo-enoates. <i>Synlett</i> , 2011, 2011, 1717-1722.	1.0	5
211	1,2-Addition of Phenylacetylene to Aldimines Catalyzed by $\text{InCl}_3/\text{CuCl}$ in Water under Barbier Conditions. <i>Synlett</i> , 2011, 2011, 627-630.	1.0	13
212	(R)-3,3'-BIS(9-PHENANTHRYL)-1,1'-BINAPHTHALENE-2,2'-DIYL HYDROGEN PHOSPHATE. <i>Organic Syntheses</i> , 2011, 88, 406.	1.0	6
213	ENANTIOSELECTIVE THREE-COMPONENT REACTION FOR THE PREPARATION OF β -AMINO- α -HYDROXY ESTERS. <i>Organic Syntheses</i> , 2011, 88, 418.	1.0	2
214	Cooperative Catalysis in Multicomponent Reactions: Highly Enantioselective Synthesis of β -Hydroxyketones with a Quaternary Carbon Stereocenter. <i>Angewandte Chemie - International Edition</i> , 2010, 49, 2190-2192.	7.2	127
215	Design, synthesis, and antibacterial activity of 2,5-dihydropyrrole formyl hydroxyamino derivatives as novel peptide deformylase inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 3592-3595.	1.0	21
216	Rh(II) Catalyzed Three-Component Reactions of Diazoacetates with Benzenemethanol and Indane-1, 2, 3-Triones. <i>Letters in Organic Chemistry</i> , 2010, 7, 106-109.	0.2	7

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217	A Strategy to Synthesize Taxol Side Chain and (â ²)-epi<i>/i> Cytosazone via Chiral Brønsted Acid-Rh₂(OAc)₄ Co-catalyzed Enantioselective Three-Component Reactions. <i>Journal of Organic Chemistry</i> , 2010, 75, 7483-7486.	1.7	82
218	Copper(ii)-catalyzed highly diastereoselective three-component reactions of aryl diazoacetates with alcohols and chalcones: an easy access to furan derivatives. <i>Chemical Communications</i> , 2010, 46, 2865.	2.2	41
219	Rh₂(OAc)₄-AgOTf Cooperative Catalysis in Cyclization/Three-Component Reactions for Concise Synthesis of 1,2-Dihydroisoquinolines. <i>Organic Letters</i> , 2010, 12, 652-655.	2.4	54
220	Trapping of an Ammonium Ylide with Activated Ketones: Synthesis of Î ² -Hydroxy-Î±-Amino Esters with Adjacent Quaternary Stereocenters. <i>Synlett</i> , 2009, 2009, 2109-2114.	1.0	3
221	Novel C-S Bond Formation Through Î±-Elimination of tert-Butyl Sulfoxonium Ylides: A Facile Approach to Chiral Sulfoxides. <i>Synlett</i> , 2009, 2009, 2183-2187.	1.0	3
222	Highly Diastereoselective Synthesis of Fully Substituted Tetrahydrofurans by a One-pot Cascade Reaction of Aryldiazoacetates with Allyl Alcohols. <i>Chemistry - A European Journal</i> , 2009, 15, 12604-12607.	1.7	28
223	Rhodium(II) catalyzed multi-component reactions of aryl diazoacetates with titanium(IV) isopropoxide and imines. <i>Tetrahedron</i> , 2009, 65, 8277-8282.	1.0	7
224	Rh(II) and Zn(II) co-catalyzed multi-component reaction for the synthesis of vicinal diols. <i>Chinese Chemical Letters</i> , 2009, 20, 1299-1302.	4.8	6
225	One-pot three-component tandem reaction of diazo compounds with anilines and unsaturated ketoesters: a novel synthesis of 2,3-dihydropyrrole derivatives. <i>Chemical Communications</i> , 2009, , 1362.	2.2	52
226	Component match in rhodium catalyzed three-component reactions of ethyl diazoacetate, H ₂ O and aryl imines: a highly diastereoselective one-step synthesis of Î ² -aryl isoserine derivatives. <i>Organic and Biomolecular Chemistry</i> , 2009, 7, 5028.	1.5	27
227	Copper(II) hexafluorophosphate: a dual functional catalyst for three-component reactions of methyl phenyldiazoacetate with alcohols and aldehydes or Î±-ketoesters. <i>Tetrahedron Letters</i> , 2008, 49, 6862-6865.	0.7	26
228	Catalytic Enantioselective Trapping of an Alcoholic Oxonium Ylide with Aldehydes: Rh^{II}/Zr^{IV}-Catalyzed Three-Component Reactions of Aryl Diazoacetates, Benzyl Alcohol, and Aldehydes. <i>Angewandte Chemie - International Edition</i> , 2008, 47, 6647-6649.	7.2	83
229	Cooperative Catalysis with Chiral Brønsted Acid-Rh₂(OAc)₄: Highly Enantioselective Three-Component Reactions of Diazo Compounds with Alcohols and Imines. <i>Journal of the American Chemical Society</i> , 2008, 130, 7782-7783.	6.6	349
230	Selectivity control in enantioselective four-component reactions of aryl diazoacetates with alcohols, aldehydes and amines: an efficient approach to synthesizing chiral Î ² -amino-Î±-hydroxyesters. <i>Chemical Communications</i> , 2008, , 6564.	2.2	93
231	Rhodium-Catalyzed Reaction of Diazoacetates, Thiols and Azodicarboxylates: An Unusual 1,2-Aza Shift from a Sulfonyl Ylide. <i>Synlett</i> , 2007, 2007, 1314-1316.	1.0	2
232	Trapping of Oxonium Ylide with Isatins: Efficient and Stereoselective Construction of Adjacent Quaternary Carbon Centers. <i>Organic Letters</i> , 2007, 9, 4721-4723.	2.4	72
233	Efficient Trapping of Oxonium Ylides with Imines: A Highly Diastereoselective Three-Component Reaction for the Synthesis of Î ² -Amino-Î±-hydroxyesters with Quaternary Stereocenters. <i>Angewandte Chemie - International Edition</i> , 2007, 46, 1337-1339.	7.2	104
234	An efficient route for the construction of cyclopenta[b]quinoline derivatives via intramolecular cyclopropanation. <i>Tetrahedron</i> , 2007, 63, 11850-11855.	1.0	8

#	ARTICLE	IF	CITATIONS
235	Regioselectivity in Lewis acids catalyzed C-H (O, S, N) insertions of methyl styryldiazoacetate with benzyl alcohol, benzyl thiol, and aniline. <i>Tetrahedron Letters</i> , 2007, 48, 3975-3977.	0.7	65
236	Facile Synthesis of Aryl α -Keto Esters via the Reaction of Aryl Diazoacetate with H ₂ O and DEAD. <i>Synlett</i> , 2006, 2006, 2486-2488.	1.0	2
237	Dirhodium catalyzed intramolecular enantioselective C-H insertion reaction of N-cumyl-N-(2-p-anisylethyl)diazoacetamide: synthesis of (α)-Rolipram. <i>Tetrahedron: Asymmetry</i> , 2005, 16, 1693-1698.	1.8	27
238	The synthesis of baclofen and GABOB via Rh(II) catalyzed intramolecular C-H insertion of α -diazoacetamides. <i>Tetrahedron</i> , 2005, 61, 1579-1586.	1.0	35
239	Rhodium-Catalyzed, Three-Component Reaction of Diazo Compounds with Amines and Azodicarboxylates. <i>Advanced Synthesis and Catalysis</i> , 2005, 347, 531-534.	2.1	50
240	Novel C-C Bond Formation Through Addition of Ammonium Ylides to Arylaldehydes: A Facile Approach to α -Aryl- β -hydroxy γ -Amino Acid Frameworks.. <i>ChemInform</i> , 2005, 36, no.	0.1	0
241	Stereoselective Synthesis of Bicyclic Pyrrolidines by a Rhodium-Catalyzed Cascade Process.. <i>ChemInform</i> , 2005, 36, no.	0.1	0
242	Three-Component Reaction of Aryl Diazoacetates, Alcohols, and Aldehydes (or Imines): Evidence of Alcoholic Oxonium Ylide Intermediates.. <i>ChemInform</i> , 2005, 36, no.	0.1	0
243	The Synthesis of Baclofen and GABOB via Rh(II) Catalyzed Intramolecular C-H Insertion of α -Diazoacetamides.. <i>ChemInform</i> , 2005, 36, no.	0.1	0
244	The Rhodium Catalyzed Three-Component Reaction of Diazoacetates, Titanium(IV) Alkoxides and Aldehydes.. <i>ChemInform</i> , 2005, 36, no.	0.1	0
245	Rhodium Acetate-Mediated Epoxidation from the Reaction of Diazobenzylphosphonates with Aldehydes and a Ketone: A Convenient and Highly Stereoselective Synthetic Method of cis-1,2-Epoxypropylphosphonates. <i>Synlett</i> , 2005, 2005, 1711-1715.	1.0	0
246	The rhodium catalyzed three-component reaction of diazoacetates, titanium(IV) alkoxides and aldehydes. <i>Chemical Communications</i> , 2005, , 2624.	2.2	38
247	Three-Component Reaction of Aryl Diazoacetates, Alcohols, and Aldehydes (or Imines): Evidence of Alcoholic Oxonium Ylide Intermediates. <i>Organic Letters</i> , 2005, 7, 83-86.	2.4	108
248	Cumyl: A Better N-Protecting Group of α -Diazo Acetamides for Intramolecular C-H Insertion Reaction and its Application in the Synthesis of Pregabalin and 3-Benzoyloxy Pyrrolidine. <i>Synlett</i> , 2004, 2004, 1763-1764.	1.0	1
249	Influence of the Diene in the Hetero-Diels-Alder Reaction Catalyzed by Dirhodium(II) Carboxamides. <i>Synlett</i> , 2004, 2004, 2425-2428.	1.0	6
250	DNA binding ligands targeting drug-resistant Gram-positive bacteria. Part 1: Internal benzimidazole derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004, 14, 1253-1257.	1.0	119
251	Stereoselective Synthesis of Bicyclic Pyrrolidines by a Rhodium-Catalyzed Cascade Process. <i>Angewandte Chemie - International Edition</i> , 2004, 43, 6713-6716.	7.2	30
252	A Novel Three-Component Reaction Catalyzed by Dirhodium(II) Acetate: Decomposition of Phenyl diazoacetate with Arylamine and Imine for Highly Diastereoselective Synthesis of 1,2-Diamines.. <i>ChemInform</i> , 2004, 35, no.	0.1	0

#	ARTICLE	IF	CITATIONS
253	A Facile Three-Component One-Pot Synthesis of Structurally Constrained Tetrahydrofurans that Are t-RNA Synthetase Inhibitor Analogues.. ChemInform, 2004, 35, no.	0.1	0
254	Highly Chemoselective 2,4,5-Triaryl-1,3-dioxolane Formation from Intermolecular 1,3-Dipolar Addition of Carbonyl Ylide with Aryl Aldehydes.. ChemInform, 2004, 35, no.	0.1	0
255	Dirhodium-catalyzed enantioselective C-H insertion of N-(2-benzyloxyethyl)-N-(tert-butyl)diazoacetamide and its application for the synthesis of chiral GABOB. Chirality, 2004, 16, 516-519.	1.3	8
256	DNA binding ligands with in vivo efficacy in murine models of bacterial infection: optimization of internal aromatic amino acids. Bioorganic and Medicinal Chemistry Letters, 2004, 14, 2067-2072.	1.0	21
257	DNA Binding Ligands with Improved in Vitro and in Vivo Potency against Drug-Resistant Staphylococcus aureus. Journal of Medicinal Chemistry, 2004, 47, 4352-4355.	2.9	27
258	Novel C=C bond formation through addition of ammonium ylides to arylaldehydes: a facile approach to β -aryl- β -hydroxy α -amino acid frameworks. Chemical Communications, 2004, , 2486-2487.	2.2	53
259	A Facile Three-Component One-Pot Synthesis of Structurally Constrained Tetrahydrofurans That Are t-RNA Synthetase Inhibitor Analogues. Journal of Organic Chemistry, 2004, 69, 4856-4859.	1.7	50
260	Highly Chemoselective 2,4,5-Triaryl-1,3-dioxolane Formation from Intermolecular 1,3-Dipolar Addition of Carbonyl Ylide with Aryl Aldehydes. Organic Letters, 2004, 6, 3071-3074.	2.4	57
261	Influences of Catalyst Configuration and Catalyst Loading on Selectivities in Reactions of Diazoacetamides. Barrier to Equilibrium between Diastereomeric Conformations. Organic Letters, 2003, 5, 407-410.	2.4	34
262	Comparative enantiocontrol with allyl phenyldiazoacetates in asymmetric catalytic intramolecular cyclopropanation. Chirality, 2003, 15, 369-373.	1.3	14
263	DNA Binding Ligands Targeting Drug-Resistant Bacteria: Structure, Activity, and Pharmacology. Journal of Medicinal Chemistry, 2003, 46, 3914-3929.	2.9	67
264	Highly Selective Catalyst-Directed Pathways to Dihydropyrroles from Vinyl diazoacetates and Imines. Journal of the American Chemical Society, 2003, 125, 4692-4693.	6.6	126
265	A Novel Three-Component Reaction Catalyzed by Dirhodium(II) Acetate: Decomposition of Phenyldiazoacetate with Arylamine and Imine for Highly Diastereoselective Synthesis of 1,2-Diamines. Organic Letters, 2003, 5, 3923-3926.	2.4	94
266	A Concise Synthesis of Gabapentin via Intramolecular C-CH Insertion Reaction. Synlett, 2003, 2003, 1965-1966.	1.0	2
267	Enantioselectivity for catalytic cyclopropanation with diazomalones. Arkivoc, 2003, 2003, 15-22.	0.3	31
268	In Search of High Stereocontrol for the Construction of cis-Disubstituted Cyclopropane Compounds. Total Synthesis of a Cyclopropane-Configured Urea-PETT Analogue That Is a HIV-1 Reverse Transcriptase Inhibitor. Organic Letters, 2002, 4, 901-904.	2.4	51
269	Total Synthesis of (S)-(+)-Imperanene. Effective Use of Regio- and Enantioselective Intramolecular Carbon-Hydrogen Insertion Reactions Catalyzed by Chiral Dirhodium(II) Carboxamidates. Journal of Organic Chemistry, 2002, 67, 2954-2959.	1.7	56
270	Enantioselective carbon-hydrogen insertion is an effective and efficient methodology for the synthesis of (r)-(-)-baclofen. Chirality, 2002, 14, 169-172.	1.3	38

#	ARTICLE	IF	CITATIONS
271	In Search of High Stereocontrol for the Construction of cis-Disubstituted Cyclopropane Compounds. Total Synthesis of a Cyclopropane-Configured Urea-PETT Analogue that Is a HIV-1 Reverse Transcriptase Inhibitor.. ChemInform, 2002, 33, 73-73.	0.1	0
272	Epoxides and Aziridines from Diazoacetates via Ylide Intermediates. Organic Letters, 2001, 3, 933-935.	2.4	162
273	Highly Stereoselective Syntheses of Five- and Seven-Membered Ring Heterocycles from Ylides Generated by Catalytic Reactions of Styryldiazoacetates with Aldehydes and Imines. Organic Letters, 2001, 3, 3741-3744.	2.4	74
274	A New Class of Chiral Lewis Acid Catalysts for Highly Enantioselective Hetero-Diels-Alder Reactions: Exceptionally High Turnover Numbers from Dirhodium(II) Carboxamidates. Journal of the American Chemical Society, 2001, 123, 5366-5367.	6.6	104
275	Reactivity Enhancement for Chiral Dirhodium(II) Tetrakis(Carboxamidates). Advanced Synthesis and Catalysis, 2001, 343, 112-117.	2.1	40
276	A New Enantioselective Synthesis of Milnacipran and an Analogue by Catalytic Asymmetric Cyclopropanation. Advanced Synthesis and Catalysis, 2001, 343, 299-302.	2.1	44
277	Macrocycle Formation from Catalytic Metal Carbene Transformations. Synlett, 2001, 2001, 1364-1370.	1.0	33
278	Enantiocontrol in Macrocycle Formation from Catalytic Metal Carbene Transformations. Chinese Journal of Chemistry, 2001, 19, 22-29.	2.6	4
279	Reactivities and selectivities in macrocyclic addition reactions with diazoacetates using copper(I) and rhodium(II) catalysts. Tetrahedron Letters, 2000, 41, 6265-6269.	0.7	33
280	Dirhodium(II) Tetrakis[methyl 2-oxaazetidone-4-carboxylate]: A Chiral Dirhodium(II) Carboxamidate of Exceptional Reactivity and Selectivity. Organic Letters, 2000, 2, 1145-1147.	2.4	142
281	Selectivity in Reactions of Allyl Diazoacetates as a Function of Catalyst and Ring Size from β^3 -Lactones to Macrocyclic Lactones. Journal of Organic Chemistry, 2000, 65, 8839-8847.	1.7	61
282	A New Approach to Macrocyclization via Alkene Formation in Catalytic Diazo Decomposition. Synthesis of Patulolides A and B. Organic Letters, 2000, 2, 1777-1779.	2.4	61
283	Optimization of enantiocontrol in cis-selective cyclopropanation reactions catalyzed by dirhodium(ii) tetrakis[alkyl 2-oxaazetidone-4(S)-carboxylates]. Chemical Communications, 2000, , 867-868.	2.2	42
284	Enantiocontrolled Macrocycle Formation by Catalytic Intramolecular Cyclopropanation. Journal of the American Chemical Society, 2000, 122, 5718-5728.	6.6	63
285	A highly effective rhodium spirocyclic phosphinite catalyst for the asymmetric hydrogenation of enamides. Tetrahedron Letters, 1999, 40, 973-976.	0.7	45
286	Highly Effective Soluble Polymer-Supported Catalysts for Asymmetric Hydrogenation. Journal of the American Chemical Society, 1999, 121, 7407-7408.	6.6	156
287	Catalytic Intramolecular Addition of Metal Carbenes to Remote Furans. Organic Letters, 1999, 1, 1327-1329.	2.4	35
288	Synthesis of a chiral pyridylphosphine ligand and a comparison of its rhodium complex with the structurally similar arylphosphine rhodium catalysts in the asymmetric hydrogenation of 2-(6-methoxy-2-naphthyl)propenoic acid. Tetrahedron: Asymmetry, 1998, 9, 3241-3246.	1.8	20

#	ARTICLE	IF	CITATIONS
289	Synthesis of (âˆ—)-(4R,5R)-4,5-bis[di-3â€²-(2â€²,6â€²-dimethoxypyridyl)phosphinomethyl]-2,2-dimethyl-1,3-dioxolane and its application in the Rh-catalyzed asymmetric hydrogenation reactions. <i>Tetrahedron: Asymmetry</i> , 1998, 9, 4183-4192.	1.8	27
290	Asymmetric Synthesis XXVIII: Novel Chiral Aminophosphine Phosphinite Ligand and Its Application in Homogenous Catalytic Asymmetric Hydrogenation of Dehydroamino Acid Derivatives. <i>Synlett</i> , 1998, 1998, 847-848.	1.0	22
291	Novel Spiro Phosphinite Ligands and Their Application in Homogeneous Catalytic Hydrogenation Reactions. <i>Journal of the American Chemical Society</i> , 1997, 119, 9570-9571.	6.6	205
292	Salen-Ti(OR) ₄ complex catalysed trimethylsilylcyanation of aldehydes. <i>Tetrahedron</i> , 1997, 53, 14327-14338.	1.0	74
293	Complete Diastereofacial Selective Cycloaddition of New Sulfinyl Dienophiles (Ss)-and (Rs)-(-)-Menthyl Î±-(2-Methoxyphenylsulfoxyl)acylate with Anthracene Under Chelation Control. <i>Synthetic Communications</i> , 1996, 26, 1867-1874.	1.1	7
294	Asymmetric Synthesis XXIV: Chiral Salen-Titanium Complexes-Efficient Catalysts for Asymmetric Trimethyl Silylcyanation of Benzaldehyde. <i>Synlett</i> , 1996, 1996, 337-338.	1.0	39
295	Synthesis of spiro[4.4]nonane-1,6-diols via the hydrogenation of spiro[4.4]nonane-1,6-dione: the profound effect of hydrogenating agents. <i>Tetrahedron: Asymmetry</i> , 1995, 6, 2953-2959.	1.8	23
296	Asymmetric Synthesis. XX. Asymmetric Oxidative Coupling Reaction of Imine Carbanions via (+)-Camphor Chiral Template. <i>Synthetic Communications</i> , 1994, 24, 3115-3122.	1.1	6
297	Diastereoselective alkylation of the (+)â€²ketopinic acid ketimine derived from benzylamine. <i>Chinese Journal of Chemistry</i> , 1990, 8, 542-548.	2.6	0