

Li-Xin Wang

List of Publications by Year in descending order

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2,287
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186265
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1909
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#	ARTICLE	IF	CITATIONS
1	New scaffold organocatalysts of chiral 3,2- ϵ^2 -pyrrolidiny spirooxindoles promoted enantioselective aldol condensation between isatins and acetone. <i>Tetrahedron Letters</i> , 2022, 97, 153780.	1.4	4
2	Enantioselective Organocatalyzed Mannich Reaction between Benzothiazolimines and β -Benzylidene Succinimides for the Preparation of Chiral Benzothiazol Succinimides. <i>Asian Journal of Organic Chemistry</i> , 2022, 11, .	2.7	3
3	Organocatalytic enantioselective aza-Friedel-Crafts reaction between benzothiazolimines and 2-naphthols for the preparation of chiral 2-aminobenzothiazolomethyl naphthols. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 7690-7694.	2.8	14
4	Organocatalytic enantioselective Diels-Alder reaction between hydroxymaleimides and <i>in situ</i> generated nitrosoalkenes for direct preparation of chiral hemiketals with 1,2-oxazine skeleton. <i>Organic Chemistry Frontiers</i> , 2021, 8, 6215-6219.	4.5	2
5	3-Amino Oxindole Schiff Base as Synthon for Enantioselective Preparation of Spiro[oxindole-3,2- ϵ^2 -pyrrol] from a Michael/Cyclization Reaction Catalyzed by a Bifunctional Cinchona. <i>Organic Letters</i> , 2021, 23, 2227-2231.	4.6	6
6	Organocatalyst-promoted Diastereoselective and Enantioselective Michael Addition/Hemiketalization Reaction between Hydroxymaleimide and Quinone. <i>Asian Journal of Organic Chemistry</i> , 2021, 10, 1713-1717.	2.7	3
7	Spiro Scaffold Chiral Organocatalyst of 3,2- ϵ^2 -Pyrrolidiny Spiro-oxindole Amine and Its Catalytic Evaluation in the Enantioselective Aldol Condensation between 3-(3-Hydroxy-1- <i>H</i> -pyrazol-1-yl)-Oxindole and Paraformaldehyde. <i>Journal of Organic Chemistry</i> , 2021, 86, 17371-17379.	3.2	6
8	A base-catalyzed domino reaction between isoindigos and β -alkylidene succinimides-convenient preparation of highly steric bispirooxindoles. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 9511-9515.	2.8	5
9	Organocatalytic and enantioselective [4+2] cyclization between hydroxymaleimides and <i>ortho</i> -hydroxyphenyl <i>para</i> -quinone methide-selective preparation of chiral hemiketals. <i>Chemical Communications</i> , 2020, 56, 14825-14828.	4.1	27
10	Organocatalytic Enantioselective Michael Addition between 3-(3-hydroxy-1- <i>H</i> -pyrazol-1-yl)Oxindole and β -Nitrostyrene for the Preparation of Chiral Disubstituted Oxindoles. <i>Journal of Organic Chemistry</i> , 2020, 85, 9290-9300.	3.2	11
11	Base Catalyzed Abnormal [3 + 2]-Cycloaddition between Isatin <i>N</i> , <i>N</i> - ϵ^2 -Cyclic Azomethine Imine 1,3-Dipole and 3-Methyleneoxindole for the One-Step Construction of Tetracyclic Bispirooxindoles. <i>Journal of Organic Chemistry</i> , 2020, 85, 3921-3928.	3.2	23
12	Substituted (E)-2-Methylene-3,4-cyclohexenones through Direct and Convenient Synthesis from Cyclohexenone-MBH Alcohol in the Presence of DMAP. <i>European Journal of Organic Chemistry</i> , 2019, 2019, 715-719.	2.4	3
13	Effective and diastereoselective preparation of dispiro[cyclopent-3- ϵ^2 -ene]bisoxindoles <i>via</i> novel [3 + 2] annulation of isoindigos and MBH carbonates. <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 1297-1304.	2.8	12
14	Isatin <i>N</i> , <i>N</i> - ϵ^2 -Cyclic Azomethine Imine 1,3-Dipole and Abnormal [3 + 2]-Cycloaddition with Maleimide in the Presence of 1,4-Diazabicyclo[2.2.2]octane. <i>Organic Letters</i> , 2017, 19, 646-649.	4.6	48
15	An efficient and enantioselective Michael addition of aromatic oximes to β -unsaturated aldehydes promoted by a chiral diamine catalyst derived from β - <i>phenyl</i> , β - <i>phenyl</i> prolinol. <i>Chirality</i> , 2017, 29, 369-375.	2.6	3
16	Organocatalytic Asymmetric Annulation between Hydroxymaleimides and Nitrosoarenes: Stereoselective Preparation of Chiral Quaternary <i>N</i> -Hydroxyindolines. <i>Organic Letters</i> , 2017, 19, 2805-2808.	4.6	27
17	Isatin <i>N</i> , <i>N</i> - ϵ^2 -Cyclic Azomethine Imine 1,3-Dipole and Base Catalyzed Michael Addition with β -Nitrostyrene <i>via</i> C3 Umpolung of Oxindole. <i>Organic Letters</i> , 2017, 19, 3051-3054.	4.6	35
18	Organocatalytic Enantioselective Michael/Cyclization Domino Reaction between 3-Amideoxindoles and β -Unsaturated Aldehydes: One-Pot Preparation of Chiral Spirocyclic Oxindole- β -lactams. <i>Journal of Organic Chemistry</i> , 2017, 82, 3908-3916.	3.2	29

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19	An enantioselective synthesis of spiro-oxindole-based 3,4-dihydropyrroles via a Michael/cyclization cascade of 3-aminooxindoles with 2-enoylpyridines. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 8518-8522.	2.8	25
20	Optimized Synthetic Route for Enantioselective Preparation of (S)-Metolachlor from Commercially Available (R)-Propylene Oxide. <i>Organic Process Research and Development</i> , 2017, 21, 1682-1688.	2.7	4
21	Enantioselective Nitroso Aldol Intramolecular Transesterification Cyclization Domino Reaction for Highly Effective Construction of Chiral Spirooxindoles. <i>European Journal of Organic Chemistry</i> , 2016, 2016, 5437-5444.	2.4	13
22	An unexpected metal-free DMAP catalyzed Michael addition-elimination domino reaction between 2-naphthols and bromomaleimides for the effective construction of 3-arylmaleimides. <i>Tetrahedron Letters</i> , 2016, 57, 1261-1264.	1.4	8
23	An Improved and Enantioselective Preparation of the Telaprevir Bicyclic [3.3.0] Proline Intermediate and Reuse of Unwanted Enantiomer. <i>Organic Process Research and Development</i> , 2016, 20, 320-324.	2.7	14
24	A cinchona alkaloid catalyzed enantioselective sulfa-Michael/aldol cascade reaction of isoindigos: construction of chiral bispirooxindole tetrahydrothiophenes with vicinal quaternary spirocenters. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 6371-6379.	2.8	56
25	An unprecedented base-promoted domino reaction of methyleneindolinones and N-tosyloxycarbamates for the construction of bispirooxindoles and spiroaziridine oxindoles. <i>Chemical Communications</i> , 2015, 51, 10726-10729.	4.1	37
26	Direct enantioselective amination of β -ketoester catalyzed by tertiary amine thiourea: a new approach to chiral β -hydroxy- β -amino acid. <i>Tetrahedron Letters</i> , 2015, 56, 4220-4223.	1.4	6
27	Asymmetric Synthesis of 3,3- α^2 -Spirooxindoles Fused with Cyclobutanes through Organocatalytic Formal [2 + 2] Cycloadditions under H-Bond-Directing Dienamine Activation. <i>Organic Letters</i> , 2014, 16, 6436-6439.	4.6	77
28	Efficient asymmetric Michael reaction of 2-oxindole-3-carboxylate esters with maleimides catalyzed by cinchonidine. <i>Tetrahedron</i> , 2014, 70, 3478-3484.	1.9	25
29	An organocatalytic domino Michael-alkylation reaction: highly enantioselective construction of spiro-cyclopentanoneoxindoles and tetrone acid scaffolds. <i>Chemical Communications</i> , 2014, 50, 14601-14604.	4.1	44
30	Organocatalytic direct asymmetric vinylogous Mannich reaction of β -butenolides with isatin-derived ketimines. <i>RSC Advances</i> , 2014, 4, 27286.	3.6	30
31	Organocatalytic asymmetric cascade Michael/hemiketalization/retro-aldol reaction of 3-acetyl-oxindole with β,β -unsaturated ketoesters catalyzed by bifunctional amino-squaramides. <i>Tetrahedron</i> , 2014, 70, 8665-8671.	1.9	16
32	A New Cyclization/Decarboxylation Reaction of Isatins with Acyl Chlorides for the Facile Synthesis of β -Alkenyl-Oxindoles. <i>Chinese Journal of Chemistry</i> , 2014, 32, 844-852.	4.9	10
33	An organocatalytic asymmetric sequential allylic alkylation-cyclization of Morita-Baylis-Hillman carbonates and 3-hydroxyoxindoles. <i>Chemical Communications</i> , 2013, 49, 9422.	4.1	68
34	A novel asymmetric organocatalytic Michael-aldol-dehydration domino reaction for the construction of spirocyclic benzofuranones. <i>Tetrahedron</i> , 2013, 69, 9303-9308.	1.9	12
35	An Improved and Economical Process for the Manufacture of the Key Intermediate of Aliskiren, a New Potent Renin Inhibitor. <i>Organic Process Research and Development</i> , 2013, 17, 1458-1462.	2.7	10
36	Highly enantioselective direct vinylogous Michael addition of β -substituted deconjugated butenolides to maleimides catalyzed by chiral squaramides. <i>RSC Advances</i> , 2013, 3, 16973.	3.6	34

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37	Enantioselective Diels-Alder reaction of anthrone and maleimide catalyzed by a simple chiral tertiary amine. <i>Tetrahedron</i> , 2013, 69, 1229-1233.	1.9	17
38	Chiral β -Arylethanamines: An Organocatalyst for the Enantioselective α -Amination of Branched Aldehydes. <i>European Journal of Organic Chemistry</i> , 2013, 2013, 2864-2868.	2.4	8
39	Asymmetric Michael/cyclization tandem reaction of 4-hydroxycoumarin with β -nitroalkenes catalyzed by chiral bifunctional thioureas. <i>Organic and Biomolecular Chemistry</i> , 2013, 11, 1286.	2.8	24
40	Construction of Quaternary Stereocenters: Asymmetric α -Amination of Branched Aldehydes Catalyzed by Monoimide Substituted Cyclohexane-1,2-Diamines. <i>Chirality</i> , 2013, 25, 668-672.	2.6	6
41	Asymmetric hydroxyamination of oxindoles catalyzed by chiral bifunctional tertiary aminethiourea: construction of 3-amino-2-oxindoles with quaternary stereocenters. <i>Organic and Biomolecular Chemistry</i> , 2012, 10, 236-239.	2.8	48
42	Organocatalytic Asymmetric Double Michael Reaction of Benzofuranone with Dienones to Construct Spirocyclic Benzofuranones. <i>Chinese Journal of Chemistry</i> , 2012, 30, 2703-2706.	4.9	11
43	Asymmetric Michael Addition of β -Substituted Isocynoacetates with Maleimides Catalyzed by Chiral Tertiary Amine Thiourea. <i>Journal of Organic Chemistry</i> , 2012, 77, 2947-2953.	3.2	85
44	Direct Asymmetric Vinylogous Mannich Reaction of 3,4-Dihalofuran-2(5H)-one with Aldimine Catalyzed by Quinine. <i>Journal of Organic Chemistry</i> , 2012, 77, 8338-8343.	3.2	24
45	Organocatalytic stereocontrolled synthesis of 3,3-dipyrrolidinyl spirooxindoles by [3+2] annulation of isocyanoesters with methyleneindolinones. <i>Chemical Communications</i> , 2012, 48, 5175.	4.1	123
46	Asymmetric Double Michael Reaction Catalyzed by Simple Primary Amine Catalysts: A Straightforward Approach to Construct Spirocyclic Oxindoles. <i>Chinese Journal of Chemistry</i> , 2012, 30, 1185-1188.	4.9	9
47	New 1,3-dipolar cycloaddition/dehydrogenation of azomethines ylides and azodicarboxylates: direct and effective construction of unsaturated 1,2,4-triazolines. <i>Tetrahedron Letters</i> , 2012, 53, 2985-2988.	1.4	5
48	Highly enantioselective aldol reaction of acetone with β,β -unsaturated β -keto esters promoted by simple chiral primary-tertiary diamine catalysts. <i>Organic and Biomolecular Chemistry</i> , 2011, 9, 4774.	2.8	24
49	A highly asymmetric direct aldol reaction catalyzed by chiral proline amide-thiourea bifunctional catalysts. <i>Canadian Journal of Chemistry</i> , 2011, 89, 1312-1318.	1.1	10
50	Enantioselective α -Amination of Branched Aldehydes Promoted by Simple Chiral Primary Amino Acids. <i>Journal of Organic Chemistry</i> , 2011, 76, 4661-4664.	3.2	43
51	A highly organocatalytic stereoselective double Michael reaction: efficient construction of optically enriched spirocyclic oxindoles. <i>Chemical Communications</i> , 2011, 47, 5593-5595.	4.1	107
52	Metal-Free Asymmetric 1,3-Dipolar Cycloaddition of β -Aryl maleimides to Azomethine Ylides Catalyzed by Chiral Tertiary Amine Thiourea. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 4472-4478.	2.4	42
53	Highly effective and enantioselective Phospho-Aldol reaction of diphenyl phosphite with N-alkylated isatins catalyzed by quinine. <i>Tetrahedron Letters</i> , 2011, 52, 1157-1160.	1.4	38
54	Highly effective and enantioselective Michael addition of 4-hydroxycoumarin to β,β -unsaturated ketones promoted by simple chiral primary amine thiourea bifunctional catalysts. <i>Tetrahedron Letters</i> , 2011, 52, 1566-1568.	1.4	43

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55	Facile Catalyst-Free Allylation of Isatins Under Mild Conditions in Dimethylformamide. <i>Letters in Organic Chemistry</i> , 2011, 8, 352-357.	0.5	2
56	Novel Preparation of H1 Receptor Antagonist Fexofenadine. <i>Organic Process Research and Development</i> , 2010, 14, 1464-1468.	2.7	6
57	Effective construction of quaternary stereocenters by highly enantioselective $\hat{1}\pm$ -amination of branched aldehydes. <i>Organic and Biomolecular Chemistry</i> , 2010, 8, 4524.	2.8	44
58	Noyori's Ts $\hat{1}$ DPEN Ligand: Simple yet Effective Catalyst for the Highly Enantioselective Michael Addition of Acetone to Nitroalkenes. <i>European Journal of Organic Chemistry</i> , 2010, 2010, 1849-1853.	2.4	55
59	Chiral primary amine thiourea promoted highly enantioselective Michael reactions of isobutylaldehyde with maleimides. <i>Tetrahedron</i> , 2010, 66, 8928-8932.	1.9	70
60	Highly asymmetric Michael additions of $\hat{1}\pm, \hat{1}\pm$ -disubstituted aldehydes to $\hat{1}^2$ -nitroalkenes promoted by chiral pyrrolidine $\hat{1}$ thiourea bifunctional catalysts. <i>Tetrahedron Letters</i> , 2010, 51, 2803-2805.	1.4	56
61	Highly effective and enantioselective $\hat{1}\pm$ -amination of aldehydes promoted by chiral proline amide $\hat{1}$ thiourea bifunctional catalysts. <i>Tetrahedron Letters</i> , 2010, 51, 4870-4873.	1.4	30
62	Asymmetric Michael Addition of Aromatic Ketones to Nitroolefins Catalyzed by Simple Chiral Bifunctional Primary Amine-Thioureas. <i>Letters in Organic Chemistry</i> , 2010, 7, 367-372.	0.5	17
63	Highly organocatalytic asymmetric Michael $\hat{1}$ ketone aldol $\hat{1}$ dehydration domino reaction: straightforward approach to construct six-membered spirocyclic oxindoles. <i>Chemical Communications</i> , 2010, 46, 8064.	4.1	117
64	Effective asymmetric Michael addition of acetone to nitroalkenes promoted by chiral proline amide-thiourea bifunctional catalysts. <i>Arkivoc</i> , 2010, 2010, 340-351.	0.5	18
65	Improved Preparation of Tyramine by Curtius Rearrangement. <i>Chinese Journal of Chemistry</i> , 2009, 27, 433-436.	4.9	8
66	Preparation of key intermediates of adrenergic receptor agonists: Highly enantioselective production of (R)- $\hat{1}\pm$ -halohydrins with <i>Saccharomyces cerevisiae</i> CGMCC 2.396. <i>Journal of Molecular Catalysis B: Enzymatic</i> , 2009, 57, 1-5.	1.8	32
67	Preparation the Key Intermediate of Angiotensin $\hat{1}$ Converting Enzyme (ACE) Inhibitors: High Enantioselective Production of Ethyl (<i>R</i>)- $\hat{1}\pm$ -Hydroxy $\hat{1}\pm$ -Phenylbutyrate with <i>Candida boidinii</i> CIOC21. <i>Advanced Synthesis and Catalysis</i> , 2008, 350, 426-430.	4.3	42
68	Synthetic Improvements in the Preparation of Clopidogrel. <i>Organic Process Research and Development</i> , 2007, 11, 487-489.	2.7	64
69	Enantiocomplementary preparation of (S)- and (R)-mandelic acid derivatives via $\hat{1}\pm$ -hydroxylation of 2-arylacetic acid derivatives and reduction of $\hat{1}\pm$ -ketoester using microbial whole cells. <i>Tetrahedron: Asymmetry</i> , 2007, 18, 2537-2540.	1.8	25
70	Synthesis of Spiro Diphosphines and Their Application in Asymmetric Hydrogenation of Ketones. <i>Journal of the American Chemical Society</i> , 2003, 125, 4404-4405.	13.7	275