

Renhua Fan

List of Publications by Year in descending order

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236833

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docs citations

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times ranked

1532
citing authors

#	ARTICLE	IF	CITATIONS
1	\hat{I}^+ and \hat{I}^\pm SP ³ C-H Bond Oxidation of Sulfonamides with PhI(OAc) ₂ /I ₂ under Metal-Free Conditions. <i>Journal of Organic Chemistry</i> , 2007, 72, 8994-8997.	1.7	147
2	Transition-Metal-Free Intermolecular Amination of sp ³ C-H Bonds with Sulfonamides. <i>Organic Letters</i> , 2009, 11, 1425-1428.	2.4	127
3	Solvent-Controlled Oxidative Cyclization for Divergent Synthesis of Highly Functionalized Oxetanes and Cyclopropanes. <i>Organic Letters</i> , 2009, 11, 3156-3159.	2.4	76
4	Gold(III) Chloride/Silver Triflate: A Highly Efficient Catalyst for Ring-Opening Reaction of Aziridines with Electron-Rich Arenes. <i>Advanced Synthesis and Catalysis</i> , 2007, 349, 2151-2155.	2.1	66
5	Aqueous Iodine(III)-Mediated Stereoselective Oxidative Cyclization for the Synthesis of Functionalized Fused Dihydrofuran Derivatives. <i>Journal of Organic Chemistry</i> , 2010, 75, 1760-1763.	1.7	58
6	Efficient Stereoselective Synthesis of Nitrocyclopropanes by the Oxidative Cyclization of Michael Adducts of Nitroolefins with Activated Methylene Compounds. <i>Advanced Synthesis and Catalysis</i> , 2008, 350, 2488-2492.	2.1	57
7	Construction of 3-oxyindoles via hypervalent iodine mediated tandem cyclization-acetoxylation of o-acyl anilines. <i>Chemical Communications</i> , 2010, 46, 6834.	2.2	53
8	PhI(OAc) ₂ induced intramolecular oxidative bromocyclization of homoallylic sulfonamides with KBr as the bromine source. <i>Tetrahedron Letters</i> , 2007, 48, 7444-7447.	0.7	49
9	Dearomatization Strategy and Palladium-Catalyzed Domino Reaction: Construction of Azepino[5,4,3-cd]indoles from 2-Alkynylanilines. <i>Organic Letters</i> , 2014, 16, 816-819.	2.4	49
10	Stereoselective Construction of Highly Functionalized Azetidines via a [2 + 2]-Cycloaddition. <i>Organic Letters</i> , 2010, 12, 2802-2805.	2.4	47
11	Recent Advances in Phenol Dearomatization and Its Application in Complex Syntheses. <i>Synthesis</i> , 2012, 45, 1-16.	1.2	47
12	Iodine(III)-Mediated Tandem Acetoxylation-Cyclization of o-Acyl Phenols for the Facile Construction of \hat{I}^\pm -Acetoxy Benzofuranones. <i>Organic Letters</i> , 2009, 11, 5174-5177.	2.4	46
13	Palladium-Catalyzed Regioselective Cross-Coupling Reactions of 3-Bromo-4-tosyloxyquinolin-2(1H)-one with Arylboronic Acids. A Facile and Convenient Route to 3,4-Disubstituted Quinolin-2(1H)-ones. <i>Advanced Synthesis and Catalysis</i> , 2007, 349, 1943-1948.	2.1	44
14	Direct Assembly of 3,4-Difunctionalized Benzofurans and Polycyclic Benzofurans by Phenol Dearomatization and Palladium-Catalyzed Domino Reaction. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 6805-6809.	7.2	42
15	Metal- and solvent-free conditions for the acylation reaction catalyzed by carbon tetrabromide (CBr ₄). <i>Green Chemistry</i> , 2007, 9, 1022.	4.6	40
16	Efficient Three-Component One-Pot Benzoylation and Allylation of Aldehydes and Amines for Synthesis of Homobenzylamines and Homoallylamines. <i>Journal of Organic Chemistry</i> , 2007, 72, 3149-3151.	1.7	37
17	A Facile Synthesis of N-Sulfonyl and N-Sulfinyl Aldimines under Barbier-Type Conditions. <i>Journal of Organic Chemistry</i> , 2008, 73, 3623-3625.	1.7	37
18	PhI(OAc) ₂ /I ₂ induced aziridination of alkenes with TsNH ₂ under mild conditions. <i>Tetrahedron Letters</i> , 2008, 49, 4925-4928.	0.7	32

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19	One-Pot Synthesis of Highly Substituted 4-Acetyloxyindoles via Sequential Dearomatization and Silver-Catalyzed Domino Reaction. <i>Organic Letters</i> , 2014, 16, 3600-3603.	2.4	32
20	Divergent Construction of Nitrogen-Containing Polycyclic Compounds with a Dearomatization Strategy. <i>Organic Letters</i> , 2012, 14, 3596-3599.	2.4	31
21	Aniline Dearomatization and Silver-Catalyzed [3+3] Dipolar Cycloaddition: Efficient Construction of Oxocino[4,3,2- <i>bcd</i>]indoles from 2-Alkynylanilines and 2-Alkynylbenzaldoximes. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 14013-14016.	7.2	30
22	One-Pot Oxidative Heteroannulations of <i>N</i> -Sulfonylanilines with Styrenes for the Construction of 5-Aminocoumaran Derivatives. <i>Advanced Synthesis and Catalysis</i> , 2008, 350, 1531-1536.	2.1	28
23	Iodine(III)-Mediated Oxidative Cross-Coupling of Enamines and Propargylamines under Metal-Free Conditions: An Alternative Way to Prepare Highly Substituted 3-Pyrrolines. <i>Organic Letters</i> , 2015, 17, 916-919.	2.4	28
24	Tandem Knoevenagel-Michael Addition of Aryl Sulfonylimines with Diethyl Malonate for Synthesis of Arylidene Dimalonates. <i>Journal of Organic Chemistry</i> , 2007, 72, 5905-5907.	1.7	27
25	Iodobenzene Diacetate/Tetrabutylammonium Iodide-Induced Aziridination of <i>N</i> -Tosylimines with Activated Methylene Compounds under Mild Conditions. <i>Advanced Synthesis and Catalysis</i> , 2008, 350, 1526-1530.	2.1	27
26	A one-pot oxidative decarboxylation-Friedel-Crafts reaction of acyclic α -amino acid derivatives activated by the combination of iodobenzene diacetate/iodine and iron dust. <i>Organic and Biomolecular Chemistry</i> , 2008, 6, 4615.	1.5	24
27	Accessing <i>N</i> -heteroarylated indoles and benzimidazoles from 2-alkynyl cyclohexadienimines and cyclohexadienones through metal-catalyzed tandem reactions. <i>Chemical Communications</i> , 2012, 48, 11775.	2.2	24
28	Application of Dearomatization Strategy on the Synthesis of Furoquinolinone and Angelicin Derivatives. <i>Organic Letters</i> , 2012, 14, 2114-2117.	2.4	23
29	syn Additions to 4-Epoxyfuranosides: Synthesis of 1-Idopyranosides. <i>Organic Letters</i> , 2007, 9, 4849-4852.	2.4	22
30	1,2- and 1,4-Additions of 2-Alkynylcyclohexadienimines with Aromatic Amines To Access 4-Amino- <i>N</i> -aryloxyindoles and -azepinoindoles. <i>Organic Letters</i> , 2012, 14, 6076-6079.	2.4	22
31	Selective C3-C3 Oxidative Cross-Coupling between Unactivated Anilines and Indoles. <i>Advanced Synthesis and Catalysis</i> , 2010, 352, 3230-3234.	2.1	21
32	Amine-Mediated Transimination and Aromatization-Triggered Domino Reaction in the Synthesis of Polyfunctionalized 4-Aminoquinolines. <i>Organic Letters</i> , 2016, 18, 5328-5331.	2.4	21
33	Facile iodine(III)-induced oxidative cycloaddition of <i>N</i> -sulfonyl imines with methylene compounds under neutral conditions. <i>Tetrahedron Letters</i> , 2009, 50, 3857-3859.	0.7	20
34	PhIO/Bu ₄ NI mediated oxidative cyclization of amidoalkylation adducts for the synthesis of <i>N</i> -benzoyl aziridines and oxazolines. <i>Tetrahedron Letters</i> , 2010, 51, 453-456.	0.7	20
35	Facile Construction of Oxa-Aza Spirobicycles via a Tandem Carbon-Hydrogen Bond Oxidation. <i>Advanced Synthesis and Catalysis</i> , 2011, 353, 1735-1740.	2.1	20
36	Destruction and Construction: Application of Dearomatization Strategy in Aromatic Carbon-Nitrogen Bond Functionalization. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 13655-13658.	7.2	19

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37	Hypervalent iodine-mediated regioselective cyclization of acetylenic malonates: facile synthesis of 1-diiodomethylene indane and cyclopentane derivatives. <i>Chemical Communications</i> , 2011, 47, 12221.	2.2	18
38	Five-Membered Ring Systems. <i>Progress in Heterocyclic Chemistry</i> , 2013, 25, 183-215.	0.5	17
39	Metal-Controlled Cycloaddition of 2-Alkynyl-1,4-benzoquinones and Styrenyl Systems: Lewis Acid versus I ⁺ Acid. <i>Organic Letters</i> , 2013, 15, 2482-2485.	2.4	17
40	Synthesis of 4-Alkylindoles from 2-Alkynylanilines via Dearomatization- and Aromatization-Triggered Alkyl Migration. <i>Organic Letters</i> , 2021, 23, 2130-2134.	2.4	16
41	FeCl ₃ -Catalyzed Aza-Diels-Alder Reactions of Methylene-cyclopropanes with Imines. <i>Synthetic Communications</i> , 2007, 37, 4425-4437.	1.1	15
42	Base-promoted selective β^2 -fragmentation of homoallylamines. <i>Tetrahedron Letters</i> , 2010, 51, 4275-4277.	0.7	15
43	Iodobenzene-Catalyzed <i>Ortho</i> -Dearomatization and Aromatization-Triggered Rearrangement of 2-Allylanilines: Construction of Indolin-3-ylmethanols with High Diastereoselectivities. <i>Organic Letters</i> , 2017, 19, 6478-6481.	2.4	15
44	Advances in the development of HIV integrase strand transfer inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2021, 225, 113787.	2.6	15
45	Dy(OTf) ₃ -mediated selective substitution of N-(β -benzotriazolyl-alkyl)amides with active methylene compounds for synthesis of benzotriazole derivatives. <i>Tetrahedron Letters</i> , 2009, 50, 5536-5538.	0.7	14
46	Dearomatization Strategy of β^2 -Enamino Ester: Construction of Indenoazepines via Tandem Michael Addition/Polycyclization. <i>Organic Letters</i> , 2013, 15, 3464-3467.	2.4	14
47	Formal group insertion into aryl C-N bonds through an aromaticity destruction-reconstruction process. <i>Nature Communications</i> , 2018, 9, 3423.	5.8	13
48	Conversion of anilines to chiral benzylic amines via formal one-carbon insertion into aromatic C-N bonds. <i>Nature Communications</i> , 2020, 11, 4805.	5.8	13
49	Dearomatization-Induced Cycloaddition and Aromatization-Triggered Rearrangement: Synthesis of Vertically Expanded Five-Ring Fused Benzofurans. <i>Organic Letters</i> , 2016, 18, 4690-4693.	2.4	12
50	Tandem Palladium Catalysis for Rapid Construction of 3,4-Fused Tricyclic Indoles. <i>Advanced Synthesis and Catalysis</i> , 2020, 362, 1281-1285.	2.1	11
51	Synthesis of 4,7-Difunctionalized Indoles via Imino Exchange and Sulfinyl Migration. <i>Organic Letters</i> , 2020, 22, 823-826.	2.4	7
52	Facile synthesis of 4-acetoxyindoles <i>via</i> PhI(OAc) ₂ -mediated dearomatization of 2-alkynylanilines. <i>Organic Chemistry Frontiers</i> , 2021, 8, 3004-3007.	2.3	7
53	Design and synthesis of novel desfluoroquinolone-aminopyrimidine hybrids as potent anti-MRSA agents with low hERG activity. <i>Bioorganic Chemistry</i> , 2020, 103, 104176.	2.0	6
54	Synthesis of ¹⁵ N-labeled heterocycles <i>via</i> the cleavage of C-N bonds of anilines and glycine- ¹⁵ N. <i>Chemical Communications</i> , 2021, 57, 5442-5445.	2.2	6

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55	Imino exchange reaction in a dearomatization strategy: synthesis of N-acyl diarylamines and phenothiazines from two anilines. <i>Organic Chemistry Frontiers</i> , 2014, 1, 1055-1057.	2.3	5
56	PhI(OAc) ₂ -mediated dialkoxylation of 4-aminostyrenes through a dearomatization process under metal-free conditions. <i>Organic Chemistry Frontiers</i> , 2017, 4, 2156-2158.	2.3	5
57	Accessing bridged bicyclic compounds or meta carbon-functionalized anilines from the dearomatization of anilines. <i>RSC Advances</i> , 2013, 3, 5775.	1.7	4
58	Iodine(III)-induced regioselective carbocyclization of terminal alkynes: a facile approach to prepare 1,1-diodomethylene substituted cyclic compounds. <i>Organic Chemistry Frontiers</i> , 2017, 4, 1005-1010.	2.3	4
59	Synthesis of chiral N-alkylated indoles through replacement of aniline nitrogen by natural amino acids. <i>Green Synthesis and Catalysis</i> , 2022, 3, 282-286.	3.7	4
60	Iodobenzene-Catalyzed Oxidative Cyclization for the Synthesis of Highly Functionalized Cyclopropanes. <i>Synthesis</i> , 2020, 52, 928-932.	1.2	3
61	Synthesis of N-indolated amino acids or peptides from 2-alkynylanilines via a dearomatization process. <i>Organic Chemistry Frontiers</i> , 2021, 8, 6869-6873.	2.3	3
62	Synthesis of C7-Functionalized Indoles through an Aromaticity Destruction–Reconstruction Process. <i>Organic Letters</i> , 2022, 24, 2665-2669.	2.4	3
63	Anodic dearomatization of 2-alkynylanilines for the synthesis of multi-functionalized indoles. <i>Chemical Communications</i> , 2022, 58, 6797-6800.	2.2	2
64	Divergent synthesis of 4-amino indoles with free amine groups <i>via</i> tandem reaction of 2-alkynylanilines. <i>Organic Chemistry Frontiers</i> , 2022, 9, 4146-4150.	2.3	1
65	Stereoselective Synthesis of Acyclic Tetrasubstituted Alkenes from Anilines by Dearomatization and Trimethylenemethane Cycloaddition. <i>Organic Letters</i> , 2022, 24, 314-318.	2.4	0