

# Jinbao Xiang

## List of Publications by Year in descending order

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34  
papers

630  
citations

759233

12  
h-index

580821

25  
g-index

37  
all docs

37  
docs citations

37  
times ranked

738  
citing authors

#	ARTICLE	IF	CITATIONS
1	Hindered dialkyl ether synthesis with electrogenerated carbocations. <i>Nature</i> , 2019, 573, 398-402.	27.8	240
2	Nickel-Catalyzed Electrochemical Phosphorylation of Aryl Bromides. <i>Organic Letters</i> , 2019, 21, 6835-6838.	4.6	66
3	A Cascade Reaction Consisting of Pictet-Spengler-Type Cyclization and Smiles Rearrangement: Application to the Synthesis of Novel Pyrrole-Fused Dihydropteridines. <i>Organic Letters</i> , 2007, 9, 765-767.	4.6	47
4	Synthesis of Pyrido[2,3-e]pyrrolo[1,2-a]pyrazine Derivatives via Tandem Iminium Cyclization and Smiles Rearrangement. <i>Journal of Organic Chemistry</i> , 2008, 73, 3281-3283.	3.2	28
5	Electroselective and Controlled Reduction of Cyclic Imides to Hydroxylactams and Lactams. <i>Organic Letters</i> , 2021, 23, 2298-2302.	4.6	27
6	Electrochemical Cross-Dehydrogenative Coupling of <i>N</i> -Aryl-tetrahydroisoquinolines with Phosphites and Indole. <i>European Journal of Organic Chemistry</i> , 2019, 2019, 2498-2501.	2.4	22
7	Discovery of Novel Tricyclic Thiazepine Derivatives as Anti-Drug-Resistant Cancer Agents by Combining Diversity-Oriented Synthesis and Converging Screening Approach. <i>ACS Combinatorial Science</i> , 2016, 18, 230-235.	3.8	18
8	Stereoselective Synthesis of 3-Carboxy-4,5-dihydropyrroles via an Intramolecular Iminium Ion Cyclization Reaction. <i>Organic Letters</i> , 2015, 17, 3818-3821.	4.6	15
9	Synthesis of highly substituted 2,3-dihydropyrimido[4,5-d]pyrimidin-4(1H)-ones from 4,6-dichloro-5-formylpyrimidine, amines and aldehydes. <i>Molecular Diversity</i> , 2011, 15, 839-847.	3.9	14
10	Electrochemical Regioselective Bromination of Electron-Rich Aromatic Rings Using <i>n</i> Bu <sub>4</sub> NBr. <i>Synlett</i> , 2019, 30, 1313-1316.	1.8	13
11	Modular and Stereoselective Approach to Highly Substituted Indole/Pyrrole-Fused Diazepanones. <i>Journal of Organic Chemistry</i> , 2021, 86, 6458-6466.	3.2	13
12	Synthesis of Novel 8,9-Dihydro-5H-pyrimido[4,5-e][1,4]diazepin-7(6H)-ones. <i>ACS Combinatorial Science</i> , 2010, 12, 503-509.	3.3	12
13	The Construction of Hydrangea-like Vanadium-Doped Iron Nickel Phosphide as an Enhanced Bifunctional Electrocatalyst for Overall Water Splitting. <i>ACS Applied Energy Materials</i> , 2020, 3, 9449-9458.	5.1	12
14	Synthesis of novel 4H-pyrimido[1,6-a]pyrimidines via a one-pot three-component condensation. <i>Molecular Diversity</i> , 2012, 16, 173-181.	3.9	11
15	Stereochemistry as a Tool in Deciphering the Processes of a Tandem Iminium Cyclization and Smiles Rearrangement. <i>Journal of Organic Chemistry</i> , 2010, 75, 8147-8154.	3.2	10
16	Single-Electron Oxidation/Alterable C3- and C10-Arylation of 9-MeO-phenanthrene. <i>Organic Letters</i> , 2018, 20, 3591-3595.	4.6	10
17	Synthesis of Isoxazolidine-Fused Eight-Membered Heterocycles via an Intramolecular Nitroene-Alkene Cycloaddition. <i>Synlett</i> , 2015, 26, 238-242.	1.8	9
18	A one-pot procedure for ring enlargement of $\beta$ -chloromethyl-N-containing heterocycles. <i>Journal of Heterocyclic Chemistry</i> , 2006, 43, 321-324.	2.6	7

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19	Electronic Effects on the cis/trans Selectivity in Formation of Isoxazolidine-Fused Eight-Membered Ring via an Intramolecular Nitron-alkene Cycloaddition. <i>Chemistry of Heterocyclic Compounds</i> , 2016, 52, 601-608.	1.2	6
20	Intramolecular Cycloaddition of Azomethine Ylides Activated by Aromatic Rings: Scope and Limitations. <i>Chemistry of Heterocyclic Compounds</i> , 2016, 52, 484-492.	1.2	6
21	A Highly Stereocontrolled Intramolecular Cycloaddition Reaction of Azomethine Ylide Activated by a Pyrimidine Ring: Access to Novel Tricyclic Hexahydro-1H-pyrrolo[2,3-d]pyrido[2,3-d]pyrimidines. <i>Synlett</i> , 2012, 23, 585-588.	1.8	5
22	Iron and nitrogen co-functionalized porous 3D graphene frameworks as an efficient oxygen reduction catalyst. <i>RSC Advances</i> , 2016, 6, 74886-74894.	3.6	5
23	New electrotriggers: <i>p</i> -methoxycarbonylbenzyl (pMCB) as an electroremovable protecting group for carboxylic acids, phosphoric acids and alcohols. <i>Green Chemistry</i> , 2022, 24, 5632-5636.	9.0	5
24	Synthesis of pyrido[2,3- <i>b</i> ][1,4]benzoxazepines via a Friedel-Crafts cyclization. <i>Chemistry of Heterocyclic Compounds</i> , 2016, 52, 326-330.	1.2	4
25	Synthesis and Evaluation of 2-Alkylthio-4-( <i>N</i> -substituted sulfonamide)pyrimidine Hydroxamic Acids as Anti-Myeloma Agents. <i>Chemical Biology and Drug Design</i> , 2016, 87, 472-477.	3.2	4
26	Diethyl Phosphite Promoted Electrochemical Oxidation of Tetrahydroisoquinolines to 3,4-Dihydroisoquinolin-1(2H)-ones. <i>Synlett</i> , 2019, 30, 2077-2080.	1.8	4
27	Direct C(sp <sup>3</sup> )-H allylation of 2-alkylpyridines with Morita-Baylis-Hillman carbonates via a tandem nucleophilic substitution/aza-Cope rearrangement. <i>Beilstein Journal of Organic Chemistry</i> , 2021, 17, 2505-2510.	2.2	4
28	Synthesis of novel tricyclic 4-chloro-7,8,10,11-tetrahydro-5H-benzo[ <i>e</i> ]pyrimido[4,5- <i>b</i> ][1,4]diazepin-9(6 <i>H</i> )-ones. <i>Journal of Heterocyclic Chemistry</i> , 2010, 47, 990-993.	2.6	3
29	Synthesis of novel 6,7-dihydro-5H-pyrimido[4,5- <i>e</i> ][1,4]diazepin-8(9 <i>H</i> )-ones. <i>Journal of Heterocyclic Chemistry</i> , 2011, 48, 1091-1094.	2.6	3
30	Highly selective electroreductive linear dimerization of electron-deficient vinylarenes. <i>Tetrahedron</i> , 2021, 102, 132535.	1.9	3
31	The discovery of kinase inhibitors by a combination of diversity-oriented synthesis and selective screening. <i>MedChemComm</i> , 2016, 7, 1946-1951.	3.4	2
32	Catalyst-Controlled Regiodivergent Synthesis of $\alpha,\beta$ -Dipeptide Derivatives via <i>N</i> -Allylic Alkylation of <i>O</i> -Alkyl Hydroxamates with MBH Carbonates. <i>Chemistry - an Asian Journal</i> , 2022, 17, .	3.3	2
33	Trifluoroacetic Acid-Mediated Nucleophilic Substitution/Smiles Rearrangement Cascade Reaction: An Alternative Approach to Constructing Pyrrole-Fused Dihydropteridines. <i>Chemistry of Heterocyclic Compounds</i> , 2016, 52, 831-835.	1.2	0
34	CC102528: A Novel Histone Deacetylase Inhibitor in the Hydroxamate Family Demonstrates Potent Anti-Myeloma Activity.. <i>Blood</i> , 2009, 114, 4927-4927.	1.4	0