

Dong Wang

List of Publications by Year in descending order

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826
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#	ARTICLE	IF	CITATIONS
1	Recent Advances in the Synthesis of C2-Functionalized Pyridines and Quinolines Using <i>N</i> -Oxide Chemistry. <i>Advanced Synthesis and Catalysis</i> , 2021, 363, 2-39.	2.1	95
2	Natural Prenylchalconaringenins and Prenylnaringenins as Antidiabetic Agents: α -Glucosidase and α -Amylase Inhibition and in Vivo Antihyperglycemic and Antihyperlipidemic Effects. <i>Journal of Agricultural and Food Chemistry</i> , 2017, 65, 1574-1581.	2.4	86
3	Synthesis of 6-hydroxyaurone analogues and evaluation of their α -glucosidase inhibitory and glucose consumption-promoting activity: Development of highly active 5,6-disubstituted derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 3226-3230.	1.0	41
4	Prohibitin ligands: a growing armamentarium to tackle cancers, osteoporosis, inflammatory, cardiac and neurological diseases. <i>Cellular and Molecular Life Sciences</i> , 2020, 77, 3525-3546.	2.4	40
5	Inhibitory activity evaluation and mechanistic studies of tetracyclic oxindole derivatives as α -glucosidase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2016, 123, 365-378.	2.6	37
6	A practical and mild chlorination of fused heterocyclic N-oxides. <i>Tetrahedron Letters</i> , 2014, 55, 7130-7132.	0.7	32
7	Strategic C-C Bond-Forming Dearomatization of Pyridines and Quinolines. <i>Organic Letters</i> , 2019, 21, 4459-4463.	2.4	30
8	Strategic Approach to 8-Azacoumarins. <i>Organic Letters</i> , 2017, 19, 984-987.	2.4	28
9	Targeting prohibitin with small molecules to promote melanogenesis and apoptosis in melanoma cells. <i>European Journal of Medicinal Chemistry</i> , 2018, 155, 880-888.	2.6	28
10	A General and Efficient Synthesis of 2-Pyridones, 2-Quinolones, and 1-Isoquinolinones from Azine <i>N</i> -Oxides. <i>Asian Journal of Organic Chemistry</i> , 2016, 5, 1442-1446.	1.3	27
11	A highly practical and convenient halogenation of fused heterocyclic N-oxides. <i>Tetrahedron</i> , 2016, 72, 5762-5768.	1.0	27
12	Exploring the pH dependence of viologen reduction by α -carbon radicals derived from Hcy and Cys. <i>Chemical Communications</i> , 2009, , 1876.	2.2	26
13	Access to 8-Azachromones via Activation of C-H in <i>N</i> -Oxides. <i>Journal of Organic Chemistry</i> , 2017, 82, 11275-11287.	1.7	21
14	Metal- and base-free regioselective thiolation of the methyl C(sp ³)-H bond in 2-picoline <i>N</i> -oxides. <i>Green Chemistry</i> , 2019, 21, 157-163.	4.6	21
15	One-Carbon Bridge Stereocontrol in Robinson Annulations Leading to Bicyclo[3.3.1]nonanes. <i>Organic Letters</i> , 2010, 12, 1232-1235.	2.4	15
16	One-Pot Selective Saturation and Functionalization of Heteroaromatics Leading to Dihydropyridines and Dihydroquinolines. <i>Journal of Organic Chemistry</i> , 2020, 85, 5027-5037.	1.7	13
17	SFPH proteins as therapeutic targets for a myriad of diseases. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020, 30, 127600.	1.0	12
18	Access to Furo[2,3- <i>b</i>]pyridines by Transition-Metal-Free Intramolecular Cyclization of C3-Substituted Pyridine <i>N</i> -Oxides. <i>Asian Journal of Organic Chemistry</i> , 2018, 7, 879-882.	1.3	10

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19	Catalyst-free three-component synthesis of highly functionalized 2,3-dihydropyrroles. <i>Green Chemistry</i> , 2018, 20, 2775-2780.	4.6	10
20	Transition-metal-free access to 7-azaindoles. <i>Tetrahedron</i> , 2018, 74, 4100-4110.	1.0	10
21	Recent advances in the synthesis of 2,3-dihydropyrroles. <i>Chemical Communications</i> , 2020, 56, 5584-5592.	2.2	10
22	The prohibitin-binding compound fluorizoline affects multiple components of the translational machinery and inhibits protein synthesis. <i>Journal of Biological Chemistry</i> , 2020, 295, 9855-9867.	1.6	9
23	A One-Pot Dearomative Approach to C4-Alkylated Tetrahydropyridines and Tetrahydroquinolines. <i>Asian Journal of Organic Chemistry</i> , 2020, 9, 1571-1575.	1.3	8
24	A facile approach to tricyclo[6.4.0.0 ^{4,9}]-dodecane framework. <i>Chinese Chemical Letters</i> , 2015, 26, 238-242.	4.8	6
25	Accessing 1,8-Naphthyridones by Metal-Free Regioselective Amination of Pyridine N-oxides/Acid-Mediated Cyclization. <i>Advanced Synthesis and Catalysis</i> , 2020, 362, 3841-3845.	2.1	6
26	Stereoselective Four-Component Synthesis of Functionalized 2,3-Dihydro-4-Nitropyrroles. <i>Frontiers in Chemistry</i> , 2019, 7, 810.	1.8	5
27	Transition-Metal-Free Regioselective Direct C2, C4 Difunctionalization and C2, C4, C6 Trifunctionalization of Pyridines. <i>Advanced Synthesis and Catalysis</i> , 2022, 364, 2720-2728.	2.1	5
28	Scalable synthesis of a tetrasubstituted 7-azabenzofuran as a key intermediate for a class of potent HCV NS5B inhibitors. <i>Tetrahedron</i> , 2020, 76, 131642.	1.0	4
29	Rapid Generation of Tetrahydropyridines and Tetrahydroquinolines by Dearomative Cyanation/Grignard Addition. <i>Chemistry - an Asian Journal</i> , 2022, 17, .	1.7	3
30	Facile and Efficient Synthesis of Tri- and Tetrasubstituted 7-Azabenzofuran Derivatives. <i>Asian Journal of Organic Chemistry</i> , 2020, 9, 749-752.	1.3	2
31	Mechanism and origin of stereoselectivity in Robinson annulations leading to bicyclo[3.3.1]nonanes: a rare Curtin-Hammett scenario. <i>Journal of Physical Organic Chemistry</i> , 2017, 30, e3595.	0.9	1
32	Flavaglines: Discovery From Plants Used in Traditional Chinese Medicine, Synthesis and Drug Development against Cancer and Immune Disorders. <i>Current Chinese Chemistry</i> , 2021, 01, .	0.3	0