

Xiaowei Wu

List of Publications by Year in descending order

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Version: 2024-02-01

25
papers

807
citations

516710

16
h-index

610901

24
g-index

25
all docs

25
docs citations

25
times ranked

640
citing authors

#	ARTICLE	IF	CITATIONS
1	Design, synthesis and biological evaluation of pyrazolo[3,4-d]pyridazinone derivatives as covalent FGFR inhibitors. <i>Acta Pharmaceutica Sinica B</i> , 2021, 11, 781-794.	12.0	16
2	Additive-Controlled Divergent Synthesis of Tetrasubstituted 1,3-Enynes and Alkynylated 3- <i>H</i> -Pyrrolo[1,2- <i>a</i>]indol-3-ones via Rhodium Catalysis. <i>Organic Letters</i> , 2021, 23, 727-733.	4.6	46
3	Small-molecule inhibitor of AF9/ENL-DOT1L/AF4/AFF4 interactions suppresses malignant gene expression and tumor growth. <i>Theranostics</i> , 2021, 11, 8172-8184.	10.0	17
4	Synthesis, Structure-Activity Relationships, and Antiviral Activity of Allosteric Inhibitors of Flavivirus NS2B-NS3 Protease. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 2777-2800.	6.4	24
5	Redox-Neutral Rhodium(III)-Catalyzed Chemospecific and Regiospecific [4+1] Annulation between Indoles and Alkenes for the Synthesis of Functionalized Imidazo[1,5- <i>a</i>]indoles. <i>Journal of Organic Chemistry</i> , 2021, 86, 10591-10607.	3.2	11
6	Rh(III)-Catalyzed Divergent Synthesis of Alkynylated Imidazo[1,5- <i>a</i>]indoles and <i>1,1</i> -Difluoromethylene Tetrasubstituted Alkenes. <i>Organic Letters</i> , 2021, 23, 5766-5771.	4.6	22
7	Chemo- and Regioselective Synthesis of Functionalized 1- <i>H</i> -imidazo[1,5- <i>a</i>]indol-3(2- <i>H</i>)-ones via a Redox-Neutral Rhodium(III)-Catalyzed [4+1] Annulation between Indoles and Alkynes. <i>Advanced Synthesis and Catalysis</i> , 2021, 363, 4380-4389.		9
8	Chemo-, Regio-, and Stereoselective Assembly of Polysubstituted Furan-2(5- <i>H</i>)-ones Enabled by Rh(III)-Catalyzed Domino C-H Alkenylation/Directing Group Migration/Lactonization: A Combined Experimental and Computational Study. <i>ACS Catalysis</i> , 2021, 11, 13921-13934.	11.2	20
9	Rhodium(III)-Catalyzed C-H Alkenylation/Directing Group Migration for the Regio- and Stereoselective Synthesis of Tetrasubstituted Alkenes. <i>Organic Letters</i> , 2020, 22, 9163-9168.	4.6	37
10	Rhodium-Catalyzed Cascade Reactions of Indoles with 4-Hydroxyalkynoates for the Synthesis of Indole-Fused Polyheterocycles. <i>Advanced Synthesis and Catalysis</i> , 2020, 362, 2953-2960.	4.3	31
11	Discovery, Structure-Activity Relationship, and Biological Activity of Histone-Competitive Inhibitors of Histone Acetyltransferases P300/CBP. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 4716-4731.	6.4	17
12	Discovery and Development of a Series of Pyrazolo[3,4- <i>d</i>]pyridazinone Compounds as the Novel Covalent Fibroblast Growth Factor Receptor Inhibitors by the Rational Drug Design. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 7473-7488.	6.4	28
13	Rhodium-Catalyzed [4 + 1] Cyclization via C-H Activation for the Synthesis of Divergent Heterocycles Bearing a Quaternary Carbon. <i>Journal of Organic Chemistry</i> , 2018, 83, 4650-4656.	3.2	60
14	Ruthenium(II)-Catalyzed Regio- and Stereoselective C-H Allylation of Indoles with Allyl Alcohols. <i>Organic Letters</i> , 2018, 20, 2224-2227.	4.6	44
15	Ruthenium-Catalyzed C-H Allylation of Alkenes with Allyl Alcohols via C-H Bond Activation in Aqueous Solution. <i>Journal of Organic Chemistry</i> , 2018, 83, 12094-12102.	3.2	22
16	Rhodium-catalyzed C-H allylation of indoles with allyl alcohols via β -hydroxide elimination. <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 5691-5698.	2.8	22
17	Regio-selective and stereo-selective hydrosilylation of internal alkynes catalyzed by ruthenium complexes. <i>RSC Advances</i> , 2018, 8, 28261-28265.	3.6	8
18	Propargyl Alcohols as One-Carbon Synthons: Redox-Neutral Rhodium(III)-Catalyzed C-H Bond Activation for the Synthesis of Isoindolinones Bearing a Quaternary Carbon. <i>Organic Letters</i> , 2017, 19, 1294-1297.	4.6	106

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19	Ruthenium-Catalyzed Redox-Neutral [4 + 1] Annulation of Benzamides and Propargyl Alcohols via C-H Bond Activation. <i>ACS Catalysis</i> , 2017, 7, 2494-2499.	11.2	118
20	Ruthenium(II)-Catalyzed Redox-Neutral [3+2] Annulation of Indoles with Internal Alkynes via C-H Bond Activation: Accessing a Pyrroloindolone Scaffold. <i>Journal of Organic Chemistry</i> , 2017, 82, 5263-5273.	3.2	45
21	Rh(III)-Catalyzed C-H Cyclization of Arylnitrones with Diazo Compounds: Access to 3-Carboxylate Substituted N-Hydroxyindoles. <i>Journal of Organic Chemistry</i> , 2017, 82, 8984-8994.	3.2	42
22	Site-specific indolation of proline-based peptides via copper-catalyzed oxidative coupling of tertiary amine N-oxides. <i>Chemical Communications</i> , 2015, 51, 12571-12573.	4.1	19
23	Design, synthesis and biological evaluation of isoquinoline-based derivatives as novel histone deacetylase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 5881-5890.	3.0	17
24	Design, synthesis and biological evaluation of 4-anilinothieno[2,3-d]pyrimidine-based hydroxamic acid derivatives as novel histone deacetylase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 6146-6155.	3.0	24
25	Temperature-Controlled Divergent Synthesis of Tetrasubstituted Alkenes and Pyrrolo[1,2-a]indole Derivatives via Iridium Catalysis. <i>Asian Journal of Organic Chemistry</i> , 0, , .	2.7	2