

Nageswara Rao Desaboini

List of Publications by Year in descending order

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Version: 2024-02-01

21
papers

648
citations

687363

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all docs

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23
times ranked

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citing authors

#	ARTICLE	IF	CITATIONS
1	Deciphering the Molecular Mechanism of HCV Protease Inhibitor Fluorination as a General Approach to Avoid Drug Resistance. <i>Journal of Molecular Biology</i> , 2022, 434, 167503.	4.2	6
2	Silicon functionalization expands the repertoire of Si-rhodamine fluorescent probes. <i>Chemical Science</i> , 2022, 13, 6081-6088.	7.4	11
3	Seralite SRC-120 resin catalyzed synthesis of bis(indolyl)methanes using indoles and low/high boiling point carbonyl compounds under solvent free conditions. <i>Synthetic Communications</i> , 2021, 51, 139-150.	2.1	6
4	Drug Design Strategies to Avoid Resistance in Direct-Acting Antivirals and Beyond. <i>Chemical Reviews</i> , 2021, 121, 3238-3270.	47.7	40
5	Discovery of Quinoxaline-Based P1â€‘P3 Macrocyclic NS3/4A Protease Inhibitors with Potent Activity against Drug-Resistant Hepatitis C Virus Variants. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 11972-11989.	6.4	15
6	Advances in Carbonâ€‘Element Bond Construction under Chanâ€‘Lam Cross-Coupling Conditions: A Second Decade. <i>Synthesis</i> , 2021, 53, 805-847.	2.3	43
7	Structural Analysis of Potent Hybrid HIV-1 Protease Inhibitors Containing Bis-tetrahydrofuran in a Pseudosymmetric Dipeptide Isostere. <i>Journal of Medicinal Chemistry</i> , 2020, 63, 8296-8313.	6.4	6
8	Avoiding Drug Resistance by Substrate Envelope-Guided Design: Toward Potent and Robust HCV NS3/4A Protease Inhibitors. <i>MBio</i> , 2020, 11, .	4.1	15
9	Avoiding Drug Resistance by Substrate Envelope Guided Design: Toward Potent and Robust HCV NS3/4A Protease Inhibitors. <i>FASEB Journal</i> , 2020, 34, 1-1.	0.5	0
10	Cobalt-Catalyzed Regioselective Ortho C(sp ²)-H Bond Nitration of Aromatics through Proton-Coupled Electron Transfer Assistance. <i>Journal of Organic Chemistry</i> , 2017, 82, 7234-7244.	3.2	32
11	Copper(II)-catalyzed Chanâ€‘Lam cross-coupling: chemoselective N-arylation of aminophenols. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 801-806.	2.8	52
12	Copper-Catalyzed C(sp ²)-NH ₂ Arylation of 2-Aminobenzimidazoles and Related C ₂ -Amino-NH ₂ azoles. <i>Advanced Synthesis and Catalysis</i> , 2016, 358, 2126-2133.	4.3	43
13	Cobalt(II)-Catalyzed C-N Bond Formation: Synthesis of Aryl-Substituted Imidazo[1,2-a]pyridines. <i>Asian Journal of Organic Chemistry</i> , 2016, 5, 1213-1218.	2.7	13
14	Nickel-catalyzed Chanâ€‘Lam cross-coupling: chemoselective N-arylation of 2-aminobenzimidazoles. <i>Organic and Biomolecular Chemistry</i> , 2016, 14, 8989-8997.	2.8	39
15	Nickel-Catalyzed C(Aryl)-O Bond Activation/Cross-Coupling Reaction of Carbazoles with Methyl Grignard: Synthesis of Ellipticine. <i>Asian Journal of Organic Chemistry</i> , 2016, 5, 1499-1507.	2.7	5
16	Palladium/Silver Synergistic Catalysis in Direct Aerobic Carbonylation of C(sp ²)-C-H Bonds Using DMF as a Carbon Source: Synthesis of Pyrido-Fused Quinazolinones and Phenanthridinones. <i>Organic Letters</i> , 2016, 18, 3142-3145.	4.6	89
17	Sulphuric acid immobilized on silica gel (H ₂ SO ₄ â€‘SiO ₂) as an eco-friendly catalyst for transamidation. <i>RSC Advances</i> , 2015, 5, 10567-10574.	3.6	48
18	Copper-Catalyzed Inter- and Intramolecular C-N Bond Formation: Synthesis of Benzimidazole-Fused Heterocycles. <i>Journal of Organic Chemistry</i> , 2015, 80, 9321-9327.	3.2	74

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19	Câ€N bond formation via Cu-catalyzed cross-coupling with boronic acids leading to methyl carbazole-3-carboxylate: synthesis of carbazole alkaloids. RSC Advances, 2014, 4, 4960.	3.6	26
20	Hypervalent iodine(<i>iii</i>) catalyzed oxidative Câ€N bond formation in water: synthesis of benzimidazole-fused heterocycles. RSC Advances, 2014, 4, 25600-25604.	3.6	48
21	Base and ligand free copper-catalyzed N-arylation of 2-amino-N-heterocycles with boronic acids in air. RSC Advances, 2013, 3, 11472.	3.6	37