

Hiteshkumar Jalani

List of Publications by Year in descending order

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papers

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#	ARTICLE	IF	CITATIONS
1	Novel thiazole–thiophene conjugates as adenosine receptor antagonists: Synthesis, biological evaluation and docking studies. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 1306-1309.	2.2	25
2	Synthesis of Substituted 1,2,3-Triazoles via Metal-Free Click Cycloaddition Reactions and Alternative Cyclization Methods. <i>Synthesis</i> , 2016, 49, 29-41.	2.3	25
3	A convenient synthesis of di- and trisubstituted 2-aminoimidazoles from 1-amidino-3-trityl-thioureas. <i>Tetrahedron Letters</i> , 2009, 50, 3955-3958.	1.4	23
4	Copper and cobalt co-catalyzed aerobic oxidative cross-dehydrogenative coupling reaction of (benzo)azoles. <i>Green Chemistry</i> , 2019, 21, 5797-5802.	9.0	23
5	Improvement of Aqueous Solubility of Lapatinib-Derived Analogues: Identification of a Quinolinimine Lead for Human African Trypanosomiasis Drug Development. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 665-687.	6.4	23
6	An efficient synthesis of 2-aminopyrroles from enamino–amidine adduct and phenacyl/benzyl/heteroalkyl-halides. <i>Tetrahedron Letters</i> , 2011, 52, 6331-6335.	1.4	21
7	A convenient one-pot synthesis of trisubstituted 1,3,5-triazines through intermediary amidinothioureas. <i>Tetrahedron Letters</i> , 2010, 51, 1486-1489.	1.4	19
8	Iridium-Catalyzed Aryl C–H Sulfonamidation and Amide Formation Using a Bifunctional Nitrogen Source. <i>Organic Letters</i> , 2018, 20, 4828-4832.	4.6	19
9	Synergistic combination of visible-light photo-catalytic electron and energy transfer facilitating multicomponent synthesis of β -functionalized β,β -diarylethylamines. <i>Chemical Communications</i> , 2019, 55, 6405-6408.	4.1	19
10	Scaffold and Parasite Hopping: Discovery of New Protozoal Proliferation Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 249-257.	2.8	17
11	A concise, greener, solvent-free novel one-pot synthesis of trisubstituted thiophenes. <i>Tetrahedron Letters</i> , 2012, 53, 6927-6930.	1.4	16
12	Copper–Catalyzed One–pot Synthesis of Pyrimidines from Amides, <i>N,N</i> -dimethylformamide dimethylacetal, and Enamines. <i>Advanced Synthesis and Catalysis</i> , 2017, 359, 2509-2513.	4.3	16
13	An efficient one-pot synthesis of functionally diverse 2-aminothiazoles from isothiocyanates, amidines/guanidines and halomethylenes. <i>Tetrahedron Letters</i> , 2013, 54, 5403-5406.	1.4	15
14	Oxidative Cascade Reaction of <i>N</i> -Aryl-3-alkylideneazetidines and Carboxylic Acids: Access to Fused Pyridines. <i>Organic Letters</i> , 2018, 20, 3833-3837.	4.6	15
15	Rh-Catalyzed Chemoselective [4 + 1] Cycloaddition Reaction toward Diverse 4-Methyleneprolines. <i>Journal of Organic Chemistry</i> , 2019, 84, 10877-10891.	3.2	15
16	Iridium-Catalyzed C–H Amination/Cyclization for Medium to Large <i>N</i> -Heterocycle-Fused Dihydroquinazolinones. <i>Organic Letters</i> , 2019, 21, 3706-3710.	4.6	15
17	A versatile one-pot multicomponent synthesis of novel quinazolinon-2-yl-tetrasubstituted thiophenes. <i>Tetrahedron Letters</i> , 2010, 51, 5686-5689.	1.4	14
18	An efficient, greener, and solvent-free one-pot multicomponent synthesis of 3-substituted quinazolin-4(3H)ones and thienopyrimidin-4(3H)ones. <i>Tetrahedron Letters</i> , 2012, 53, 4062-4064.	1.4	13

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19	Discovery of enantioselectivity of urea inhibitors of soluble epoxide hydrolase. <i>European Journal of Medicinal Chemistry</i> , 2016, 117, 113-124.	5.5	12
20	Design and synthesis of sulfonamidophenylethylureas as novel cardiac myosin activator. <i>European Journal of Medicinal Chemistry</i> , 2018, 143, 1869-1887.	5.5	10
21	Iridium-Catalyzed C-H Amination of Weinreb Amides: A Facile Pathway toward Anilines and Quinazolin-2,4-diones. <i>Journal of Organic Chemistry</i> , 2020, 85, 13096-13107.	3.2	10
22	Stereospecific Electrophilic Fluorocyclization of β,β -Unsaturated Amides with Selectfluor. <i>Organic Letters</i> , 2020, 22, 2651-2656.	4.6	10
23	Exploration of flexible phenylpropylurea scaffold as novel cardiac myosin activators for the treatment of systolic heart failure. <i>European Journal of Medicinal Chemistry</i> , 2017, 134, 379-391.	5.5	9
24	A Diversified Assembly of 1,2,4-Oxadiazol-3-amines: Metallic Thiophile Catalyzed Chemoselective One-Pot Reaction of Aryl Isothiocyanates, Amidines/Guanidines, and Hydroxylamine. <i>Synthesis</i> , 2012, 44, 3378-3386.	2.3	7
25	Discovery of a novel series of N-hydroxypyridone derivatives protecting astrocytes against hydrogen peroxide-induced toxicity via improved mitochondrial functionality. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 1394-1405.	3.0	7
26	Iodine-Promoted One-Pot Synthesis of Highly Substituted 4-Aminopyrroles and Bis-4-Aminopyrrole from Aryl Methyl Ketones, Arylamines, and Enamines. <i>Advanced Synthesis and Catalysis</i> , 2018, 360, 4073-4079.	4.3	7
27	Stability-indicating assay method for determination of actarit, its process related impurities and degradation products: Insight into stability profile and degradation pathways. <i>Journal of Pharmaceutical Analysis</i> , 2014, 4, 374-383.	5.3	6
28	<i>p</i> -Toluenesulfonic acid catalyzed, isocyanide-free, Groebke-Blackburn-Bienayme (GBB) type multicomponent synthesis of 3-anilino-1,2,4-diazopyridines. <i>Journal of Heterocyclic Chemistry</i> , 2022, 59, 1266-1271.	2.6	6
29	Exploration and Optimization of an Efficient One-Pot Sequential Synthesis of Di/tri-substituted Thiazoles from β -Bromoketones, Thioacids Salt, and Ammonium Acetate. <i>Journal of Heterocyclic Chemistry</i> , 2016, 53, 1449-1456.	2.6	5
30	Discovery of 2-aminoimidazole and 2-aminoimidazolyl-thiazoles as non-xanthine human adenosine A3 receptor antagonists: SAR and molecular modeling studies. <i>MedChemComm</i> , 2018, 9, 676-684.	3.4	4
31	Sequential One-Pot Synthesis of Imidazoles and 2H-imidazolones from β -Ketoamines, Acylating Agents and Ammonium Acetate. <i>Bulletin of the Korean Chemical Society</i> , 2016, 37, 1966-1970.	1.9	3
32	Synthesis of Selenopyrano[2,3-c]pyrazol-4(1H)-ones and Their C-H Activation. <i>Synlett</i> , 2021, 32, 321-325.	1.8	3
33	Exploration of diphenylalkyloxadiazoles as novel cardiac myosin activator. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018, 28, 2369-2374.	2.2	2
34	Structure-Bioactivity Relationships of Lapatinib Derived Analogs against <i>Schistosoma mansoni</i> . <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 258-265.	2.8	2
35	Catalyst-Free One-Pot Multi-Component Synthesis of 2-Substituted Quinazolin-4-carboxamides from 2-Aminophenyl-2-oxoacetamides, Aldehydes, and Ammonium Acetate. <i>ChemistrySelect</i> , 2021, 6, 5446-5450.	1.5	2
36	Design and synthesis of sulfonamidophenylethylamides as novel cardiac myosin activator. <i>Bioorganic and Medicinal Chemistry</i> , 2019, 27, 4110-4123.	3.0	0

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37	Synthesis of selenopyrano[2,3-b]indol-4(9H)-ones and their C-alkylation. Journal of Heterocyclic Chemistry, 0, , .	2.6	0