

Ganesh Chandra Nandi

List of Publications by Year in descending order

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46
papers

2,065
citations

279798

23
h-index

233421

45
g-index

65
all docs

65
docs citations

65
times ranked

1858
citing authors

#	ARTICLE	IF	CITATIONS
1	Direct Synthesis of Sulfonylimidoyl Guanidines from Sulfonylimidoyl Azides under Dual (Cobalt and Tj ETQq1 1 0.784314 rgBT /Qverlock	4.3	10
2	Advances in the photoredox catalysis of S(VI) compounds. Tetrahedron, 2022, 111, 132711.	1.9	10
3	Advances in the Synthesis and Applications of Three Membered Sila, Silaâ€Aza/â€Phospha/â€Oxa/â€Thia Cyclopropanes. European Journal of Organic Chemistry, 2021, 2021, 587-606.	2.4	11
4	A metal-free Petasis reaction towards the synthesis of <i>N</i>-(\pm-substituted)alkyl sulfoximines/sulfonylimidamides. Organic and Biomolecular Chemistry, 2021, 19, 7061-7065.	2.8	12
5	Recent Advances in the Preparations and Synthetic Applications of Oxaziridines and Diaziridines. Advanced Synthesis and Catalysis, 2021, 363, 1756-1781.	4.3	13
6	Applications of Carbon Dots (CDs) in Latent Fingerprints Imaging. Chemistry - an Asian Journal, 2021, 16, 1057-1072.	3.3	23
7	Design and Synthesis of Triphenylamine Based Cyano Stilbenes for Picric Acid Sensing and Two Photon Absorption Applications. ChemistrySelect, 2021, 6, 12300-12308.	1.5	4
8	Mild and Metal-Free Protocol toward the Synthesis of Triarylmethanes by Reactions of (Hetero)Arylboronic Acids and <i>ortho</i>-Hydroxyarylaldehydes. Journal of Organic Chemistry, 2020, 85, 3000-3009.	3.2	18
9	Sulfonylimidamide as a directing agent for Pd-catalyzed regioselective oxidative Câ€H acyloxylation of arenes. Tetrahedron, 2019, 75, 130622.	1.9	12
10	Recent Advances in the A³ Coupling Reactions and their Applications. European Journal of Organic Chemistry, 2019, 2019, 2704-2720.	2.4	99
11	Sulfonylimidoyl Azide: A Novel Precursor for the Direct and Rapid Access to <i>N</i>-â€Aryl Sulfonylimidamides <i>via</i> Cuâ€Catalyzed Chanâ€Evansâ€Lam Reaction with Boronic Acids under Mild And Efficient Condition. ChemistrySelect, 2019, 4, 14004-14006.	1.5	5
12	Catalystâ€Controlled Dual Reactivity of Sulfonylimidamides: Synthesis of Propargylamines and <i>N</i>-â€Propargyl Sulfonylimidamides. Chemistry - A European Journal, 2019, 25, 743-749.	3.3	12
13	Sulfonylimidamides: Synthesis and Applications in Preparative Organic Chemistry. Advanced Synthesis and Catalysis, 2018, 360, 2976-3001.	4.3	77
14	Direct Synthesis of <i>N</i>-â€Acyl Sulfonylimidamides and <i>N</i>-â€Sulfonylimidoyl Amidines from Sulfonylimidoyl Azides. Advanced Synthesis and Catalysis, 2018, 360, 2465-2469.	4.3	15
15	CuBr/TBHP-mediated synthesis of N-acyl sulfonylimidamides via the oxidative cross-coupling of sulfonylimidamides and aldehydes. Organic and Biomolecular Chemistry, 2017, 15, 2234-2239.	2.8	15
16	An efficient Cu-catalyzed microwave-assisted synthesis of diaryl sulfones. Synthetic Communications, 2017, 47, 319-323.	2.1	17
17	Cuâ€Catalysed Mild Synthesis of <i>N</i>-imidoyl and <i>N</i>-Oximidoyl Sulfonylimidamides through the Threeâ€Component Coupling of Sulfonylimidamides, Azides, and Alkynes. European Journal of Organic Chemistry, 2017, 2017, 6633-6638.	2.4	18
18	Catalyst-Controlled Straightforward Synthesis of Highly Substituted Pyrroles/Furans via Propargylation/Cycloisomerization of \pm-Oxoketene-N,S-acetals. Journal of Organic Chemistry, 2016, 81, 11909-11915.	3.2	30

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19	p-TSA/Base-Promoted Propargylation/Cyclization of \hat{I}^2 -Ketothioamides for the Regioselective Synthesis of Highly Substituted (Hydro)thiophenes. <i>Journal of Organic Chemistry</i> , 2016, 81, 5824-5836.	3.2	35
20	An Efficient Protecting-Group-Free Synthesis of Vinylic Sulfoximines via Horner-Wadsworth-Emmons Reaction. <i>Synlett</i> , 2016, 27, 1423-1427.	1.8	9
21	Pd-catalyzed C-N coupling of vinylbromides and sulfonimidamides: a facile synthesis of N^{ϵ^2} -vinylsulfonimidamides. <i>RSC Advances</i> , 2015, 5, 62084-62090.	3.6	12
22	Cu(OAc) ₂ -Catalysed Oxidative Dual C-H/N-H Activation of Terminal Alkynes and Deprotected Sulfonimidamides: An Easy Access to Alkynylated Sulfonimidamides. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 2861-2867.	2.4	27
23	Iron-Promoted Domino Annulation of \hat{I}^{\pm} -Enolic Dithioesters with Ninhydrin under Solvent-Free Conditions: Chemoselective Direct Access to Indeno[1,2-b]thiophenes. <i>European Journal of Organic Chemistry</i> , 2014, 2014, 5501-5508.	2.4	12
24	Cu(OAc) ₂ promoted Chan-Evans-Lam C-N cross coupling reactions on the N- and N^{ϵ^2} -nitrogen atoms of sulfonimidamides with aryl boronic acids. <i>Tetrahedron</i> , 2014, 70, 5428-5433.	1.9	23
25	Selectfluor-mediated mild oxidative halogenation and thiocyanation of 1-aryl-allenes with TMSX (X=Cl, Br, I, NCS) and NH ₄ SCN. <i>Tetrahedron Letters</i> , 2014, 55, 2401-2405.	1.4	34
26	\hat{I}^2 -Oxodithioesters: a new frontier for diverse heterocyclic architectures. <i>RSC Advances</i> , 2013, 3, 14183.	3.6	53
27	Y(OTf) ₃ catalyzed substitution dependent oxidative C(sp ³)-C(sp ³) cleavage and regioselective dehydration of \hat{I}^2 -allyl- \hat{I}^2 -hydroxydithioesters: alternate route to \hat{I}^{\pm} , \hat{I}^2 -unsaturated ketones and functionalized dienes. <i>Tetrahedron</i> , 2013, 69, 8899-8903.	1.9	12
28	Electrophilic Addition of Propargylic Cations to Allenes: Formation of Crowded Chloro- and Azido-Enynes by Trapping of the Resulting Allylic Cations with TMSX (X = Cl, N ₃): A Synthetic and Computational Study. <i>European Journal of Organic Chemistry</i> , 2013, 2013, 5455-5463.	2.4	7
29	Mild conversion of propargylic alcohols to \hat{I}^{\pm} , \hat{I}^2 -unsaturated enones in ionic liquids (ILs); a new metal free life for the Rupe rearrangement. <i>Tetrahedron Letters</i> , 2013, 54, 6258-6263.	1.4	19
30	Schmidt reaction in ionic liquids: highly efficient and selective conversion of aromatic and heteroaromatic aldehydes to nitriles with [BMIM(SO ₃ H)][OTf] as catalyst and [BMIM][PF ₆] as solvent. <i>Tetrahedron Letters</i> , 2013, 54, 2177-2179.	1.4	34
31	Organocatalyzed straightforward synthesis of highly fluorescent 3,5-disubstituted 2,6-dicyanoanilines via domino annulation of \hat{I}^{\pm} -enolicdithioesters with malononitrile. <i>RSC Advances</i> , 2013, 3, 5345.	3.6	12
32	DABCO-Promoted three-component regioselective synthesis of functionalized chromen-5-ones and pyrano[3,2-c]chromen-5-ones via direct annulation of \hat{I}^{\pm} -oxoketene-N,S-arylaminoacetals under solvent-free conditions. <i>Green Chemistry</i> , 2012, 14, 447.	9.0	67
33	Highly convergent one-pot four-component regioselective synthesis of 4H-benzo[f]chromenes via annulation of \hat{I}^2 -oxodithioesters. <i>Tetrahedron</i> , 2012, 68, 1247-1252.	1.9	33
34	Highly Regioselective One-Pot, Three-Component Synthesis of 1-Aryl-3,4-Substituted/Annulated-5-(Cycloamino)/(Alkylamino)pyrazoles from \hat{I}^2 -Oxodithioesters. <i>European Journal of Organic Chemistry</i> , 2012, 2012, 967-974.	2.4	54
35	One-Pot Two-Component [3 + 2] Cycloaddition/Annulation Protocol for the Synthesis of Highly Functionalized Thiophene Derivatives. <i>Journal of Organic Chemistry</i> , 2011, 76, 8009-8014.	3.2	90
36	Regioselective Synthesis of Tetrahydrothiochromen-5-ones via a One-Pot Three-Component Solvent-Free Domino Protocol. <i>Organic Letters</i> , 2011, 13, 3762-3765.	4.6	67

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37	Silica-Gel Catalyzed Efficient Synthesis of Quinoxaline Derivatives Under Solvent-Free Conditions. <i>Synthetic Communications</i> , 2011, 41, 417-425.	2.1	25
38	l-Proline catalyzed synthesis of densely functionalized pyrido[2,3-d]pyrimidines via three-component one-pot domino Knoevenagel aza-Diels-Alder reaction. <i>Tetrahedron</i> , 2011, 67, 5935-5941.	1.9	62
39	A facile approach for the synthesis of 14-aryl- or alkyl-14H-dibenzo[a,j]xanthenes under solvent-free condition. <i>Tetrahedron Letters</i> , 2010, 51, 442-445.	1.4	94
40	An efficient and facile one-pot synthesis of propargylamines by three-component coupling of aldehydes, amines, and alkynes via C-H activation catalyzed by NiCl ₂ . <i>Tetrahedron Letters</i> , 2010, 51, 5555-5558.	1.4	135
41	First InCl ₃ -Catalyzed, Three-Component Coupling of Aldehydes, 1 ² -Naphthol, and 6-Amino-1,3-dimethyluracil to Functionalized Naphthopyranopyrimidines. <i>Synlett</i> , 2010, 2010, 1133-1137.	1.8	10
42	Biginelli and Hantzsch-Type Reactions Leading to Highly Functionalized Dihydropyrimidinone, Thiocoumarin, and Pyridopyrimidinone Frameworks via Ring Annulation with 1 ² -Oxidithioesters. <i>Journal of Organic Chemistry</i> , 2010, 75, 7785-7795.	3.2	88
43	An efficient one-pot synthesis of tetrahydrobenzo[a]xanthene-11-one and diazabenz[a]anthracene-9,11-dione derivatives under solvent free condition. <i>Tetrahedron</i> , 2009, 65, 7129-7134.	1.9	198
44	l-Proline: an efficient catalyst for the one-pot synthesis of 2,4,5-trisubstituted and 1,2,4,5-tetrasubstituted imidazoles. <i>Tetrahedron</i> , 2009, 65, 10155-10161.	1.9	251
45	Multicomponent one-pot solvent-free synthesis of functionalized unsymmetrical dihydro-1H-indeno[1,2-b]pyridines. <i>Tetrahedron Letters</i> , 2009, 50, 7096-7098.	1.4	72
46	Atom-efficient and environment-friendly multicomponent synthesis of amidoalkyl naphthols catalyzed by P ₂ O ₅ . <i>Tetrahedron Letters</i> , 2009, 50, 7220-7222.	1.4	131