

Nagula Shankaraiah

List of Publications by Year in descending order

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156
papers

4,670
citations

81743

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155451

55
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175
all docs

175
docs citations

175
times ranked

3873
citing authors

#	ARTICLE	IF	CITATIONS
1	Recent advances in multi-component reactions and their mechanistic insights: a triennium review. <i>Organic Chemistry Frontiers</i> , 2021, 8, 4237-4287.	2.3	158
2	Synthesis of 1,2,3-triazole-linked pyrrolobenzodiazepine conjugates employing "click" chemistry: DNA-binding affinity and anticancer activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 1468-1473.	1.0	145
3	Synthesis and anticancer activity of chalcone-pyrrolobenzodiazepine conjugates linked via 1,2,3-triazole ring side-armed with alkane spacers. <i>European Journal of Medicinal Chemistry</i> , 2011, 46, 3820-3831.	2.6	124
4	Spirooxindole-derived morpholine-fused-1,2,3-triazoles: Design, synthesis, cytotoxicity and apoptosis inducing studies. <i>European Journal of Medicinal Chemistry</i> , 2015, 102, 413-424.	2.6	107
5	Water mediated Heck and Ullmann couplings by supported palladium nanoparticles: importance of surface polarity of the carbon spheres. <i>Green Chemistry</i> , 2012, 14, 2513.	4.6	91
6	DNA-binding affinity and anticancer activity of \hat{I}^2 -carboline"chalcone conjugates as potential DNA intercalators: Molecular modelling and synthesis. <i>Bioorganic Chemistry</i> , 2015, 59, 130-139.	2.0	83
7	Synthesis and biological evaluation of new benzimidazole-thiazolidinedione hybrids as potential cytotoxic and apoptosis inducing agents. <i>European Journal of Medicinal Chemistry</i> , 2016, 124, 608-621.	2.6	80
8	Design and synthesis of dithiocarbamate linked \hat{I}^2 -carboline derivatives: DNA topoisomerase II inhibition with DNA binding and apoptosis inducing ability. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 5511-5526.	1.4	79
9	Design and synthesis of C3-tethered 1,2,3-triazolo- \hat{I}^2 -carboline derivatives: Anticancer activity, DNA-binding ability, viscosity and molecular modeling studies. <i>Bioorganic Chemistry</i> , 2016, 64, 42-50.	2.0	77
10	Solid-phase synthesis of new pyrrolobenzodiazepine"chalcone conjugates: DNA-binding affinity and anticancer activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 2434-2439.	1.0	72
11	Design and Synthesis of C3"Pyrazole/Chalcone"Linked Beta"Carboline Hybrids: Antitopoisomerase...I, DNA"Interactive, and Apoptosis"Inducing Anticancer Agents. <i>ChemMedChem</i> , 2014, 9, 2084-2098.	1.6	72
12	Novel Supramolecular Palladium Catalyst for the Asymmetric Reduction of Imines in Aqueous Media. <i>Organic Letters</i> , 2009, 11, 3238-3241.	2.4	71
13	Copper Oxide Nanoparticles Supported on Graphene Oxide" Catalyzed S"arylation: An Efficient and Ligand"Free Synthesis of Aryl Sulfides. <i>Advanced Synthesis and Catalysis</i> , 2013, 355, 2297-2307.	2.1	69
14	Conventional and microwave-assisted synthesis of new 1 H -benzimidazole-thiazolidinedione derivatives: A potential anticancer scaffold. <i>European Journal of Medicinal Chemistry</i> , 2017, 138, 234-245.	2.6	66
15	Recent Advances in the Solid-Phase Combinatorial Synthetic Strategies for the Quinoxaline, Quinazoline and Benzimidazole Based Privileged Structures. <i>Mini-Reviews in Medicinal Chemistry</i> , 2006, 6, 71-89.	1.1	65
16	H ₂ O-mediated isatin spiro-epoxide ring opening with NaCN: Synthesis of novel 3-tetrazolylmethyl-3-hydroxy-oxindole hybrids and their anticancer evaluation. <i>European Journal of Medicinal Chemistry</i> , 2015, 104, 11-24.	2.6	61
17	Exploration of carbamide derived pyrimidine-thioindole conjugates as potential VEGFR-2 inhibitors with anti-angiogenesis effect. <i>European Journal of Medicinal Chemistry</i> , 2020, 200, 112457.	2.6	61
18	Design, synthesis and apoptosis inducing effect of novel (Z)-3-(3"methoxy-4"-(2-amino-2-oxoethoxy)-benzylidene)indolin-2-ones as potential antitumour agents. <i>European Journal of Medicinal Chemistry</i> , 2016, 118, 34-46.	2.6	60

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19	Structural insights of oxindole based kinase inhibitors as anticancer agents: Recent advances. <i>European Journal of Medicinal Chemistry</i> , 2021, 216, 113334.	2.6	58
20	Recent Advances in the Solid-Phase Combinatorial Synthetic Strategies for the Benzodiazepine Based Privileged Structures. <i>Mini-Reviews in Medicinal Chemistry</i> , 2006, 6, 53-69.	1.1	56
21	evaluation and apoptosis inducing studies. <i>European Journal of Medicinal Chemistry</i> , 2016, 122, 584-600.	2.6	55
22	Synthesis of podophyllotoxin linked \hat{I}^2 -carboline congeners as potential anticancer agents and DNA topoisomerase II inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2018, 144, 557-571.	2.6	55
23	Chemoselective Aromatic Azido Reduction with Concomitant Aliphatic Azide Employing Al/Gd Triflates/NaI and ESI-MS Mechanistic Studies. <i>Chemistry - A European Journal</i> , 2009, 15, 7215-7224.	1.7	53
24	Enantioselective total syntheses of ropivacaine and its analogues. <i>Tetrahedron Letters</i> , 2008, 49, 5098-5100.	0.7	52
25	Anticancer potential of spirocompounds in medicinal chemistry: A pentennial expedition. <i>European Journal of Medicinal Chemistry</i> , 2021, 215, 113263.	2.6	50
26	Targeting DNA Minor Groove by Hybrid Molecules as Anticancer Agents. <i>Current Medicinal Chemistry</i> , 2017, 24, 2887-2907.	1.2	50
27	TBAI/TBHP-catalyzed [3 + 2] cycloaddition/oxidation/aromatization cascade and online ESI-MS mechanistic studies: synthesis of pyrrolo[2,1- <i>a</i>]isoquinolines and indolino[8,7- <i>b</i>]indoles. <i>RSC Advances</i> , 2016, 6, 2671-2677.	1.7	47
28	Design and Synthesis of DNA-Interactive \hat{I}^2 -Carboline-Oxindole Hybrids as Cytotoxic and Apoptosis-Inducing Agents. <i>ChemMedChem</i> , 2018, 13, 1909-1922.	1.6	47
29	Microwave-assisted direct oxidative synthesis of \hat{I}^{\pm} -ketoamides from aryl methyl ketones and amines by a water soluble Cu(<i>scp</i>)-complex. <i>Green Chemistry</i> , 2016, 18, 3439-3447.	4.6	46
30	Synthesis and <i>in vitro</i> cytotoxicity evaluation of \hat{I}^2 -carboline-linked 2,4-thiazolidinedione hybrids: potential DNA intercalation and apoptosis-inducing studies. <i>New Journal of Chemistry</i> , 2018, 42, 16226-16236.	1.4	45
31	An insight into medicinal attributes of dithiocarbamates: Bird's eye view. <i>Bioorganic Chemistry</i> , 2020, 105, 104346.	2.0	45
32	Syntheses and Applications of Spirocyclopropyl Oxindoles: A Decade Review. <i>European Journal of Organic Chemistry</i> , 2021, 2021, 757-772.	1.2	45
33	Solid-Phase Synthesis of a Library of Pyrrolo[2,1- <i>c</i>][1,4]benzodiazepine-5,11-diones with Potential Antitubercular Activity. <i>ACS Combinatorial Science</i> , 2007, 9, 29-42.	3.3	43
34	Silver(<i>scp</i>)-catalysed domino alkyne-annulation/Diels-Alder reaction: a mild synthetic approach to tetrahydrospiro[carbazole-4,3- \hat{I}^2 -indoline] scaffolds. <i>Organic Chemistry Frontiers</i> , 2016, 3, 1503-1508.	2.3	43
35	Silver catalyzed domino aza-annulation/Diels-Alder cyclization of 2-ene-yne anilines: a facile one-pot access to carbazole, dihydrocarbazole and tetrahydrocarbazole frameworks. <i>Chemical Communications</i> , 2016, 52, 4581-4584.	2.2	43
36	Synthesis of 2,3,6,7-tetramethoxyphenanthren-9-amine: An efficient precursor to access new 4-aza-2,3-dihydropyridophenanthrenes as apoptosis inducing agents. <i>European Journal of Medicinal Chemistry</i> , 2017, 127, 305-317.	2.6	43

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37	Enantioselective total synthesis of (S)-($\hat{\alpha}$)-quinolactacin B. <i>Tetrahedron Letters</i> , 2008, 49, 4289-4291.	0.7	42
38	An efficient one-pot synthesis of benzothiazolo-4 $\hat{\beta}$ -anilino-podophyllotoxin congeners: DNA topoisomerase-II inhibition and anticancer activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 350-353.	1.0	42
39	Synthesis of 1,2,4-triazole-linked urea/thiourea conjugates as cytotoxic and apoptosis inducing agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018, 28, 1919-1924.	1.0	42
40	Design, synthesis and anticancer evaluation of tetrahydro- $\hat{\beta}$ -carboline-hydantoin hybrids. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 5413-5417.	1.0	41
41	Cinnamide derived pyrimidine-benzimidazole hybrids as tubulin inhibitors: Synthesis, in silico and cell growth inhibition studies. <i>Bioorganic Chemistry</i> , 2021, 110, 104765.	2.0	41
42	Synthesis and biological evaluation of podophyllotoxin congeners as tubulin polymerization inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 5466-5475.	1.4	40
43	An efficient one-pot decarboxylative aromatization of tetrahydro- $\hat{\beta}$ -carbolines by using N-chlorosuccinimide: total synthesis of norharmine, harmine and eudistomins. <i>RSC Advances</i> , 2015, 5, 90121-90126.	1.7	39
44	Synthesis of 2-aryl-1,2,4-oxadiazolo-benzimidazoles: Tubulin polymerization inhibitors and apoptosis inducing agents. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 4608-4623.	1.4	38
45	Selective reduction of aromatic azides in solution/solid-phase and resin cleavage by employing BF $\hat{3}$ -OEt $\hat{2}$ /EtSH. Preparation of DC-81. <i>Tetrahedron Letters</i> , 2006, 47, 4253-4257.	0.7	37
46	Palladium-catalyzed Aryl C-H Activation and Tandem ortho-hydroxylation/alkoxylation of 2-aryl benzimidazoles: Cytotoxicity and DNA-binding studies. <i>Asian Journal of Organic Chemistry</i> , 2014, 3, 68-76.	1.3	37
47	Synthesis of different heterocycles-linked chalcone conjugates as cytotoxic agents and tubulin polymerization inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 4805-4816.	1.4	36
48	Design and synthesis of substituted dihydropyrimidinone derivatives as cytotoxic and tubulin polymerization inhibitors. <i>Bioorganic Chemistry</i> , 2019, 93, 103317.	2.0	36
49	Design and synthesis of DNA-intercalative naphthalimide-benzothiazole/cinnamide derivatives: cytotoxicity evaluation and topoisomerase-III inhibition. <i>MedChemComm</i> , 2019, 10, 72-79.	3.5	36
50	Synthesis and in vitro cytotoxicity evaluation of $\hat{\beta}$ -carboline-combretastatin carboxamides as apoptosis inducing agents: DNA intercalation and topoisomerase-II inhibition. <i>Bioorganic and Medicinal Chemistry</i> , 2019, 27, 3285-3298.	1.4	34
51	Sulfamic acid promoted one-pot synthesis of phenanthrene fused-dihydrodibenzo-quinolinones: Anticancer activity, tubulin polymerization inhibition and apoptosis inducing studies. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 1996-2008.	1.4	33
52	Expedition of sulfur-containing heterocyclic derivatives as cytotoxic agents in medicinal chemistry: A decade update. <i>Medicinal Research Reviews</i> , 2022, 42, 513-575.	5.0	33
53	Enantioselective total synthesis of pyrroloquinolone as a potent PDE5 inhibitor. <i>Tetrahedron Letters</i> , 2009, 50, 520-523.	0.7	32
54	Synthesis of Combretastatin A4 Carboxamide that Mimic Sulfonyl Piperazines by a Molecular Hybridization Approach: in vitro Cytotoxicity Evaluation and Inhibition of Tubulin Polymerization. <i>ChemMedChem</i> , 2019, 14, 2052-2060.	1.6	32

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55	Syntheses and reactivity of spiro-epoxy/aziridine oxindole cores: developments in the past decade. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 8572-8596.	1.5	32
56	Synthesis of bis-1,2,3-triazolo-bridged unsymmetrical pyrrolobenzodiazepine trimers via <i>click</i> TM chemistry and their DNA-binding studies. <i>Tetrahedron</i> , 2010, 66, 5498-5506.	1.0	31
57	A one-pot <i>click</i> TM reaction from spiro-epoxides catalyzed by Cu(<i>scpd</i>)-pyrrolidinyl-oxazole-carboxamide. <i>New Journal of Chemistry</i> , 2015, 39, 3973-3981.	1.4	31
58	\hat{I}^2 -Carboline-based molecular hybrids as anticancer agents: a brief sketch. <i>RSC Medicinal Chemistry</i> , 2021, 12, 730-750.	1.7	30
59	An update on the progress of cycloaddition reactions of 3-methyleneindolinones in the past decade: versatile approaches to spirooxindoles. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 7768-7791.	1.5	30
60	Ru(II)-Catalyzed Regioselective Hydroxymethylation of \hat{I}^2 -Carbolines and Isoquinolines via C ^H Functionalization: Probing the Mechanism by Online ESI-MS/MS Screening. <i>Journal of Organic Chemistry</i> , 2019, 84, 5504-5513.	1.7	29
61	Solid-phase synthesis of fused [2,1- <i>b</i>]quinazolinone alkaloids. <i>Tetrahedron Letters</i> , 2006, 47, 9025-9028.	0.7	28
62	4 \hat{I}^2 -[4 \hat{E}^2 -(1-(Aryl)ureido)benzamide]podophyllotoxins as DNA topoisomerase I and III \pm inhibitors and apoptosis inducing agents. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 5198-5208.	1.4	28
63	Microwave-assisted one-pot synthesis of new phenanthrene fused-tetrahydrodibenzo-acridinones as potential cytotoxic and apoptosis inducing agents. <i>European Journal of Medicinal Chemistry</i> , 2018, 151, 173-185.	2.6	28
64	Design and synthesis of thiadiazolo-carboxamide bridged \hat{I}^2 -carboline-indole hybrids: DNA intercalative topo-III \pm inhibition with promising antiproliferative activity. <i>Bioorganic Chemistry</i> , 2020, 105, 104357.	2.0	28
65	Development of \hat{I}^2 -carboline-benzothiazole hybrids via carboxamide formation as cytotoxic agents: DNA intercalative topoisomerase III \pm inhibition and apoptosis induction. <i>Bioorganic Chemistry</i> , 2021, 106, 104481.	2.0	28
66	Synthesis of DNA-Interactive Pyrrolo[2,1- <i>c</i>][1,4]benzodiazepines by Employing Polymer-Supported Reagents: Preparation of DC-81. <i>Synlett</i> , 2004, 2004, 2533-2536.	1.0	27
67	Molecular iodine-catalysed oxidative CO ^H -C(alkyl) bond cleavage of aryl/heteroaryl alkyl ketones: an efficient strategy to access fused polyheterocycles. <i>New Journal of Chemistry</i> , 2018, 42, 15820-15829.	1.4	27
68	Microwave-Assisted One-Pot [3+2] Cycloaddition of Azomethine Ylides and \hat{I}^2 -Alkenyl Oxindoles: A Facile Approach to Pyrrolidine-Fused Bis-Spirooxindoles. <i>ChemistrySelect</i> , 2019, 4, 1727-1730.	0.7	27
69	Microwave-assisted multicomponent reactions in heterocyclic chemistry and mechanistic aspects. <i>Beilstein Journal of Organic Chemistry</i> , 2021, 17, 819-865.	1.3	27
70	An efficient solid-phase synthesis of biologically important DNA-interactive pyrrolo[2,1- <i>c</i>][1,4]benzodiazepine dimers (DSB-120) and their C2-fluorinated analogues. <i>Tetrahedron Letters</i> , 2006, 47, 6553-6556.	0.7	26
71	Conversion of Amines to Imines Employing Polymer-Supported Sulfoxide (PSS) and Polymer-Supported Perruthenate (PSP): Synthesis of Pyrrolo[2,1- <i>c</i>][1,4]benzodiazepines. <i>Advanced Synthesis and Catalysis</i> , 2006, 348, 249-254.	2.1	26
72	An Efficient Selective Reduction of Aromatic Azides to Amines Employing BF ₃ ·OEt ₂ /NaI: Synthesis of Pyrrolobenzodiazepines. <i>Synlett</i> , 2008, 2008, 1297-1300.	1.0	25

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73	Synthesis and biological evaluation of 4-aza-2,3-dihydropyridophenanthrolines as tubulin polymerization inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 3356-3360.	1.0	25
74	Synthesis of substituted biphenyl methylene indolinones as apoptosis inducers and tubulin polymerization inhibitors. <i>Bioorganic Chemistry</i> , 2019, 86, 210-223.	2.0	25
75	Synthesis and potential cytotoxic activity of new phenanthrylphenol-pyrrolobenzodiazepines. <i>European Journal of Medicinal Chemistry</i> , 2010, 45, 2173-2181.	2.6	24
76	Asymmetric Michael addition of ketones to nitroolefins: pyrrolidinyl-oxazole-carboxamides as new efficient organocatalysts. <i>Organic and Biomolecular Chemistry</i> , 2014, 12, 8008-8018.	1.5	24
77	One-pot synthesis of podophyllotoxin thiourea congeners by employing NH ₂ SO ₃ H/NaI: Anticancer activity, DNA topoisomerase-II inhibition, and apoptosis inducing agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 4239-4244.	1.0	24
78	Synthesis of substituted phenanthrene-9-benzimidazole conjugates: Cytotoxicity evaluation and apoptosis inducing studies. <i>European Journal of Medicinal Chemistry</i> , 2017, 140, 128-140.	2.6	24
79	Synthesis of DNA interactive C3-trans-cinnamide linked \hat{I}^2 -carboline conjugates as potential cytotoxic and DNA topoisomerase I inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 4916-4929.	1.4	24
80	Synthesis and biological evaluation of substituted N-(2-(1H-benzo[d]imidazol-2-yl)phenyl)cinnamides as tubulin polymerization inhibitors. <i>Bioorganic Chemistry</i> , 2020, 103, 104191.	2.0	24
81	Design and synthesis of 4-O-alkylamino-tethered-benzylideneindolin-2-ones as potent cytotoxic and apoptosis inducing agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 4061-4069.	1.0	23
82	Iron-Mediated One-Pot Synthesis of 3,5-Diarylpyridines from \hat{I}^2 -Nitrostyrenes. <i>Journal of Organic Chemistry</i> , 2016, 81, 2159-2165.	1.7	23
83	Reliability of Click Chemistry on Drug Discovery: A Personal Account. <i>Chemical Record</i> , 2020, 20, 253-272.	2.9	23
84	Synthesis of Novel C3-Linked \hat{I}^2 -Carboline-Pyridine Derivatives Employing Khronke Reaction: DNA-binding Ability and Molecular Modeling Studies. <i>Letters in Drug Design and Discovery</i> , 2016, 13, 335-342.	0.4	23
85	A One-Pot Azido Reductive Tandem Mono-N-Alkylation Employing Dialkylboron Triflates: Online ESI-MS Mechanistic Investigation. <i>Journal of Organic Chemistry</i> , 2011, 76, 7017-7026.	1.7	22
86	Synthesis of 1,2,3-triazolo fused tetrahydro \hat{I}^2 -carboline Derivatives via 1,3-Dipolar Cycloaddition Reaction; Cytotoxicity Evaluation and DNA Binding studies. <i>ChemistrySelect</i> , 2017, 2, 7210-7221.	0.7	22
87	Diverse Targeted Approaches to Battle Multidrug Resistance in Cancer. <i>Current Medicinal Chemistry</i> , 2019, 26, 7059-7080.	1.2	22
88	Synthesis of C8-C8-linked triazolo pyrrolobenzodiazepine dimers by employing "click" chemistry and their DNA-binding affinity. <i>Tetrahedron Letters</i> , 2008, 49, 3620-3624.	0.7	21
89	A recyclable and water soluble copper (<sc>i</sc>)-catalyst: one-pot synthesis of 1,4-disubstituted 1,2,3-triazoles and their biological evaluation. <i>RSC Advances</i> , 2016, 6, 103556-103566.	1.7	20
90	Short synthesis of noscapine, bicuculline, egenine, capnoidine, and corytensine alkaloids through the addition of 1-siloxy-isobenzofurans to imines. <i>Tetrahedron Letters</i> , 2010, 51, 1770-1773.	0.7	19

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91	Design and synthesis of 1,2,3-triazolo-phenanthrene hybrids as cytotoxic agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 2369-2376.	1.0	19
92	A Polymer-Assisted Solution-Phase Strategy for the Synthesis of Fused [2,1-b]Quinazolinones and the Preparation of Optically Active Vasicinone. <i>Synlett</i> , 2006, 2006, 2609-2612.	1.0	18
93	Enantioselective total synthesis of (S)-(+)-lennoxamine through asymmetric hydrogenation mediated by l-proline-tetrazole ruthenium catalyst. <i>Tetrahedron Letters</i> , 2012, 53, 3672-3675.	0.7	18
94	Dithiocarbamate/piperazine bridged pyrrolobenzodiazepines as DNA-minor groove binders: Synthesis, DNA-binding affinity and cytotoxic activity. <i>Bioorganic Chemistry</i> , 2015, 59, 23-30.	2.0	18
95	Iodine-mediated C-N and N-N bond formation: a facile one-pot synthetic approach to 1,2,3-triazoles under metal-free and azide-free conditions. <i>RSC Advances</i> , 2019, 9, 27021-27031.	1.7	18
96	The microwave-assisted syntheses and applications of non-fused single-nitrogen-containing heterocycles. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 9737-9761.	1.5	18
97	Efficient Solid-Phase Synthesis of a Library of Imidazo[1,2-a]pyridine-8-carboxamides. <i>ACS Combinatorial Science</i> , 2007, 9, 267-274.	3.3	17
98	Short Total Synthesis of (-)-Lupinine and (-)-Epiquinamide by Double Mitsunobu Reaction. <i>Synthesis</i> , 2011, 2011, 51-56.	1.2	17
99	$\hat{2}$ -Carboline directed regioselective hydroxylation by employing Cu(OAc) ₂ and mechanistic investigation by ESI-MS. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 2307-2311.	1.5	17
100	Application of Transition Metal-Catalyzed C-H Activation Strategies in the Synthesis and Functionalization of $\hat{2}$ -Carbolines. <i>Asian Journal of Organic Chemistry</i> , 2021, 10, 1050-1066.	1.3	17
101	Unravelling KDM4 histone demethylase inhibitors for cancer therapy. <i>Drug Discovery Today</i> , 2021, 26, 1841-1856.	3.2	17
102	TCCA-mediated oxidative rearrangement of tetrahydro- $\hat{2}$ -carbolines: facile access to spirooxindoles and the total synthesis of ($\hat{\pm}$)-coerulescine and ($\hat{\pm}$)-horsfiline. <i>RSC Advances</i> , 2021, 11, 16537-16546.	1.7	16
103	A facile intramolecular azido/amido reductive cyclization approach: synthesis of pyrrolobenzodiazepines and their dimers. <i>Tetrahedron Letters</i> , 2008, 49, 1465-1468.	0.7	15
104	Studies towards the construction of quaternary indolizidines by [2,3]-sigmatropic rearrangement cocatalyzed by ionic liquid. <i>Journal of the Brazilian Chemical Society</i> , 2009, 20, 813-819.	0.6	15
105	Facile and Efficient Solid-Phase Synthesis of DNA-Interactive Pyrrolo[2,1-c][1,4]benzodiazepines. <i>Synlett</i> , 2004, 2004, 1841-1843.	1.0	14
106	A new approach for the solid-phase synthesis of pyrrolo[2,1-c][1,4]benzodiazepines involving reductive cleavage. <i>Tetrahedron Letters</i> , 2004, 45, 7667-7669.	0.7	14
107	Asymmetric syntheses of piperidino-benzodiazepines through \hat{c} ation-pool TM host/guest supramolecular approach and their DNA-binding studies. <i>Tetrahedron: Asymmetry</i> , 2010, 21, 2625-2630.	1.8	14
108	Synthesis of enamino-2-oxindoles via conjugate addition between $\hat{\pm}$ -azido ketones and 3-alkenyl oxindoles: Cytotoxicity evaluation and apoptosis inducing studies. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2018, 28, 3564-3573.	1.0	14

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109	Synthesis and biological evaluation of novel imidazo[1,2-a]pyridine-oxadiazole hybrids as anti-proliferative agents: Study of microtubule polymerization inhibition and DNA binding. <i>Bioorganic and Medicinal Chemistry</i> , 2021, 43, 116277.	1.4	14
110	Total Synthesis of Rutaecarpine and Analogues by Tandem Azido Reductive Cyclization Assisted by Microwave Irradiation. <i>Synlett</i> , 2011, 2011, 61-64.	1.0	13
111	Î²-Carboline tethered cinnamoyl 2-aminobenzamides as class I selective HDAC inhibitors: Design, synthesis, biological activities and modelling studies. <i>Bioorganic Chemistry</i> , 2021, 117, 105461.	2.0	13
112	AlCl ₃ •NaI assisted cleavage of polymer-bound esters with concomitant amine coupling and azido-reductive cyclization: synthesis of pyrrolobenzodiazepine derivatives. <i>Tetrahedron Letters</i> , 2013, 54, 4435-4441.	0.7	12
113	Iodine-promoted one-pot synthesis of 1,3,4-oxadiazole scaffolds <i>via</i> sp ³ C-H functionalization of azaarenes. <i>New Journal of Chemistry</i> , 2019, 43, 15999-16006.	1.4	12
114	Stereoselective Aldol and Conjugate Addition Reactions Mediated by Proline-Based Catalysts and Its Analogues: A Concise Review. <i>European Journal of Organic Chemistry</i> , 2021, 2021, 5288-5311.	1.2	12
115	Transition metal-free one-pot synthesis of substituted pyrroles by employing aza-Wittig reaction. <i>RSC Advances</i> , 2019, 9, 30659-30665.	1.7	11
116	Contribution of Knoevenagel Condensation Products toward the Development of Anticancer Agents: An Updated Review. <i>ChemMedChem</i> , 2022, 17, .	1.6	11
117	Synthesis of indolo/pyrroloazepinone-oxindoles as potential cytotoxic, DNA-intercalating and Topo I inhibitors. <i>Bioorganic Chemistry</i> , 2022, 122, 105706.	2.0	11
118	H ₂ O ₂ -Mediated Epoxide Ring-Opening with Concomitant S Bond Formation: A One-Pot Method to 3-Hydroxy-oxindolino-dithiocarbamates as Cytotoxic Agents. <i>ChemistrySelect</i> , 2018, 3, 6766-6774.	0.7	10
119	Lewis-acid catalyzed dehydrative [3+2] cycloaddition reaction: A facile synthetic approach to spiro-benzoindoline oxindoles. <i>Tetrahedron Letters</i> , 2020, 61, 152007.	0.7	10
120	FeCl ₃ Catalyzed [3+2] Cycloaddition Reaction: A Mild Synthetic Approach to Spirooxindolo-2-aminothiazolidine Scaffolds. <i>ChemistrySelect</i> , 2020, 5, 2886-2891.	0.7	10
121	Design and Synthesis of 5-Morpholino-Thiophene-Indole/ Oxindole Hybrids as Cytotoxic Agents. <i>ChemistrySelect</i> , 2020, 5, 4356-4363.	0.7	10
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