

Suresh Kumar Raju

List of Publications by Year in descending order

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210
papers

3,748
citations

126907

33
h-index

189892

50
g-index

231
all docs

231
docs citations

231
times ranked

2705
citing authors

#	ARTICLE	IF	CITATIONS
1	Bio-inspired silver nanoparticles from <i>Artocarpus lakoocha</i> fruit extract and evaluation of their antibacterial activity and anticancer activity on human prostate cancer cell line. <i>Applied Nanoscience (Switzerland)</i> , 2023, 13, 3041-3051.	3.1	10
2	Characterization, antimicrobial activity and anticancer activity of <i>Pyrostegia venusta</i> leaf extract-synthesized silver nanoparticles against COS-7 cell line. <i>Applied Nanoscience (Switzerland)</i> , 2023, 13, 2303-2314.	3.1	10
3	Biomimetic synthesis of silver nanoparticles using <i>Cucumis sativus</i> var. <i>hardwickii</i> fruit extract and their characterizations, anticancer potential and apoptosis studies against Pa-1 (Human ovarian) Tj ETQq1 1 0.784314 rgBT / Overlock	3.1	10
4	Raman Spectroscopic and Electrochemical Measurements of Dynamic Shocked MnFe ₂ O ₄ Nano-crystalline Materials. <i>Journal of Inorganic and Organometallic Polymers and Materials</i> , 2022, 32, 344-352.	3.7	9
5	Synthesis, structure and biological evaluation of highly functionalized 2-azetidinone integrated spirooxindolopyrrolidine heterocyclic hybrid. <i>Journal of Molecular Structure</i> , 2022, 1250, 131745.	3.6	2
6	Cardioprotective effects of phytopigments via multiple signaling pathways. <i>Phytomedicine</i> , 2022, 95, 153859.	5.3	8
7	Bioprospection and secondary metabolites profiling of marine <i>Streptomyces levis</i> strain KS46. <i>Saudi Journal of Biological Sciences</i> , 2022, 29, 667-679.	3.8	18
8	Synthesis and antimicrobial potential of spirooxindolopyrrolidine tethered oxindole heterocyclic hybrid against multidrug resistant microbial pathogens. <i>Process Biochemistry</i> , 2022, 114, 66-70.	3.7	2
9	Stereoselective synthesis, structural determination, computational studies and antimicrobial activity of novel class of spiropyrroloquinoxaline engrafted ferrocenoindole hybrid heterocycle. <i>Journal of Molecular Structure</i> , 2022, 1252, 132131.	3.6	2
10	High Shock Resistance of Polycrystalline Sodium Sulfate Crystals at Dynamic Shocked Conditions. <i>Physica Status Solidi (B): Basic Research</i> , 2022, 259, .	1.5	1
11	Dynamic Shock Wave-Induced Amorphous-to-Crystalline Switchable Phase Transition of Lithium Sulfate. <i>Journal of Physical Chemistry C</i> , 2022, 126, 3194-3201.	3.1	16
12	Switchable crystalâ€™amorphous states of NiSO ₄ ·6H ₂ O induced by a Reddy tube. <i>New Journal of Chemistry</i> , 2022, 46, 5091-5099.	2.8	9
13	Reversible magnetic phase transitions of zirconium oxide nanoparticles induced by dynamic shock waves. <i>Applied Physics A: Materials Science and Processing</i> , 2022, 128, 1.	2.3	1
14	Investigation on crystallinity and optical properties of l-tartaric acid single crystal at dynamic shocked conditions. <i>Journal of Materials Science: Materials in Electronics</i> , 2022, 33, 10841-10850.	2.2	5
15	Switchable Phase Transition from Crystalline to Amorphous States of Cadmium Sulfate Octahydrate Single Crystals by Shock Waves. <i>Physica Status Solidi (B): Basic Research</i> , 2022, 259, .	1.5	1
16	Eco-friendly synthesis and structural determination of pyrene fused pyrroloquinolinone hybrid. <i>Journal of Molecular Structure</i> , 2022, 1259, 132714.	3.6	2
17	Label-Free Electrochemical Detection of the Cancer Biomarker Platelet-Derived Growth Factor Receptor in Human Serum and Cancer Cells. <i>ACS Biomaterials Science and Engineering</i> , 2022, 8, 826-833.	5.2	5
18	Investigation of the Optical Properties of a Novel Class of Quinoline Derivatives and Their Random Laser Properties Using ZnO Nanoparticles. <i>Molecules</i> , 2022, 27, 145.	3.8	2

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19	Discovery of spirooxindole-pyrrolidine heterocyclic hybrids with potent antifungal activity against fungal pathogens. <i>Pathogens and Disease</i> , 2022, , .	2.0	0
20	Environmentally friendly domino multicomponent strategy for the synthesis of pyrroloquinolinone hybrid heterocycles. <i>RSC Advances</i> , 2022, 12, 15440-15446.	3.6	3
21	Investigation of the one-step electrochemical deposition of graphene oxide-doped poly(3,4-ethylenedioxythiophene)â€“polyphenol oxidase as a dopamine sensor. <i>RSC Advances</i> , 2022, 12, 15575-15583.	3.6	2
22	Synthesis, computational studies and antibacterial assessment of dispirooxindolopyrrolidine integrated indandione hybrids. <i>Journal of Molecular Structure</i> , 2022, 1267, 133577.	3.6	2
23	Microwave-Assisted Copper(II)-Catalyzed Cascade Cyclization of 2-Propargylamino/Oxy-Arylaldehydes and <i>O</i> -Phenylenediamines: Access to Densely Functionalized Benzo[<i>f</i>]Imidazo[1,2- <i>d</i>][1,4]Oxazepines and Benzo[<i>f</i>]Imidazo[1,2- <i>d</i>][1,4]Diazepines. <i>Journal of Organic Chemistry</i> , 2022, 87, 8956-8969.	3.2	12
24	Design, stereoselective synthesis, computational studies and cholinesterase inhibitory activity of novel spiropyrrolidinoquinoxaline tethered indole hybrid heterocycle. <i>Journal of Molecular Structure</i> , 2021, 1225, 129165.	3.6	6
25	Synthesis, X-ray structural determination and biological evaluation of novel ferrocene grafted spiroquinoxalinopyrrolidine. <i>Journal of Molecular Structure</i> , 2021, 1226, 129348.	3.6	2
26	Ionic liquid mediated synthesis and <i>in vitro</i> mechanistic exploration of polycyclic cage-like heterocyclic hybrid. <i>Journal of Heterocyclic Chemistry</i> , 2021, 58, 580-588.	2.6	5
27	Diastereoselective synthesis and anticancer potential of a small library of cage-like heterocyclic hybrids. <i>Journal of King Saud University - Science</i> , 2021, 33, 101238.	3.5	0
28	A stereo, regioselective synthesis and discovery of antimycobacterium tuberculosis activity of novel Î²-lactam grafted spirooxindolopyrrolidine hybrid heterocycles. <i>Arabian Journal of Chemistry</i> , 2021, 14, 102938.	4.9	13
29	Cholinesterase inhibitory activity of highly functionalized fluorinated spiropyrrolidine heterocyclic hybrids. <i>Saudi Journal of Biological Sciences</i> , 2021, 28, 754-761.	3.8	13
30	Imidazolium ylide mediated tandem Knoevenagelâ€“Michaelâ€“ <i>O</i> -cyclization sequence for the synthesis of multi-substituted 4,5-dihydrofurans. <i>Synthetic Communications</i> , 2021, 51, 234-244.	2.1	4
31	Switchable phase transition between crystalline and amorphous states of CuSO ₄ ·5H ₂ O by dynamic shock waves. <i>CrystEngComm</i> , 2021, 23, 7044-7048.	2.6	8
32	Biogenic synthesis, characterization and antimicrobial activity of <i>Ixora brachypoda</i> (DC) leaf extract mediated silver nanoparticles. <i>Journal of King Saud University - Science</i> , 2021, 33, 101296.	3.5	42
33	Stereo- and regioselective synthesis of novel Î²-lactam tethered spiropyrrolidone/pyrrolothiazole heterocyclic hybrids. <i>Tetrahedron</i> , 2021, 84, 132026.	1.9	3
34	Assessment of crystallographic and magnetic phase stabilities of cubic copper ferrite at shocked conditions. <i>Journal of Materials Science: Materials in Electronics</i> , 2021, 32, 12732-12742.	2.2	1
35	Assessment of Structural Stability of Dye-Doped Potassium Dihydrogen Phosphate Under Shocked Conditions. <i>Journal of Electronic Materials</i> , 2021, 50, 4215-4221.	2.2	3
36	Stereoselective synthesis and discovery of novel spirooxindolopyrrolidine engrafted indandione heterocyclic hybrids as antimycobacterial agents. <i>Bioorganic Chemistry</i> , 2021, 110, 104798.	4.1	20

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37	Assessment of shock wave resistance on brookite TiO ₂ . Journal of Materials Science: Materials in Electronics, 2021, 32, 15134-15142.	2.2	4
38	Assessment of shock resistance of barium ferrite at dynamic shocked conditions. Journal of Materials Science: Materials in Electronics, 2021, 32, 22429-22439.	2.2	2
39	Dynamic Shock Wave-Induced Switchable Phase Transition of Magnesium Sulfate Heptahydrate. Crystal Growth and Design, 2021, 21, 5050-5057.	3.0	11
40	Stereoselective synthesis, structure and DFT studies on fluoro- and nitro- substituted spirooxindole-pyrrolidine heterocyclic hybrids. Journal of Molecular Structure, 2021, 1237, 130396.	3.6	8
41	The switchable phase transition of sodium sulfate crystals activated by shock waves. New Journal of Chemistry, 2021, 45, 16529-16536.	2.8	19
42	Spectroscopic Assessment of Shock Wave Resistance on ZnO Nanorods for Aerospace Applications. Journal of Inorganic and Organometallic Polymers and Materials, 2021, 31, 2553-2559.	3.7	8
43	Phase Transformation of Amorphous to Crystalline of Multiwall Carbon Nanotubes by Shock Waves. Crystal Growth and Design, 2021, 21, 1617-1624.	3.0	33
44	Synthesis of Fused Quinoline Derivatives from Easily Accessible N-(2-aminobenzylidene)-4-methylanilines under Catalyst-Free Conditions in Water. ChemistrySelect, 2021, 6, 10436-10439.	1.5	1
45	Purification of Colon Carcinoma Cells from Primary Colon Tumor Using a Filtration Method via Porous Polymeric Filters. Polymers, 2021, 13, 3411.	4.5	0
46	Antimicrobial activities of novel class of dispirooxindolopyrrolidine grafted indanedione hybrid heterocycles against carbapenemase producing Klebsiella pneumoniae (CKP). Journal of Infection and Public Health, 2021, 14, 1870-1874.	4.1	6
47	Pyrano[2,3-f]pyrazolo[3,4-b]quinoline-3-carbonitriles: A three-component synthesis and AChE inhibitory studies. Synthetic Communications, 2021, 51, 1058-1065.	2.1	3
48	Shock Wave Induced Crystallographic Structural Phase Transitions (Tri-states) of L-Threonine Crystal. Journal of Physical Chemistry C, 2021, 125, 25217-25226.	3.1	8
49	Antimicrobial activities of spirooxindolopyrrolidine tethered dicarbonitrile heterocycles against multidrug resistant nosocomial pathogens. Journal of Infection and Public Health, 2021, 14, 1810-1814.	4.1	6
50	Graphene oxide-rhodamine nanocomposite for picomolar detection of chromium(III) by fluorimetry and its biofilm inhibition. Mikochimica Acta, 2021, 188, 414.	5.0	2
51	Multicomponent Domino Synthesis of Highly Functionalized Aryl and Heteroaryl Fused Pyrroloquinolinone Ring Systems via Environmentally Benign Solid-State Melt Reaction. ChemistrySelect, 2021, 6, 12001-12006.	1.5	0
52	Sustainability of the crystallographic phase stability of the barium carbonate nanoparticles at dynamic shocked conditions. Applied Physics A: Materials Science and Processing, 2021, 127, 1.	2.3	2
53	Shock wave induced phase transition from crystalline to the amorphous state of lead nitrate crystals. CrystEngComm, 2021, 24, 52-56.	2.6	7
54	Design, synthesis and cholinesterase inhibitory activity of novel spiropyrrolidine tethered imidazole heterocyclic hybrids. Bioorganic and Medicinal Chemistry Letters, 2020, 30, 126789.	2.2	23

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55	Synthesis, anticancer and molecular docking studies of new class of benzoisoxazolyl-piperidinyl-1, 2, 3-triazoles. <i>Journal of King Saud University - Science</i> , 2020, 32, 3286-3292.	3.5	1
56	A simple, rapid, expedient and sustainable green strategy for the synthesis of benz-/naphthimidazoles. <i>Journal of King Saud University - Science</i> , 2020, 32, 3153-3158.	3.5	3
57	Functionalized N-Pyridinylmethyl Engrafted Bisarylmethylidenepyridinones as Anticancer Agents. <i>Processes</i> , 2020, 8, 1154.	2.8	5
58	Regio- and stereoselective synthesis of novel β -lactam engrafted spiroheterocyclic hybrids via one-pot three component cycloaddition strategy. <i>Tetrahedron Letters</i> , 2020, 61, 152661.	1.4	6
59	In Vitro Molecular Biology Studies of Spirooxindole Heterocyclic Hybrids. <i>Processes</i> , 2020, 8, 1473.	2.8	2
60	Highly functionalized N-1-(2-pyridinylmethyl)-3,5-bis[(E)-arylmethylidene]tetrahydro-4(1H)-pyridinones: Synthesis, characterization, crystal structure and DFT studies. <i>Journal of Molecular Structure</i> , 2020, 1222, 128940.	3.6	11
61	An efficient, sustainable approach to the chemo and regioselective synthesis of novel spiroindenoquinoxaline grafted piperidone hybrid heterocycles. <i>Journal of King Saud University - Science</i> , 2020, 32, 3059-3064.	3.5	3
62	A One-Pot Three-Component Synthesis and Investigation of the In Vitro Mechanistic Anticancer Activity of Highly Functionalized Spirooxindole-Pyrrolidine Heterocyclic Hybrids. <i>Molecules</i> , 2020, 25, 5581.	3.8	6
63	Caspase dependent apoptotic activity of polycyclic cage-like heterocyclic hybrids. <i>Saudi Journal of Biological Sciences</i> , 2020, 27, 3290-3300.	3.8	4
64	Broad spectrum antimicrobial activity of dispirooxindolopyrrolidine fused acenaphthenone heterocyclic hybrid against healthcare associated microbial pathogens (HAMPS). <i>Journal of Infection and Public Health</i> , 2020, 13, 2001-2008.	4.1	11
65	[Bmim]Br Accelerated One-Pot Three-Component Cascade Protocol for the Construction of Spirooxindole-Pyrrolidine Heterocyclic Hybrids. <i>Molecules</i> , 2020, 25, 4779.	3.8	5
66	In vitro Mechanistic Exploration of Novel Spiropyrrrolidine Heterocyclic Hybrids as Anticancer Agents. <i>Frontiers in Chemistry</i> , 2020, 8, 465.	3.6	8
67	In vitro mechanistic investigation of polycyclic cage-like heterocyclic hybrid possessing diverse pharmacophoric units. <i>Journal of King Saud University - Science</i> , 2020, 32, 2406-2413.	3.5	3
68	Substitution induced switch between Pictet-Spengler and Eschweiler-Clarke reactions: Selective synthesis of spiro acenaphthylene pyrrolo[1,2-b]-isoquinoline/pyrrolidine hybrids. <i>Tetrahedron Letters</i> , 2020, 61, 151606.	1.4	5
69	Anti-tubercular activity of novel class of spiropyrrolidine tethered indenoquinoxaline heterocyclic hybrids. <i>Bioorganic Chemistry</i> , 2020, 99, 103799.	4.1	24
70	Dispiropyrrrolidine tethered piperidone heterocyclic hybrids with broad-spectrum antifungal activity against <i>Candida albicans</i> and <i>Cryptococcus neoformans</i> . <i>Bioorganic Chemistry</i> , 2020, 100, 103865.	4.1	13
71	A facile ionic liquid-accelerated, four-component cascade reaction protocol for the regioselective synthesis of biologically interesting ferrocene engrafted spiropyrrolidine hybrid heterocycles. <i>Journal of King Saud University - Science</i> , 2020, 32, 2500-2504.	3.5	8
72	Regio- and diastereoselective synthesis of spiropyrroloquinoxaline grafted indole heterocyclic hybrids and evaluation of their anti- <i>Mycobacterium tuberculosis</i> activity. <i>RSC Advances</i> , 2020, 10, 23522-23531.	3.6	21

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73	Discovery of novel cage-like heterocyclic hybrids as anti-inflammatory agents through the inhibition of nitrite, PGE2 and TNF- α . <i>Bioorganic Chemistry</i> , 2019, 91, 103180.	4.1	3
74	Broad-spectrum antifungal activity of spirooxindolo-pyrrolidine tethered indole/imidazole hybrid heterocycles against fungal pathogens. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2019, 29, 2059-2063.	2.2	29
75	Design, Synthesis and In Vitro Mechanistic Investigation of Novel Hexacyclic Cage-Like Hybrid Heterocycles. <i>Molecules</i> , 2019, 24, 3820.	3.8	7
76	A New Class of β -Pyrrolidino-1,2,3-Triazole Derivatives as β -Adrenergic Receptor Inhibitors: Synthesis, Pharmacological, and Docking Studies. <i>Molecules</i> , 2019, 24, 3501.	3.8	0
77	Domino Multicomponent Approach for the Synthesis of Functionalized Spiro-Indeno[1,2-b]quinoxaline Heterocyclic Hybrids and Their Antimicrobial Activity, Synergistic Effect and Molecular Docking Simulation. <i>Molecules</i> , 2019, 24, 1962.	3.8	16
78	Design and synthesis of A- and D ring-modified analogues of luotonin A with reduced planarity. <i>Tetrahedron Letters</i> , 2019, 60, 1514-1517.	1.4	8
79	Spirooxindole-pyrrolidine heterocyclic hybrids promotes apoptosis through activation of caspase-3. <i>Bioorganic and Medicinal Chemistry</i> , 2019, 27, 2487-2498.	3.0	26
80	Glucosamine-6-phosphate synthase inhibiting C3- β -cholesterol tethered spiro heterocyclic conjugates: Synthesis and their insight of DFT and docking study. <i>Bioorganic Chemistry</i> , 2019, 88, 102920.	4.1	4
81	Dispiropyrrolidinyl-piperidone embedded indeno[1,2-b]quinoxaline heterocyclic hybrids: Synthesis, cholinesterase inhibitory activity and their molecular docking simulation. <i>Bioorganic and Medicinal Chemistry</i> , 2019, 27, 2621-2628.	3.0	38
82	D-Ring-Modified Analogues of Luotonin A with Reduced Planarity: Design, Synthesis, and Evaluation of Their Topoisomerase Inhibition-Associated Cytotoxicity. <i>BioMed Research International</i> , 2019, 2019, 1-12.	1.9	5
83	Discovery of diazahexa/hepta cyclic cage-like compounds with broad-spectrum antifungal activity against <i>Candida</i> and <i>Cryptococcus</i> species. <i>RSC Advances</i> , 2019, 9, 29909-29916.	3.6	2
84	Design of New Amino Tf-Amide Organocatalysts: Environmentally Benign Approach to Asymmetric Aldol Synthesis. <i>Synlett</i> , 2019, 30, 401-404.	1.8	12
85	Multicomponent domino protocol for the stereoselective synthesis of novel pyrrolo[3,2-c]quinolinone hybrid heterocycles. <i>Tetrahedron Letters</i> , 2019, 60, 602-605.	1.4	12
86	Benzimidazole tethered pyrrolo[3,4-b]quinoline with broad-spectrum activity against fungal pathogens. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2019, 29, 729-733.	2.2	22
87	Regio- and diastereoselective synthesis of anticancer spirooxindoles derived from tryptophan and histidine via three-component 1,3-dipolar cycloadditions in an ionic liquid. <i>Tetrahedron</i> , 2018, 74, 5358-5366.	1.9	44
88	Ionic liquid-enabled synthesis, cholinesterase inhibitory activity, and molecular docking study of highly functionalized tetrasubstituted pyrrolidines. <i>Bioorganic Chemistry</i> , 2018, 77, 263-268.	4.1	29
89	Functionalized spirooxindole-indolizine hybrids: Stereoselective green synthesis and evaluation of anti-inflammatory effect involving TNF- α and nitrite inhibition. <i>European Journal of Medicinal Chemistry</i> , 2018, 152, 417-423.	5.5	38
90	ACI/EG eutectic mixture mediated synthesis, characterization and <i>in vitro</i> osteoblast differentiation assessment of spiropyrrolo[1,2- <i>b</i>]isoquinoline analogues. <i>RSC Advances</i> , 2018, 8, 16303-16313.	3.6	9

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91	Spiropyrrolidine/spiroindolizino[6,7-b]indole heterocyclic hybrids: Stereoselective synthesis, cholinesterase inhibitory activity and their molecular docking study. <i>Bioorganic Chemistry</i> , 2018, 79, 64-71.	4.1	37
92	Stereoselective green synthesis and molecular structures of highly functionalized spirooxindole-pyrrolidine hybrids – A combined experimental and theoretical investigation. <i>Journal of Molecular Structure</i> , 2018, 1152, 266-275.	3.6	13
93	Highly functionalized pyrrolidine analogues: stereoselective synthesis and caspase-dependent apoptotic activity. <i>RSC Advances</i> , 2018, 8, 41226-41236.	3.6	18
94	Synthesis of indole-cycloalkylpyridine hybrids via a four-component six-step tandem process. <i>Beilstein Journal of Organic Chemistry</i> , 2018, 14, 2907-2915.	2.2	11
95	Synthesis of spiro-linked quinolinone-pyrrolidine/pyrrolo[1,2-c]thiazole-oxindole/acenaphthalene hybrids via multi-component [3+2] cycloaddition. <i>Tetrahedron Letters</i> , 2018, 59, 4086-4089.	1.4	10
96	Unsupported nanoporous palladium-catalyzed chemoselective hydrogenation of quinolines: Heterolytic cleavage of H ₂ molecule. <i>Chinese Journal of Catalysis</i> , 2018, 39, 1746-1752.	14.0	11
97	Regio and stereoselective synthesis of anticancer spirooxindolopyrrolidine embedded piperidone heterocyclic hybrids derived from one-pot cascade protocol. <i>Chemistry Central Journal</i> , 2018, 12, 95.	2.6	15
98	Practical synthesis of four different pseudoenantiomeric organocatalysts with both cis- and trans-substituted 1,2-cis-cyclohexanediamine structures from a common intermediate. <i>Tetrahedron</i> , 2018, 74, 5263-5269.	1.9	2
99	Enantioselective Alkylation of Arylhydrazones Derived from Keto Esters and Isatin Derivatives through Asymmetric Phase-Transfer Catalysis. <i>Chemistry - an Asian Journal</i> , 2018, 13, 1780-1783.	3.3	8
100	Dipolar cycloaddition based multi-component reaction: Synthesis of spiro tethered acenaphthylene-indolizine-pyridinone hybrids. <i>Tetrahedron Letters</i> , 2018, 59, 3336-3340.	1.4	8
101	A one-pot access to pyridine/benzo fused cyclododecanes via multi-component tandem reactions. <i>Tetrahedron</i> , 2018, 74, 4569-4577.	1.9	9
102	Multicomponent Domino Synthesis, Anticancer Activity and Molecular Modeling Simulation of Complex Dispirooxindolopyrrolidines. <i>Molecules</i> , 2018, 23, 1094.	3.8	12
103	Highly functionalized 2-amino-4H-pyrans as potent cholinesterase inhibitors. <i>Bioorganic Chemistry</i> , 2018, 81, 134-143.	4.1	24
104	Carboxylative Suzuki coupling reactions of benzyl chlorides with allyl pinacolborate catalyzed by palladium nanoparticles. <i>Chinese Journal of Catalysis</i> , 2018, 39, 1258-1262.	14.0	5
105	Stereoselective Synthesis of Vinyl Iodides through Copper(I)-Catalyzed Finkelstein-Type Halide-Exchange Reaction. <i>Synthesis</i> , 2017, 49, 2727-2732.	2.3	12
106	Practical Synthesis of both Enantiomeric Amino Acid, Mannich, and Aldol Derivatives by Asymmetric Organocatalysis. <i>Chemical Record</i> , 2017, 17, 1059-1069.	5.8	11
107	Copper(II)-Catalyzed and Chelation-Induced Remote C-H Halogenation of Quinolines under Neutral Conditions. <i>ChemistrySelect</i> , 2017, 2, 3414-3418.	1.5	13
108	Practical Synthesis of Two Different Pseudoenantiomeric Organocatalysts with cis-cyclohexanediamine Structure from a Common Chiral Source. <i>Asian Journal of Organic Chemistry</i> , 2017, 6, 1226-1230.	2.7	1

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109	Unsupported Nanoporous Gold-Catalyzed Chemoselective Reduction of α,β -Unsaturated Aldehydes Using Formic Acid as Hydrogen Source. <i>Asian Journal of Organic Chemistry</i> , 2017, 6, 867-872.	2.7	12
110	A One-Pot Multicomponent 1,3-Dipolar Cycloaddition Strategy: Combinatorial Synthesis of Dihydrothiophenone-Engrafted Dispiro Hybrid Heterocycles. <i>ACS Combinatorial Science</i> , 2017, 19, 308-314.	3.8	27
111	Synthesis and cholinesterase inhibitory activity study of new piperidone grafted spiropyrrolidines. <i>Bioorganic Chemistry</i> , 2017, 75, 210-216.	4.1	21
112	Synthesis of penta- and tetra-cyclic cage-like compounds and dispiro heterocycles through microwave-assisted solvent-free multi-component domino reactions. <i>New Journal of Chemistry</i> , 2017, 41, 11009-11015.	2.8	8
113	Synthesis, theoretical studies and molecular docking of a novel chlorinated tetracyclic: (Z/E)-3-(1,8-dichloro-9,10-dihydro-9,10-ethanoanthracen-11-yl)acrylaldehyde. <i>Journal of Molecular Structure</i> , 2017, 1150, 358-365.	3.6	7
114	Design, synthesis and antiproliferative activity of decarbonyl luotonin analogues. <i>European Journal of Medicinal Chemistry</i> , 2017, 138, 932-941.	5.5	36
115	Hypervalent iodine(III) catalyzed radical hydroacylation of chiral alkylidenemalonates with aliphatic aldehydes under photolysis. <i>Tetrahedron</i> , 2017, 73, 5841-5846.	1.9	11
116	A Sustainable Approach to the Stereoselective Synthesis of Diazaheptacyclic Cage Systems Based on a Multicomponent Strategy in an Ionic Liquid. <i>Molecules</i> , 2016, 21, 165.	3.8	2
117	Synthesis, Spectroscopic, X-ray Diffraction and DFT Studies of Novel Benzimidazole Fused-1,4-Oxazepines. <i>Molecules</i> , 2016, 21, 724.	3.8	11
118	Multicomponent Dipolar Cycloaddition Strategy: Combinatorial Synthesis of Novel Spiro-Tethered Pyrazolo[3,4- <i>b</i>]quinoline Hybrid Heterocycles. <i>ACS Combinatorial Science</i> , 2016, 18, 262-270.	3.8	41
119	Highly functionalized dispiro oxindole-pyrrolo[1,2- <i>c</i>]thiazole-piperidone hybrid: Synthesis, characterization and theoretical investigations on the regiochemistry. <i>Journal of Molecular Structure</i> , 2016, 1121, 93-103.	3.6	6
120	Synthesis of highly functionalized 2-thiaspiro[4.5]deca-6,8-dienes via atom efficient tandem Michael addition/Thorpe-Ziegler cyclization. <i>RSC Advances</i> , 2016, 6, 40585-40592.	3.6	10
121	Rhodium-Catalyzed Oxidative Benzannulation of <i>N</i> -Pivaloylanilines with Internal Alkynes through Dual C-H Bond Activation: Synthesis of Highly Substituted Naphthalenes. <i>Chemistry - an Asian Journal</i> , 2016, 11, 3241-3250.	3.3	10
122	Rhodium-Catalyzed Oxidative Benzannulation of <i>N</i> -Adamantyl-1-naphthylamines with Internal Alkynes via Dual C-H Bond Activation: Synthesis of Substituted Anthracenes. <i>Organic Letters</i> , 2016, 18, 4246-4249.	4.6	43
123	1-Naphthol Synthesis through Base-Promoted S_NAr Reactions of <i>ortho</i> -Haloacetophenones Followed by Lewis-Acid-Catalyzed Cyclization. <i>Asian Journal of Organic Chemistry</i> , 2016, 5, 699-704.	2.7	8
124	Selective synthesis of γ -lactone via palladium nanoparticles-catalyzed telomerization of CO ₂ with 1,3-butadiene. <i>Tetrahedron Letters</i> , 2016, 57, 3163-3166.	1.4	23
125	Synthesis of cycloalkano[<i>b</i>]pyridines by multicomponent strategy: ring-size mediated product selectivity, substitution-induced axial chirality and influence of the 14N quadrupole-relaxation. <i>Tetrahedron</i> , 2016, 72, 4582-4592.	1.9	9
126	Palladium-catalyzed propargylative and allenylative dearomatization of 2-(chloromethyl)thiophenes: remarkable effect of solvents. <i>Tetrahedron</i> , 2016, 72, 170-175.	1.9	4

#	ARTICLE	IF	CITATIONS
127	Ī-Conjugated carbocycles and heterocycles via annulation through C-H and X-Y activation across CC triple bonds. <i>Arkivoc</i> , 2016, 2016, 9-41.	0.5	2
128	A Novel One-Pot Green Synthesis of Dispirooxindolo-pyrrolidines via 1,3-Dipolar Cycloaddition Reactions of Azomethine Ylides. <i>Molecules</i> , 2015, 20, 780-791.	3.8	31
129	A Facile Ionic Liquid Promoted Synthesis, Cholinesterase Inhibitory Activity and Molecular Modeling Study of Novel Highly Functionalized Spiropyrrrolidines. <i>Molecules</i> , 2015, 20, 2296-2309.	3.8	37
130	An Expedient Regio- and Diastereoselective Synthesis of Hybrid Frameworks with Embedded Spiro[9,10]dihydroanthracene [9,3- β^2]-pyrrolidine and Spiro[oxindole-3,2- α^2 -pyrrolidine] Motifs via an Ionic Liquid-Mediated Multicomponent Reaction. <i>Molecules</i> , 2015, 20, 16142-16153.	3.8	18
131	An Expedient Synthesis, Acetylcholinesterase Inhibitory Activity, and Molecular Modeling Study of Highly Functionalized Hexahydro-1,6-naphthyridines. <i>BioMed Research International</i> , 2015, 2015, 1-9.	1.9	14
132	Regioselective synthesis of novel dispiro oxindole- α^2 -pyrrolizine- α^2 -thiazolidine-2,4-dione hybrids. <i>Tetrahedron Letters</i> , 2015, 56, 4374-4376.	1.4	16
133	Applications of Metal Nanopore Catalysts in Organic Synthesis. <i>Synlett</i> , 2015, 26, 2355-2380.	1.8	21
134	Dipolar Cycloaddition-Based Multicomponent Reactions in Ionic Liquids: A Green, Fully Stereoselective Synthesis of Novel Polycyclic Cage Systems with the Generation of Two New Azaheterocyclic Rings. <i>Synthesis</i> , 2015, 47, 2721-2730.	2.3	18
135	Straightforward synthesis of pyrrolo[3,4-b]quinolines through intramolecular Povarov reactions. <i>Tetrahedron Letters</i> , 2015, 56, 6900-6903.	1.4	20
136	Carboxylative coupling reaction of five-membered (chloromethyl)heteroarenes with allyltributylstannane catalyzed by palladium nanoparticles. <i>Tetrahedron Letters</i> , 2015, 56, 6747-6750.	1.4	16
137	Domino four-component synthesis of novel cycloocta[b]pyridines. <i>Tetrahedron Letters</i> , 2015, 56, 179-181.	1.4	12
138	Ionic Liquid-Promoted Synthesis and Cholinesterase Inhibitory Activity of Highly Functionalized Spiropyrrrolidines. <i>Australian Journal of Chemistry</i> , 2015, 68, 863.	0.9	22
139	Crystal structure of 8-(2-methylphenyl)-11-[(E)-(2-methylphenyl)-methylidene]-14-hydroxy-3,13-diazaheptacyclo-[13.7.1.9,13.02,9.02,14.03,7.019,23]tetracos-1(22),15,17,19(23),20-pentaen-10-one methanol monosolvate, C ₃₇ H ₃₄ N ₂ O ₂ ·CH ₃ OH, C ₃₈ H ₃₈ N ₂ O ₃ . <i>Zeitschrift Fur Kristallographie - New Crystal Structures</i> , 2014, 229, 175-177.	0.3	2
140	Facile, Regio- and Diastereoselective Synthesis of Spiro-Pyrrolidine and Pyrrolizine Derivatives and Evaluation of Their Antiproliferative Activities. <i>Molecules</i> , 2014, 19, 10033-10055.	3.8	35
141	An expedient pseudo four-component synthesis of dispiroindandione fused indeno-N-methylmorpholine, spectroscopic, X-ray diffraction and DFT studies. <i>Journal of Molecular Structure</i> , 2014, 1063, 283-288.	3.6	4
142	Cholinesterase inhibitory activity versus aromatic core multiplicity: A facile green synthesis and molecular docking study of novel piperidone embedded thiazolopyrimidines. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 906-916.	3.0	19
143	Ionic liquid mediated synthesis of mono- and bis-spirooxindole-hexahydropyrrolidines as cholinesterase inhibitors and their molecular docking studies. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 1318-1328.	3.0	43
144	Ultrasound-Assisted Sequential Multicomponent Strategy for the Combinatorial Synthesis of Novel Coumarin Hybrids. <i>ACS Combinatorial Science</i> , 2014, 16, 566-572.	3.8	30

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145	Microwave-assisted chemoselective synthesis of novel pyrazolo[3,4-b]thieno[3,4-e]pyridines: substitution induced axial chirality. <i>Tetrahedron Letters</i> , 2014, 55, 5805-5807.	1.4	13
146	Synthesis and discovery of highly functionalized mono- and bis-spiro-pyrrolidines as potent cholinesterase enzyme inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 1815-1819.	2.2	50
147	An Expedient Synthesis and Screening for Antiacetylcholinesterase Activity of Piperidine Embedded Novel Pentacyclic Cage Compounds. <i>Medicinal Chemistry</i> , 2014, 10, 228-236.	1.5	13
148	An Efficient Ionic Liquid Mediated Synthesis, Cholinesterase Inhibitory Activity and Molecular Modeling Study of Novel Piperidone Embedded α ,β-Unsaturated Ketones. <i>Medicinal Chemistry</i> , 2014, 10, 512-520.	1.5	3
149	An expedient, ionic liquid mediated multi-component synthesis of novel piperidone grafted cholinesterase enzymes inhibitors and their molecular modeling study. <i>European Journal of Medicinal Chemistry</i> , 2013, 67, 221-229.	5.5	43
150	â€™On-waterâ€™ one-pot pseudo four-component domino protocol for the synthesis of novel benzo[a]cyclooctenes. <i>Tetrahedron Letters</i> , 2013, 54, 4800-4802.	1.4	13
151	Microwave assisted synthesis, cholinesterase enzymes inhibitory activities and molecular docking studies of new pyridopyrimidine derivatives. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 3022-3031.	3.0	31
152	Three-Component Synthesis and 1,3-Dipolar Cycloaddition of Highly Functionalized Pyrans with Nitrile Oxides: Easy Access to 1,2,4-Oxadiazoles. <i>Synthetic Communications</i> , 2013, 43, 2763-2772.	2.1	14
153	An atom economic synthesis and AChE inhibitory activity of novel dispiro 7-aryltetrahydro-1H-pyrrolo[1,2-c][1,3]thiazole and 4-aryloctahydroindolizine N-methylpiperidin-4-one hybrid heterocycles. <i>European Journal of Medicinal Chemistry</i> , 2013, 65, 240-248.	5.5	30
154	A facile chemo-, regio- and stereoselective synthesis and cholinesterase inhibitory activity of spirooxindoleâ€“pyrrolizineâ€“piperidine hybrids. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013, 23, 2979-2983.	2.2	66
155	Synthesis and discovery of novel piperidone-grafted mono- and bis-spirooxindole-hexahydropyrrolizines as potent cholinesterase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2013, 21, 1696-1707.	3.0	87
156	Synthesis of novel 16-spiro steroids: Spiro-7â€“(aryl)tetrahydro-1H-pyrrolo[1,2-c][1,3]thiazolo-trans-androsterone hybrid heterocycles. <i>Steroids</i> , 2013, 78, 409-417.	1.8	22
157	A 1,3-dipolar cycloadditionâ€“annulation protocol for the expedient regio-, stereo- and product-selective construction of novel hybrid heterocycles comprising seven rings and seven contiguous stereocentres. <i>Tetrahedron Letters</i> , 2013, 54, 2515-2519.	1.4	29
158	A facile three-component [3+2]-cycloaddition for the regioselective synthesis of highly functionalised dispiropyrrolidines acting as antimycobacterial agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013, 23, 1383-1386.	2.2	43
159	Antituberculosis: Synthesis and Antimycobacterial Activity of Novel Benzimidazole Derivatives. <i>BioMed Research International</i> , 2013, 2013, 1-6.	1.9	16
160	(3E,5E)-1-Allyl-3,5-bis(4-methoxybenzylidene)piperidin-4-one. <i>Acta Crystallographica Section E: Structure Reports Online</i> , 2013, 69, o1071-o1071.	0.2	3
161	9-(2-Chlorobenzylidene)anthracen-10(9H)-one. <i>Acta Crystallographica Section E: Structure Reports Online</i> , 2013, 69, o627-o627.	0.2	0
162	Multicomponent 1,3-Dipolar Cycloaddition Reactions in the Construction of Hybrid Spiroheterocycles. <i>Current Organic Chemistry</i> , 2013, 17, 1929-1956.	1.6	22

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163	Antimycobacterial Activity: Synthesis and Biological Evaluation of Novel Substituted (3E,5E)-3,5-diarylidene-1-phenethylpiperidine-4-one Derivatives. Letters in Drug Design and Discovery, 2013, 10, 471-476.	0.7	3
164	Synthesis of Novel and Highly Functionalized 4-hydroxycoumarin Chalcone and their Pyrazoline Derivatives as Anti-Tuberculosis Agents. Letters in Drug Design and Discovery, 2013, 11, 222-230.	0.7	4
165	5-Chlorospiro[indoline-3,7-6H,7H,8H-pyrano[3,2-c:5,6-câ€™²]di[1]benzopyran]-2,6-8â€™²-trione. Acta Crystallographica Section E: Structure Reports Online, 2012, 68, o1172-o1172.	0.2	0
166	5-Methylspiro[indoline-3,7-6H,7H,8H-pyrano[3,2-c:5,6-câ€™²]di[1]benzopyran]-2,6-8â€™²-trione chloroform hemisolvate. Acta Crystallographica Section E: Structure Reports Online, 2012, 68, o1194-o1194.	0.2	0
167	11-[(E)-Benzylidene]-14-hydroxy-8-phenyl-6-thia-3,13-diazaheptacyclo[13.7.1.1^{9,13}.0^{2,9}.0^{2,1}]heptacosan-1(20)-one. Acta Crystallographica Section E: Structure Reports Online, 2012, 68, o2013-o2014.	0.2	0
168	2-Hydroxy-5-[(E)-2-methylbenzylidene]-8-(2-methylphenyl)-9-phenyl-3,10-diazahexacyclo[10.7.1.13,7.02,11.07,11.016,20]heptacosan-1(20)-one. Acta Crystallographica Section E: Structure Reports Online, 2012, 68, o3375-o3375.	0.2	0
169	5-Fluoro-6-8H,7-8H-spiro[indoline-3,7-6H-pyrano[3,2-c:5,6-câ€™²]di-1-benzopyran]-2,6-8â€™²-trione. Acta Crystallographica Section E: Structure Reports Online, 2012, 68, o744-o744.	0.2	1
170	11-[(E)-2-Fluorobenzylidene]-8-(2-fluorophenyl)-14-hydroxy-6-thia-3,13-diazaheptacyclo[13.7.1.19,13.02,9.02,14.03,7.019,23]tetracosan-1(20)-one. Acta Crystallographica Section E: Structure Reports Online, 2012, 68, o2094-o2095.	0.2	1
171	Antimycobacterial activity: A facile three-component [3+2]-cycloaddition for the regioselective synthesis of highly functionalised dispiropyrrolidines. Bioorganic and Medicinal Chemistry Letters, 2012, 22, 4930-4933.	2.2	40
172	A solvent free, four-component synthesis and 1,3-dipolar cycloaddition of 4(H)-pyrans with nitrile oxides: Synthesis and discovery of antimycobacterial activity of enantiomerically pure 1,2,4-oxadiazoles. European Journal of Medicinal Chemistry, 2012, 53, 416-423.	5.5	31
173	AChE inhibitor : A regio- and stereo-selective 1,3-dipolar cycloaddition for the synthesis of novel substituted 5,6-dimethoxy spiro[5.3]-oxindole-spiro-[6.3]-2,3-dihydro-1H-inden-1-one-7-(substituted) Tj. ETQq1 1 0,784314 mg 508-511.	2.2	51
174	Antimycobacterial Agents: Synthesis and Biological Evaluation of Novel 4-(Substituted-phenyl)-6-methyl-2-oxo-N-(pyridin-2-yl)-1,2,3,4-tetrahydropyrimidine-5-carboxamide Derivatives by Using One-pot Multicomponent Method. Letters in Drug Design and Discovery, 2012, 9, 953-957.	0.7	0
175	Antimycobacterial activity of novel 1,2,4-oxadiazole-pyranopyridine/chromene hybrids generated by chemoselective 1,3-dipolar cycloadditions of nitrile oxides. Bioorganic and Medicinal Chemistry, 2011, 19, 3444-3450.	3.0	51
176	A green expedient synthesis of pyridopyrimidine-2-thiones and their antitubercular activity. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 3012-3016.	2.2	50
177	Synthesis and discovery of novel hexacyclic cage compounds as inhibitors of acetylcholinesterase. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 3997-4000.	2.2	21
178	A facile three-component [3+2]-cycloaddition/annulation domino protocol for the regio- and diastereoselective synthesis of novel penta- and hexacyclic cage systems, involving the generation of two heterocyclic rings and five contiguous stereocenters. Tetrahedron, 2011, 67, 3132-3139.	1.9	25
179	5-[(E)-Benzylidene]-2-hydroxy-8,9-diphenyl-3,10-diazahexacyclo[10.7.1.13,7.02,11.07,11.016,20]heptacosan-1(19),12(20),13,15,17-pentacosan-1(20)-one. Acta Crystallographica Section E: Structure Reports Online, 2011, 67, o2877-o2878.	0.2	0
180	5-[(E)-2-Fluorobenzylidene]-8-(2-fluorophenyl)-2-hydroxy-10-methyl-3,10-diazahexacyclo[10.7.1.13,7.02,11.07,11.016,20]heptacosan-1(20)-one. Acta Crystallographica Section E: Structure Reports Online, 2011, 67, o211-o212.	0.2	2

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182	16-[(E)-4-Bromobenzylidene]-13-(4-bromophenyl)-2-hydroxy-11-methyl-1,11-diazapentacyclo[12.3.1.02,10.03,8.010,14]octadeca-3(8),4 Acta Crystallographica Section E: Structure Reports Online, 2011, 67, o137-o138.	0.2	1
183	14-Hydroxy-11-[(E)-4-methoxybenzylidene]-8-(4-methoxyphenyl)-5-thia-3,13-diazaheptacyclo[13.7.1.19,13.02,9.02,14.03,7.019,23]tetra Acta Crystallographica Section E: Structure Reports Online, 2011, 67, o2881-o2882.	0.2	1
184	A Facile Synthesis and Discovery of Highly Functionalized Tetrahydro-pyridines and Pyridines as Antimycobacterial Agents. Chemical and Pharmaceutical Bulletin, 2010, 58, 602-610.	1.3	10
185	1,3-Dipolar cycloaddition of C-aryl-N-phenylnitrones to (R)-1-(1-phenylethyl)-3-[(E)-arylmethylidene]tetrahydro-4(1H)-pyridinones: Synthesis and antimycobacterial evaluation of enantiomerically pure spiroisoxazolidines. European Journal of Medicinal Chemistry, 2010, 45, 124-133.	5.5	37
186	1,3-Dipolar cycloaddition of nitrile oxides to (R)-1-(1-phenylethyl)-3,5-bis[(E)-arylmethylidene]tetrahydro-4(1H)-pyridinones: synthesis and antimycobacterial evaluation of novel enantiomerically pure di- and trispiroheterocycles. Tetrahedron: Asymmetry, 2010, 21, 1315-1327.	1.8	24
187	Substituted spiro [2.3] oxindolespiro [3.2]-5,6-dimethoxy-indane-1-one-pyrrolidine analogue as inhibitors of acetylcholinesterase. Bioorganic and Medicinal Chemistry Letters, 2010, 20, 7064-7066.	2.2	129
188	Novel three-component domino reactions of ketones, isatin and amino acids: Synthesis and discovery of antimycobacterial activity of highly functionalised novel dispiropyrrrolidines. European Journal of Medicinal Chemistry, 2010, 45, 411-422.	5.5	129
189	19-[(E)-4-Chlorobenzylidene]-16-(4-chlorophenyl)-2-hydroxy-1,11-diazahexacyclo[15.3.1.02,10.03,8.010,17.011,15]henicosa-3(8),4,6 Acta Crystallographica Section E: Structure Reports Online, 2010, 66, o1444-o1445.	0.2	5
190	16-[(E)-Benzylidene]-13-hydroxy-4-methyl-2-phenyl-4,14-diazapentacyclo-[12.3.1.01,5.05,13.07,12]octadeca-7(12),8,10-triene-6,17-dione Acta Crystallographica Section E: Structure Reports Online, 2010, 66, o1540-o1541.	0.2	9
191	16-[(E)-Benzylidene]-2-hydroxy-12,13-diphenyl-1,11-diazapentacyclo[12.3.1.02,10.03,8.010,14]octadeca-3(8),4,6-triene-9,15-dione. Acta Crystallographica Section E: Structure Reports Online, 2010, 66, o2084-o2085.	0.2	1
192	2-Hydroxy-16-[(E)-4-methylbenzylidene]-13-(4-methylphenyl)-12-phenyl-1,11-diazapentacyclo[12.3.1.02,10.03,8.010,14]octadeca-3(8), Acta Crystallographica Section E: Structure Reports Online, 2010, 66, o2086-o2087.	0.2	0
193	2-Hydroxy-11-methyl-16-[(E)-4-methylbenzylidene]-13-(4-methylphenyl)-1,11-diazapentacyclo[12.3.1.02,10.03,8.010,14]octadeca-3(8), Acta Crystallographica Section E: Structure Reports Online, 2010, 66, o2370-o2371.	0.2	1
194	5-[(E)-2-Bromobenzylidene]-8-(2-bromophenyl)-2-hydroxy-10-methyl-3,10-diazahexacyclo[10.7.1.13,7.02,11.07,11.016,20]henicosa-1(20) Acta Crystallographica Section E: Structure Reports Online, 2010, 66, o2376-o2377.	0.2	3
195	5-[(E)-Benzylidene]-2-hydroxy-10-methyl-8-phenyl-3,10-diazahexacyclo[10.7.1.13,7.02,11.07,11.016,20]henicosa-1(19),12(20),13,15 ethanol 0.25-solvate 0.6-hydrate. Acta Crystallographica Section E: Structure Reports Online, 2010, 66, o2736-o2737.	0.2	1
196	11-[(E)-4-Bromobenzylidene]-8-(4-bromophenyl)-14-hydroxy-3,13-diazaheptacyclo[13.7.1.19,13.02,9.02,14.03,7.019,23]tetracos-1(22) Acta Crystallographica Section E: Structure Reports Online, 2010, 66, o2926-o2927.	0.2	1
197	11-[(E)-Benzylidene]-14-hydroxy-8-phenyl-3,13-diazaheptacyclo[13.7.1.19,13.02,9.02,14.03,7.019,23]tetracos-1(22),15,17,19(23),20 Acta Crystallographica Section E: Structure Reports Online, 2010, 66, o3045-o3045.	0.2	1
198	2-Hydroxy-16-[(E)-4-hydroxy-3-methoxybenzylidene]-13-(4-hydroxy-3-methoxyphenyl)-11-methyl-1,11-diazapentacyclo[12.3.1.02,10.03, Acta Crystallographica Section E: Structure Reports Online, 2010, 66, o1370-o1371.	0.2	5

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199	A facile synthesis and antimycobacterial evaluation of novel spiro-pyrido-pyrrolizines and pyrrolidines. <i>European Journal of Medicinal Chemistry</i> , 2009, 44, 3821-3829.	5.5	154
200	An atom economic synthesis and antitubercular evaluation of novel spiro-cyclohexanones. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009, 19, 3461-3465.	2.2	75
201	A highly atom economic, chemo-, regio- and stereoselective synthesis, and discovery of spiro-pyrido-pyrrolizines and pyrrolidines as antimycobacterial agents. <i>Tetrahedron</i> , 2008, 64, 2962-2971.	1.9	71
202	Discovery of Antimycobacterial Spiro-piperidin-4-ones: An Atom Economic, Stereoselective Synthesis, and Biological Intervention. <i>Journal of Medicinal Chemistry</i> , 2008, 51, 5731-5735.	6.4	216
203	Four-component tandem protocol for the stereoselective synthesis of highly functionalized [1,4]-thiazines. <i>Tetrahedron</i> , 2007, 63, 1411-1416.	1.9	37
204	A facile synthesis and highly atom economic 1,3-dipolar cycloaddition of hexahydropyrido[3,4-c][1,5]benzothiazepines with nitrile oxide: stereoselective formation of hexahydro[1,2,4]oxadiazolo[5,4-d]pyrido[3,4-c][1,5]benzothiazepines. <i>Tetrahedron</i> , 2007, 63, 7850-7857.	1.9	38
205	Sacrificial azomethine ylide cycloaddition controlled chemoselective nitrile oxide cycloaddition to 1-methyl-3,5-bis[(E)-arylmethylidene]tetrahydro-4(1H)-pyridinones: formation of mono-spiro-isoxazolines. <i>Tetrahedron</i> , 2007, 63, 12220-12231.	1.9	55
206	1,3-Dipolar cycloaddition of nitrile oxides to (R)-1-(1-phenylethyl)-3-[(E)-arylmethylidene]tetrahydro-4(1H)-pyridinones: synthesis of enantiomerically pure spiro heterocycles. <i>Tetrahedron: Asymmetry</i> , 2007, 18, 170-180.	1.8	23
207	Three-component tandem reactions of (2-arylsulfanyl-3-aryl-2-oxiranyl)(aryl)methanones and o-phenylenediamine: formation of quinoxalines. <i>Tetrahedron Letters</i> , 2007, 48, 2155-2158.	1.4	34
208	Novel three-component tandem reactions of cyclic mono ketones, isatin and sarcosine: formation of dispiropyrrrolidines. <i>Tetrahedron Letters</i> , 2007, 48, 7164-7168.	1.4	52
209	An atom efficient, solvent-free, green synthesis and antimycobacterial evaluation of 2-amino-6-methyl-4-aryl-8-[(E)-arylmethylidene]-5,6,7,8-tetrahydro-4H-pyrano[3,2-c]pyridine-3-carbonitriles. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007, 17, 6459-6462.	2.2	125
210	Chemo-, regio- and stereoselective 1,3-dipolar cycloaddition of C-aryl-N-phenylnitrones over 3,5-bis(arylidene)-1-methylpiperidin-4-ones: synthesis of highly substituted novel spiro-isoxazolidines. <i>Tetrahedron</i> , 2006, 62, 12380-12391.	1.9	21