

Rashmi Venugopala

List of Publications by Year in descending order

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Version: 2024-02-01

20
papers

1,008
citations

567281

15
h-index

794594

19
g-index

20
all docs

20
docs citations

20
times ranked

1542
citing authors

#	ARTICLE	IF	CITATIONS
1	Antitubercular, Cytotoxicity, and Computational Target Validation of Dihydroquinazolinone Derivatives. <i>Antibiotics</i> , 2022, 11, 831.	3.7	5
2	4-Aryl-1,4-Dihydropyridines as Potential Enoyl-Acyl Carrier Protein Reductase Inhibitors: Antitubercular Activity and Molecular Docking Study. <i>Current Topics in Medicinal Chemistry</i> , 2021, 21, 295-306.	2.1	8
3	Crystallography, Molecular Modeling, and COX-2 Inhibition Studies on Indolizine Derivatives. <i>Molecules</i> , 2021, 26, 3550.	3.8	10
4	Cytotoxicity and Antimycobacterial Properties of Pyrrolo[1,2-a]quinoline Derivatives: Molecular Target Identification and Molecular Docking Studies. <i>Antibiotics</i> , 2020, 9, 233.	3.7	30
5	Synthesis and characterization of pyrrolo[1,2-a]quinoline derivatives for their larvicidal activity against <i>Anopheles arabiensis</i> . <i>Structural Chemistry</i> , 2020, 31, 1533-1543.	2.0	22
6	Larvicidal Activities of 2-Aryl-2,3-Dihydroquinazolin-4-ones against Malaria Vector <i>Anopheles arabiensis</i> , In Silico ADMET Prediction and Molecular Target Investigation. <i>Molecules</i> , 2020, 25, 1316.	3.8	16
7	In silico Design and Synthesis of Tetrahydropyrimidinones and Tetrahydropyrimidinethiones as Potential Thymidylate Kinase Inhibitors Exerting Anti-TB Activity Against <i>Mycobacterium tuberculosis</i> . <i>Drug Design, Development and Therapy</i> , 2020, Volume 14, 1027-1039.	4.3	26
8	Crystallography, in Silico Studies, and In Vitro Antifungal Studies of 2,4,5 Trisubstituted 1,2,3-Triazole Analogues. <i>Antibiotics</i> , 2020, 9, 350.	3.7	13
9	Novel Series of Methyl 3-(Substituted Benzoyl)-7-Substituted-2-Phenylindolizine-1-Carboxylates as Promising Anti-Inflammatory Agents: Molecular Modeling Studies. <i>Biomolecules</i> , 2019, 9, 661.	4.0	21
10	Computational, crystallographic studies, cytotoxicity and anti-tubercular activity of substituted 7-methoxy-indolizine analogues. <i>PLoS ONE</i> , 2019, 14, e0217270.	2.5	29
11	Anti-Tubercular Activity of Substituted 7-Methyl and 7-Formylindolizines and In Silico Study for Prospective Molecular Target Identification. <i>Antibiotics</i> , 2019, 8, 247.	3.7	32
12	Anti-tubercular Potency and Computationally assessed Drug-likeness and Toxicology of Diversely Substituted Indolizines. <i>Indian Journal of Pharmaceutical Education and Research</i> , 2019, 53, 545-552.	0.6	25
13	Efficient synthesis and characterization of novel indolizines: exploration of <i>in vitro</i> COX-2 inhibitory activity and molecular modelling studies. <i>New Journal of Chemistry</i> , 2018, 42, 4893-4901.	2.8	32
14	One-pot microwave assisted synthesis and structural elucidation of novel ethyl 3-substituted-7-methylindolizine-1-carboxylates with larvicidal activity against <i>Anopheles arabiensis</i> . <i>Journal of Molecular Structure</i> , 2018, 1156, 377-384.	3.6	36
15	Design and Synthesis of Novel Indolizine Analogues as COX-2 Inhibitors: Computational Perspective and in vitro Screening. <i>Indian Journal of Pharmaceutical Education and Research</i> , 2017, 51, 452-460.	0.6	23
16	Silica-Sulfuric Acid: Novel, Simple, Efficient and Reusable Catalyst for Hydration of Nitrile to Amide. <i>Asian Journal of Chemistry</i> , 2016, 28, 2177-2180.	0.3	0
17	Synthesis and Characterization of Ethyl 7-Acetyl-2-substituted 3-(substituted) Tj ETQq1 1 0.784314 rgBT /Overlock 10 Tf 50 107 Td (ber 1043-1048.	0.3	33
18	Greener synthesis of indolizine analogues using water as a base and solvent: study for larvicidal activity against <i>Anopheles arabiensis</i> . <i>Chemical Biology and Drug Design</i> , 2016, 88, 899-904.	3.2	40

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19	Design, synthesis, and computational studies on dihydropyrimidine scaffolds as potential lipoyxygenase inhibitors and cancer chemopreventive agents. <i>Drug Design, Development and Therapy</i> , 2015, 9, 911.	4.3	20
20	Review on Natural Coumarin Lead Compounds for Their Pharmacological Activity. <i>BioMed Research International</i> , 2013, 2013, 1-14.	1.9	587