

Hayat Ullah

List of Publications by Year in descending order

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38
papers

1,916
citations

201658

27
h-index

345203

36
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docs citations

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times ranked

1093
citing authors

#	ARTICLE	IF	CITATIONS
1	Isatin based Schiff bases as inhibitors of α -glucosidase: Synthesis, characterization, in vitro evaluation and molecular docking studies. <i>Bioorganic Chemistry</i> , 2015, 60, 42-48.	4.1	147
2	Triazinoindole analogs as potent inhibitors of α -glucosidase: Synthesis, biological evaluation and molecular docking studies. <i>Bioorganic Chemistry</i> , 2015, 58, 81-87.	4.1	126
3	Synthesis, molecular docking, acetylcholinesterase and butyrylcholinesterase inhibitory potential of thiazole analogs as new inhibitors for Alzheimer disease. <i>Bioorganic Chemistry</i> , 2015, 62, 106-116.	4.1	114
4	Synthesis, in vitro evaluation and molecular docking studies of thiazole derivatives as new inhibitors of α -glucosidase. <i>Bioorganic Chemistry</i> , 2015, 62, 15-21.	4.1	109
5	Synthesis of 4-thiazolidinone analogs as potent in vitro anti-urease agents. <i>Bioorganic Chemistry</i> , 2015, 63, 123-131.	4.1	85
6	Synthesis and in vitro acetylcholinesterase and butyrylcholinesterase inhibitory potential of hydrazide based Schiff bases. <i>Bioorganic Chemistry</i> , 2016, 68, 30-40.	4.1	82
7	Synthesis and study of the α -amylase inhibitory potential of thiadiazole quinoline derivatives. <i>Bioorganic Chemistry</i> , 2017, 74, 179-186.	4.1	80
8	Synthesis of alpha amylase inhibitors based on privileged indole scaffold. <i>Bioorganic Chemistry</i> , 2017, 72, 248-255.	4.1	75
9	Biology-oriented drug synthesis (BIODS) of 2-(2-methyl-5-nitro-1H-imidazol-1-yl)ethyl aryl ether derivatives, in vitro α -amylase inhibitory activity and in silico studies. <i>Bioorganic Chemistry</i> , 2017, 74, 1-9.	4.1	75
10	Synthesis, α -glucosidase inhibitory activity and in silico study of tris-indole hybrid scaffold with oxadiazole ring: As potential leads for the management of type-II diabetes mellitus. <i>Bioorganic Chemistry</i> , 2017, 74, 30-40.	4.1	72
11	Synthesis, β -glucuronidase inhibition and molecular docking studies of hybrid bisindole-thiosemicarbazides analogs. <i>Bioorganic Chemistry</i> , 2016, 68, 56-63.	4.1	66
12	Synthesis of bis-indolylmethanes as new potential inhibitors of β -glucuronidase and their molecular docking studies. <i>European Journal of Medicinal Chemistry</i> , 2018, 143, 1757-1767.	5.5	65
13	Synthesis of 2-acylated and sulfonated 4-hydroxycoumarins: In vitro urease inhibition and molecular docking studies. <i>Bioorganic Chemistry</i> , 2016, 66, 111-116.	4.1	60
14	Bisindolylmethane thiosemicarbazides as potential inhibitors of urease: Synthesis and molecular modeling studies. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 152-160.	3.0	59
15	Synthesis and biological evaluation of novel N-arylidenequinoline-3-carbohydrazides as potent β -glucuronidase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 3696-3704.	3.0	58
16	Development of bis-thiobarbiturates as successful urease inhibitors and their molecular modeling studies. <i>Chinese Chemical Letters</i> , 2016, 27, 693-697.	9.0	53
17	Synthesis, in vitro alpha-glucosidase inhibitory potential of benzimidazole bearing bis-Schiff bases and their molecular docking study. <i>Bioorganic Chemistry</i> , 2020, 94, 103394.	4.1	51
18	Synthesis of Bis-indolylmethane sulfonylhydrazides derivatives as potent α -Glucosidase inhibitors. <i>Bioorganic Chemistry</i> , 2018, 80, 112-120.	4.1	49

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19	Synthesis, in vitro urease inhibitory potential and molecular docking study of Benzimidazole analogues. <i>Bioorganic Chemistry</i> , 2019, 89, 103024.	4.1	45
20	New triazinoindole bearing thiazole/oxazole analogues: Synthesis, $\hat{\alpha}$ -amylase inhibitory potential and molecular docking study. <i>Bioorganic Chemistry</i> , 2019, 92, 103284.	4.1	38
21	In vitro $\hat{\alpha}$ -glucosidase and $\hat{\alpha}$ -amylase inhibitory potential and molecular docking studies of benzohydrazide based imines and thiazolidine-4-one derivatives. <i>Journal of Molecular Structure</i> , 2022, 1251, 132058.	3.6	34
22	Synthesis, molecular docking study and in vitro thymidine phosphorylase inhibitory potential of oxadiazole derivatives. <i>Bioorganic Chemistry</i> , 2018, 78, 58-67.	4.1	33
23	Synthesis of new arylhydrazide bearing Schiff bases/thiazolidinone: $\hat{\alpha}$ -Amylase, urease activities and their molecular docking studies. <i>Bioorganic Chemistry</i> , 2019, 91, 103112.	4.1	33
24	Synthesis, in vitro urease inhibitory potential and molecular docking study of benzofuran-based-thiazolidinone analogues. <i>Scientific Reports</i> , 2020, 10, 10673.	3.3	33
25	Synthesis, In vitro $\hat{\alpha}$ -Glucosidase Inhibitory Potential and Molecular Docking Studies of 2-Amino-1,3,4-Oxadiazole Derivatives. <i>Medicinal Chemistry</i> , 2020, 16, 724-734.	1.5	31
26	Synthesis, in vitro alpha glucosidase, urease activities and molecular docking study of bis-indole bearing Schiff base analogs. <i>Chemical Data Collections</i> , 2020, 28, 100396.	2.3	29
27	Synthesis, In Vitro $\hat{\alpha}$ -Amylase Activity, and Molecular Docking Study of New Benzimidazole Derivatives. <i>Russian Journal of Organic Chemistry</i> , 2021, 57, 968-975.	0.8	29
28	The immunomodulation potential of the synthetic derivatives of benzothiazoles: Implications in immune system disorders through in vitro and in silico studies. <i>Bioorganic Chemistry</i> , 2016, 64, 21-28.	4.1	28
29	Aryl-oxadiazole Schiff bases: Synthesis, $\hat{\alpha}$ -glucosidase in vitro inhibitory activity and their in silico studies. <i>Arabian Journal of Chemistry</i> , 2020, 13, 4904-4915.	4.9	27
30	Synthesis, in-vitro and in-silico studies of triazinoindole bearing bis-Schiff base as $\hat{\alpha}$ -glucuronidase inhibitors. <i>Journal of Molecular Structure</i> , 2021, 1244, 131003.	3.6	25
31	Synthesis and in vitro study of benzofuran hydrazone derivatives as novel alpha-amylase inhibitor. <i>Bioorganic Chemistry</i> , 2017, 75, 78-85.	4.1	24
32	Synthesis of Novel Triazinoindole-Based Thiourea Hybrid: A Study on $\hat{\alpha}$ -Glucosidase Inhibitors and Their Molecular Docking. <i>Molecules</i> , 2019, 24, 3819.	3.8	18
33	Synthesis of substituted benzohydrazide derivatives: In vitro urease activities and their molecular docking studies. <i>Chemical Data Collections</i> , 2021, 36, 100778.	2.3	15
34	Recent Progress in Nanoparticles Based Sensors for the Detection of Mercury (II) Ions in Environmental and Biological Samples. <i>Critical Reviews in Analytical Chemistry</i> , 2024, 54, 44-60.	3.5	15
35	Recent Development in Coordination Compounds as a Sensor for Cyanide Ions in Biological and Environmental Segments. <i>Critical Reviews in Analytical Chemistry</i> , 0, , 1-21.	3.5	12
36	Synthesis of triazinoindole bearing sulfonamide derivatives, in vitro $\hat{\alpha}$ -amylase activity and their molecular docking study. <i>Chemical Data Collections</i> , 2022, 39, 100875.	2.3	11

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37	Synthesis, in vitro thymidine phosphorylase activity and molecular docking study of thiadiazole bearing isatin analogs. Chemical Papers, 2022, 76, 213-224.	2.2	10
38	Antimicrobial Susceptibility of Pseudomonas aeruginosa Isolated from Hospital Environment. Abasyn Journal of Life Sciences, 2021, , 40-50.	0.1	0