

Nasimul Hoda

List of Publications by Year in descending order

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Version: 2024-02-01

28
papers

1,107
citations

516710

16
h-index

501196

28
g-index

29
all docs

29
docs citations

29
times ranked

1766
citing authors

#	ARTICLE	IF	CITATIONS
1	Pivotal role of glycogen synthase kinase-3: A therapeutic target for Alzheimer's disease. <i>European Journal of Medicinal Chemistry</i> , 2016, 107, 63-81.	5.5	191
2	A comprehensive review of monoamine oxidase inhibitors as Anti-Alzheimer's disease agents: A review. <i>European Journal of Medicinal Chemistry</i> , 2020, 206, 112787.	5.5	123
3	Coumarin: A Privileged Scaffold for the Design and Development of Antineurodegenerative Agents. <i>Chemical Biology and Drug Design</i> , 2016, 87, 21-38.	3.2	100
4	Curcumin specifically binds to the human calcium-calmodulin-dependent protein kinase IV: fluorescence and molecular dynamics simulation studies. <i>Journal of Biomolecular Structure and Dynamics</i> , 2016, 34, 572-584.	3.5	68
5	Synthesis and screening of triazolopyrimidine scaffold as multi-functional agents for Alzheimer's disease therapies. <i>European Journal of Medicinal Chemistry</i> , 2016, 119, 260-277.	5.5	55
6	Rational design, synthesis and biological screening of triazine-triazolopyrimidine hybrids as multitarget anti-Alzheimer agents. <i>European Journal of Medicinal Chemistry</i> , 2017, 136, 36-51.	5.5	55
7	GSK3 Inhibitors in the Therapeutic Development of Diabetes, Cancer and Neurodegeneration: Past, Present and Future. <i>Current Pharmaceutical Design</i> , 2017, 23, 4332-4350.	1.9	54
8	A multifunctional therapeutic approach: Synthesis, biological evaluation, crystal structure and molecular docking of diversified 1H-pyrazolo[3,4-b]pyridine derivatives against Alzheimer's disease. <i>European Journal of Medicinal Chemistry</i> , 2019, 175, 2-19.	5.5	53
9	Dihydroorotate dehydrogenase: A drug target for the development of antimalarials. <i>European Journal of Medicinal Chemistry</i> , 2017, 125, 640-651.	5.5	52
10	Development of cyanopyridine-triazine hybrids as lead multitarget anti-Alzheimer agents. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 2777-2788.	3.0	48
11	Discovery of new phenyl sulfonyl-pyrimidine carboxylate derivatives as the potential multi-target drugs with effective anti-Alzheimer's action: Design, synthesis, crystal structure and in-vitro biological evaluation. <i>European Journal of Medicinal Chemistry</i> , 2021, 215, 113224.	5.5	37
12	Structure guided design of potential inhibitors of human calcium-calmodulin dependent protein kinase IV containing pyrimidine scaffold. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 782-788.	2.2	34
13	Alzheimer's Disease: A Systemic Review of Substantial Therapeutic Targets and the Leading Multi-functional Molecules. <i>Current Topics in Medicinal Chemistry</i> , 2018, 17, 3370-3389.	2.1	31
14	Design, synthesis, and biological evaluation of pyrimidine derivatives as potential inhibitors of human calcium/calmodulin-dependent protein kinase IV. <i>Chemical Biology and Drug Design</i> , 2017, 89, 741-754.	3.2	28
15	Pyrimidine-Triazolopyrimidine and Pyrimidine-Pyridine Hybrids as Potential Acetylcholinesterase Inhibitors for Alzheimer's Disease. <i>ChemistrySelect</i> , 2018, 3, 736-747.	1.5	24
16	Novel triazole-sulfonamide bearing pyrimidine moieties with carbonic anhydrase inhibitory action: Design, synthesis, computational and enzyme inhibition studies. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021, 48, 128249.	2.2	20
17	Naphthalene-triazolopyrimidine hybrid compounds as potential multifunctional anti-Alzheimer's agents. <i>Bioorganic and Medicinal Chemistry</i> , 2019, 27, 3156-3166.	3.0	16
18	An insight into the recent development of the clinical candidates for the treatment of malaria and their target proteins. <i>European Journal of Medicinal Chemistry</i> , 2021, 210, 112955.	5.5	15

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19	Recent advancements in chromone as a privileged scaffold towards the development of small molecules for neurodegenerative therapeutics. <i>RSC Medicinal Chemistry</i> , 2022, 13, 258-279.	3.9	15
20	Selective inhibitors of phosphodiesterases: therapeutic promise for neurodegenerative disorders. <i>MedChemComm</i> , 2015, 6, 2063-2080.	3.4	14
21	Quinoline carboxamide core moiety-based compounds inhibit <i>P. falciparum</i> falcipain-2: Design, synthesis and antimalarial efficacy studies. <i>Bioorganic Chemistry</i> , 2021, 108, 104514.	4.1	14
22	Dual targeting of acetylcholinesterase and tau aggregation: Design, synthesis and evaluation of multifunctional deoxyvasicinone analogues for Alzheimer's disease. <i>Bioorganic Chemistry</i> , 2021, 116, 105354.	4.1	13
23	New amyloid beta-disaggregating agents: synthesis, pharmacological evaluation, crystal structure and molecular docking of <i>N</i> -(4-((7-chloroquinolin-4-yl)oxy)-3-ethoxybenzyl)amines. <i>MedChemComm</i> , 2018, 9, 1891-1904.	3.4	10
24	Identification of lead BAY60-7550 analogues as potential inhibitors that utilize the hydrophobic groove in PDE2A: a molecular dynamics simulation study. <i>Journal of Molecular Modeling</i> , 2017, 23, 7.	1.8	9
25	Diphenyl triazine hybrids inhibit α -synuclein fibrillogenesis: Design, synthesis and <i>in vitro</i> efficacy studies. <i>European Journal of Medicinal Chemistry</i> , 2020, 207, 112705.	5.5	9
26	De novo lead optimization of triazine derivatives identifies potent antimalarials. <i>Journal of Molecular Graphics and Modelling</i> , 2017, 71, 96-103.	2.4	8
27	Deciphering the robustness of pyrazolo-pyridine carboxylate core structure-based compounds for inhibiting α -synuclein in transgenic <i>C. elegans</i> model of Synucleinopathy. <i>Bioorganic and Medicinal Chemistry</i> , 2020, 28, 115640.	3.0	6
28	Unravelling the potency of triazole analogues for inhibiting α -synuclein fibrillogenesis and <i>in vitro</i> disaggregation. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 1589-1603.	2.8	5