

# Mehdi Asadi

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6978527/publications.pdf>

Version: 2024-02-01

41  
papers

618  
citations

516710

16  
h-index

642732

23  
g-index

47  
all docs

47  
docs citations

47  
times ranked

621  
citing authors

#	ARTICLE	IF	CITATIONS
1	Recent Developments in Arylation of N-Nucleophiles via Chan-Lam Reaction: Updates from 2012 Onwards. <i>Current Organic Synthesis</i> , 2022, 19, 16-30.	1.3	6
2	Thebaine Derivatives as a New Regulator of Tumor Angiogenesis. <i>Polycyclic Aromatic Compounds</i> , 2022, 42, 4501-4519.	2.6	0
3	Synthesis and in vitro urease inhibitory activity of 5-nitrofuranyl-thiadiazole linked to different cyclohexyl-2-(phenylamino)acetamides, in silico and kinetic studies. <i>Bioorganic Chemistry</i> , 2022, 120, 105592.	4.1	14
4	Design and synthesis of novel nitrothiazolacetamide conjugated to different thioquinazolinone derivatives as anti-urease agents. <i>Scientific Reports</i> , 2022, 12, 2003.	3.3	21
5	Synthesis, molecular docking, and cytotoxicity of quinazolinone and dihydroquinazolinone derivatives as cytotoxic agents. <i>BMC Chemistry</i> , 2022, 16, 35.	3.8	1
6	Isoindolin-1-ones Fused to Barbiturates: From Design and Molecular Docking to Synthesis and Urease Inhibitory Evaluation. <i>ACS Omega</i> , 2022, 7, 19401-19411.	3.5	5
7	Novel (thio)barbituric-phenoxy-N-phenylacetamide derivatives as potent urease inhibitors: synthesis, in vitro urease inhibition, and in silico evaluations. <i>Structural Chemistry</i> , 2021, 32, 37-48.	2.0	19
8	Design, synthesis, and evaluation of novel racecadotril-tetrazole-amino acid derivatives as new potent analgesic agents. <i>Research in Pharmaceutical Sciences</i> , 2021, 16, 341.	1.8	3
9	Design, synthesis, and evaluation of metronidazole-1,2,3-triazole derivatives as potent urease inhibitors. <i>Chemical Papers</i> , 2021, 75, 4217-4226.	2.2	12
10	Synthesis, molecular docking, and biological evaluation of nitroimidazole derivatives as potent urease inhibitors. <i>Medicinal Chemistry Research</i> , 2021, 30, 1220-1229.	2.4	18
11	Arylmethylene hydrazine derivatives containing 1,3-dimethylbarbituric moiety as novel urease inhibitors. <i>Scientific Reports</i> , 2021, 11, 10607.	3.3	19
12	Recent Opportunities and Challenges in Selective C-H Functionalization of Methyl Azaarenes: a Highlight from 2010 to 2020 Literatures. <i>Current Organic Synthesis</i> , 2021, 18, 761-789.	1.3	0
13	Design, synthesis, docking study and urease inhibitory activity evaluation of novel 2-((5-amino-1,3,4-thiadiazol-2-yl)thio)-N-arylacetamide derivatives. <i>Medicinal Chemistry Research</i> , 2021, 30, 729-742.	2.4	14
14	Multicomponent reaction of amine, carbon disulfide, and fluoronitrobenzene via nucleophilic attack on the fluorinated carbon for the synthesis of nitrophenyl methylcarbomodithioates. <i>Journal of the Chinese Chemical Society</i> , 2020, 67, 160-164.	1.4	8
15	Novel fused 1,2,3-triazolo-benzodiazepine derivatives as potent anticonvulsant agents: design, synthesis, in vivo, and in silico evaluations. <i>Molecular Diversity</i> , 2020, 24, 179-189.	3.9	19
16	Novel N,N-dimethylbarbituric-pyridinium derivatives as potent urease inhibitors: Synthesis, in vitro, and in silico studies. <i>Bioorganic Chemistry</i> , 2020, 95, 103529.	4.1	21
17	Synthesis and biological evaluation of new benzimidazole-1,2,3-triazole hybrids as potential $\alpha$ -glucosidase inhibitors. <i>Bioorganic Chemistry</i> , 2020, 95, 103482.	4.1	50
18	Amine-carbon disulfide promoted synthesis of novel benzo[e][1,3]thiazepin-5(1H)-one derivatives. <i>Journal of Heterocyclic Chemistry</i> , 2020, 57, 413-418.	2.6	2

#	ARTICLE	IF	CITATIONS
19	Synthesis and Evaluation of Anti-Epileptic Properties of New Phthalimide-4,5-Dihydrothiazole-Amide Derivatives. <i>Polycyclic Aromatic Compounds</i> , 2020, , 1-11.	2.6	3
20	Synthesis and labeling of p-NH <sub>2</sub> -Bn-DTPA-(DabcyL-Lys6,Phe7)-pHBSP with <sup>99m</sup> Tc as a radiopeptide scintigraphic agent to detect cardiac ischemia. <i>Journal of Radioanalytical and Nuclear Chemistry</i> , 2020, 324, 635-646.	1.5	0
21	New 1,2,3-triazole (thio)barbituric acid hybrids as urease inhibitors: Design, synthesis, in vitro urease inhibition, docking study, and molecular dynamic simulation. <i>Archiv Der Pharmazie</i> , 2020, 353, e2000023.	4.1	29
22	Synthesis, in Vivo and in Silico Studies of N-Aryl-4-(1,3-Dioxisoindolin-2-Yl)Benzamides as an Anticonvulsant Agent. <i>Pharmaceutical Sciences</i> , 2020, 26, 38-44.	0.2	4
23	Design, Synthesis, Molecular Docking, and Cholinesterase Inhibitory Potential of Phthalimide-Dithiocarbamate Hybrids as New Agents for Treatment of Alzheimer's Disease. <i>Chemistry and Biodiversity</i> , 2019, 16, e1900370.	2.1	15
24	Design, Synthesis, and Cholinesterase Inhibition Assay of Coumarin-3-carboxamide-N-morpholine Hybrids as New Anti-Alzheimer Agents. <i>Chemistry and Biodiversity</i> , 2019, 16, e1900144.	2.1	28
25	Synthesis, molecular docking, and antiepileptic activity of novel phthalimide derivatives bearing amino acid conjugated anilines. <i>Research in Pharmaceutical Sciences</i> , 2019, 14, 534.	1.8	11
26	Design, synthesis, and biological evaluation of selective and potent Carbazole-based butyrylcholinesterase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 4952-4962.	3.0	17
27	Synthesis and Urease Inhibitory Activity of Some 5-Aminomethylene Barbituric/Thiobarbituric Acid Derivatives. <i>Letters in Drug Design and Discovery</i> , 2018, 15, 428-436.	0.7	8
28	Synthesis and Biological Evaluation of 1,3,4-Thiadiazole Linked Phthalimide Derivatives as Anticancer Agents. <i>Letters in Drug Design and Discovery</i> , 2017, 14, .	0.7	15
29	CuBr/Et <sub>3</sub> N-Promoted Reactions of 2-Aminobenzamides and Isothiocyanates: Efficient Synthesis of Novel Quinazolin-4(3H)-ones. <i>Helvetica Chimica Acta</i> , 2016, 99, 378-383.	1.6	8
30	Iodine-Mediated Synthesis of Novel Pyrazole Derivatives. <i>Synthesis</i> , 2016, 48, 541-546.	2.3	10
31	N-(2-(Piperazin-1-yl)phenyl)arylamide Derivatives as $\beta$ -Secretase (BACE1) Inhibitors: Simple Synthesis by Ugi Four-Component Reaction and Biological Evaluation. <i>Archiv Der Pharmazie</i> , 2015, 348, 330-337.	4.1	23
32	Synthesis and Evaluation of Chroman-4-one Linked to N-Benzyl Pyridinium Derivatives as New Acetylcholinesterase Inhibitors. <i>Archiv Der Pharmazie</i> , 2015, 348, 643-649.	4.1	22
33	Convenient and sequential one-pot route for synthesis of 2-thioxoquinazolinone and quinazolinobenzothiazinedione derivatives. <i>Monatshefte für Chemie</i> , 2014, 145, 497-504.	1.8	15
34	Green and Catalyst-Free One-Pot Synthesis of Anthranilamide Schiff Bases: An Approach Toward Sirtinol. <i>Synthetic Communications</i> , 2014, 44, 665-673.	2.1	10
35	Imidazo[2,1-b]thiazole derivatives as new inhibitors of 15-lipoxygenase. <i>European Journal of Medicinal Chemistry</i> , 2014, 87, 759-764.	5.5	30
36	Green Synthesis of New Boron-Containing Quinazolines: Preparation of Benzo[1,3,2]diazaborinin-4(1H)-one Derivatives. <i>Synthetic Communications</i> , 2013, 43, 2936-2942.	2.1	19

#	ARTICLE	IF	CITATIONS
37	Reaction of Isatoic Anhydride, Amine, and <i>N,N</i> -Dialkyl Carbodiimides Under Solvent-Free Conditions: New and Efficient Synthesis of 3-Alkyl-2-(alkylamino)quinazolin-4(3 <i>H</i> )-ones. <i>Synthetic Communications</i> , 2013, 43, 2385-2392.	2.1	27
38	Synthesis of Novel 1,4-Benzodiazepine-3,5-dione Derivatives: Reaction of 2-Aminobenzamides under Bargellini Reaction Conditions. <i>Synlett</i> , 2012, 23, 2521-2525.	1.8	33
39	One-Pot, Four-Component Synthesis of Novel Imidazo[2,1- <i>b</i> ]thiazol-5-amine Derivatives. <i>Synthesis</i> , 2012, 44, 3649-3654.	2.3	27
40	Synthesis of 2,3-diaryl-5 <i>H</i> -imidazo[2,1- <i>a</i> ]isoindol-5-ones via the one-pot reaction of 1,2-diketones, 2-formylbenzoic acids, and ammonium acetate. <i>Tetrahedron Letters</i> , 2012, 53, 3448-3451.	1.4	31
41	Synthesis, molecular docking, and antiepileptic activity of new <i>N</i> -phthaloylglycine derivatives. <i>Journal of the Iranian Chemical Society</i> , 0, , 1.	2.2	1