

Latifeh Navidpour

List of Publications by Year in descending order

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26
papers

535
citations

840776
11
h-index

642732
23
g-index

26
all docs

26
docs citations

26
times ranked

832
citing authors

#	ARTICLE	IF	CITATIONS
1	Design, synthesis, and biological evaluation of substituted 3-alkylthio-4,5-diaryl-4H-1,2,4-triazoles as selective COX-2 inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2006, 14, 2507-2517.	3.0	139
2	Design, synthesis, and biological evaluation of substituted 2-alkylthio-1,5-diarylimidazoles as selective COX-2 inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2007, 15, 1976-1982.	3.0	78
3	Synthesis, antibacterial activity, and quantitative structure–activity relationships of new (Z)-2-(nitroimidazolylmethylene)-3(-)-benzofuranone derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007, 17, 6354-6363.	2.2	68
4	Lipophilic 4-imidazolyl-1,4-dihydropyridines: synthesis, calcium channel antagonist activity and protection against pentylenetetrazole-induced seizure. <i>Il Farmaco</i> , 2004, 59, 261-269.	0.9	40
5	Design and synthesis of new water-soluble tetrazolidine derivatives of celecoxib and rofecoxib as selective cyclooxygenase-2 (COX-2) inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2006, 16, 4483-4487.	2.2	34
6	Bioassay-guided fractionation and identification of wound healing active compound from <i>Pistacia vera</i> L. hull extract. <i>Journal of Ethnopharmacology</i> , 2020, 248, 112335.	4.1	25
7	Chemodiversity of <i>Nepeta menthoides</i> Boiss. & Bohse. essential oil from Iran and antimicrobial, acetylcholinesterase inhibitory and cytotoxic properties of 1,8-cineole chemotype. <i>Natural Product Research</i> , 2018, 32, 2745-2748.	1.8	22
8	Design and synthesis of niflumic acid-based N-acylhydrazone derivatives as novel anti-inflammatory and analgesic agents. <i>Medicinal Chemistry Research</i> , 2013, 22, 2411-2420.	2.4	19
9	Characterization of arginine preventive effect on heat-induced aggregation of insulin. <i>International Journal of Biological Macromolecules</i> , 2020, 145, 1039-1048.	7.5	16
10	Synthesis, anti-inflammatory and analgesic activities of arylidene-2-(3-chloroanilino)nicotinic acid hydrazides. <i>Medicinal Chemistry Research</i> , 2014, 23, 2793-2802.	2.4	14
11	Synthesis, receptor affinity and effect on pentylenetetrazole-induced seizure threshold of novel benzodiazepine analogues: 3-Substituted 5-(2-phenoxybenzyl)-4H-1,2,4-triazoles and 2-amino-5-(phenoxybenzyl)-1,3,4-oxadiazoles. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 1929-1937.	3.0	13
12	A convenient synthesis of 5-alkylthio-3,4-diarylisoaxazoles by palladium-catalyzed coupling reactions. <i>Journal of Heterocyclic Chemistry</i> , 2007, 44, 449-453.	2.6	10
13	Amino Acids as Additives against Amorphous Aggregation: In Vitro and In Silico Study on Human Lysozyme. <i>Applied Biochemistry and Biotechnology</i> , 2019, 189, 305-317.	2.9	9
14	Syntheses of 5-alkylthio-1,3-diaryl-1,2,4-triazoles. <i>Journal of Heterocyclic Chemistry</i> , 2004, 41, 201-204.	2.6	8
15	5-[Aryloxy-pyridyl (or nitrophenyl)]-4H-1,2,4-triazoles as novel flexible benzodiazepine analogues: Synthesis, receptor binding affinity and lipophilicity-dependent anti-seizure onset of action. <i>Bioorganic Chemistry</i> , 2021, 106, 104504.	4.1	8
16	Syntheses of 4,5-Diaryl-1,2,3-thiadiazoles. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 2005, 180, 1593-1600.	1.6	7
17	Synthetic approaches towards the sulfonamide substituted 4,5-diaryl-1,2,4-triazole-3-thiones. <i>Journal of Heterocyclic Chemistry</i> , 2007, 44, 1323-1331.	2.6	6
18	Monocyclic phenolic compounds stabilize human insulin and suppress its amorphous aggregation: In vitro and in vivo study. <i>Biochemical and Biophysical Research Communications</i> , 2019, 518, 362-367.	2.1	5

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19	Synthetic Approaches towards the Sulfonamide Substituted 1,5-Diarylimidazole-2-thiones as Selective Cyclooxygenase-2 inhibitors. <i>Journal of Heterocyclic Chemistry</i> , 2014, 51, 71-79.	2.6	4
20	Antimalarial Activities of (Z)-2-(Nitroheteroarylmethylene)-3(2H)-Benzofuranone Derivatives: <i>In Vitro</i> and <i>In Vivo</i> Assessment and β -Hematin Formation Inhibition Activity. <i>Antimicrobial Agents and Chemotherapy</i> , 2021, 65, e0268320.	3.2	4
21	Synthesis, conformational assignment, and anti-inflammatory activities of N-arylidene-2-(4-chloro-2-(2-substituted phenoxy)phenyl)acetic acid hydrazides. <i>Medicinal Chemistry Research</i> , 2016, 25, 2220-2236.	2.4	3
22	Synthesis of aryl-substituted or aryl-fused N-hydroxyethyl and N-hydroxymethylpyrazole derivatives as potential ligands for the estrogen receptor. <i>Journal of the Iranian Chemical Society</i> , 2013, 10, 43-53.	2.2	2
23	Design, synthesis and apoptosis inducing activity of nonsteroidal flavone-methanesulfonate derivatives on MCF-7 cell line as potential sulfatase inhibitor. <i>Medicinal Chemistry Research</i> , 2021, 30, 1677-1687.	2.4	1
24	Synthesis and Calcium Channel Antagonist Activity of New 1,4-Dihydropyridine Derivatives Containing Lipophilic 4-Imidazolyl Substituents. <i>Arzneimittelforschung</i> , 2004, 54, 499-504.	0.4	0
25	COX inhibition: Catalepsy and Striatum Dopaminergic-GABAergic-Glutamatergic Neurotransmission. <i>Nature Precedings</i> , 2008, , .	0.1	0
26	Phenylalanine and indole effects on the pathogenicity of human lysozyme amorphous aggregates. <i>Enzyme and Microbial Technology</i> , 2022, 158, 110036.	3.2	0