

Mohammad Mahdavi

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4393171/publications.pdf>

Version: 2024-02-01

332
papers

6,720
citations

81743

39
h-index

143772

57
g-index

381
all docs

381
docs citations

381
times ranked

5233
citing authors

#	ARTICLE	IF	CITATIONS
1	A review on tacrine-based scaffolds as multi-target drugs (MTDLs) for Alzheimer's disease. <i>European Journal of Medicinal Chemistry</i> , 2017, 128, 332-345.	2.6	147
2	Design and synthesis of novel quinazolinone-1,2,3-triazole hybrids as new anti-diabetic agents: In vitro α -glucosidase inhibition, kinetic, and docking study. <i>Bioorganic Chemistry</i> , 2019, 83, 161-169.	2.0	119
3	Dimethyl Sulfoxide: Yesterday's Solvent, Today's Reagent. <i>Advanced Synthesis and Catalysis</i> , 2020, 362, 65-86.	2.1	112
4	Catalyst-free three-component reaction between 2-aminopyridines (or 2-aminothiazoles), aldehydes, and isocyanides in water. <i>Tetrahedron Letters</i> , 2007, 48, 7263-7265.	0.7	105
5	Novel tacrine-coumarin hybrids linked to 1,2,3-triazole as anti-Alzheimer's compounds: In vitro and in vivo biological evaluation and docking study. <i>Bioorganic Chemistry</i> , 2019, 83, 303-316.	2.0	94
6	Potent acetylcholinesterase inhibitors: Design, synthesis, biological evaluation, and docking study of acridone linked to 1,2,3-triazole derivatives. <i>European Journal of Medicinal Chemistry</i> , 2015, 92, 799-806.	2.6	91
7	C ³ -Functionalization of Imidazo[1,2-a]pyridines. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 269-284.	1.2	90
8	Novel tacrine-1,2,3-triazole hybrids: In vitro, in vivo biological evaluation and docking study of cholinesterase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2017, 125, 1200-1212.	2.6	88
9	Synthesis, in vitro cytotoxicity and apoptosis inducing study of 2-aryl-3-nitro-2H-chromene derivatives as potent anti-breast cancer agents. <i>European Journal of Medicinal Chemistry</i> , 2014, 86, 562-569.	2.6	84
10	Transition-Metal-Catalyzed Acyloxylation: Activation of C(sp ²)-H and C(sp ³)-H Bonds. <i>European Journal of Organic Chemistry</i> , 2016, 2016, 3282-3299.	1.2	82
11	Design, synthesis and anti-Alzheimer's activity of novel 1,2,3-triazole-chromenone carboxamide derivatives. <i>Bioorganic Chemistry</i> , 2019, 83, 391-401.	2.0	77
12	Hybrid Bionanocomposite Containing Magnesium Hydroxide Nanoparticles Embedded in a Carboxymethyl Cellulose Hydrogel Plus Silk Fibroin as a Scaffold for Wound Dressing Applications. <i>ACS Applied Materials & Interfaces</i> , 2021, 13, 33840-33849.	4.0	77
13	Design, synthesis, pharmacological evaluation, and docking study of new acridone-based 1,2,4-oxadiazoles as potential anticonvulsant agents. <i>European Journal of Medicinal Chemistry</i> , 2016, 112, 91-98.	2.6	75
14	New 6-amino-pyrido[2,3-d]pyrimidine-2,4-diones as novel agents to treat type 2 diabetes: A simple and efficient synthesis, α -glucosidase inhibition, molecular modeling and kinetic study. <i>European Journal of Medicinal Chemistry</i> , 2018, 155, 353-363.	2.6	75
15	Microwave-assisted efficient, one-pot, three-component synthesis of 3,5-disubstituted 1,2,4-oxadiazoles under solvent-free conditions. <i>Tetrahedron Letters</i> , 2006, 47, 2965-2967.	0.7	74
16	Synthesis and anti-leishmanial activity of 5-(5-nitrofuran-2-yl)-1,3,4-thiadiazol-2-amines containing N-[(1-benzyl-1H-1,2,3-triazol-4-yl)methyl] moieties. <i>European Journal of Medicinal Chemistry</i> , 2012, 50, 124-128.	2.6	71
17	Biscoumarin-1,2,3-triazole hybrids as novel anti-diabetic agents: Design, synthesis, in vitro α -glucosidase inhibition, kinetic, and docking studies. <i>Bioorganic Chemistry</i> , 2019, 92, 103206.	2.0	70
18	Multifunctional iminochromene-2H-carboxamide derivatives containing different aminomethylene triazole with BACE1 inhibitory, neuroprotective and metal chelating properties targeting Alzheimer's disease. <i>European Journal of Medicinal Chemistry</i> , 2017, 141, 690-702.	2.6	69

#	ARTICLE	IF	CITATIONS
19	Design, synthesis and in vitro α -glucosidase inhibition of novel dihydropyrano[3,2-c]quinoline derivatives as potential anti-diabetic agents. <i>Bioorganic Chemistry</i> , 2018, 77, 280-286.	2.0	68
20	Synthesis and anticancer activity of N-substituted 2-arylquinazolinones bearing trans-stilbene scaffold. <i>European Journal of Medicinal Chemistry</i> , 2015, 95, 492-499.	2.6	65
21	Large-scale virtual screening for the identification of new <i>Helicobacter pylori</i> urease inhibitor scaffolds. <i>Journal of Molecular Modeling</i> , 2012, 18, 2917-2927.	0.8	63
22	Synthesis of novel chromenones linked to 1,2,3-triazole ring system: Investigation of biological activities against Alzheimer's disease. <i>Bioorganic Chemistry</i> , 2017, 70, 86-93.	2.0	61
23	Synthesis of novel fused 4,5-dihydro-1,2,3-triazolo[1,5-a][1,4]benzodiazepine derivatives via four-component Ugi-Smiles-type reaction. <i>Tetrahedron</i> , 2013, 69, 3506-3510.	1.0	58
24	Design, Synthesis, Biological Evaluation, and Docking Study of Acetylcholinesterase Inhibitors: New Acridone-1,2,4-oxadiazole-1,2,3-triazole Hybrids. <i>Chemical Biology and Drug Design</i> , 2015, 86, 1425-1432.	1.5	58
25	Benzofuran-derived benzylpyridinium bromides as potent acetylcholinesterase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2015, 93, 196-201.	2.6	57
26	Design and synthesis of novel anti-Alzheimer's agents: Acridine-chromenone and quinoline-chromenone hybrids. <i>Bioorganic Chemistry</i> , 2016, 67, 84-94.	2.0	55
27	Synthesis, biological evaluation and docking study of 3-aryl-1-(4-sulfamoylphenyl)thiourea derivatives as 15-lipoxygenase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2014, 82, 308-313.	2.6	51
28	Design, synthesis and in vitro α -glucosidase inhibition of novel coumarin-pyridines as potent antidiabetic agents. <i>New Journal of Chemistry</i> , 2018, 42, 17268-17278.	1.4	51
29	Synthesis of New Benzimidazole-1,2,3-triazole Hybrids as Tyrosinase Inhibitors. <i>Chemistry and Biodiversity</i> , 2018, 15, e1800120.	1.0	50
30	Design, synthesis, docking study, α -glucosidase inhibition, and cytotoxic activities of acridine linked to thioacetamides as novel agents in treatment of type 2 diabetes. <i>Bioorganic Chemistry</i> , 2018, 80, 288-295.	2.0	50
31	Synthesis and biological evaluation of new benzimidazole-1,2,3-triazole hybrids as potential α -glucosidase inhibitors. <i>Bioorganic Chemistry</i> , 2020, 95, 103482.	2.0	50
32	Synthesis, characterization, molecular docking, and biological activities of coumarin-1,2,3-triazole-acetamide hybrid derivatives. <i>Archiv Der Pharmazie</i> , 2020, 353, e2000109.	2.1	50
33	Synthesis and structure-activity relationship study of benzofuran-based chalconoids bearing benzylpyridinium moiety as potent acetylcholinesterase inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2015, 103, 361-369.	2.6	48
34	Ullmann-Goldberg and Buchwald-Hartwig C-N Cross Couplings: Synthetic Methods to Pharmaceutically Potential Heterocycles. <i>Asian Journal of Organic Chemistry</i> , 2021, 10, 1319-1344.	1.3	46
35	Design and synthesis of novel coumarin-pyridinium hybrids: In vitro cholinesterase inhibitory activity. <i>Bioorganic Chemistry</i> , 2018, 77, 311-319.	2.0	44
36	Pectin-cellulose hydrogel, silk fibroin and magnesium hydroxide nanoparticles hybrid nanocomposites for biomedical applications. <i>International Journal of Biological Macromolecules</i> , 2021, 192, 7-15.	3.6	44

#	ARTICLE	IF	CITATIONS
37	Discovery of imidazopyridines containing isoindoline-1,3-dione framework as a new class of BACE1 inhibitors: Design, synthesis and SAR analysis. <i>European Journal of Medicinal Chemistry</i> , 2017, 138, 729-737.	2.6	42
38	Design, synthesis, in vitro cytotoxic activity evaluation, and apoptosis-induction study of new 9(10H)-acridinone-1,2,3-triazoles. <i>Molecular Diversity</i> , 2015, 19, 787-795.	2.1	41
39	New tacrine-derived AChE/BuChE inhibitors: Synthesis and biological evaluation of 5-amino-2-phenyl-4H-pyrano[2,3-b]quinoline-3-carboxylates. <i>European Journal of Medicinal Chemistry</i> , 2017, 128, 237-246.	2.6	41
40	Design, synthesis, biological evaluation, and docking study of novel dual-acting thiazole-pyridiniums inhibiting acetylcholinesterase and β -amyloid aggregation for Alzheimer's disease. <i>Bioorganic Chemistry</i> , 2020, 103, 104186.	2.0	41
41	Design, synthesis, characterization, enzymatic inhibition evaluations, and docking study of novel quinazolinone derivatives. <i>International Journal of Biological Macromolecules</i> , 2021, 170, 1-12.	3.6	40
42	A review on synthesis, mechanism of action, and structure-activity relationships of 1,2,3-triazole-based α -glucosidase inhibitors as promising anti-diabetic agents. <i>Journal of Molecular Structure</i> , 2022, 1255, 132469.	1.8	40
43	Palladium catalyst supported on N-aminoguanidine functionalized magnetic graphene oxide as a robust water-tolerant and versatile nanocatalyst. <i>RSC Advances</i> , 2014, 4, 48613-48620.	1.7	39
44	Design, synthesis, molecular modeling and anticholinesterase activity of benzylidene-benzofuran-3-ones containing cyclic amine side chain. <i>Future Medicinal Chemistry</i> , 2017, 9, 659-671.	1.1	39
45	Phthalimide-1,2,3-triazole hybrid compounds as tyrosinase inhibitors; synthesis, biological evaluation and molecular docking analysis. <i>Journal of Molecular Structure</i> , 2019, 1176, 86-93.	1.8	38
46	The natural-based optimization of kojic acid conjugated to different thio-quinazolinones as potential anti-melanogenesis agents with tyrosinase inhibitory activity. <i>Bioorganic and Medicinal Chemistry</i> , 2021, 36, 116044.	1.4	38
47	Magnetic Copper Ferrite Nanoparticles Functionalized by Aromatic Polyamide Chains for Hyperthermia Applications. <i>Langmuir</i> , 2021, 37, 8847-8854.	1.6	38
48	Novel quinazolinone linked to 1,2,3-triazoles: Synthesis and anticancer activity. <i>Chemical Biology and Drug Design</i> , 2018, 92, 1373-1381.	1.5	36
49	A Novel, One-Pot, Three-Component Synthesis of 1,2,4-Oxadiazoles under Microwave Irradiation and Solvent-Free Conditions. <i>Synlett</i> , 2006, 2006, 1765-1767.	1.0	34
50	Synthesis and pharmacological properties of polysubstituted 2-amino-4H-pyran-3-carbonitrile derivatives. <i>Molecular Diversity</i> , 2020, 24, 1385-1431.	2.1	34
51	Review: the latest advances in biomedical applications of chitosan hydrogel as a powerful natural structure with eye-catching biological properties. <i>Journal of Materials Science</i> , 2022, 57, 3855-3891.	1.7	34
52	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.	1.0	33
53	Efficient multi-component synthesis of 1,4-benzodiazepine-3,5-diones: a Petasis-based approach. <i>Tetrahedron</i> , 2015, 71, 6272-6275.	1.0	33
54	Synthesis, docking study and neuroprotective effects of some novel pyrano[3,2-c]chromene derivatives bearing morpholine/phenylpiperazine moiety. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 3980-3988.	1.4	33

#	ARTICLE	IF	CITATIONS
55	Design and synthesis of novel pyrazole-phenyl semicarbazone derivatives as potential α -glucosidase inhibitor: Kinetics and molecular dynamics simulation study. <i>International Journal of Biological Macromolecules</i> , 2021, 166, 1082-1095.	3.6	33
56	Copper supported β -cyclodextrin grafted magnetic nanoparticles as an efficient recyclable catalyst for one-pot synthesis of 1-benzyl-1H-1,2,3-triazole-dibenzodiazepinone derivatives via click reaction. <i>RSC Advances</i> , 2016, 6, 28838-28843.	1.7	32
57	Design and synthesis of new fused carbazole-imidazole derivatives as anti-diabetic agents: In vitro α -glucosidase inhibition, kinetic, and in silico studies. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2019, 29, 713-718.	1.0	32
58	Synthesis of 2,3-diaryl-5H-imidazo[2,1-a]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.	0.7	31
59	A green one-pot synthesis of N-alkyl-2-(2-oxoazepan-1-yl)-2-arylacamide derivatives via an Ugi four-center, three-component reaction in water. <i>Tetrahedron Letters</i> , 2012, 53, 7088-7092.	0.7	30
60	Imidazo[2,1-b]thiazole derivatives as new inhibitors of 15-lipoxygenase. <i>European Journal of Medicinal Chemistry</i> , 2014, 87, 759-764.	2.6	30
61	Sulfonic acid-functionalized poly(4-styrenesulfonic acid) mesoporous graphene oxide hybrid for one-pot preparation of coumarin-based pyrido[2,3-d]pyrimidine-dione derivatives. <i>Research on Chemical Intermediates</i> , 2020, 46, 491-507.	1.3	30
62	Recent advances in biological activities of rhodium complexes: Their applications in drug discovery research. <i>European Journal of Medicinal Chemistry</i> , 2021, 216, 113308.	2.6	30
63	Synthesis of Novel 1,2,3-Triazole-dihydro[3,2-c<i>/i>]chromenones as Acetylcholinesterase Inhibitors. <i>Synthetic Communications</i> , 2015, 45, 2311-2318.	1.1	29
64	Design, synthesis, in vitro, and in silico studies of novel diarylimidazole-1,2,3-triazole hybrids as potent α -glucosidase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2019, 27, 115148.	1.4	29
65	Anti-cancer, anti-oxidant and molecular docking studies of thiosemicarbazone indole-based derivatives. <i>Research on Chemical Intermediates</i> , 2019, 45, 2827-2854.	1.3	29
66	New Biscoumarin Derivatives as Potent α -Glucosidase Inhibitors: Synthesis, Biological Evaluation, Kinetic Analysis, and Docking Study. <i>Polycyclic Aromatic Compounds</i> , 2020, 40, 915-926.	1.4	29
67	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.	2.1	29
68	Anti-melanogenesis and anti-tyrosinase properties of aryl-substituted acetamides of phenoxy methyl triazole conjugated with thiosemicarbazide: Design, synthesis and biological evaluations. <i>Bioorganic Chemistry</i> , 2021, 114, 104979.	2.0	29
69	Ionic liquid-functionalized magnetic nanostructures as an efficient catalyst for the synthesis of 6H-chromeno[4,3-b]quinolin-6-ones. <i>Molecular Diversity</i> , 2017, 21, 597-609.	2.1	29
70	Design, Synthesis, and Cholinesterase Inhibition Assay of Coumarin-3-carboxamide<i>-N</i>-morpholine Hybrids as New Anti-Alzheimer Agents. <i>Chemistry and Biodiversity</i> , 2019, 16, e1900144.	1.0	28
71	Design and Synthesis of Selective Acetylcholinesterase Inhibitors: Arylisoxazole-Phenylpiperazine Derivatives. <i>Chemistry and Biodiversity</i> , 2019, 16, e1800433.	1.0	28
72	Design and synthesis of novel quinazolinone-pyrazole derivatives as potential α -glucosidase inhibitors: Structure-activity relationship, molecular modeling and kinetic study. <i>Bioorganic Chemistry</i> , 2021, 114, 105127.	2.0	28

#	ARTICLE	IF	CITATIONS
73	One-Pot, Four-Component Synthesis of Novel Imidazo[2,1-b]thiazol-5-amine Derivatives. <i>Synthesis</i> , 2012, 44, 3649-3654.	1.2	27
74	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.	1.1	27
75	Synthesis and Evaluation of Coumarin-Resveratrol Hybrids as 15-Lipoxygenase Inhibitors. <i>Synthetic Communications</i> , 2015, 45, 741-749.	1.1	27
76	Copper-catalyzed four-component synthesis of imidazo[1,2-a]pyridines via sequential reductive amination, condensation, and cyclization. <i>Tetrahedron Letters</i> , 2017, 58, 121-124.	0.7	27
77	Copper-supported β -cyclodextrin-functionalized magnetic nanoparticles: Efficient multifunctional catalyst for one-pot green synthesis of 1,2,3-triazolylquinazolinone derivatives. <i>Applied Organometallic Chemistry</i> , 2018, 32, e4212.	1.7	27
78	An efficient and targeted synthetic approach towards new highly substituted 6-amino-pyrazolo[1,5-a]pyrimidines with α -glucosidase inhibitory activity. <i>Scientific Reports</i> , 2020, 10, 2595.	1.6	27
79	Preparation of an improved sulfonated carbon-based solid acid as a novel, efficient, and reusable catalyst for chemoselective synthesis of 2-oxazolines and bis-oxazolines. <i>Monatshefte für Chemie</i> , 2009, 140, 1489-1494.	0.9	26
80	Synthesis of Novel Benzo[6,7][1,4]oxazepino[4,5-a]quinazolinone Derivatives via Transition-Metal-Free Intramolecular Hydroamination. <i>Synlett</i> , 2014, 25, 385-388.	1.0	26
81	Synthesis and characterization of β -Fe ₂ O ₃ @SiO ₂ -(CH ₂) ₃ -PDTCA-Pd magnetic nanoparticles: a new and highly active catalyst for the Heck/Sonogashira coupling reactions. <i>New Journal of Chemistry</i> , 2019, 43, 8930-8938.	1.4	26
82	Design and synthesis of new imidazo[1,2-b]pyrazole derivatives, in vitro α -glucosidase inhibition, kinetic and docking studies. <i>Molecular Diversity</i> , 2020, 24, 69-80.	2.1	26
83	Recent strategies in the synthesis of thiophene derivatives: highlights from the 2012-2020 literature. <i>Molecular Diversity</i> , 2021, 25, 2571-2604.	2.1	26
84	Design and Synthesis of Novel Arylisoxazole-Chromenone Carboxamides: Investigation of Biological Activities Associated with Alzheimer's Disease. <i>Chemistry and Biodiversity</i> , 2020, 17, e1900746.	1.0	26
85	Palladium functionalized phosphinite polyethyleneimine grafted magnetic silica nanoparticles as an efficient catalyst for the synthesis of isoquinolino[1,2-b]quinazolin-8-ones. <i>New Journal of Chemistry</i> , 2018, 42, 5499-5507.	1.4	25
86	Novel tetrahydrocarbazole benzyl pyridine hybrids as potent and selective butyryl cholinesterase inhibitors with neuroprotective and β -secretase inhibition activities. <i>European Journal of Medicinal Chemistry</i> , 2018, 155, 49-60.	2.6	25
87	A new series of Schiff base derivatives bearing 1,2,3-triazole: Design, synthesis, molecular docking, and α -glucosidase inhibition. <i>Archiv Der Pharmazie</i> , 2019, 352, e1900034.	2.1	25
88	New thiosemicarbazide-1,2,3-triazole hybrids as potent α -glucosidase inhibitors: Design, synthesis, and biological evaluation. <i>Journal of Molecular Structure</i> , 2019, 1192, 192-200.	1.8	25
89	A review on α -glucosidase inhibitory activity of first row transition metal complexes: a futuristic strategy for treatment of type 2 diabetes. <i>RSC Advances</i> , 2022, 12, 12011-12052.	1.7	25
90	A solvent-free reaction between acetophenone oximes and epoxy styrenes: an efficient synthesis of 2,4,6-triarylpyridines under neutral conditions. <i>Tetrahedron Letters</i> , 2014, 55, 3844-3846.	0.7	24

#	ARTICLE	IF	CITATIONS
91	Vilsmeier Reagent: An Efficient Reagent for the Transformation of 2-Aminobenzamides into Quinazolin-4(3H)-one Derivatives. <i>Synthetic Communications</i> , 2014, 44, 481-487.	1.1	24
92	Isoindolin-1-one derivatives as urease inhibitors: Design, synthesis, biological evaluation, molecular docking and in-silico ADME evaluation. <i>Bioorganic Chemistry</i> , 2019, 87, 1-11.	2.0	24
93	N-Cyclohexylimidazo[1,2-a]pyridine derivatives as multi-target-directed ligands for treatment of Alzheimer's disease. <i>Bioorganic Chemistry</i> , 2020, 103, 104146.	2.0	24
94	Synthesis, in vitro and in silico screening of 2-amino-4-aryl-6-(phenylthio) pyridine-3,5-dicarbonitriles as novel α -glucosidase inhibitors. <i>Bioorganic Chemistry</i> , 2020, 100, 103879.	2.0	24
95	Reaction between isocyanides and nitrostyrenes in water: a novel and efficient synthesis of 5-(alkylamino)-4-aryl-3-isoxazolecarboxamides. <i>Tetrahedron Letters</i> , 2009, 50, 7246-7248.	0.7	23
96	Novel Four-Step Synthesis of Thioxo-quinazolino[3,4-a]quinazolinone Derivatives. <i>Synthetic Communications</i> , 2014, 44, 215-221.	1.1	23
97	N-(2-(Piperazin-1-yl)phenyl)arylamide Derivatives as β -Secretase (BACE1) Inhibitors: Simple Synthesis by Ugi Four-Component Reaction and Biological Evaluation. <i>Archiv Der Pharmazie</i> , 2015, 348, 330-337.	2.1	23
98	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.	2.1	22
99	Copper (II)-supported polyethylenimine-functionalized magnetic graphene oxide as a catalyst for the green synthesis of 2-arylquinazolin-4(3H)-ones. <i>Research on Chemical Intermediates</i> , 2018, 44, 5241-5253.	1.3	22
100	Iodine-catalyzed tandem oxidative coupling reaction: A one-pot strategy for the synthesis of new coumarin-fused pyrroles. <i>Tetrahedron Letters</i> , 2018, 59, 94-98.	0.7	22
101	Synthesis, evaluation, and molecular docking studies of aryl urea-triazole-based derivatives as anti-urease agents. <i>Archiv Der Pharmazie</i> , 2018, 351, e1800005.	2.1	22
102	C-N cross-coupling reaction catalysed by efficient and reusable CuO/SiO ₂ nanoparticles under ligand-free conditions. <i>Applied Organometallic Chemistry</i> , 2014, 28, 809-813.	1.7	21
103	Simple and efficient syntheses of novel benzo[4,5]imidazo[1,2-a]pyridine derivatives. <i>Tetrahedron Letters</i> , 2015, 56, 743-746.	0.7	21
104	2-Imino 2H-chromene and 2-(phenylimino) 2H-chromene 3-aryl carboxamide derivatives as novel cytotoxic agents: synthesis, biological assay, and molecular docking study. <i>Journal of the Iranian Chemical Society</i> , 2016, 13, 2163-2171.	1.2	21
105	Synthesis and anticholinesterase activity of new substituted benzo[d]oxazole-based derivatives. <i>Chemical Biology and Drug Design</i> , 2017, 89, 783-789.	1.5	21
106	DABCO-modified superparamagnetic nanoparticles as an efficient and water-compatible catalyst for the synthesis of pyrano[3,2-c:5,6-c']dichromene-6,8-dione derivatives under mild reaction conditions. <i>Applied Organometallic Chemistry</i> , 2018, 32, e4561.	1.7	21
107	Novel N,N-dimethylbarbituric-pyridinium derivatives as potent urease inhibitors: Synthesis, in vitro, and in silico studies. <i>Bioorganic Chemistry</i> , 2020, 95, 103529.	2.0	21
108	Design and synthesis of 4,5-diphenyl-imidazol-1,2,3-triazole hybrids as new anti-diabetic agents: in vitro α -glucosidase inhibition, kinetic and docking studies. <i>Molecular Diversity</i> , 2021, 25, 877-888.	2.1	21

#	ARTICLE	IF	CITATIONS
109	Design and synthesis of novel nitrothiazolacetamide conjugated to different thioquinazolinone derivatives as anti-urease agents. <i>Scientific Reports</i> , 2022, 12, 2003.	1.6	21
110	Synthesis and cytotoxic activity of novel poly-substituted imidazo[2,1- <i>b</i>][1,2,4]triazin-6-amines. <i>Molecular Diversity</i> , 2015, 19, 273-281.	2.1	20
111	Synthesis and evaluation of novel oxoisindoline derivatives as acetylcholinesterase inhibitors. <i>Monatshefte für Chemie</i> , 2015, 146, 637-643.	0.9	20
112	Phthalimide-Derived Benzylpyridinium Halides Targeting Cholinesterases: Synthesis and Bioactivity of New Potential Anti-Alzheimer's Disease Agents. <i>Archiv Der Pharmazie</i> , 2016, 349, 293-301.	2.1	20
113	Novel morpholine containing cinnamoyl amides as potent tyrosinase inhibitors. <i>International Journal of Biological Macromolecules</i> , 2019, 135, 978-985.	3.6	20
114	CuBr ₂ -catalysed one-pot multicomponent synthesis of 3-substituted 2-thioxo-2,3-dihydroquinazolin-4(1 <i>H</i>)-one derivatives. <i>Applied Organometallic Chemistry</i> , 2019, 33, 1-7.	3.7	20
115	Recent Advances in Alkyne Hydroamination as a Powerful Tool for the Construction of C-N Bonds. <i>Asian Journal of Organic Chemistry</i> , 2020, 9, 969-991.	1.3	20
116	1,2,3-Triazole-Isoxazole Based Acetylcholinesterase Inhibitors: Synthesis, Biological Evaluation and Docking Study. <i>Letters in Drug Design and Discovery</i> , 2016, 14, 58-65.	0.4	20
117	Functionalized graphene oxide nanosheets with folic acid and silk fibroin as a novel nanobiocomposite for biomedical applications. <i>Scientific Reports</i> , 2022, 12, 6205.	1.6	20
118	Green Synthesis of New Boron-Containing Quinazolines: Preparation of Benzo[<i>d</i>][1,3,2]diazaborin-4(1 <i>H</i>)-one Derivatives. <i>Synthetic Communications</i> , 2013, 43, 2936-2942.	1.1	19
119	Combined isocyanide-based multi-component Ullmann-type reaction: an efficient access to novel nitrogen-containing pentacyclic compounds. <i>Molecular Diversity</i> , 2015, 19, 797-805.	2.1	19
120	Experimental and computational evidence for KOt-Bu-promoted synthesis of oxopyrazino[1,2- <i>a</i>]indoles. <i>RSC Advances</i> , 2015, 5, 101353-101361.	1.7	19
121	Synthesis of new benzo[<i>f</i>]imidazo[1,2- <i>d</i>][1,4]oxazepines: AgNO ₃ -mediated intramolecular hydroamination. <i>Tetrahedron Letters</i> , 2015, 56, 7082-7084.	0.7	19
122	Synthesis of Novel Tacrine Analogs as Acetylcholinesterase Inhibitors. <i>Journal of Heterocyclic Chemistry</i> , 2017, 54, 384-390.	1.4	19
123	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.	2.1	19
124	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.	1.0	19
125	Arylmethylene hydrazine derivatives containing 1,3-dimethylbarbituric moiety as novel urease inhibitors. <i>Scientific Reports</i> , 2021, 11, 10607.	1.6	19
126	Synthesis of Two Novel 3-Amino-5-[4-chloro-2-phenoxyphenyl]-4 <i>H</i> -1,2,4-triazoles with Anticonvulsant Activity. <i>Iranian Journal of Pharmaceutical Research</i> , 2010, 9, 265-9.	0.3	19

#	ARTICLE	IF	CITATIONS
127	Synthesis and Biological Investigation of some Novel Sulfonamide and Amide Derivatives Containing Coumarin Moieties. <i>Iranian Journal of Pharmaceutical Research</i> , 2014, 13, 881-92.	0.3	19
128	Novel Tacrine-Based Pyrano[3,4:5,6]pyrano[2,3- <i>b</i>]quinolinones: Synthesis and Cholinesterase Inhibitory Activity. <i>Archiv Der Pharmazie</i> , 2016, 349, 915-924.	2.1	18
129	An efficient four-component reaction for the synthesis of chromeno[4,3- <i>b</i>]quinolone derivatives. <i>Journal of the Iranian Chemical Society</i> , 2017, 14, 771-775.	1.2	18
130	Design, synthesis and biological evaluation of novel phthalimide-Schiff base-coumarin hybrids as potent α -glucosidase inhibitors. <i>Chemical Papers</i> , 2020, 74, 4379-4388.	1.0	18
131	α -Glucosidase and α -amylase inhibition, molecular modeling and pharmacokinetic studies of new quinazolinone-1,2,3-triazole-acetamide derivatives. <i>Medicinal Chemistry Research</i> , 2021, 30, 702-711.	1.1	18
132	Design, synthesis, in vitro α -glucosidase inhibition, docking, and molecular dynamics of new phthalimide-benzenesulfonamide hybrids for targeting type 2 diabetes. <i>Scientific Reports</i> , 2022, 12, .	1.6	18
133	Synthesis of novel indolo[2,3- <i>c</i>]quinolinones via Ugi-4CR/palladium-catalyzed arylation. <i>Tetrahedron</i> , 2014, 70, 3931-3934.	1.0	17
134	Cu(II)-cyclodextrin-catalyzed synthesis of spiro[indoline-3,4-pyrano[3,2- <i>c</i>]chromene]-3-carbonitrile derivatives. <i>Synthetic Communications</i> , 2017, 47, 2324-2329.	1.1	17
135	Design, synthesis, and biological evaluation of selective and potent Carbazole-based butyrylcholinesterase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 4952-4962.	1.4	17
136	Synthesis and Biological Activity of Some Benzochromenoquinolinones: Tacrine Analogs as Potent Anti-Alzheimer's Agents. <i>Chemistry and Biodiversity</i> , 2019, 16, e1800488.	1.0	17
137	One-pot multi-component synthesis of novel chromeno[4,3- <i>b</i>]pyrrol-3-yl derivatives as α -glucosidase inhibitors. <i>Molecular Diversity</i> , 2022, 26, 2393-2405.	2.1	17
138	Novel N-benzylpyridinium moiety linked to arylisoxazole derivatives as selective butyrylcholinesterase inhibitors: Synthesis, biological evaluation, and docking study. <i>Bioorganic Chemistry</i> , 2019, 92, 103192.	2.0	16
139	Thieno[2,3- <i>b</i>]pyridine amines: Synthesis and evaluation of tacrine analogs against biological activities related to Alzheimer's disease. <i>Archiv Der Pharmazie</i> , 2020, 353, 2000101.	2.1	16
140	Design, Synthesis and Cytotoxicity of Novel Coumarin-1,2,3-triazole-1,2,4- Oxadiazole Hybrids as Potent Anti-breast Cancer Agents. <i>Letters in Drug Design and Discovery</i> , 2019, 16, 818-824.	0.4	16
141	Novel magnetic organic-inorganic hybrids based on aromatic polyamides and ZnFe ₂ O ₄ nanoparticles with biological activity. <i>Scientific Reports</i> , 2021, 11, 20310.	1.6	16
142	Reaction between anthranilic acids, salicylaldehydes and isocyanides in water: an efficient synthesis of 2-[[2-(alkylimino)-1-benzofuran-3-ylidene]amino]benzoic acids. <i>Tetrahedron Letters</i> , 2010, 51, 27-29.	0.7	15
143	Synthesis of Isoindolo[2,1- <i>a</i>]quinazoline-5,11-dione Derivatives via the Reductive One-Pot Reaction of <i>N</i> -Substituted <i>N</i> -Nitrobenzamides and <i>N</i> -Formylbenzoic Acids. <i>Helvetica Chimica Acta</i> , 2013, 96, 419-423.	1.0	15
144	Convenient and sequential one-pot route for synthesis of 2-thioxoquinazolinone and quinazolinobenzothiazinedione derivatives. <i>Monatshefte für Chemie</i> , 2014, 145, 497-504.	0.9	15

#	ARTICLE	IF	CITATIONS
145	Synthesis of novel 5-phenylimidazo[1,2-c]quinazolin-3-amine derivatives via Groebke-B Blackburn-Bienaym multicomponent reaction. <i>Monatshefte für Chemie</i> , 2014, 145, 1483-1487.	9.9	15
146	Quinoline-based imidazole-fused heterocycles as new inhibitors of 15-lipoxygenase. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016, 31, 205-209.	2.5	15
147	Novel cinnamic acid-tryptamine hybrids as potent butyrylcholinesterase inhibitors: Synthesis, biological evaluation, and docking study. <i>Archiv Der Pharmazie</i> , 2018, 351, e1800115.	2.1	15
148	Synthesis and cholinesterase inhibitory activity of new 2-benzofuran carboxamide-benzylpyridinium salts. <i>Bioorganic Chemistry</i> , 2018, 80, 180-188.	2.0	15
149	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.	1.0	15
150	New benzyl pyridinium derivatives bearing 2,4-dioxochroman moiety as potent agents for treatment of Alzheimer's disease: Design, synthesis, biological evaluation, and docking study. <i>Bioorganic Chemistry</i> , 2019, 87, 506-515.	2.0	15
151	N-sulfonyl ketenimine as a versatile intermediate for the synthesis of heteroatom containing compounds. <i>Journal of Organometallic Chemistry</i> , 2021, 939, 121773.	0.8	15
152	Synthesis, in vitro, and in silico evaluation of Indazole Schiff bases as potential α -glucosidase inhibitors. <i>Journal of Molecular Structure</i> , 2021, 1242, 130826.	1.8	15
153	Design, synthesis, biological evaluation, and molecular modeling studies of pyrazole-benzofuran hybrids as new α -glucosidase inhibitor. <i>Scientific Reports</i> , 2021, 11, 20776.	1.6	15
154	An Efficient and Direct Solvent-Free Synthesis of Naphtho[1,2-b]furans, Naphtho[2,1-b]furans, and Furo[3,2-c]chromenes. <i>Synlett</i> , 2009, 2009, 2542-2544.	1.0	14
155	Potassium tert-Butoxide Promoted Intramolecular Amination of 1-Aryl-2-(2-nitrobenzylidene)hydrazines: Efficient Synthesis of 1-Aryl-1H-indazoles. <i>Synlett</i> , 2014, 25, 2605-2608.	1.0	14
156	An Efficient Synthesis of 2,4,6-Triarylpyridines via Solvent-Free Reaction between Acetophenoneoximes and Aldehydes. <i>Synlett</i> , 2014, 25, 1299-1301.	1.0	14
157	Synthesis of novel 5-arylidene (thio)barbituric acid and evaluation of their urease inhibitory activity. <i>Journal of the Iranian Chemical Society</i> , 2015, 12, 1487-1491.	1.2	14
158	Sulfamic acid-functionalized hydroxyapatite-encapsulated β -Fe ₂ O ₃ nanoparticles as a magnetically recoverable catalyst for synthesis of N-fused imidazole-quinoline conjugates under solvent-free conditions. <i>RSC Advances</i> , 2015, 5, 83530-83537.	1.7	14
159	Synthesis and Cytotoxic Activity of Some Novel Dihydrobenzo[<i>h</i>]pyrano[3,2- <i>c</i>]chromene Derivatives. <i>Journal of Heterocyclic Chemistry</i> , 2015, 52, 97-104.	1.4	14
160	Synthesis and In Vitro Cytotoxic Activity of Novel Triazole-Isoxazole Derivatives. <i>Journal of Heterocyclic Chemistry</i> , 2015, 52, 1743-1747.	1.4	14
161	A novel and efficient synthesis of 2-substituted quinazolin-4(3H)-ones by the reaction of (het)arylmethanamines with isoic anhydride. <i>Tetrahedron Letters</i> , 2016, 57, 3770-3772.	0.7	14
162	Design and Synthesis of Novel Cytotoxic Indole-Thiosemicarbazone Derivatives: Biological Evaluation and Docking Study. <i>Chemistry and Biodiversity</i> , 2019, 16, e1800470.	1.0	14

#	ARTICLE	IF	CITATIONS
163	Mo (CO) ₆ -assisted Pd-supported magnetic graphene oxide-catalyzed carbonylation-cyclization as an efficient way for the synthesis of 4(3 <i>H</i>)-quinazolinones. <i>Applied Organometallic Chemistry</i> , 2019, 33, e4769.	1.7	14
164	Novel and efficient synthesis of triazolobenzodiazepine analogues through the sequential Ugi 4CR-click-N-arylation reactions. <i>Tetrahedron Letters</i> , 2019, 60, 583-585.	0.7	14
165	Magnetic silica nanoparticle-supported copper complex as an efficient catalyst for the synthesis of novel triazolopyrazinylacetamides with improved antibacterial activity. <i>Chemistry of Heterocyclic Compounds</i> , 2020, 56, 488-494.	0.6	14
166	Design, Synthesis, and Molecular Docking of Some Novel Tacrine Based Cyclopentapyranopyridine and Tetrahydropyranoquinoline-Kojic Acid Derivatives as Anti-Acetylcholinesterase Agents. <i>Chemistry and Biodiversity</i> , 2021, 18, e2000924.	1.0	14
167	Triflic Anhydride (Tf) ₂ O: An Efficient Catalyst for Electrophilic Activation of Amides. <i>ChemistrySelect</i> , 2021, 6, 5320-5328.	0.7	14
168	Pyrano[3,2- <i>c</i>]quinoline Derivatives as New Class of α -glucosidase Inhibitors to Treat Type 2 Diabetes: Synthesis, in vitro Biological Evaluation and Kinetic Study. <i>Medicinal Chemistry</i> , 2019, 15, 8-16.	0.7	14
169	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.	2.0	14
170	Design, synthesis, and in silico studies of benzimidazole bearing phenoxyacetamide derivatives as α -glucosidase and α -amylase inhibitors. <i>Journal of Molecular Structure</i> , 2022, 1268, 133650.	1.8	14
171	Efficient Synthesis of 2-Methylenethiazolo[2,3- <i>b</i>]quinazolinone Derivatives. <i>Synlett</i> , 2015, 26, 173-176.	1.0	13
172	Hetero-annulated coumarins as new AChE/BuChE inhibitors: synthesis and biological evaluation. <i>Medicinal Chemistry Research</i> , 2016, 25, 1831-1841.	1.1	13
173	Synthesis and anti-acetylcholinesterase activity of benzotriazinone-triazole systems. <i>Journal of Chemical Sciences</i> , 2016, 128, 1445-1449.	0.7	13
174	Synthesis and Evaluation of Novel Quinazolinone-1,2,3-Triazoles as Inhibitors of Lipoxygenase. <i>Journal of Chemical Research</i> , 2016, 40, 188-191.	0.6	13
175	The use of magnetic starch as a support for an ionic liquid- β -cyclodextrin based catalyst for the synthesis of imidazothiadiazolamine derivatives. <i>International Journal of Biological Macromolecules</i> , 2019, 135, 453-461.	3.6	13
176	Design, synthesis, and evaluation of novel cinnamic acid-tryptamine hybrid for inhibition of acetylcholinesterase and butyrylcholinesterase. <i>DARU, Journal of Pharmaceutical Sciences</i> , 2020, 28, 463-477.	0.9	13
177	Efficient One Pot Synthesis of Phenylimidazo[1,2- <i>a</i>]pyridine Derivatives using Multifunctional Copper Catalyst Supported on β -Cyclodextrin Functionalized Magnetic Graphene oxide. <i>Applied Organometallic Chemistry</i> , 2020, 34, e5913.	1.7	13
178	Recent Advances in the Synthesis of Heterocycles by the Aza-Wittig Reaction. <i>Synthesis</i> , 2021, 53, 2342-2366.	1.2	13
179	Catalytic and non-catalytic amidation of carboxylic acid substrates. <i>Molecular Diversity</i> , 2022, 26, 1311-1344.	2.1	13
180	A review on the latest progress of C-S cross-coupling in diaryl sulfide synthesis: Update from 2012 to 2021. <i>Applied Organometallic Chemistry</i> , 2022, 36, e6482.	1.7	13

#	ARTICLE	IF	CITATIONS
181	Solvent-Free Reaction between Anthranilic Acids and Isocyanides: A Novel Approach for the Synthesis of 2-Unsubstituted 4(3H)-Quinazolinones. <i>Synlett</i> , 2011, 2011, 834-836.	1.0	12
182	Synthesis of novel fused quinazolinone derivatives. <i>Molecular Diversity</i> , 2016, 20, 677-685.	2.1	12
183	Synthesis of highly functionalized organic compounds through Ugi post-transformations started from propiolic acids. <i>Molecular Diversity</i> , 2020, 24, 855-887.	2.1	12
184	New phthalimide-benzamide-1,2,3-triazole hybrids; design, synthesis, \pm -glucosidase inhibition assay, and docking study. <i>Medicinal Chemistry Research</i> , 2020, 29, 868-876.	1.1	12
185	Novel <i>N</i> -benzylpiperidine derivatives of 5-arylisoxazole-carboxamides as anti-Alzheimer's agents. <i>Archiv Der Pharmazie</i> , 2021, 354, e2000258.	2.1	12
186	Design, synthesis, and evaluation of metronidazole-1,2,3-triazole derivatives as potent urease inhibitors. <i>Chemical Papers</i> , 2021, 75, 4217-4226.	1.0	12
187	Quinazolinone-dihydropyrano[3,2-b]pyran hybrids as new \pm -glucosidase inhibitors: Design, synthesis, enzymatic inhibition, docking study and prediction of pharmacokinetic. <i>Bioorganic Chemistry</i> , 2021, 109, 104703.	2.0	12
188	New quinoxalin-1,3,4-oxadiazole derivatives: Synthesis, characterization, in vitro biological evaluations, and molecular modeling studies. <i>Archiv Der Pharmazie</i> , 2021, 354, e2000471.	2.1	12
189	Synthesis and evaluation of novel arylisoxazoles linked to tacrine moiety: in vitro and in vivo biological activities against Alzheimer's disease. <i>Molecular Diversity</i> , 2022, 26, 409-428.	2.1	12
190	Design and synthesis of phenoxymethylbenzimidazole incorporating different aryl thiazole-triazole acetamide derivatives as \pm -glycosidase inhibitors. <i>Molecular Diversity</i> , 2021, , 1.	2.1	12
191	Efficient Solvent-Free Synthesis of Benzothiazine-Fused Pyrrolo[3,4-c]coumarins: Cycloaddition Reactions between Coumarin-Based Dihydrobenzothiazoles and Isocyanides. <i>Helvetica Chimica Acta</i> , 2014, 97, 847-853.	1.0	11
192	Novel 1,2,3,4-tetrahydroquinazolinones via Reaction of 2-Amino-substituted Benzamides and Dimethyl Acetylenedicarboxylate. <i>Helvetica Chimica Acta</i> , 2015, 98, 1028-1033.	1.0	11
193	Synthesis of Novel Pyrazino[2,1-a]isoindoliones via Intramolecular Hydroamination of 2,3-Dihydro-3-oxo-2-(prop-2-yn-1-yl)-1H-isoindole-1-carboxamides. <i>Helvetica Chimica Acta</i> , 2016, 99, 187-190.	1.0	11
194	Synthesis of Novel Isoindolo[2,1-a]quinazolinone Derivatives Containing a 1,2,3-Triazole Ring System. <i>Helvetica Chimica Acta</i> , 2016, 99, 37-40.	1.0	11
195	Synthesis and biological evaluation of novel imidazopyrimidinamines as anticancer agents. <i>Chemical Biology and Drug Design</i> , 2017, 89, 797-805.	1.5	11
196	Palladium supported aminobenzamide modified silica coated superparamagnetic iron oxide as an applicable nanocatalyst for Heck cross-coupling reaction. <i>Journal of Organometallic Chemistry</i> , 2021, 936, 121711.	0.8	11
197	New β -phenylpiperazine-carbodithioate- <i>N</i> -phenylacetamide hybrids: Synthesis, in vitro and in silico evaluations against cholinesterase and \pm -glucosidase enzymes. <i>Archiv Der Pharmazie</i> , 2022, 355, e2100313.	2.1	11
198	Novel phenylurea-pyridinium derivatives as potent urease inhibitors: Synthesis, in vitro, and in silico studies. <i>Journal of Molecular Structure</i> , 2022, 1263, 133078.	1.8	11

#	ARTICLE	IF	CITATIONS
199	An efficient one-pot synthesis of 3-aryl-1,2,4-oxadiazol-5-amines under solvent-free conditions. <i>Mendeleev Communications</i> , 2010, 20, 50-51.	0.6	10
200	Green and Catalyst-Free One-Pot Synthesis of Anthranilamide Schiff Bases: An Approach Toward Sirtinol. <i>Synthetic Communications</i> , 2014, 44, 665-673.	1.1	10
201	Palladium-Catalyzed Regioselective Direct Cyanation of Acetanilide Derivatives with $K_4[Fe(CN)_6]$ by C-H Bond Activation. <i>European Journal of Organic Chemistry</i> , 2016, 2016, 4269-4274.	1.2	10
202	One-pot synthesis of oxoisoindoline-1,2,3-triazole hybrid by a Ugi-click reaction. <i>Synthetic Communications</i> , 2016, 46, 1708-1712.	1.1	10
203	Efficient three-step synthesis of benzo[<i>c</i>]imidazo[1,2- <i>c</i>][1,2,3]triazines. <i>Synthetic Communications</i> , 2016, 46, 563-567.	1.1	10
204	Iodine-Mediated Synthesis of Novel Pyrazole Derivatives. <i>Synthesis</i> , 2016, 48, 541-546.	1.2	10
205	Green synthesis of 2-((2-aryl-3-oxoisoindolin-1-yl)methyl)quinazolin-4(3H)-ones via sequential condensation, sp ³ C-H bond functionalization and cyclization. <i>Tetrahedron Letters</i> , 2018, 59, 1555-1559.	0.7	10
206	SBA-15-SO ₃ H-assisted preparation of 4-aza-phenanthrene-3,10-dione derivatives via a one-pot, four-component reaction. <i>Research on Chemical Intermediates</i> , 2018, 44, 739-747.	1.3	10
207	Anticancer properties of N-alkyl-2, 4-diphenylimidazo [1, 2-a] quinoxalin-1-amine derivatives; kinase inhibitors. <i>Bioorganic Chemistry</i> , 2019, 90, 103055.	2.0	10
208	New ciprofloxacin-dithiocarbamate-benzyl hybrids: design, synthesis, antibacterial evaluation, and molecular modeling studies. <i>Research on Chemical Intermediates</i> , 2019, 45, 223-236.	1.3	10
209	New acridine-9-carboxamide linked to 1,2,3-triazole-N-phenylacetamide derivatives as potent α -glucosidase inhibitors: design, synthesis, in vitro, and in silico biological evaluations. <i>Medicinal Chemistry Research</i> , 2020, 29, 1836-1845.	1.1	10
210	Design, synthesis, and α -glucosidase-inhibitory activity of phenoxy-bis coumarin-N-phenylacetamide hybrids. <i>Archiv Der Pharmazie</i> , 2021, 354, e2100179.	2.1	10
211	Synthesis, and in vitro biological evaluations of novel naphthoquinone conjugated to aryl triazole acetamide derivatives as potential anti-Alzheimer agents. <i>Journal of Molecular Structure</i> , 2022, 1255, 132229.	1.8	10
212	Synthesis and characterization of cellulose, β -cyclodextrin, silk fibroin-based hydrogel containing copper-doped cobalt ferrite nanospheres and exploration of its biocompatibility. <i>Journal of Nanostructure in Chemistry</i> , 2023, 13, 103-113.	5.3	10
213	Photochemical regioselective C-H arylation of imidazo[1,2- <i>a</i>]pyridine derivatives using chlorophyll as a biocatalyst and diazonium salts. <i>New Journal of Chemistry</i> , 2022, 46, 10814-10819.	1.4	10
214	Synthesis of Novel 2-Oxoquinoline Derivatives via Ugi-Four-Component-Heck Reaction. <i>Journal of Heterocyclic Chemistry</i> , 2015, 52, 386-391.	1.4	9
215	Synthesis of Novel Phthalazino[1,2- <i>b</i>]quinazolin-4(1H)-one Derivatives: Efficient and Practical Reaction of 2-Amino-N-arylbisbenzohydrazides and 2-Formylbenzoic Acids. <i>Helvetica Chimica Acta</i> , 2016, 99, 539-542.	1.0	9
216	Novel quinazolin-sulfonamid derivatives: synthesis, characterization, biological evaluation, and molecular docking studies. <i>Journal of Biomolecular Structure and Dynamics</i> , 2020, , 1-12.	2.0	9

#	ARTICLE	IF	CITATIONS
217	Synthesis and biological evaluation of new dihydroindolizino[8,7-b]indole derivatives as novel α -glucosidase inhibitors. <i>Journal of Molecular Structure</i> , 2021, 1224, 129290.	1.8	9
218	Electrochemical synthesis of three-dimensional flower-like Ni/Co-BTC bimetallic organic framework as heterogeneous catalyst for solvent-free and green synthesis of substituted chromeno[4,3-b]quinolones. <i>Journal of the Chinese Chemical Society</i> , 2021, 68, 620-629.	0.8	9
219	Copper Supported onto Magnetic Nanoparticles as an Efficient Catalyst for the Synthesis of Triazolobenzodiazepino[7,1-b]quinazolin-1(9H)-ones via Click N-Arylation Reactions. <i>ChemistrySelect</i> , 2021, 6, 1385-1392.	0.7	9
220	Design and synthesis of a novel nanocomposite based on magnetic dopamine nanoparticles for purification of α -amylase from the bovine milk. <i>Scientific Reports</i> , 2021, 11, 13428.	1.6	9
221	Isatoic Anhydride: A Fascinating and Basic Molecule for the Synthesis of Substituted Quinazolinones and Benzo di/triazepines. <i>Current Organic Chemistry</i> , 2019, 23, 1090-1130.	0.9	9
222	A novel, bioactive and antibacterial scaffold based on functionalized graphene oxide with lignin, silk fibroin and ZnO nanoparticles. <i>Scientific Reports</i> , 2022, 12, .	1.6	9
223	Highly Efficient Synthesis of 14-Aryl-14H-dibenzo[a,j]xanthenes Catalyzed by Carbon-Based Solid Acid Under Solvent-Free Conditions. <i>Synthetic Communications</i> , 2009, 39, 4328-4340.	1.1	8
224	A Highly Efficient Method for the Synthesis of Novel 1 ^H -spiro[indene-2,2'-quinazoline]-1,3,4'-trione Derivatives. <i>Journal of Chemical Research</i> , 2015, 39, 495-498.	0.6	8
225	CuBr/Et ₃ N-Promoted Reactions of α -Aminobenzamides and Isothiocyanates: Efficient Synthesis of Novel Quinazolin-4(3H)-ones. <i>Helvetica Chimica Acta</i> , 2016, 99, 378-383.	1.0	8
226	Biology-Oriented Drug Synthesis (BIODS) Approach towards Synthesis of Ciprofloxacin-Dithiocarbamate Hybrids and Their Antibacterial Potential both <i>in Vitro</i> and <i>in Silico</i> . <i>Chemistry and Biodiversity</i> , 2018, 15, e1800273.	1.0	8
227	Design, synthesis, and biological evaluation of novel 4-oxobenzo[d]1,2,3-triazin-benzylpyridinium derivatives as potent anti-Alzheimer agents. <i>Bioorganic and Medicinal Chemistry</i> , 2019, 27, 2914-2922.	1.4	8
228	Facile Non-Transition Metal-Catalyzed Synthesis of 2-Thioxo-2,3-dihydroquinazolin-4(1H)-one Derivatives via One-Pot Multicomponent Reactions. <i>ChemistrySelect</i> , 2019, 4, 100-104.	0.7	8
229	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.	0.8	8
230	Benzoylquinazolinone derivatives as new potential antidiabetic agents: α -glucosidase inhibition, kinetic, and docking studies. <i>Journal of the Chinese Chemical Society</i> , 2020, 67, 856-863.	0.8	8
231	New 4,5-diphenylimidazole-acetamide-1,2,3-triazole hybrids as potent α -glucosidase inhibitors: synthesis, <i>in vitro</i> and <i>in silico</i> enzymatic and toxicity evaluations. <i>Monatshefte für Chemie</i> , 2021, 152, 679-693.	0.9	8
232	C1-Functionalization of 1,2,3,4-Tetrahydroisoquinolines (THIQs). <i>Asian Journal of Organic Chemistry</i> , 2021, 10, 2421-2439.	1.3	8
233	Novel Indole-Isoxazole Hybrids: Synthesis and <i>In Vitro</i> Anti-Cholinesterase Activity. <i>Letters in Drug Design and Discovery</i> , 2017, 14, .	0.4	8
234	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.	0.4	8

#	ARTICLE	IF	CITATIONS
235	Rational Design, Synthesis, <i>in Vitro</i> , and <i>in Silico</i> Studies of Chlorophenylquinazolin-4(3H)-one Containing Different Aryl Acetohydrazides as Tyrosinase Inhibitors. <i>Chemistry and Biodiversity</i> , 2022, 19, .	1.0	8
236	Design, synthesis, and biological evaluation of new series of 2-amido-1,3,4-thiadiazole derivatives as cytotoxic agents. <i>Zeitschrift Fur Naturforschung - Section B Journal of Chemical Sciences</i> , 2016, 71, 205-210.	0.3	7
237	Improvement of the Van Leusen reaction in the presence of β -cyclodextrin: a green and efficient synthesis of oxazoles in water. <i>Zeitschrift Fur Naturforschung - Section B Journal of Chemical Sciences</i> , 2017, 72, 923-926.	0.3	7
238	Copper-catalyzed efficient synthesis of 5-arylindazolo[3,2-b]quinazolin-7(5H)-ones from 2-nitrobenzaldehydes. <i>Tetrahedron</i> , 2018, 74, 2197-2201.	1.0	7
239	Novel Coumarin Containing Dithiocarbamate Derivatives as Potent α -Glucosidase Inhibitors for Management of Type 2 Diabetes. <i>Medicinal Chemistry</i> , 2021, 17, 264-272.	0.7	7
240	Utilizing Amines and Carbon Disulfide to Obtain Nitrogen- and Sulfur-containing Compounds under Green Conditions: A Review. <i>Current Organic Chemistry</i> , 2018, 22, 2315-2380.	0.9	7
241	Characteristics of published/registered clinical trials on COVID-19 treatment: A systematic review. <i>DARU, Journal of Pharmaceutical Sciences</i> , 2021, 29, 449-467.	0.9	7
242	In silico and in vitro studies of thiosemicarbazone-indole hybrid compounds as potent α -glycosidase inhibitors. <i>Computational Biology and Chemistry</i> , 2022, 97, 107642.	1.1	7
243	A novel and efficient route for the synthesis of 5-nitrobenzo[d]oxazole derivatives. <i>Journal of Fluorine Chemistry</i> , 2014, 161, 83-86.	0.9	6
244	Synthesis, antileishmanial activity and QSAR study of (1,3,4-thiadiazol-2-ylthio) acetamides derived from 5-nitrofurans. <i>Medicinal Chemistry Research</i> , 2015, 24, 891-900.	1.1	6
245	Straightforward Approach Toward Dihydrothiazoles via Intramolecular Bromocyclization. <i>Synthetic Communications</i> , 2015, 45, 2142-2147.	1.1	6
246	Three-component one-pot synthesis of dihydrochromeno[4,3-b]pyrazolo[4,3-e]pyridines. <i>Heterocyclic Communications</i> , 2016, 22, 247-250.	0.6	6
247	An efficient approach to the synthesis of coumarin-fused dihydropyridinones. <i>Heterocyclic Communications</i> , 2017, 23, 305-308.	0.6	6
248	Metal-free, air-promoted, radical-mediated arylation of benzoquinone with phenylhydrazines. <i>Zeitschrift Fur Naturforschung - Section B Journal of Chemical Sciences</i> , 2018, 73, 703-706.	0.3	6
249	4-Oxobenzo[d]1,2,3-triazin-pyridinium-phenylacetamide derivatives as new anti-Alzheimer agents: design, synthesis, in vitro evaluation, molecular modeling, and molecular dynamic study. <i>Structural Chemistry</i> , 2020, 31, 999-1012.	1.0	6
250	The possible effect of microRNA-155 (miR-155) and BACE1 inhibitors in the memory of patients with down syndrome and Alzheimer's disease: Design, synthesis, virtual screening, molecular modeling and biological evaluations. <i>Journal of Biomolecular Structure and Dynamics</i> , 2022, 40, 5803-5814.	2.0	6
251	Recent Developments in Arylation of N-Nucleophiles via Chan-Lam Reaction: Updates from 2012 Onwards. <i>Current Organic Synthesis</i> , 2022, 19, 16-30.	0.7	6
252	Copper-catalyzed one-pot synthesis of amide linked 1,2,3-triazoles bearing aryloxy skeletons. <i>Tetrahedron Letters</i> , 2021, 65, 152765.	0.7	6

#	ARTICLE	IF	CITATIONS
253	Palladium-coated thiourea core-shell nanocomposite as a new, efficient, and magnetic responsive nanocatalyst for the Suzuki-Miyaura coupling reactions. <i>Materials Research Express</i> , 2021, 8, 026102.	0.8	6
254	Synthesis of novel tetracyclic coumarin-fused furo-pyridone scaffolds via sequential N-arylation and intramolecular amidation reactions. <i>Tetrahedron Letters</i> , 2021, 68, 152904.	0.7	6
255	Synthesis, in vitro and in silico enzymatic inhibition assays, and toxicity evaluations of new 4,5-diphenylimidazole-N-phenylacetamide derivatives as potent α -glucosidase inhibitors. <i>Medicinal Chemistry Research</i> , 2021, 30, 1273-1283.	1.1	6
256	Synthesis of Thiazolone Derivatives as Novel Soybean 15-LOX Inhibitors. <i>Letters in Organic Chemistry</i> , 2017, 14, 186-191.	0.2	6
257	Design, Synthesis, In vitro Cytotoxic Activity Evaluation, and Study of Apoptosis Inducing Effect of New Styrylimidazo[1,2-a]Pyridines as Potent Anti-Breast Cancer Agents. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2019, 19, 265-275.	0.9	6
258	Synthesis and biological evaluation of a new series of benzofuran-1,3,4-oxadiazole containing 1,2,3-triazole-acetamides as potential α -glucosidase inhibitors. <i>Journal of Biochemical and Molecular Toxicology</i> , 2021, 35, e22688.	1.4	6
259	Pd@Py2PZ@MSN as a Novel and Efficient Catalyst for C-C Bond Formation Reactions. <i>Frontiers in Chemistry</i> , 2022, 10, 838294.	1.8	6
260	Sodium Azide: An Inorganic Nitrogen Source for the Synthesis of Organic N-Compounds. <i>ChemistrySelect</i> , 2021, 6, 13419-13433.	0.7	6
261	An Efficient Synthesis of Novel Dihydrothiazol-2-yl-amides via Cyclisation of Propargylic Carbamthioyl-amides. <i>Journal of Chemical Research</i> , 2014, 38, 131-133.	0.6	5
262	Efficient Synthesis of Novel Thiazol-2-ylidene-amides Using Carbonylthiourea Building Blocks. <i>Journal of Heterocyclic Chemistry</i> , 2015, 52, 1150-1153.	1.4	5
263	Efficient Synthesis of Polyfunctionalized Pyrimidine Derivatives. <i>Synlett</i> , 2016, 27, 1689-1692.	1.0	5
264	Synthesis and Cytotoxic Evaluation of Novel 1,2,3-Triazole-4-Linked (2-E,6-E)-2-Benzylidene-6-(4-nitrobenzylidene)cyclohexanones. <i>Helvetica Chimica Acta</i> , 2016, 99, 175-180.	1.0	5
265	Copper-catalyzed synthesis of 2,3-disubstituted quinazolin-4(3H)-ones from benzyl-substituted anthranilamides. <i>Heterocyclic Communications</i> , 2018, 24, 267-271.	0.6	5
266	Synthesis and Characterization of Novel Phthalimide-pyrano[3,2-c]chromene and Phthalimide-pyrano-2-one Hybrids. <i>Journal of Heterocyclic Chemistry</i> , 2018, 55, 1678-1684.	1.4	5
267	Preparation of some novel imidazopyridine derivatives of indole as anticancer agents: one-pot multicomponent synthesis, biological evaluation and docking studies. <i>Research on Chemical Intermediates</i> , 2019, 45, 5261-5290.	1.3	5
268	Facile access to new pyrido[2,3-d]pyrimidine derivatives. <i>Molecular Diversity</i> , 2019, 23, 333-340.	2.1	5
269	Synthesis of Arylidene Isoquinolinones bearing Combretastatin Skeleton by Cyclocarbopalladation/cross coupling Tandem Heck-Suzuki Miyaura Reactions using nano catalyst Pd@Py@SPION. <i>Applied Organometallic Chemistry</i> , 2020, 34, e5279.	1.7	5
270	Design, synthesis, biological evaluation, and docking study of new acridine-9-carboxamide linked to 1,2,3-triazole derivatives as antidiabetic agents targeting α -glucosidase. <i>Journal of Heterocyclic Chemistry</i> , 2020, 57, 4348-4357.	1.4	5

#	ARTICLE	IF	CITATIONS
271	Synthesis, in vitro, and in silico studies of newly functionalized quinazolinone analogs for the identification of potent β -glucosidase inhibitors. <i>Journal of the Iranian Chemical Society</i> , 2021, 18, 2017-2034.	1.2	5
272	Sulfonic Acid Functionalized Magnetic Starch as an Efficient Catalyst for the Synthesis of Chromeno[4,3-b]quinoline-6,8-dione Derivatives. <i>Starch/Staerke</i> , 2021, 73, 2000257.	1.1	5
273	Efficient synthesis of chromeno[4,3-b]pyrano[3,4-e]pyridine-6,8-dione derivatives via multicomponent one-pot reaction under mild reaction conditions in water. <i>Research on Chemical Intermediates</i> , 2021, 47, 4101-4112.	1.3	5
274	Bi Metal-Organic Framework (Ce/Ni-BTC) as Heterogeneous Catalyst for the Green Synthesis of Substituted Chromeno[4,3-b]quinolone under Solvent Free Condition. <i>Current Organic Synthesis</i> , 2021, 18, 475-482.	0.7	5
275	Identification of Essential 2D and 3D Chemical Features for Discovery of the Novel Tubulin Polymerization Inhibitors. <i>Current Topics in Medicinal Chemistry</i> , 2019, 19, 1092-1120.	1.0	5
276	In vitro cell-based models of drug-induced hepatotoxicity screening: progress and limitation. <i>Drug Metabolism Reviews</i> , 2022, 54, 161-193.	1.5	5
277	Synthesis and cytotoxicity of novel chromenone derivatives bearing 4-nitrophenoxy phenyl acryloyl moiety. <i>Journal of the Iranian Chemical Society</i> , 2016, 13, 1139-1144.	1.2	4
278	A Novel Copper-Catalyzed Preparation of Pyrido[1,2-a]pyrimidine Derivatives. <i>Synlett</i> , 2016, 27, 1359-1362.	1.0	4
279	Sulfonic Acid Supported Phosphonium Based Ionic Liquid Functionalized SBA-15 for the Synthesis of 2-Amino-3-cyano-4,6-diarylpyridines. <i>Synthesis and Reactivity in Inorganic, Metal Organic, and Nano Metal Chemistry</i> , 2016, 46, 306-310.	0.6	4
280	A green and efficient synthesis of 2-thioxoquinazolinone derivatives in water using potassium thiocyanate. <i>Journal of Sulfur Chemistry</i> , 2017, 38, 519-529.	1.0	4
281	An efficient, four-component reaction for the synthesis of novel carbamodithioates. <i>Journal of Sulfur Chemistry</i> , 2017, 38, 43-51.	1.0	4
282	Synthesis and biological evaluation of chalcone-triazole hybrid derivatives as 15-LOX inhibitors. <i>Zeitschrift Fur Naturforschung - Section B Journal of Chemical Sciences</i> , 2018, 73, 77-83.	0.3	4
283	Efficient copper-catalyzed synthesis of 2-arylbenzimidazole derivatives by reaction of 1-fluoro-2-nitrobenzene with benzamidine hydrochlorides. <i>Chemistry of Heterocyclic Compounds</i> , 2018, 54, 351-354.	0.6	4
284	Catalyst-free three-component synthesis of 2-amino-4,6-diarylpyridine-3-carbonitriles under solvent-free conditions. <i>Chemistry of Heterocyclic Compounds</i> , 2019, 55, 725-728.	0.6	4
285	Synthesis of quinazolin-4(3H)-ones via the reaction of isatoic anhydride with benzyl azides in the presence of potassium tert-butoxide in DMSO. <i>Chemistry of Heterocyclic Compounds</i> , 2019, 55, 964-967.	0.6	4
286	A one-pot and three-component synthetic approach for the preparation of asymmetric and multi-substituted 1,4-dihydropyrazines. <i>Tetrahedron Letters</i> , 2019, 60, 151257.	0.7	4
287	Design, synthesis, in vivo and in vitro studies of 1,2,3,4-tetrahydro-9H-carbazole derivatives, highly selective and potent butyrylcholinesterase inhibitors. <i>Molecular Diversity</i> , 2020, 24, 211-223.	2.1	4
288	Regio- and Diastereoselective $\text{KMnO}_4/\text{RCO}_2\text{H}$ Mediated Acyloxyarylation of Chalcones - An Indirect β -Arylation of Chalcones. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 2045-2051.	1.2	4

#	ARTICLE	IF	CITATIONS
289	Comparison of serologic status of <i>Toxoplasma gondii</i> infection in pre- and post-heart transplantation in a pediatric population: A preliminary study. <i>Transplant Infectious Disease</i> , 2020, 22, e13339.	0.7	4
290	Synthesis and In Vitro Biological Activity Evaluation of Novel Imidazo [2,1-B][1,3,4] Thiadiazole as Anti-Alzheimer Agents. <i>Letters in Drug Design and Discovery</i> , 2020, 17, 610-617.	0.4	4
291	Design and synthesis of new benzofuran-1,2,3-triazole hybrid preservatives and the evaluation of their antifungal potential against white and brown-rot fungi. <i>BioResources</i> , 2020, 15, 7828-7843.	0.5	4
292	Novel aryl(4-phenylpiperazin-1-yl)methanethione derivatives as new anti-Alzheimer agents: Design, synthesis, in vitro and in silico assays. <i>Journal of Molecular Structure</i> , 2022, 1262, 132945.	1.8	4
293	N-Arylation Reaction of 2-Amino-N-phenylbenzamide with Phenyl Boronic Acid via Chan-Evans-Lam (CEL) Type Reaction Using Cu@Phen@MGO Catalyst. <i>Catalysis Letters</i> , 2023, 153, 805-813.	1.4	4
294	6-Methoxy-4-chlorotetralone Derivatives Bearing an Arylpyridinium Moiety as Cholinesterase Inhibitors: Design, Synthesis, Biological Evaluation, and Molecular Docking Study. <i>ChemistrySelect</i> , 2022, 7, .	0.7	4
295	Efficient and Ecofriendly Route for the Solvent-Free Synthesis of 4-Alkoxy-5H-chromen[2,3-d]pyrimidines Using Phosphonic Acid Functionalized KIT-6 Confined Ionic Liquid as Recoverable Catalyst. <i>Synthetic Communications</i> , 2014, 44, 2826-2837.	1.1	3
296	Synthesis and biological evaluation of 1,3,4,5-tetrasubstituted pyrazole derivatives. <i>Zeitschrift Fur Naturforschung - Section B Journal of Chemical Sciences</i> , 2016, 71, 973-977.	0.3	3
297	An efficient access to 2,3-diarylimidazo[1,2-a]pyridines via silver(I)-catalyzed C-H bond functionalization. <i>Monatshefte Für Chemie</i> , 2017, 148, 1817-1821.	0.9	3
298	Appel reagent as novel promoter for the synthesis of polysubstituted imidazoles. <i>Arkivoc</i> , 2017, 2017, 343-352.	0.3	3
299	Synthesis and cytotoxicity of novel thioxo-quinazolino[3,4-b]quinazolinones. <i>Turkish Journal of Chemistry</i> , 2017, 41, 125-134.	0.5	3
300	Design, Synthesis, and Biological Evaluation of New Indole-Acrylamide-1,2,3-Triazole Derivatives as Potential α -Glucosidase Inhibitors. <i>Polycyclic Aromatic Compounds</i> , 2022, 42, 3157-3165.	1.4	3
301	Design, synthesis and antibacterial activity evaluation of novel 2-((1-aryl-1H)-1H-tetrazol-5-yl)ethanol derivatives. <i>Chemistry</i> , 2020, 57, 4254-4261.	1.4	3
302	β -Fe ₂ O ₃ @SiO ₂ (CH ₂) ₃ -HPBM-Pd as a versatile boosted nanocatalyst for carbon-carbon bond formation. <i>Materials Today Communications</i> , 2021, 26, 101913.	0.9	3
303	Design and Synthesis of Novel 5-Arylisoxazole-1,3,4-thiadiazole Hybrids as α -Glucosidase Inhibitors. <i>Letters in Drug Design and Discovery</i> , 2021, 18, 436-444.	0.4	3
304	Copper-Mediated Direct Cyanation of Benzamides: A New Approach to the Synthesis of Quinazolinones. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 708-713.	1.2	3
305	Design, Synthesis and In vitro Cytotoxicity of New 1,2,3-triazol- and Nitrostyrene Hybrids as Potent Anticancer Agents. <i>Letters in Drug Design and Discovery</i> , 2018, 16, 213-219.	0.4	3
306	Synthesis of novel 1,2,3-triazole derivatives of 2,3-dihydroquinazolin-4(1H)-one. <i>Monatshefte Für Chemie</i> , 2016, 147, 2151-2156.	0.9	2

#	ARTICLE	IF	CITATIONS
307	A simple one-pot synthesis of 2,4-diaryl-9H-pyrido[2,3-b]indoles under solvent-free conditions. <i>Heterocyclic Communications</i> , 2017, 23, 293-296.	0.6	2
308	The synthesis of 2,3-dihydroquinazoline-4(1H)-one and dihydroisoindolo[2,1-a]quinazoline-5,11-dione derivatives in the presence of imidazolium ionic liquid sulfonic acid functionalized SBA-15: a novel feature of SBA-15. <i>Arkivoc</i> , 2019, 2018, 302-314.	0.3	2
309	Synthesis and Anticancer Activity of N-(di/trimethoxyaryl)-5-arylisoxazole-3-carboxamide. <i>Polycyclic Aromatic Compounds</i> , 2020, 40, 1568-1580.	1.4	2
310	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.	1.4	2
311	Vinylazides: versatile synthons and magical precursors for the construction of N-heterocycles. <i>Molecular Diversity</i> , 2021, 25, 2533-2570.	2.1	2
312	An Efficient and Convenient Approach for Synthesizing Iodohydrin and Iodoether from Aromatic Alkenes Using Hg(BF ₄) ₂ ·SiO ₂ and I ₂ . <i>Polycyclic Aromatic Compounds</i> , 2022, 42, 3975-3983.	1.4	2
313	A Convenient Method for the Synthesis of Chromeno[4,3-b]pyridines Via Three-component Reaction. <i>Combinatorial Chemistry and High Throughput Screening</i> , 2018, 21, 344-348.	0.6	2
314	Four-Component Heterocyclization Reaction for the One-Pot Synthesis of 2,4-Dichloro-Substituted Pyrano/Furo[2,3-d]pyrimidines in an Environmentally Benign Procedure Mediated by Ceric Ammonium Nitrate in Phosphorus Ionic Liquid. <i>Polycyclic Aromatic Compounds</i> , 0, , 1-9.	1.4	2
315	Synthesis of novel derivatives of chromenone bearing an N-carbamothioyl moiety as soybean 15-LOX inhibitors. <i>Turkish Journal of Chemistry</i> , 2017, 41, 335-344.	0.5	1
316	Efficient three-component synthesis of N-alkyl-3,6-diaryl-[1,2,4]triazolo[4,3-b][1,2,4]triazin-7-amines under solvent-free condition. <i>Arkivoc</i> , 2017, 2017, 293-300.	0.3	1
317	Tandem synthesis of benzo[d]naphtho[2,3-g][1,3]oxazocine-8,13(6H,14H)-dione derivatives. <i>Monatshefte für Chemie</i> , 2019, 150, 347-352.	0.9	1
318	Efficient synthesis of novel 2-(2-chloroquinolin-3-yl)imidazo[1,2-a]pyridin-3-amine derivatives. <i>Journal of the Chinese Chemical Society</i> , 2021, 68, 1328-1333.	0.8	1
319	Stage-Specific Oligonucleotide Primers for the Diagnosis of Toxoplasmosis Among Iranian Pediatric Heart Transplant Recipients; Evaluation of Cotrimoxazole as a Preventive Therapy. <i>Archives of Pediatric Infectious Diseases</i> , 2021, 9, .	0.1	1
320	Nickel Supported MCM-Functionalized 1,2,3-Triazol-4-ylmethanamine: An Efficient Nano-particle-Heterogeneous Catalyst Activate for Suzuki Reaction. <i>Catalysis Letters</i> , 2022, 152, 2186-2199.	1.4	1
321	2,4-Dioxochroman Moiety Linked to 1,2,3-triazole Derivatives as Novel Î±-glucosidase Inhibitors: Synthesis, In vitro Biological Evaluation, and Docking Study. <i>Current Organic Chemistry</i> , 2020, 24, 2019-2027.	0.9	1
322	Anticholinesterase Activity of Cinnamic Acids Derivatives: In Vitro, In Vivo Biological Evaluation, and Docking Study. <i>Letters in Drug Design and Discovery</i> , 2020, 17, 965-982.	0.4	1
323	Design, Synthesis, <i>in Vitro</i> , and <i>in Silico</i> Evaluation of <i>N</i> -[Phenylacetamide- <i>O</i> -indole- <i>Î</i> -thiosemicarbazide Hybrids as New Potential Tyrosinase Inhibitors. <i>Chemistry and Biodiversity</i> , 2022, , .	1.0	1
324	Synthesis, molecular docking, and cytotoxicity of quinazolinone and dihydroquinazolinone derivatives as cytotoxic agents. <i>BMC Chemistry</i> , 2022, 16, 35.	1.6	1

#	ARTICLE	IF	CITATIONS
325	Efficient one-pot synthesis of novel 6,9-dihydro-2H,7H-spiro[pyrimidine-5,8-[1,3]dioxolo[4,5-f]quinoline]-2,4,6(1H,3H)-trione derivatives under mild and "green" reaction conditions. <i>Journal of Heterocyclic Chemistry</i> , 2020, 57, 3161-3166.	1.4	0
326	Design and synthesis of 2,4-dioxochromanopyridiniumphenylacetamide derivatives as new anti-Alzheimer agents: in vitro and in silico studies. <i>Journal of the Chinese Chemical Society</i> , 2020, 67, 1910-1928.	0.8	0
327	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.	0.7	0
328	Aminoimidazo[1,2-a]pyridine Bearing Different Pyrazole Moieties as the Structural Scaffold for the Development of BACE1 Inhibitor; Synthesis, Structural Characterization, In vitro and In silico Studies. <i>Current Organic Synthesis</i> , 2022, 19, .	0.7	0
329	Synthesis of Novel Pyrazino[2,1-a]isoindoliones via Intramolecular Hydroamination of 2,3-Dihydro-3-oxo-2-(prop-2-yn-1-yl)-1H-isoindole-1-carboxamides. <i>Helvetica Chimica Acta</i> , 2016, , n/a-n/a.	1.0	0
330	Copper Supported Imidazolylpyridine Modified SPION as an Efficient Catalyst for Eco-friendly, One-Pot and Green Synthesis of Novel (3-Cyanothiophen-2-yl)-N-(arylsulfonyl)acetimidamide Derivatives. <i>Current Organic Synthesis</i> , 2022, 19, .	0.7	0
331	New imidazo[1,2-a]pyridin-2-yl derivatives as AChE, BChE and 15-LOX inhibitors; design, synthesis, and biological evaluation. <i>Letters in Drug Design and Discovery</i> , 2022, 19, .	0.4	0
332	Synthesis and Evaluation of 6-Ethoxy-2-mercaptobenzothiazole Scaffolds as Potential α -Glucosidase Inhibitors. <i>ChemistrySelect</i> , 2022, 7, .	0.7	0