

List of Publications by Year in descending order

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319
papers

6,696
citations

93792

39
h-index

169272

56
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368
all docs

368
docs citations

368
times ranked

5592
citing authors

#	ARTICLE	IF	CITATIONS
1	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
2	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
3	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
4	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
5	One-pot multi-component synthesis of novel chromeno[4,3-b]pyrrol-3-yl derivatives as alpha-glucosidase inhibitors. <i>Molecular Diversity</i> , 2022, 26, 2393-2405.	2.1	17
6	Synthesis and characterization of 1-amidino-O-alkylureas metal complexes as α -glucosidase Inhibitors: Structure-activity relationship, molecular docking, and kinetic studies. <i>Journal of Molecular Structure</i> , 2022, 1250, 131726.	1.8	17
7	Design, synthesis, biological evaluation, and molecular docking study of thioxo-2,3-dihydroquinazolinone derivative as tyrosinase inhibitors. <i>Journal of Molecular Structure</i> , 2022, 1253, 132283.	1.8	11
8	Green synthesized silver nanoparticles obtained from <i>Stachys schtschegleevii</i> extract: ct-DNA interaction and in silico and in vitro investigation of antimicrobial activity. <i>Journal of Biomolecular Structure and Dynamics</i> , 2022, , 1-14.	2.0	0
9	Magnetic graphene oxide-lignin nanobiocomposite: a novel, eco-friendly and stable nanostructure suitable for hyperthermia in cancer therapy. <i>RSC Advances</i> , 2022, 12, 3593-3601.	1.7	21
10	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
11	Synthesis, molecular dynamic, and in silico study of new ethyl 4-arylpyrimido[1,2-b]indazole-2-carboxylate: Potential inhibitors of α -glucosidase. <i>Journal of Molecular Structure</i> , 2022, 1257, 132507.	1.8	4
12	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
13	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
14	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
15	New 4-phenylpiperazine-carbodithioate-N-phenylacetamide hybrids: Synthesis, in vitro and in silico evaluations against cholinesterase and α -glucosidase enzymes. <i>Archiv Der Pharmazie</i> , 2022, 355, e2100313.	2.1	11
16	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
17	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
18	Can polyoxometalates (POMs) prevent of coronavirus 2019-nCoV cell entry? Interaction of POMs with TMPRSS2 and spike receptor domain complexed with ACE2 (ACE2-RBD): Virtual screening approaches. <i>Informatics in Medicine Unlocked</i> , 2022, 29, 100902.	1.9	8

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19	Design, Synthesis, <i>in Vitro</i> , and <i>in Silico</i> Evaluation of <i>N</i> -Phenylacetamide- <i>O</i> -indole- <i>T</i> -thiosemicarbazide Hybrids as New Potential Tyrosinase Inhibitors. <i>Chemistry and Biodiversity</i> , 2022, , .	1.0	1
20	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
21	Novel aryl(4-phenylpiperazin-1-yl)methanethione derivatives as new anti-Alzheimer agents: Design, synthesis, <i>in vitro</i> and <i>in silico</i> assays. <i>Journal of Molecular Structure</i> , 2022, 1262, 132945.	1.8	4
22	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
23	Novel phenylurea-pyridinium derivatives as potent urease inhibitors: Synthesis, <i>in vitro</i> , and <i>in silico</i> studies. <i>Journal of Molecular Structure</i> , 2022, 1263, 133078.	1.8	11
24	<i>In vitro</i> cell-based models of drug-induced hepatotoxicity screening: progress and limitation. <i>Drug Metabolism Reviews</i> , 2022, 54, 161-193.	1.5	5
25	Modern metal-catalyzed and organocatalytic methods for synthesis of coumarin derivatives: a review. <i>Organic and Biomolecular Chemistry</i> , 2022, 20, 4846-4883.	1.5	6
26	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
27	Evaluating the effects of disubstituted 3-hydroxy-1H-pyrrol-2(5H)-one analog as novel tyrosinase inhibitors. <i>Bioorganic Chemistry</i> , 2022, 126, 105876.	2.0	13
28	Synthesis, molecular docking, and cytotoxicity of quinazolinone and dihydroquinazolinone derivatives as cytotoxic agents. <i>BMC Chemistry</i> , 2022, 16, 35.	1.6	1
29	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
30	Magnetic carboxymethyl cellulose/silk fibroin hydrogel embedded with halloysite nanotubes as a biocompatible nanobiocomposite with hyperthermia application. <i>Materials Chemistry and Physics</i> , 2022, 287, 126347.	2.0	19
31	Design, synthesis, <i>in vitro</i> 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
32	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
33	Biocompatibility and Antimicrobial Investigation of Agar-Tannic Acid Hydrogel Reinforced with Silk Fibroin and Zinc Manganese Oxide Magnetic Microparticles. <i>Journal of Inorganic and Organometallic Polymers and Materials</i> , 2022, 32, 4057-4069.	1.9	9
34	Design, synthesis, and <i>in silico</i> 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
35	6-Methoxy- α -tetralone Derivatives Bearing an <i>N</i> -Arylpyridinium Moiety as Cholinesterase Inhibitors: Design, Synthesis, Biological Evaluation, and Molecular Docking Study. <i>ChemistrySelect</i> , 2022, 7, .	0.7	4
36	Synthesis and Evaluation of 6-Ethoxy-2-mercaptobenzothiazole Scaffolds as Potential α -glucosidase Inhibitors. <i>ChemistrySelect</i> , 2022, 7, .	0.7	0

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37	Recent advances on biomedical applications of pectin-containing biomaterials. <i>International Journal of Biological Macromolecules</i> , 2022, 217, 1-18.	3.6	28
38	Vinylazides: versatile synthons and magical precursors for the construction of N-heterocycles. <i>Molecular Diversity</i> , 2021, 25, 2533-2570.	2.1	2
39	Multispectroscopic analysis, atomic force microscopy, molecular docking and molecular dynamic simulation studies of the interaction between [SnMe ₂ Cl ₂ (Me ₂ phen)] complex and ct-DNA in the presence of glucose. <i>Journal of Biomolecular Structure and Dynamics</i> , 2021, 39, 5068-5082.	2.0	4
40	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
41	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
42	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
43	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
44	Novel N-benzylpiperidine derivatives of 5-arylisoxazole-3-carboxamides as anti-Alzheimer's agents. <i>Archiv Der Pharmazie</i> , 2021, 354, e2000258.	2.1	12
45	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
46	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
47	Synthesis of the new tri-amide derivatives as novel α -glucosidase inhibitors by Ugi four-component reaction. <i>Journal of Molecular Structure</i> , 2021, 1227, 129531.	1.8	5
48	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
49	Synthesis, in vitro, and in silico studies of newly functionalized quinazolinone analogs for the identification of potent α -glucosidase inhibitors. <i>Journal of the Iranian Chemical Society</i> , 2021, 18, 2017-2034.	1.2	5
50	Investigation of the biological activity, mechanical properties and wound healing application of a novel scaffold based on lignin-agarose hydrogel and silk fibroin embedded zinc chromite nanoparticles. <i>RSC Advances</i> , 2021, 11, 17914-17923.	1.7	68
51	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
52	Inhibitory activity of FDA-approved drugs cetilistat, abiraterone, diiodohydroxyquinoline, bexarotene, remdesivir, and hydroxychloroquine on COVID-19 main protease and human ACE2 receptor: A comparative in silico approach. <i>Informatics in Medicine Unlocked</i> , 2021, 26, 100745.	1.9	11
53	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
54	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

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55	Copper Supported onto Magnetic Nanoparticles as an Efficient Catalyst for the Synthesis of Triazolobenzodiazepino[7,1-b]quinazolin-1(9H)-ones. <i>Click N-Arylation Reactions. ChemistrySelect</i> , 2021, 6, 1385-1392.	0.7	9
56	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
57	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
58	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
59	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
60	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
61	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
62	Quinazolinone-dihydropyrano[3,2-b]pyran hybrids as new α -glucosidase inhibitors: Design, synthesis, enzymatic inhibition, docking study and prediction of pharmacokinetic. <i>Bioorganic Chemistry</i> , 2021, 109, 104703.	2.0	12
63	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
64	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
65	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
66	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
67	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
68	Arylmethylene hydrazine derivatives containing 1,3-dimethylbarbituric moiety as novel urease inhibitors. <i>Scientific Reports</i> , 2021, 11, 10607.	1.6	19
69	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
70	New 4,5-diphenylimidazole-acetamide-1,2,3-triazole hybrids as potent α -glucosidase inhibitors: synthesis, in vitro and in silico enzymatic and toxicity evaluations. <i>Monatshefte Für Chemie</i> , 2021, 152, 679-693.	0.9	8
71	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
72	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

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73	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
74	Magnetic Copper Ferrite Nanoparticles Functionalized by Aromatic Polyamide Chains for Hyperthermia Applications. <i>Langmuir</i> , 2021, 37, 8847-8854.	1.6	38
75	Design, synthesis, and α -glucosidase inhibitory activity of phenoxy-bis coumarin-phenylacetamide hybrids. <i>Archiv Der Pharmazie</i> , 2021, 354, e2100179.	2.1	10
76	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
77	Functionalized magnetic nanoparticles for the separation and purification of proteins and peptides. <i>TrAC - Trends in Analytical Chemistry</i> , 2021, 141, 116291.	5.8	70
78	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
79	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
80	Design and synthesis of phenoxy-methylbenzimidazole incorporating different aryl thiazole-triazole acetamide derivatives as α -glucosidase inhibitors. <i>Molecular Diversity</i> , 2021, , 1.	2.1	12
81	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
82	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
83	Design, synthesis, in vitro and in silico biological assays of new quinazolinone-2-thio-metronidazole derivatives. <i>Journal of Molecular Structure</i> , 2021, 1244, 130889.	1.8	9
84	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
85	Synthesis of Chromene-Fused Heterocycles by the Intramolecular Diels-Alder Reaction: An Overview. <i>Tetrahedron</i> , 2021, 102, 132524.	1.0	12
86	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
87	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
88	Sodium Azide: An Inorganic Nitrogen Source for the Synthesis of Organic N-Compounds. <i>ChemistrySelect</i> , 2021, 6, 13419-13433.	0.7	6
89	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
90	Multicomponent reaction of amine, carbon disulfide, and fluoronitrobenzene via nucleophilic attack on the fluorinated carbon for the synthesis of nitrophenyl methylcarbamodithioates. <i>Journal of the Chinese Chemical Society</i> , 2020, 67, 160-164.	0.8	8

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91	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
92	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
93	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
94	Synthesis and Anticancer Activity of N-(di/trimethoxyaryl)-5-arylisoxazole-3-carboxamide. <i>Polycyclic Aromatic Compounds</i> , 2020, 40, 1568-1580.	1.4	2
95	Synthesis and pharmacological properties of polysubstituted 2-amino-4H-pyran-3-carbonitrile derivatives. <i>Molecular Diversity</i> , 2020, 24, 1385-1431.	2.1	34
96	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
97	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
98	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
99	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
100	Synthesis of Arylidene α -Isoquinolinones bearing Combretastatin Skeleton by Cyclocarbopalladation/cross coupling Tandem Heck-Suzuki-Miyaura Reactions using nano catalyst Pd@Py@SiO ₂ . <i>Applied Organometallic Chemistry</i> , 2020, 34, e5279.	1.7	5
101	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
102	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
103	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
104	Novel quinazolin-2-sulfonamide derivatives: synthesis, characterization, biological evaluation, and molecular docking studies. <i>Journal of Biomolecular Structure and Dynamics</i> , 2020, , 1-12.	2.0	9
105	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
106	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
107	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
108	Design, synthesis and antibacterial activity evaluation of novel 2-arylamino-1H-benzotriazole derivatives. <i>Journal of Heterocyclic Chemistry</i> , 2020, 57, 4254-4261.	1.4	3

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109	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
110	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
111	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
112	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
113	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
114	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
115	New phthalimide-benzamide-1,2,3-triazole hybrids; design, synthesis, α -glucosidase inhibition assay, and docking study. <i>Medicinal Chemistry Research</i> , 2020, 29, 868-876.	1.1	12
116	Design and Synthesis of Novel Arylisoazolechromenone Carboxamides: Investigation of Biological Activities Associated with Alzheimer's Disease. <i>Chemistry and Biodiversity</i> , 2020, 17, e1900746.	1.0	26
117	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
118	Synthesis, characterization, molecular docking, and biological activities of coumarin-1,2,3-triazoleacetamide hybrid derivatives. <i>Archiv Der Pharmazie</i> , 2020, 353, e2000109.	2.1	50
119	Thieno[2,3-b]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
120	Efficient One Pot Synthesis of Phenylimidazo[1,2-a]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
121	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
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128	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
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158	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
159	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
160	Facile access to new pyrido[2,3-d]pyrimidine derivatives. <i>Molecular Diversity</i> , 2019, 23, 333-340.	2.1	5
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174	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
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180	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

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182	Copper-catalyzed intramolecular domino synthesis of 6H-chromeno[4,3-b]quinolines in green condition. <i>Arkivoc</i> , 2018, 2018, 20-28.	0.3	9
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185	Synthesis of New Benzimidazole-1,2,3-Triazole Hybrids as Tyrosinase Inhibitors. <i>Chemistry and Biodiversity</i> , 2018, 15, e1800120.	1.0	50
186	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
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188	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
189	Synthesis and Characterization of Novel Phthalimide–pyrano[3,2 <i>c</i>]chromene and Phthalimide–pyrano–one Hybrids. <i>Journal of Heterocyclic Chemistry</i> , 2018, 55, 1678-1684.	1.4	5
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196	Synthesis of Novel Tacrine Analogs as Acetylcholinesterase Inhibitors. <i>Journal of Heterocyclic Chemistry</i> , 2017, 54, 384-390.	1.4	19
197	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
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200	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
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211	Synthesis and biological evaluation of novel imidazopyrimidinamines as anticancer agents. <i>Chemical Biology and Drug Design</i> , 2017, 89, 797-805.	1.5	11
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218	Novel Indole-Isoxazole Hybrids: Synthesis and In Vitro Anti-Cholinesterase Activity. <i>Letters in Drug Design and Discovery</i> , 2017, 14, .	0.4	8
219	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, .	0.4	15
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236	Synthesis of Novel Pyrazino[2,1-a]isoindolones 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
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256	Synthesis and cytotoxic activity of novel poly-substituted imidazo[2,1- <i>c</i>][1,2,4]triazin-6-amines. <i>Molecular Diversity</i> , 2015, 19, 273-281.	2.1	20
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258	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
259	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
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267	Synthesis of Novel 1,2,3-Triazole-dihydro[3,2- <i>c</i>]chromenones as Acetylcholinesterase Inhibitors. <i>Synthetic Communications</i> , 2015, 45, 2311-2318.	1.1	29
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