Mohammad Mahdavi

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

316 papers

4,248 citations

32 h-index

47 g-index

381 ext. papers

5,509 ext. citations

2.9 avg, IF

5.92 L-index

| # | Paper | IF | Citations |
|-----|--|-----|-----------|
| 316 | 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 | 4.3 | 4 |
| 315 | Synthesis and in vitro urease inhibitory activity of 5-nitrofuran-2-yl-thiadiazole linked to different cyclohexyl-2-(phenylamino)acetamides, in silico and kinetic studies <i>Bioorganic Chemistry</i> , 2022 , 120, 105592 | 5.1 | 1 |
| 314 | A review on synthesis, mechanism of action, and structure-activity relationships of 1,2,3-triazole-based Eglucosidase inhibitors as promising anti-diabetic agents. <i>Journal of Molecular Structure</i> , 2022 , 1255, 132469 | 3.4 | 5 |
| 313 | 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 | 3.4 | 3 |
| 312 | New 4-phenylpiperazine-carbodithioate-N-phenylacetamide hybrids: Synthesis, in vitro and in silico evaluations against cholinesterase and Eglucosidase enzymes <i>Archiv Der Pharmazie</i> , 2022 , e2100313 | 4.3 | O |
| 311 | Design and synthesis of novel nitrothiazolacetamide conjugated to different thioquinazolinone derivatives as anti-urease agents <i>Scientific Reports</i> , 2022 , 12, 2003 | 4.9 | 4 |
| 310 | In silico and in vitro studies of thiosemicarbazone-indole hybrid compounds as potent Eglycosidase inhibitors <i>Computational Biology and Chemistry</i> , 2022 , 97, 107642 | 3.6 | O |
| 309 | Pd@PyPZ@MSN as a Novel and Efficient Catalyst for C-C Bond Formation Reactions <i>Frontiers in Chemistry</i> , 2022 , 10, 838294 | 5 | 0 |
| 308 | 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 | 3.4 | 1 |
| 307 | Functionalized graphene oxide nanosheets with folic acid and silk fibroin as a novel nanobiocomposite for biomedical applications <i>Scientific Reports</i> , 2022 , 12, 6205 | 4.9 | O |
| 306 | Novel phenylurea-pyridinium derivatives as potent urease inhibitors: Synthesis, in vitro, and in silico studies. <i>Journal of Molecular Structure</i> , 2022 , 133078 | 3.4 | 1 |
| 305 | In vitro cell-based models of drug-induced hepatotoxicity screening: progress and limitation <i>Drug Metabolism Reviews</i> , 2022 , 1-76 | 7 | O |
| 304 | A review on Eglucosidase 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 | 3.7 | 2 |
| 303 | Synthesis, molecular docking, and cytotoxicity of quinazolinone and dihydroquinazolinone derivatives as cytotoxic agents <i>BMC Chemistry</i> , 2022 , 16, 35 | 3.7 | |
| 302 | Vinylazides: versatile synthons and magical precursors for the construction of N-heterocycles. <i>Molecular Diversity</i> , 2021 , 25, 2533-2570 | 3.1 | 1 |
| 301 | Characteristics of published/registered clinical trials on COVID-19 treatment: A systematic review. <i>DARU, Journal of Pharmaceutical Sciences</i> , 2021 , 29, 449-467 | 3.9 | 2 |
| 300 | One-pot multi-component synthesis of novel chromeno[4,3-b]pyrrol-3-yl derivatives as alpha-glucosidase inhibitors. <i>Molecular Diversity</i> , 2021 , 1 | 3.1 | 3 |

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| 299 | Novel magnetic organic-inorganic hybrids based on aromatic polyamides and ZnFeO nanoparticles with biological activity. <i>Scientific Reports</i> , 2021 , 11, 20310 | 4.9 | 1 |
|-----|---|-----|----|
| 298 | Design, synthesis, biological evaluation, and molecular modeling studies of pyrazole-benzofuran hybrids as new Eglucosidase inhibitor. <i>Scientific Reports</i> , 2021 , 11, 20776 | 4.9 | 2 |
| 297 | Synthesis and biological evaluation of a new series of benzofuran-1,3,4-oxadiazole containing 1,2,3-triazole-acetamides as potential lglucosidase inhibitors. <i>Journal of Biochemical and Molecular Toxicology</i> , 2021 , 35, e22688 | 3.4 | 2 |
| 296 | Recent strategies in the synthesis of thiophene derivatives: highlights from the 2012-2020 literature. <i>Molecular Diversity</i> , 2021 , 25, 2571-2604 | 3.1 | 8 |
| 295 | Recent advances in biological activities of rhodium complexes: Their applications in drug discovery research. <i>European Journal of Medicinal Chemistry</i> , 2021 , 216, 113308 | 6.8 | 13 |
| 294 | Design, synthesis, and evaluation of metronidazole-1,2,3-triazole derivatives as potent urease inhibitors. <i>Chemical Papers</i> , 2021 , 75, 4217-4226 | 1.9 | 2 |
| 293 | Quinazolinone-dihydropyrano[3,2-b]pyran hybrids as new lglucosidase inhibitors: Design, synthesis, enzymatic inhibition, docking study and prediction of pharmacokinetic. <i>Bioorganic Chemistry</i> , 2021 , 109, 104703 | 5.1 | 7 |
| 292 | Ullmann-Goldberg and Buchwald-Hartwig CN Cross Couplings: Synthetic Methods to Pharmaceutically Potential N-Heterocycles. <i>Asian Journal of Organic Chemistry</i> , 2021 , 10, 1319-1344 | 3 | 17 |
| 291 | 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 | 3.4 | 13 |
| 290 | Synthesis, in vitro and in silico enzymatic inhibition assays, and toxicity evaluations of new 4,5-diphenylimidazole-N-phenylacetamide derivatives as potent Eglucosidase inhibitors. <i>Medicinal Chemistry Research</i> , 2021 , 30, 1273-1283 | 2.2 | 1 |
| 289 | 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 | 2.5 | 2 |
| 288 | 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 | 4.3 | 3 |
| 287 | N-sulfonyl ketenimine as a versatile intermediate for the synthesis of heteroatom containing compounds. <i>Journal of Organometallic Chemistry</i> , 2021 , 939, 121773 | 2.3 | 6 |
| 286 | Arylmethylene hydrazine derivatives containing 1,3-dimethylbarbituric moiety as novel urease inhibitors. <i>Scientific Reports</i> , 2021 , 11, 10607 | 4.9 | 6 |
| 285 | Sulfonic Acid Functionalized Magnetic Starch as an Efficient Catalyst for the Synthesis of Chromeno[4,3-b]quinoline-6,8(9H)-dione Derivatives. <i>Starch/Staerke</i> , 2021 , 73, 2000257 | 2.3 | 2 |
| 284 | New 4,5-diphenylimidazole-acetamide-1,2,3-triazole hybrids as potent ⊞lucosidase inhibitors: synthesis, in vitro and in silico enzymatic and toxicity evaluations. <i>Monatshefte F□ Chemie</i> , 2021 , 152, 679 | 1.4 | O |
| 283 | Catalytic and non-catalytic amidation of carboxylic acid substrates. <i>Molecular Diversity</i> , 2021 , 1 | 3.1 | 2 |
| 282 | Design and synthesis of a novel nanocomposite based on magnetic dopamine nanoparticles for purification of the mylase from the bovine milk. <i>Scientific Reports</i> , 2021 , 11, 13428 | 4.9 | 1 |

| 281 | Triflic Anhydride (Tf2O): An Efficient Catalyst for Electrophilic Activation of Amides. <i>ChemistrySelect</i> , 2021 , 6, 5320-5328 | 1.8 | 2 |
|-------------|--|-----|----|
| 2 80 | Recent Opportunities and Challenges in Selective C-H Functionalization of Methyl Azaarenes: A Highlight from 2010 to 2020 Studies. <i>Current Organic Synthesis</i> , 2021 , 18, 761-789 | 1.9 | |
| 279 | Design and Synthesis of Novel 5-Arylisoxazole-1,3,4-thiadiazole Hybrids as EGlucosidase Inhibitors. Letters in Drug Design and Discovery, 2021 , 18, 436-444 | 0.8 | 0 |
| 278 | 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 | 2.8 | 1 |
| 277 | Design and synthesis of 4,5-diphenyl-imidazol-1,2,3-triazole hybrids as new anti-diabetic agents: in vitro Eglucosidase inhibition, kinetic and docking studies. <i>Molecular Diversity</i> , 2021 , 25, 877-888 | 3.1 | 7 |
| 276 | Design and synthesis of novel pyrazole-phenyl semicarbazone derivatives as potential ⊞glucosidase inhibitor: Kinetics and molecular dynamics simulation study. <i>International Journal of Biological Macromolecules</i> , 2021 , 166, 1082-1095 | 7.9 | 13 |
| 275 | Synthesis and biological evaluation of new dihydroindolizino[8,7-b]indole derivatives as novel Eglucosidase inhibitors. <i>Journal of Molecular Structure</i> , 2021 , 1224, 129290 | 3.4 | 3 |
| 274 | Electrochemical synthesis of three-dimensional flower-like Ni/CoBTC bimetallic organic framework as heterogeneous catalyst for solvent-free and green synthesis of substituted chromeno[4,3B]quinolones. <i>Journal of the Chinese Chemical Society</i> , 2021 , 68, 620-629 | 1.5 | 2 |
| 273 | Novel N-benzylpiperidine derivatives of 5-arylisoxazole-3-carboxamides as anti-Alzheimer's agents. <i>Archiv Der Pharmazie</i> , 2021 , 354, e2000258 | 4.3 | 3 |
| 272 | Fe2O3@SiO2(CH2)3-HPBM-Pd as a versatile boosted nanocatalyst for carboncarbon bond f ormation. <i>Materials Today Communications</i> , 2021 , 26, 101913 | 2.5 | 2 |
| 271 | Design, synthesis, characterization, enzymatic inhibition evaluations, and docking study of novel quinazolinone derivatives. <i>International Journal of Biological Macromolecules</i> , 2021 , 170, 1-12 | 7.9 | 20 |
| 270 | 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 | 1.8 | 7 |
| 269 | Synthesis, in vitro, and in silico studies of newly functionalized quinazolinone analogs for the identification of potent glucosidase inhibitors. <i>Journal of the Iranian Chemical Society</i> , 2021 , 18, 2017 | 2 | 0 |
| 268 | EGlucosidase and Emylase inhibition, molecular modeling and pharmacokinetic studies of new quinazolinone-1,2,3-triazole-acetamide derivatives. <i>Medicinal Chemistry Research</i> , 2021 , 30, 702-711 | 2.2 | 3 |
| 267 | 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> , 2021 , 1-13 | 3.6 | 1 |
| 266 | Novel Coumarin Containing Dithiocarbamate Derivatives as Potent Educosidase Inhibitors for Management of Type 2 Diabetes. <i>Medicinal Chemistry</i> , 2021 , 17, 264-272 | 1.8 | 3 |
| 265 | Copper-catalyzed one-pot synthesis of amide linked 1,2,3-triazoles bearing aryloxy skeletons. <i>Tetrahedron Letters</i> , 2021 , 65, 152765 | 2 | 2 |
| 264 | Copper Supported onto Magnetic Nanoparticles as an Efficient Catalyst for the Synthesis of Triazolobenzodiazepino[7,1-b]quinazolin-11(9H)-ones via Click N-Arylation Reactions. <i>ChemistrySelect</i> , 2021 , 6, 1385-1392 | 1.8 | 2 |

| 263 | Efficient synthesis of novel 2-(2-chloroquinolin-3-yl)imidazo[1,2-a]pyridin-3-amine derivatives. Journal of the Chinese Chemical Society, 2021 , 68, 1328-1333 | 1.5 | О |
|-----|---|---------------------|----|
| 262 | 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 | 1.7 | 4 |
| 261 | 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 | 2.3 | 4 |
| 260 | Synthesis of novel tetracyclic coumarin-fused furo-pyridone scaffolds via sequential N-arylation and intramolecular amidation reactions. <i>Tetrahedron Letters</i> , 2021 , 68, 152904 | 2 | 3 |
| 259 | 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 & Dressing Applications</i> 13, 33840-33849 | 9.5 | 18 |
| 258 | 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> , 2021 , 1 | 3.1 | 2 |
| 257 | Magnetic Copper Ferrite Nanoparticles Functionalized by Aromatic Polyamide Chains for Hyperthermia Applications. <i>Langmuir</i> , 2021 , 37, 8847-8854 | 4 | 14 |
| 256 | Design, synthesis, and Eglucosidase-inhibitory activity of phenoxy-biscoumarin-N-phenylacetamide hybrids. <i>Archiv Der Pharmazie</i> , 2021 , 354, e2100179 | 4.3 | 1 |
| 255 | 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 | 1.9 | 1 |
| 254 | Design and synthesis of novel quinazolinone-pyrazole derivatives as potential ঘ lucosidase inhibitors: Structure-activity relationship, molecular modeling and kinetic study. <i>Bioorganic Chemistry</i> , 2021 , 114, 105127 | 5.1 | 9 |
| 253 | 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 | 5.1 | 6 |
| 252 | Design and synthesis of phenoxymethybenzoimidazole incorporating different aryl thiazole-triazole acetamide derivatives as Eglycosidase inhibitors. <i>Molecular Diversity</i> , 2021 , 1 | 3.1 | Ο |
| 251 | 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 | 7.9 | 7 |
| 250 | Synthesis, in vitro, and in silico evaluation of Indazole Schiff bases as potential ⊞glucosidase inhibitors. <i>Journal of Molecular Structure</i> , 2021 , 1242, 130826 | 3.4 | 3 |
| 249 | Sodium Azide: An Inorganic Nitrogen Source for the Synthesis of Organic N -Compounds. <i>ChemistrySelect</i> , 2021 , 6, 13419-13433 | 1.8 | О |
| 248 | Synthesis, in vitro and in silico screening of 2-amino-4-aryl-6-(phenylthio) pyridine-3,5-dicarbonitriles as novel Eglucosidase inhibitors. <i>Bioorganic Chemistry</i> , 2020 , 100, 103879 | 5.1 | 11 |
| 247 | 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 | 1.4 | 6 |
| 246 | 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, 46 | 53 ³ 497 | 5 |

| 245 | Efficient one-pot synthesis of novel 6?,9?-dihydro-2H,7?H-spiro[pyrimidine-5,8?-[1,3]dioxolo[4,5-f]quinoline]-2,4,6(1H,3H)-trione derivatives under mild and greenDeaction conditions. <i>Journal of Heterocyclic Chemistry</i> , 2020 , 57, 3161-3 | 1.9 3166 | |
|-----|--|------------------|----|
| 244 | Design, synthesis and biological evaluation of novel phthalimide-Schiff base-coumarin hybrids as potent 🗄 glucosidase inhibitors. <i>Chemical Papers</i> , 2020 , 74, 4379-4388 | 1.9 | 4 |
| 243 | Design and synthesis of 2,4-dioxochroman-pyridinium-phenylacetamide derivatives as new anti-Alzheimer agents: in vitro and in silico studies. <i>Journal of the Chinese Chemical Society</i> , 2020 , 67, 1910-1928 | 1.5 | |
| 242 | New phthalimide-benzamide-1,2,3-triazole hybrids; design, synthesis, Eglucosidase inhibition assay, and docking study. <i>Medicinal Chemistry Research</i> , 2020 , 29, 868-876 | 2.2 | 4 |
| 241 | 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 | 6 ^{2.5} | 13 |
| 240 | Regio- and Diastereoselective KMnO4/RCO2H Mediated Acyloxyarylation of Chalcones An Indirect Arylation of Chalcones. <i>European Journal of Organic Chemistry</i> , 2020 , 2020, 2045-2051 | 3.2 | 2 |
| 239 | New 1,2,3-triazole-(thio)barbituric acid hybrids as urease inhibitors: Design, synthesis, in vitro urease inhibition, docking study, and molecular dynamic simulation. <i>Archiv Der Pharmazie</i> , 2020 , 353, e2000023 | 4.3 | 11 |
| 238 | Synthesis, characterization, molecular docking, and biological activities of coumarin-1,2,3-triazole-acetamide hybrid derivatives. <i>Archiv Der Pharmazie</i> , 2020 , 353, e2000109 | 4.3 | 27 |
| 237 | 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, e2000101 | 4.3 | 9 |
| 236 | Efficient One Pot Synthesis of Phenylimidazo[1,2-a]pyridine Derivatives using Multifunctional Copper Catalyst Supported on Ecyclodextrin Functionalized Magnetic Graphene oxide. <i>Applied Organometallic Chemistry</i> , 2020 , 34, e5913 | 3.1 | 7 |
| 235 | An efficient and targeted synthetic approach towards new highly substituted 6-amino-pyrazolo[1,5-a]pyrimidines with lglucosidase inhibitory activity. <i>Scientific Reports</i> , 2020 , 10, 2595 | 4.9 | 8 |
| 234 | C3-Functionalization of Imidazo[1,2-a]pyridines. European Journal of Organic Chemistry, 2020 , 2020, 269 | -3.84 | 56 |
| 233 | Comparison of serologic status of Toxoplasma gondii infection in pre- and post-heart transplantation in a pediatric population: A preliminary study. <i>Transplant Infectious Disease</i> , 2020 , 22, e13339 | 2.7 | O |
| 232 | 2,4-Dioxochroman Moiety Linked to 1,2,3-triazole Derivatives as Novel Eglucosidase Inhibitors: Synthesis, In vitro Biological Evaluation, and Docking Study. <i>Current Organic Chemistry</i> , 2020 , 24, 2019-2 | d27 | 1 |
| 231 | 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 | 0.8 | 1 |
| 230 | 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 | 0.8 | |
| 229 | 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 | 1.3 | 2 |
| 228 | Dimethyl Sulfoxide: Yesterday's Solvent, Today's Reagent. <i>Advanced Synthesis and Catalysis</i> , 2020 , 362, 65-86 | 5.6 | 53 |

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| 227 | 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 | 1.8 | 4 |
|-----|---|-----|----|
| 226 | Novel N,N-dimethylbarbituric-pyridinium derivatives as potent urease inhibitors: Synthesis, in vitro, and in silico studies. <i>Bioorganic Chemistry</i> , 2020 , 95, 103529 | 5.1 | 6 |
| 225 | Synthesis and biological evaluation of new benzimidazole-1,2,3-triazole hybrids as potential glucosidase inhibitors. <i>Bioorganic Chemistry</i> , 2020 , 95, 103482 | 5.1 | 25 |
| 224 | Synthesis of Arylidene Isoquinolinones bearing Combretastatin Skeleton by Cyclocarbopalladation/cross coupling Tandem Heck-Suzuki Miaura Reactions using nano catalyst Pd@Py-IL-SPION. <i>Applied Organometallic Chemistry</i> , 2020 , 34, e5279 | 3.1 | 2 |
| 223 | Amine-carbon disulfide promoted synthesis of novel benzo[e][1,3]thiazepin-5(1H)-one derivatives. Journal of Heterocyclic Chemistry, 2020 , 57, 413-418 | 1.9 | 2 |
| 222 | Benzoylquinazolinone derivatives as new potential antidiabetic agents: Ælucosidase inhibition, kinetic, and docking studies. <i>Journal of the Chinese Chemical Society</i> , 2020 , 67, 856-863 | 1.5 | 5 |
| 221 | Design, synthesis, biological evaluation, and docking study of novel dual-acting thiazole-pyridiniums inhibiting acetylcholinesterase and Emyloid aggregation for Alzheimer's disease. <i>Bioorganic Chemistry</i> , 2020 , 103, 104186 | 5.1 | 18 |
| 220 | Novel quinazolin-sulfonamid derivatives: synthesis, characterization, biological evaluation, and molecular docking studies. <i>Journal of Biomolecular Structure and Dynamics</i> , 2020 , 1-12 | 3.6 | 4 |
| 219 | Design, Synthesis, and Biological Evaluation of New Indole-Acrylamide-1,2,3-Triazole Derivatives as Potential Educosidase Inhibitors. <i>Polycyclic Aromatic Compounds</i> , 2020 , 1-9 | 1.3 | О |
| 218 | New acridine-9-carboxamide linked to 1,2,3-triazole-N-phenylacetamide derivatives as potent glucosidase inhibitors: design, synthesis, in vitro, and in silico biological evaluations. <i>Medicinal Chemistry Research</i> , 2020 , 29, 1836-1845 | 2.2 | 2 |
| 217 | 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 | 5.1 | 10 |
| 216 | Design, synthesis, biological evaluation, and docking study of new acridine-9-carboxamide linked to 1,2,3-triazole derivatives as antidiabetic agents targeting Eglucosidase. <i>Journal of Heterocyclic Chemistry</i> , 2020 , 57, 4348-4357 | 1.9 | 2 |
| 215 | Design, synthesis and antibacterial activity evaluation of novel 2-(4-((1-aryl-1H-1,2,3-triazol-4-yl)methoxy)phenyl)2-(2-oxoazetidin-1-yl)acetamide derivatives. <i>Journal of Heterocyclic Chemistry</i> , 2020 , 57, 4254-4261 | 1.9 | 3 |
| 214 | New Biscoumarin Derivatives as Potent Educosidase Inhibitors: Synthesis, Biological Evaluation, Kinetic Analysis, and Docking Study. <i>Polycyclic Aromatic Compounds</i> , 2020 , 40, 915-926 | 1.3 | 16 |
| 213 | 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 | 1.5 | 5 |
| 212 | 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 | 3.1 | 4 |
| 211 | Novel fused 1,2,3-triazolo-benzodiazepine derivatives as potent anticonvulsant agents: design, synthesis, in vivo, and in silico evaluations. <i>Molecular Diversity</i> , 2020 , 24, 179-189 | 3.1 | 8 |
| 210 | Design and synthesis of new imidazo[1,2-b]pyrazole derivatives, in vitro ⊞glucosidase inhibition, kinetic and docking studies. <i>Molecular Diversity</i> , 2020 , 24, 69-80 | 3.1 | 13 |

| 209 | Synthesis and Anticancer Activity of N-(di/trimethoxyaryl)-5-arylisoxazole-3-carboxamide. <i>Polycyclic Aromatic Compounds</i> , 2020 , 40, 1568-1580 | 1.3 | 2 |
|-----|---|-----|----|
| 208 | Synthesis and pharmacological properties of polysubstituted 2-amino-4H-pyran-3-carbonitrile derivatives. <i>Molecular Diversity</i> , 2020 , 24, 1385-1431 | 3.1 | 11 |
| 207 | Synthesis of highly functionalized organic compounds through Ugi post-transformations started from propiolic acids. <i>Molecular Diversity</i> , 2020 , 24, 855-887 | 3.1 | 6 |
| 206 | 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 | 2.8 | 23 |
| 205 | Recent Advances in Alkyne Hydroamination as a Powerful Tool for the Construction of CN Bonds. <i>Asian Journal of Organic Chemistry</i> , 2020 , 9, 969-991 | 3 | 11 |
| 204 | Copper-Mediated Direct Cyanatation of Benzamides: A New Approach to the Synthesis of Quinazolinediones. <i>European Journal of Organic Chemistry</i> , 2020 , 2020, 708-713 | 3.2 | 2 |
| 203 | Design, Synthesis, Molecular Docking, and Cholinesterase Inhibitory Potential of Phthalimide-Dithiocarbamate Hybrids as New Agents for Treatment of Alzheimer's Disease. <i>Chemistry and Biodiversity</i> , 2019 , 16, e1900370 | 2.5 | 12 |
| 202 | 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 | 5.1 | 11 |
| 201 | Synthesis and Biological Activity of Some Benzochromenoquinolinones: Tacrine Analogs as Potent Anti-Alzheimer's Agents. <i>Chemistry and Biodiversity</i> , 2019 , 16, e1800488 | 2.5 | 10 |
| 200 | Design, Synthesis, and Cholinesterase Inhibition Assay of Coumarin-3-carboxamide-N-morpholine Hybrids as New Anti-Alzheimer Agents. <i>Chemistry and Biodiversity</i> , 2019 , 16, e1900144 | 2.5 | 15 |
| 199 | Anticancer properties of N-alkyl-2, 4-diphenylimidazo [1, 2-a] quinoxalin-1-amine derivatives; kinase inhibitors. <i>Bioorganic Chemistry</i> , 2019 , 90, 103055 | 5.1 | 7 |
| 198 | The use of magnetic starch as a support for an ionic liquid-Etyclodextrin based catalyst for the synthesis of imidazothiadiazolamine derivatives. <i>International Journal of Biological Macromolecules</i> , 2019 , 135, 453-461 | 7.9 | 10 |
| 197 | Novel morpholine containing cinnamoyl amides as potent tyrosinase inhibitors. <i>International Journal of Biological Macromolecules</i> , 2019 , 135, 978-985 | 7.9 | 12 |
| 196 | Synthesis and characterization of Fe2O3@SiO2(CH2)3PDTCPd magnetic nanoparticles: a new and highly active catalyst for the Heck/Sonogashira coupling reactions. <i>New Journal of Chemistry</i> , 2019 , 43, 8930-8938 | 3.6 | 18 |
| 195 | Design, synthesis, and biological evaluation of novel 4-oxobenzo[d]1,2,3-triazin-benzylpyridinum derivatives as potent anti-Alzheimer agents. <i>Bioorganic and Medicinal Chemistry</i> , 2019 , 27, 2914-2922 | 3.4 | 7 |
| 194 | New thiosemicarbazide-1,2,3-triazole hybrids as potent Eglucosidase inhibitors: Design, synthesis, and biological evaluation. <i>Journal of Molecular Structure</i> , 2019 , 1192, 192-200 | 3.4 | 17 |
| 193 | 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 | 5.1 | 11 |
| 192 | Design and Synthesis of Novel Cytotoxic Indole-Thiosemicarbazone Derivatives: Biological Evaluation and Docking Study. <i>Chemistry and Biodiversity</i> , 2019 , 16, e1800470 | 2.5 | 7 |

| 191 | Anti-cancer, anti-oxidant and molecular docking studies of thiosemicarbazone indole-based derivatives. <i>Research on Chemical Intermediates</i> , 2019 , 45, 2827-2854 | 2.8 | 19 |
|-----|--|------------------|----|
| 190 | Mo (CO)6-assisted Pd-supported magnetic graphene oxide-catalyzed carbonylation-cyclization as an efficient way for the synthesis of 4(3H)-quinazolinones. <i>Applied Organometallic Chemistry</i> , 2019 , 33, e4769 | 3.1 | 11 |
| 189 | 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 | 5.1 | 12 |
| 188 | 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 | 3.4 | 25 |
| 187 | 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 | 1.4 | 1 |
| 186 | Biscoumarin-1,2,3-triazole hybrids as novel anti-diabetic agents: Design, synthesis, in vitro Eglucosidase inhibition, kinetic, and docking studies. <i>Bioorganic Chemistry</i> , 2019 , 92, 103206 | 5.1 | 40 |
| 185 | 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 | 2.8 | 3 |
| 184 | A new series of Schiff base derivatives bearing 1,2,3-triazole: Design, synthesis, molecular docking, and 🗄 lucosidase inhibition. <i>Archiv Der Pharmazie</i> , 2019 , 352, e1900034 | 4.3 | 10 |
| 183 | Design, synthesis, in vitro, and in silico studies of novel diarylimidazole-1,2,3-triazole hybrids as potent Eglucosidase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2019 , 27, 115148 | 3.4 | 17 |
| 182 | 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 | 1.4 | 2 |
| 181 | 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 | 2 | 4 |
| 180 | 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 | 1.7 | 4 |
| 179 | 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 | 3 | 3 |
| 178 | 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 | 0.8 | 9 |
| 177 | Pyrano[3,2-c]quinoline Derivatives as New Class of Eglucosidase Inhibitors to Treat Type 2 Diabetes: Synthesis, in vitro Biological Evaluation and Kinetic Study. <i>Medicinal Chemistry</i> , 2019 , 15, 8-16 | 1.8 | 10 |
| 176 | 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 | 2.2 | 5 |
| 175 | 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 | 1.8 | 6 |
| 174 | CuBr-catalysed one-pot multicomponent synthesis of 3-substituted 2-thioxo-2,3-dihydroquinazolin-4(1H)-one derivatives. <i>Applied Organometallic Chemistry</i> , 2019 , 33, e463 | 5 ^{3.1} | 12 |

| 173 | Design and Synthesis of Selective Acetylcholinesterase Inhibitors: Arylisoxazole-Phenylpiperazine Derivatives. <i>Chemistry and Biodiversity</i> , 2019 , 16, e1800433 | 2.5 | 17 |
|-----|--|-------------------|----|
| 172 | Novel and efficient synthesis of triazolobenzodiazepine analogues through the sequential Ugi 4CR-click-N-arylation reactions. <i>Tetrahedron Letters</i> , 2019 , 60, 583-585 | 2 | 13 |
| 171 | Tandem synthesis of benzo[d]naphtho[2,3-g][1,3]oxazocine-8,13(6H,14H)-dione derivatives. <i>Monatshefte F</i> ☐ <i>Chemie</i> , 2019 , 150, 347-352 | 1.4 | 1 |
| 170 | Design and synthesis of new fused carbazole-imidazole derivatives as anti-diabetic agents: In vitro Eglucosidase inhibition, kinetic, and in silico studies. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2019 , 29, 713-718 | 2.9 | 22 |
| 169 | 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 | 5.1 | 55 |
| 168 | Design, synthesis and anti-Alzheimer's activity of novel 1,2,3-triazole-chromenone carboxamide derivatives. <i>Bioorganic Chemistry</i> , 2019 , 83, 391-401 | 5.1 | 50 |
| 167 | Design and synthesis of novel quinazolinone-1,2,3-triazole hybrids as new anti-diabetic agents: In vitro 🗄 lucosidase inhibition, kinetic, and docking study. <i>Bioorganic Chemistry</i> , 2019 , 83, 161-169 | 5.1 | 74 |
| 166 | Facile access to new pyrido[2,3-d]pyrimidine derivatives. <i>Molecular Diversity</i> , 2019 , 23, 333-340 | 3.1 | 2 |
| 165 | New ciprofloxacindithiocarbamateBenzyl hybrids: design, synthesis, antibacterial evaluation, and molecular modeling studies. <i>Research on Chemical Intermediates</i> , 2019 , 45, 223-236 | 2.8 | 6 |
| 164 | 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 | 0.9 | 1 |
| 163 | Green synthesis of 2-((2-aryl-3-oxoisoindolin-1-yl)methyl)quinazolin-4(3H)-ones via sequential condensation, sp3 C H bond functionalization and cyclization. <i>Tetrahedron Letters</i> , 2018 , 59, 1555-1559 | 2 | 6 |
| 162 | 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 | 3.6 | 19 |
| 161 | Copper-catalyzed efficient synthesis of 5-arylindazolo[3,2-b]quinazolin-7(5H)-ones from 2-nitrobenzaldehydes. <i>Tetrahedron</i> , 2018 , 74, 2197-2201 | 2.4 | 4 |
| 160 | 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- | 5 2 83 | 15 |
| 159 | Novel quinazolin-4(3H)-one linked to 1,2,3-triazoles: Synthesis and anticancer activity. <i>Chemical Biology and Drug Design</i> , 2018 , 92, 1373-1381 | 2.9 | 21 |
| 158 | Design and synthesis of novel coumarin-pyridinium hybrids: In vitro cholinesterase inhibitory activity. <i>Bioorganic Chemistry</i> , 2018 , 77, 311-319 | 5.1 | 33 |
| 157 | Synthesis and biological evaluation of chalcone-triazole hybrid derivatives as 15-LOX inhibitors. Zeitschrift Fur Naturforschung - Section B Journal of Chemical Sciences, 2018 , 73, 77-83 | 1 | 3 |
| 156 | Design, synthesis and in vitro ঘ lucosidase inhibition of novel dihydropyrano[3,2-c]quinoline derivatives as potential anti-diabetic agents. <i>Bioorganic Chemistry</i> , 2018 , 77, 280-286 | 5.1 | 48 |

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| 155 | Copper-supported Ecyclodextrin-functionalized magnetic nanoparticles: Efficient multifunctional catalyst for one-pot greenBynthesis of 1,2,3-triazolylquinazolinone derivatives. <i>Applied Organometallic Chemistry</i> , 2018 , 32, e4212 | 3.1 | 20 | |
|-----|---|-----|----|--|
| 154 | SBA-15-SO3H-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 | 2.8 | 7 | |
| 153 | Biology-Oriented Drug Synthesis (BIODS) Approach towards Synthesis of Ciprofloxacin-Dithiocarbamate Hybrids and Their Antibacterial Potential both in Vitro and in Silico. <i>Chemistry and Biodiversity</i> , 2018 , 15, e1800273 | 2.5 | 3 | |
| 152 | Synthesis and cholinesterase inhibitory activity of new 2-benzofuran carboxamide-benzylpyridinum salts. <i>Bioorganic Chemistry</i> , 2018 , 80, 180-188 | 5.1 | 10 | |
| 151 | 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 | 1.7 | 5 | |
| 150 | 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 | 0.8 | 7 | |
| 149 | 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 | 1.3 | 0 | |
| 148 | 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 | 0.8 | 1 | |
| 147 | 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 | 2 | 16 | |
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| 145 | Design, synthesis and in vitro ঘ lucosidase inhibition of novel coumarin-pyridines as potent antidiabetic agents. <i>New Journal of Chemistry</i> , 2018 , 42, 17268-17278 | 3.6 | 31 | |
| 144 | Copper-catalyzed synthesis of 2,3-disubstituted quinazolin-4(3H)-ones from benzyl-substituted anthranilamides. <i>Heterocyclic Communications</i> , 2018 , 24, 267-271 | 1.7 | 2 | |
| 143 | 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 | 1 | 6 | |
| 142 | Novel cinnamic acid-tryptamine hybrids as potent butyrylcholinesterase inhibitors: Synthesis, biological evaluation, and docking study. <i>Archiv Der Pharmazie</i> , 2018 , 351, e1800115 | 4.3 | 8 | |
| 141 | Design, synthesis, and biological evaluation of selective and potent Carbazole-based butyrylcholinesterase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2018 , 26, 4952-4962 | 3.4 | 9 | |
| 140 | New 6-amino-pyrido[2,3-d]pyrimidine-2,4-diones as novel agents to treat type 2 diabetes: A simple and efficient synthesis, ⊞glucosidase inhibition, molecular modeling and kinetic study. <i>European Journal of Medicinal Chemistry</i> , 2018 , 155, 353-363 | 6.8 | 53 | |
| 139 | Synthesis of New Benzimidazole-1,2,3-triazole Hybrids as Tyrosinase Inhibitors. <i>Chemistry and Biodiversity</i> , 2018 , 15, e1800120 | 2.5 | 29 | |
| 138 | Synthesis, evaluation, and molecular docking studies of aryl urea-triazole-based derivatives as anti-urease agents. <i>Archiv Der Pharmazie</i> , 2018 , 351, e1800005 | 4.3 | 13 | |

| 137 | Novel tetrahydrocarbazole benzyl pyridine hybrids as potent and selective butryl cholinesterase inhibitors with neuroprotective and Elecretase inhibition activities. <i>European Journal of Medicinal Chemistry</i> , 2018 , 155, 49-60 | 6.8 | 18 |
|-----|--|------------------|----|
| 136 | 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 | 1.4 | 4 |
| 135 | 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 | 1.9 | 2 |
| 134 | Design, synthesis, docking study, lglucosidase 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- | 2 9 5 | 41 |
| 133 | Synthesis of Novel Tacrine Analogs as Acetylcholinesterase Inhibitors. <i>Journal of Heterocyclic Chemistry</i> , 2017 , 54, 384-390 | 1.9 | 13 |
| 132 | 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 | 6.8 | 29 |
| 131 | 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 | 5.1 | 42 |
| 130 | A green and efficient synthesis of 2-thioxoquinazolinone derivatives in water using potassium thiocyanate. <i>Journal of Sulfur Chemistry</i> , 2017 , 38, 519-529 | 2.3 | 3 |
| 129 | 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 | 4.1 | 32 |
| 128 | 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 | 3.4 | 26 |
| 127 | An efficient four-component reaction for the synthesis of chromeno[4,3-b]quinolone derivatives. Journal of the Iranian Chemical Society, 2017 , 14, 771-775 | 2 | 15 |
| 126 | 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 | 6.8 | 46 |
| 125 | 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 | 1 | O |
| 124 | Appel reagent as novel promoter for the synthesis of polysubstituted imidazoles. <i>Arkivoc</i> , 2017 , 2017, 343-352 | 0.9 | 1 |
| 123 | Synthesis and cytotoxicity of novel thioxo-quinazolino[3,4-\$a\$]quinazolinones. <i>Turkish Journal of Chemistry</i> , 2017 , 41, 125-134 | 1 | 2 |
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| 121 | An efficient approach to the synthesis of coumarin-fused dihydropyridinones. <i>Heterocyclic Communications</i> , 2017 , 23, | 1.7 | 3 |
| 120 | Improvement of the Van Leusen reaction in the presence of Etyclodextrin: a green and efficient synthesis of oxazoles in water. <i>Zeitschrift Fur Naturforschung - Section B Journal of Chemical Sciences</i> 2017 , 72, 923-926 | 1 | 5 |

| 119 | 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- | -7 37 8 | 32 | |
|-----|--|--------------------|-----|--|
| 118 | 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, | 1.7 | 2 | |
| 117 | An efficient access to 2,3-diarylimidazo[1,2-a]pyridines via silver(I)-catalyzed C-H bond functionalization. <i>Monatshefte Fil Chemie</i> , 2017 , 148, 1817-1821 | 1.4 | 2 | |
| 116 | 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 | 6.8 | 104 | |
| 115 | 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 | 6.8 | 64 | |
| 114 | Synthesis and biological evaluation of novel imidazopyrimidin-3-amines as anticancer agents. <i>Chemical Biology and Drug Design</i> , 2017 , 89, 797-805 | 2.9 | 9 | |
| 113 | Synthesis and anticholinesterase activity of new substituted benzo[d]oxazole-based derivatives. <i>Chemical Biology and Drug Design</i> , 2017 , 89, 783-789 | 2.9 | 17 | |
| 112 | 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 | 2 | 21 | |
| 111 | An efficient, four-component reaction for the synthesis of novel carbamodithioates. <i>Journal of Sulfur Chemistry</i> , 2017 , 38, 43-51 | 2.3 | 3 | |
| 110 | Synthesis of Thiazolone Derivatives as Novel Soybean 15-LOX Inhibitors. <i>Letters in Organic Chemistry</i> , 2017 , 14, 186-191 | 0.6 | 3 | |
| 109 | Novel Indole-Isoxazole Hybrids: Synthesis and In Vitro Anti-Cholinesterase Activity. <i>Letters in Drug Design and Discovery</i> , 2017 , 14, | 0.8 | 7 | |
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| 107 | 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 | | 3 | |
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| 103 | Quinoline-based imidazole-fused heterocycles as new inhibitors of 15-lipoxygenase. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016 , 31, 205-209 | 5.6 | 8 | |
| 102 | Novel Tacrine-Based Pyrano[3',4':5,6]pyrano[2,3-b]quinolinones: Synthesis and Cholinesterase Inhibitory Activity. <i>Archiv Der Pharmazie</i> , 2016 , 349, 915-924 | 4.3 | 16 | |

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|-----|--|------------------|----|
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| 99 | Design and synthesis of novel anti-Alzheimer's agents: Acridine-chromenone and quinoline-chromenone hybrids. <i>Bioorganic Chemistry</i> , 2016 , 67, 84-94 | 5.1 | 47 |
| 98 | Synthesis of novel fused quinazolinone derivatives. <i>Molecular Diversity</i> , 2016 , 20, 677-85 | 3.1 | 10 |
| 97 | Synthesis of Novel Phthalazino[1,2-b]quinazolinedione Derivatives: Efficient and Practical Reaction of 2-Amino-N?-Arylbenzohydrazides and 2-Formylbenzoic Acids. <i>Helvetica Chimica Acta</i> , 2016 , 99, 539-54 | 12 | 6 |
| 96 | A Novel Copper-Catalyzed Preparation of Pyrido[1,2-a]pyrimidine Derivatives. <i>Synlett</i> , 2016 , 27, 1359-13 | 86.2 | 3 |
| 95 | Synthesis of Novel Isoindolo[2,1-a]quinazolinedione Derivatives Containing a 1,2,3-Triazole Ring System. <i>Helvetica Chimica Acta</i> , 2016 , 99, 37-40 | 2 | 10 |
| 94 | Efficient three-step synthesis of benzo[e]imidazo[1,2-c][1,2,3]triazines. <i>Synthetic Communications</i> , 2016 , 46, 563-567 | 1.7 | 8 |
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| 92 | Iodine-Mediated Synthesis of Novel Pyrazole Derivatives. <i>Synthesis</i> , 2016 , 48, 541-546 | 2.9 | 7 |
| 91 | 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 | 6.8 | 55 |
| 90 | Copper supported Eyclodextrin grafted magnetic nanoparticles as an efficient recyclable catalyst for one-pot synthesis of 1-benzyl-1H-1,2,3-triazoldibenzodiazepinone derivatives via click reaction. <i>RSC Advances</i> , 2016 , 6, 28838-28843 | 3.7 | 25 |
| 89 | Synthesis and Evaluation of Novel Quinazolinone-1,2,3-Triazoles as Inhibitors of Lipoxygenase. Journal of Chemical Research, 2016 , 40, 188-191 | 0.6 | 12 |
| 88 | 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 | 0.8 | 18 |
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| 85 | CuBr/Et3N-Promoted Reactions of 2-Aminobenzamides and Isothiocyanates: Efficient Synthesis of Novel Quinazolin-4(3H)-ones. <i>Helvetica Chimica Acta</i> , 2016 , 99, 378-383 | 2 | 6 |
| 84 | Hetero-annulated coumarins as new AChE/BuChE inhibitors: synthesis and biological evaluation. <i>Medicinal Chemistry Research</i> , 2016 , 25, 1831-1841 | 2.2 | 11 |

| 83 | Phthalimide-Derived N-Benzylpyridinium Halides Targeting Cholinesterases: Synthesis and Bioactivity of New Potential Anti-Alzheimer's Disease Agents. <i>Archiv Der Pharmazie</i> , 2016 , 349, 293-301 | 4.3 | 17 | |
|----|--|------------------|----|--|
| 82 | Efficient Synthesis of Polyfunctionalized Pyrimidine Derivatives. <i>Synlett</i> , 2016 , 27, 1689-1692 | 2.2 | 3 | |
| 81 | 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 | 2 | 2 | |
| 80 | Synthesis of novel 1,2,3-triazole derivatives of 2,3-dihydroquinazolin-4(1H)-one. <i>Monatshefte F</i> ☐ <i>Chemie</i> , 2016 , 147, 2151-2156 | 1.4 | 2 | |
| 79 | Three-component one-pot synthesis of dihydrochromeno[4,3-b]pyrazolo[4,3-e]pyridines. <i>Heterocyclic Communications</i> , 2016 , 22, | 1.7 | 6 | |
| 78 | 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 | 1 | 3 | |
| 77 | Palladium-Catalyzed Regioselective Direct Cyanation of Acetanilide Derivatives with K4[Fe(CN)6] by CH Bond Activation. <i>European Journal of Organic Chemistry</i> , 2016 , 2016, 4269-4274 | 3.2 | 9 | |
| 76 | 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, 799-8 | 306 ⁸ | 74 | |
| 75 | 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 | 3.1 | 18 | |
| 74 | 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, 787-95 | 3.1 | 36 | |
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| 72 | Efficient Synthesis of 2-Methylenethiazolo[2,3-b]quinazolinone Derivatives. <i>Synlett</i> , 2015 , 26, 173-176 | 2.2 | 11 | |
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| 69 | A Highly Efficient Method for the Synthesis of Novel 1?H-spiro[indene-2,2?-quinazoline]-1,3,4?(3?H)-trione Derivatives. <i>Journal of Chemical Research</i> , 2015 , 39, 495-498 | 0.6 | 7 | |
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|----|--|-----|----|
| 64 | 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, 361-9 | 6.8 | 40 |
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| 62 | Experimental and computational evidence for KOt-Bu-promoted synthesis of oxopyrazino[1,2-a]indoles. <i>RSC Advances</i> , 2015 , 5, 101353-101361 | 3.7 | 16 |
| 61 | Synthesis of new benzo[f]imidazo[1,2-d][1,4]oxazepines: AgNO3-mediated intramolecular hydroamination. <i>Tetrahedron Letters</i> , 2015 , 56, 7082-7084 | 2 | 15 |
| 60 | Simple and efficient syntheses of novel benzo[4,5]imidazo[1,2-a]pyridine derivatives. <i>Tetrahedron Letters</i> , 2015 , 56, 743-746 | 2 | 17 |
| 59 | Synthesis of Novel 2-Oxoquinoline Derivatives via Ugi-Four-Component-Heck Reaction. <i>Journal of Heterocyclic Chemistry</i> , 2015 , 52, 386-391 | 1.9 | 6 |
| 58 | Synthesis and Cytotoxic Activity of Some Novel Dihyrobenzo[h]pyrano[3,2-c]chromene Derivatives. <i>Journal of Heterocyclic Chemistry</i> , 2015 , 52, 97-104 | 1.9 | 10 |
| 57 | Efficient Synthesis of Novel Thiazol-2-ylidene-amides Using Carbonylthiourea Building Blocks. Journal of Heterocyclic Chemistry, 2015 , 52, 1150-1153 | 1.9 | 3 |
| 56 | Synthesis and Evaluation of Coumarin R esveratrol Hybrids as 15-Lipoxygenaze Inhibitors. <i>Synthetic Communications</i> , 2015 , 45, 741-749 | 1.7 | 22 |
| 55 | Synthesis and evaluation of novel oxoisoindoline derivatives as acetylcholinesterase inhibitors. <i>Monatshefte Fil Chemie</i> , 2015 , 146, 637-643 | 1.4 | 17 |
| 54 | Synthesis and In Vitro Cytotoxic Activity of Novel Triazole-Isoxazole Derivatives. <i>Journal of Heterocyclic Chemistry</i> , 2015 , 52, 1743-1747 | 1.9 | 12 |
| 53 | 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, 1425-32 | 2.9 | 50 |
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| 48 | Convenient and sequential one-pot route for synthesis of 2-thioxoquinazolinone and quinazolinobenzothiazinedione derivatives. <i>Monatshefte Fill Chemie</i> , 2014 , 145, 497-504 | 1.4 | 11 |

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| 45 | Synthesis of novel indolo[2,3-c]quinolinones via Ugi-4CR/palladium-catalyzed arylation. <i>Tetrahedron</i> , 2014 , 70, 3931-3934 | 2.4 | 13 |
| 44 | A novel and efficient route for the synthesis of 5-nitrobenzo[d]oxazole derivatives. <i>Journal of Fluorine Chemistry</i> , 2014 , 161, 83-86 | 2.1 | 3 |
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| 42 | Imidazo[2,1-b]thiazole derivatives as new inhibitors of 15-lipoxygenase. <i>European Journal of Medicinal Chemistry</i> , 2014 , 87, 759-64 | 6.8 | 26 |
| 41 | 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 | 1.7 | 3 |
| 40 | 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- | 9 ^{6.8} | 75 |
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| 38 | 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 | 1.7 | 21 |
| 37 | CN cross-coupling reaction catalysed by efficient and reusable CuO/SiO2 nanoparticles under ligand-free conditions. <i>Applied Organometallic Chemistry</i> , 2014 , 28, 809-813 | 3.1 | 18 |
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| 32 | An Efficient Synthesis of 2,4,6-Triarylpyridines via Solvent-Free Reaction between Acetophenoneoximes and Aldehydes. <i>Synlett</i> , 2014 , 25, 1299-1301 | 2.2 | 10 |
| 31 | Novel Four-Step Synthesis of Thioxo-quinazolino[3,4-a]quinazolinone Derivatives. <i>Synthetic Communications</i> , 2014 , 44, 215-221 | 1.7 | 22 |
| 30 | 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 | 2.2 | 23 |

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| 28 | Synthesis of novel fused 4,5-dihydro-1,2,3-triazolo[1,5-a][1,4]benzodiazepine derivatives via four-component UgiBmiles-type reaction. <i>Tetrahedron</i> , 2013 , 69, 3506-3510 | 2.4 | 50 |
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| 26 | Green Synthesis of New Boron-Containing Quinazolines: Preparation of Benzo[d][1,3,2]diazaborinin-4(1H)-one Derivatives. <i>Synthetic Communications</i> , 2013 , 43, 2936-2942 | 1.7 | 17 |
| 25 | Reaction of Isatoic Anhydride, Amine, and N,N?-Dialkyl Carbodiimides Under Solvent-Free Conditions: New and Efficient Synthesis of 3-Alkyl-2-(alkylamino)quinazolin-4(3H)-ones. <i>Synthetic Communications</i> , 2013 , 43, 2385-2392 | 1.7 | 26 |
| 24 | 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, 124-8 | 6.8 | 58 |
| 23 | 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 | 2 | 29 |
| 22 | A green one-pot synthesis of N-alkyl-2-(2-oxoazepan-1-yl)-2-arylacetamide derivatives via an Ugi four-center, three-component reaction in water. <i>Tetrahedron Letters</i> , 2012 , 53, 7088-7092 | 2 | 25 |
| 21 | Large-scale virtual screening for the identification of new Helicobacter pylori urease inhibitor scaffolds. <i>Journal of Molecular Modeling</i> , 2012 , 18, 2917-27 | 2 | 50 |
| 20 | 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 | 2.2 | 32 |
| 19 | One-Pot, Four-Component Synthesis of Novel Imidazo[2,1-b]thiazol-5-amine Derivatives. <i>Synthesis</i> , 2012 , 44, 3649-3654 | 2.9 | 26 |
| 18 | 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 | 2.2 | 11 |
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| 10 | Catalyst-free three-component reaction between 2-aminopyridines (or 2-aminothiazoles), aldehydes, and isocyanides in water. <i>Tetrahedron Letters</i> , 2007 , 48, 7263-7265 | 2 | 95 |
| 9 | 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 | 2.2 | 32 |
| 8 | 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 | 2 | 68 |
| 7 | 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> ,1-9 | 1.3 | 1 |
| 6 | 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> ,e6482 | 3.1 | 2 |
| 5 | Design and synthesis of novel pyrazole-benzofuran hybrids: in vitro Eglucosidase inhibitory activity, kinetic and molecular modeling study | | 2 |
| 4 | An Efficient and Convenient Approach for Synthesizing Iodohydrin and Iodoether from Aromatic Alkenes Using Hg(BF4)2.SiO2 and I2. <i>Polycyclic Aromatic Compounds</i> ,1-9 | 1.3 | O |
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| 1 | Nickel Supported MCM-Functionalized 1,2,3-Triazol-4-ylmethanamine: An Efficient Nano-particle-Heterogeneous Catalyst Activate for Suzuki Reaction. <i>Catalysis Letters</i> ,1 | 2.8 | 0 |