Yaseen A M M El-Shaier

List of Publications by Citations

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42 636 16 23 g-index

43 902 3.7 4.52 ext. papers ext. citations avg, IF L-index

| # | Paper | IF | Citations |
|----|---|-------------------|-----------|
| 42 | Design, synthesis, modeling studies and biological evaluation of thiazolidine derivatives containing pyrazole core as potential anti-diabetic PPAR-lagonists and anti-inflammatory COX-2 selective inhibitors. <i>Bioorganic Chemistry</i> , 2019 , 82, 86-99 | 5.1 | 49 |
| 41 | FDA-Approved Drugs with Potent In Vitro Antiviral Activity against Severe Acute Respiratory Syndrome Coronavirus 2. <i>Pharmaceuticals</i> , 2020 , 13, | 5.2 | 47 |
| 40 | Synthesis of new thiazolo-pyrrolidine-(spirooxindole) tethered to 3-acylindole as anticancer agents. <i>Bioorganic Chemistry</i> , 2019 , 82, 423-430 | 5.1 | 46 |
| 39 | Substituted spirooxindole derivatives as potent anticancer agents through inhibition of phosphodiesterase 1 <i>RSC Advances</i> , 2018 , 8, 14335-14346 | 3.7 | 37 |
| 38 | Chitosan-Gelatin Hydrogel Crosslinked With Oxidized Sucrose for the Ocular Delivery of Timolol Maleate. <i>Journal of Pharmaceutical Sciences</i> , 2018 , 107, 3098-3104 | 3.9 | 30 |
| 37 | New spiro-oxindole constructed with pyrrolidine/thioxothiazolidin-4-one derivatives: Regioselective synthesis, X-ray crystal structures, Hirshfeld surface analysis, DFT, docking and antimicrobial studies. <i>Journal of Molecular Structure</i> , 2018 , 1152, 101-114 | 3.4 | 29 |
| 36 | Design and synthesis of new substituted spirooxindoles as potential inhibitors of the MDM2-p53 interaction. <i>Bioorganic Chemistry</i> , 2019 , 86, 598-608 | 5.1 | 28 |
| 35 | Novel Pyrazoloquinolin-2-ones: Design, synthesis, docking studies, and biological evaluation as antiproliferative EGFR-TK inhibitors. <i>Bioorganic Chemistry</i> , 2019 , 90, 103045 | 5.1 | 28 |
| 34 | A new procedure for the preparation of 2-vinylindoles and their [4+2] cycloaddition reaction. <i>Tetrahedron</i> , 2011 , 67, 5133-5141 | 2.4 | 28 |
| 33 | Non-acidic 1,3,4-trisubstituted-pyrazole derivatives as lonazolac analogs with promising COX-2 selectivity, anti-inflammatory activity and gastric safety profile. <i>Bioorganic Chemistry</i> , 2018 , 77, 568-578 | 5.1 | 27 |
| 32 | Design, synthesis, and antihypertensive activity of new pyrimidine derivatives endowing new pharmacophores. <i>Medicinal Chemistry Research</i> , 2019 , 28, 360-379 | 2.2 | 25 |
| 31 | New 1,2,4-triazole/pyrazole hybrids linked to oxime moiety as nitric oxide donor celecoxib analogs: Synthesis, cyclooxygenase inhibition anti-inflammatory, ulcerogenicity, anti-proliferative activities, apoptosis, molecular modeling and nitric oxide release studies. <i>Bioorganic Chemistry</i> , 2020 , 98, 103752 | 5.1 | 24 |
| 30 | Design and synthesis of pyrazolo[3,4-d]pyrimidines: Nitric oxide releasing compounds targeting hepatocellular carcinoma. <i>Bioorganic and Medicinal Chemistry</i> , 2017 , 25, 2956-2970 | 3.4 | 18 |
| 29 | New N-ribosides and N-mannosides of rhodanine derivatives with anticancer activity on leukemia cell line: Design, synthesis, DFT and molecular modelling studies. <i>Carbohydrate Research</i> , 2020 , 487, 107 | 1 89 4 | 18 |
| 28 | Novel Thiourea Derivatives Bearing Sulfonamide Moiety as Anticancer Agents Through COX-2 Inhibition. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2017 , 17, 1411-1425 | 2.2 | 16 |
| 27 | An perception for newly isolated flavonoids from peach fruit as privileged avenue for a countermeasure outbreak of COVID-19 <i>RSC Advances</i> , 2020 , 10, 29983-29998 | 3.7 | 16 |
| 26 | Synthesis of Pyrazole-Thiobarbituric Acid Derivatives: Antimicrobial Activity and Docking Studies. <i>Molecules</i> , 2016 , 21, | 4.8 | 14 |

(2021-2019)

| 25 | Discovery of New S-Glycosides and N-Glycosides of Pyridine-biphenyl System with Antiviral Activity and Induction of Apoptosis in MCF7 Cells. <i>Journal of Heterocyclic Chemistry</i> , 2019 , 56, 1733-1747 | 1.9 | 13 |
|----|---|-----|----|
| 24 | Design, synthesis and biological evaluation of new 2-aminothiazole scaffolds as phosphodiesterase type 5 regulators and COX-1/COX-2 inhibitors <i>RSC Advances</i> , 2020 , 10, 29723-29736 | 3.7 | 13 |
| 23 | Molecular structure and spectroscopic investigations combined with hypoglycemic/anticancer and docking studies of a new barbituric acid derivative. <i>Journal of Molecular Structure</i> , 2017 , 1134, 99-111 | 3.4 | 12 |
| 22 | Design, synthesis and anticonvulsant activity of new imidazolidindione and imidazole derivatives. <i>Bioorganic Chemistry</i> , 2020 , 101, 104020 | 5.1 | 12 |
| 21 | Synthesis, Anticancer Activity, and Molecular Modeling of New Halogenated Spiro[pyrrolidine-thiazolo-oxindoles] Derivatives. <i>Applied Sciences (Switzerland)</i> , 2020 , 10, 2170 | 2.6 | 11 |
| 20 | Synthesis of some benzimidazole derivatives endowed with 1,2,3-triazole as potential inhibitors of hepatitis C virus. <i>Acta Pharmaceutica</i> , 2016 , 66, 219-31 | 3.2 | 11 |
| 19 | New 2-Oxopyridine/2-Thiopyridine Derivatives Tethered to a Benzotriazole with Cytotoxicity on MCF7 Cell Lines and with Antiviral Activities. <i>Letters in Drug Design and Discovery</i> , 2020 , 17, 124-137 | 0.8 | 9 |
| 18 | Bioactive constituents from Thunbergia erecta as potential anticholinesterase and anti-ageing agents: Experimental and in silico studies. <i>Bioorganic Chemistry</i> , 2021 , 108, 104643 | 5.1 | 9 |
| 17 | Arylidenes of Quinolin-2-one scaffold as Erlotinib analogues with activities against leukemia through inhibition of EGFR TK/STAT-3 pathways. <i>Bioorganic Chemistry</i> , 2020 , 96, 103628 | 5.1 | 8 |
| 16 | Design, synthesis and biological study of hybrid drug candidates of nitric oxide releasing cucurbitacin-inspired estrone analogs for treatment of hepatocellular carcinoma. <i>Bioorganic Chemistry</i> , 2019 , 85, 515-533 | 5.1 | 7 |
| 15 | Design and synthesis of pyrazolo[3,4-d]pyrimidinone derivatives: Discovery of selective phosphodiesterase-5 inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020 , 30, 127337 | 2.9 | 7 |
| 14 | Synthesis of Pyridine-Dicarboxamide-Cyclohexanone Derivatives: Anticancer and Educosidase Inhibitory Activities and In Silico Study. <i>Molecules</i> , 2019 , 24, | 4.8 | 6 |
| 13 | Structure- and Ligand-Based Studies towards the Repurposing of Marine Bioactive Compounds to Target SARS-CoV-2 <i>Arabian Journal of Chemistry</i> , 2021 , 14, 103092 | 5.9 | 6 |
| 12 | A new barbituric acid derivatives as reactive oxygen scavenger: Experimental and theoretical investigations. <i>Journal of Molecular Structure</i> , 2019 , 1175, 524-535 | 3.4 | 5 |
| 11 | Synthesis and Inhibitory Effect of Some Indole-Pyrimidine Based Hybrid Heterocycles on Eulucosidase and EAmylase as Potential Hypoglycemic Agents. <i>ChemistryOpen</i> , 2019 , 8, 1288-1297 | 2.3 | 5 |
| 10 | A review on the synthesis of heteroannulated quinolones and their biological activities. <i>Molecular Diversity</i> , 2021 , 1 | 3.1 | 5 |
| 9 | Potent Quinoline-Containing Combretastatin A-4 Analogues: Design, Synthesis, Antiproliferative, and Anti-Tubulin Activity. <i>Pharmaceuticals</i> , 2020 , 13, | 5.2 | 4 |
| 8 | Identification and molecular modeling of new quinolin-2-one thiosemicarbazide scaffold with antimicrobial urease inhibitory activity. <i>Molecular Diversity</i> , 2021 , 25, 13-27 | 3.1 | 4 |

| 7 | Discovery of novel quinoline-based analogues of combretastatin A-4 as tubulin polymerisation inhibitors with apoptosis inducing activity and potent anticancer effect. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2021 , 36, 802-818 | 5.6 | 4 | |
|---|--|--------------|---|--|
| 6 | Design and synthesis of novel pyrazolo[3,4-d]pyrimidin-4-one bearing quinoline scaffold as potent dual PDE5 inhibitors and apoptotic inducers for cancer therapy. <i>Bioorganic Chemistry</i> , 2020 , 105, 104352 | 5.1 | 2 | |
| 5 | Quinolones as prospective drugs: Their syntheses and biological applications. <i>Advances in Heterocyclic Chemistry</i> , 2021 , 147-196 | 2.4 | 1 | |
| 4 | Multifunctional Isosteric Pyridine Analogs-Based 2-Aminothiazole: Design, Synthesis, and Potential Phosphodiesterase-5 Inhibitory Activity. <i>Molecules</i> , 2021 , 26, | 4.8 | 1 | |
| 3 | Regioselective formation of new 3-S-alkylated-1,2,4-triazole-quinolones. <i>Journal of Sulfur Chemistry</i> ,1-17 | 7 2.3 | О | |
| 2 | Lipid polymer hybrid nanocarriers as a combinatory platform for different anti-SARS-CoV-2 drugs supported by computational studies <i>RSC Advances</i> , 2021 , 11, 28876-28891 | 3.7 | О | |
| 1 | Intelligent Drug Descriptors Analysis: Toward COVID-19 Drug Repurposing. Studies in Computational Intelligence, 2022, 173-191 | 0.8 | О | |