Hajjaj H M Abdu-Allah

List of Publications by Year in descending order

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| # | Article | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Design and synthesis of novel 2,3-dihydropyrazino[1,2-a]indole-1,4-dione derivatives as antiproliferative EGFR and BRAFV600E dual inhibitors. Bioorganic Chemistry, 2020, 104, 104260. | 2.0 | 50 |
| 2 | Synthesis, biological evaluation and docking study of 1,3,4-thiadiazole-thiazolidinone hybrids as anti-inflammatory agents with dual inhibition of COX-2 and 15-LOX. Bioorganic Chemistry, 2018, 80, 461-471. | 2.0 | 49 |
| 3 | Further insight into the dual COX-2 and 15-LOX anti-inflammatory activity of 1,3,4-thiadiazole-thiazolidinone hybrids: The contribution of the substituents at 5th positions is size dependent. Bioorganic Chemistry, 2020, 97, 103657. | 2.0 | 39 |
| 4 | CD22-Antagonists with nanomolar potency: The synergistic effect of hydrophobic groups at C-2 and C-9 of sialic acid scaffold. Bioorganic and Medicinal Chemistry, 2011, 19, 1966-1971. | 1.4 | 37 |
| 5 | Click chemistry synthesis, biological evaluation and docking study of some novel 2′-hydroxychalcone-triazole hybrids as potent anti-inflammatory agents. Bioorganic Chemistry, 2020, 95, 103505. | 2.0 | 37 |
| 6 | Design, Synthesis, and Structureâ^'Affinity Relationships of Novel Series of Sialosides as CD22-Specific Inhibitors. Journal of Medicinal Chemistry, 2008, 51, 6665-6681. | 2.9 | 31 |
| 7 | Inhibition of SHP2 by new compounds induces differential effects on RAS/RAF/ERK and PI3K/AKT pathways in different cancer cell types. Investigational New Drugs, 2019, 37, 252-261. | 1.2 | 27 |
| 8 | Proximity labeling of cis-ligands of CD22/Siglec-2 reveals stepwise α2,6 sialic acid-dependent and -independent interactions. Biochemical and Biophysical Research Communications, 2018, 495, 854-859. | 1.0 | 26 |
| 9 | CD22-Binding Synthetic Sialosides Regulate B Lymphocyte Proliferation Through CD22 Ligand-Dependent and Independent Pathways, and Enhance Antibody Production in Mice. Frontiers in Immunology, 2018, 9, 820. | 2.2 | 25 |
| 10 | 5-Aminosalyclic Acid (5-ASA): A Unique Anti-Inflammatory Salicylate. , 2016, 06, . | | 23 |
| 11 | Synthesis of some benzimidazole derivatives endowed with 1,2,3-triazole as potential inhibitors of hepatitis C virus. Acta Pharmaceutica, 2016, 66, 219-231. | 0.9 | 21 |
| 12 | Design and synthesis of novel 5-aminosalicylate (5-ASA)–4-thiazolinone hybrid derivatives with promising antiproliferative activity. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 1647-1650. | 1.0 | 20 |
| 13 | Synthesis and anti-inflammatory testing of some new compounds incorporating 5-aminosalicylic acid (5-ASA) as potential prodrugs. Archives of Pharmacal Research, 2005, 28, 637-647. | 2.7 | 19 |
| 14 | Potent small molecule mouse CD22-inhibitors: Exploring the interaction of the residue at C-2 of sialic acid scaffold. Bioorganic and Medicinal Chemistry Letters, 2009, 19, 5573-5575. | 1.0 | 19 |
| 15 | Novel N-substituted 5-aminosalicylamides as dual inhibitors of cyclooxygenase and 5-lipoxygenase enzymes: Synthesis, biological evaluation and docking study. Bioorganic Chemistry, 2018, 78, 80-93. | 2.0 | 18 |
| 16 | Synthesis and anti-mycobacterial activity of 4-(4-phenyl-1H-1,2,3-triazol-1-yl)salicylhydrazones: revitalizing an old drug. Archives of Pharmacal Research, 2017, 40, 168-179. | 2.7 | 15 |
| 17 | Conjugation of 4-aminosalicylate with thiazolinones afforded non-cytotoxic potent in vitro and in vivo anti-inflammatory hybrids. Bioorganic Chemistry, 2020, 94, 103378. | 2.0 | 14 |
| 18 | Induction of DNA damage, apoptosis and cell cycle perturbation mediate cytotoxic activity of new 5-aminosalicylate–4-thiazolinone hybrid derivatives. Biomedicine and Pharmacotherapy, 2020, 131, 110571. | 2.5 | 11 |

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|----|---|-------|-----------|
| 19 | Bis-(5-substituted-2-thiono-1,3,5-thiadiazinan-3-yl) butane as a scaffold of anti-proliferative activity, blended by a multicomponent process. Medicinal Chemistry Research, 2018, 27, 1103-1110. | 1.1 | 10 |
| 20 | Synthesis of biotinylated sialoside to probe CD22–ligand interactions. Tetrahedron Letters, 2009, 50, 4488-4491. | 0.7 | 9 |
| 21 | The Protein Tyrosine Phosphatase SHP-1 (PTPN6) but Not CD45 (PTPRC) Is Essential for the Ligand-Mediated Regulation of CD22 in BCR-Ligated B Cells. Journal of Immunology, 2021, 206, 2544-2551. | 0.4 | 9 |
| 22 | Synthesis of B- and C-ring-modified lithocholic acid analogues as potential sialyltransferase inhibitors. Steroids, 2016, 112, 54-61. | 0.8 | 8 |
| 23 | The inhibitory coreceptor CD22 restores B cell signaling by developmentally regulating <i> Cd45 ^{â^'/â^'} </i> immunodeficient B cells. Science Signaling, 2022, 15, eabf9570. | 1.6 | 6 |
| 24 | Ionic liquid of ketoprofen-piperine modulates the pharmaceutical and therapeutic characters of ketoprofen. International Journal of Pharmaceutics, 2022, 620, 121724. | 2.6 | 6 |
| 25 | Design, synthesis and molecular docking study of α-triazolylsialosides as non-hydrolyzable and potent CD22 ligands. European Journal of Medicinal Chemistry, 2020, 208, 112707. | 2.6 | 5 |
| 26 | Synthesis, characterization, and photophysical properties of some new thieno[2,3â€ <i>b</i>]pyridines bearing phenylethenyl moiety. Journal of Heterocyclic Chemistry, 2022, 59, 359-370. | 1.4 | 5 |
| 27 | Design and Synthesis of a Multivalent Heterobifunctional CD22 Ligand as a Potential Immunomodulator. Synthesis, 2011, 2011, 2968-2974. | 1.2 | 4 |
| 28 | Crystal structure of methyl 2-hydroxy-5-[(4-oxo-4,5-dihydro-1,3-thiazol-2-yl)amino]benzoate. Acta Crystallographica Section E: Crystallographic Communications, 2015, 71, o282-o283. | 0.2 | 3 |
| 29 | Synthesis and characterization of some new pyridines, thieno[2,3―b] pyridines and pyrido[3′,2′:4,5]thieno[3,2―d]pyrimidineâ€4(3 H)â€ones bearing styryl moiety. Journal of Heterocyclic Chemistry, 2020, 57, 2379-2388. | 1.4 | 3 |
| 30 | Stereoselective trimethylsilylation of \hat{I}_{\pm} - and \hat{I}_{\pm} -galactopyranoses. Carbohydrate Research, 2019, 474, 51-56. | 1.1 | 1 |
| 31 | Synthesis and characterization of some new S-substituted sulfanylpyridines, thieno[2,3-b]pyridines and related heterocycles. Arkivoc, 2021, 2020, 46-57. | 0.3 | 1 |
| 32 | Nature-inspired design of tetraindoles: Optimization of the core structure and evaluation of structure–activity relationship. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 4497-4503. | 1.0 | 0 |
| 33 | 5-aminosalicylate–4-thiazolinone hybrid derivatives: A potent modulator of DNA damage response and G2/M cell cycle arrest via ATM/ATR pathway and Cyclin-CDK complex. , 0, , . | | 0 |
| 34 | Crystal structure of 3-methyl-1-phenyl-6-propylamino-1H-pyrazolo[3,4-b]pyridine-5-carbonitrile. Acta Crystallographica Section E: Crystallographic Communications, 2015, 71, 0766-0767. | 0.2 | 0 |
| 35 | Crystal structure and Hirshfeld surface analysis of 5-acetyl-3-amino-6-methyl- <i>N</i> -phenyl-4-[(<i>E</i>)-2-phenylethenyl]thieno[2,3- <i>b</i>]pyridine-2-carboxan Acta Crystallographica Section E: Crystallographic Communications, 2022, 78, 225-230. | nide2 | 0 |