Ove Alexander HÃ, gmoen Ästrand

List of Publications by Year in descending order

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Ove Alexander HÃ,gmoen

| # | Article | IF | CITATIONS |
|----|--|-------------------|----------------|
| 1 | Synthesis and Preclinical Evaluation of TPA-Based Zinc Chelators as Metallo-β-lactamase Inhibitors. ACS Infectious Diseases, 2018, 4, 1407-1422. | 3.8 | 35 |
| 2 | Total synthesis and antileukemic evaluations of the phenazine 5,10-dioxide natural products iodinin, myxin and their derivatives. Bioorganic and Medicinal Chemistry, 2017, 25, 2285-2293. | 3.0 | 25 |
| 3 | Antimicrobial Activity and Cytotoxicity of Ag(I) and Au(I) Pillarplexes. Frontiers in Chemistry, 2018, 6, 584. | 3.6 | 22 |
| 4 | ZN148 Is a Modular Synthetic Metallo-β-Lactamase Inhibitor That Reverses Carbapenem Resistance in Gram-Negative Pathogens <i>In Vivo</i> . Antimicrobial Agents and Chemotherapy, 2020, 64, . | 3.2 | 22 |
| 5 | Synthesis of a novel legumain-cleavable colchicine prodrug with cell-specific toxicity. Bioorganic and Medicinal Chemistry, 2014, 22, 3309-3315. | 3.0 | 18 |
| 6 | Synthesis and biological evaluation of zinc chelating compounds as metallo-β-lactamase inhibitors. MedChemComm, 2019, 10, 528-537. | 3.4 | 13 |
| 7 | Synthesis and biological evaluation of new dipicolylamine zinc chelators as metallo-β-lactamase inhibitors. Tetrahedron, 2019, 75, 1525-1540. | 1.9 | 10 |
| 8 | Regulation of liver X receptor target genes by 22-functionalized oxysterols. Synthesis, in silico and in vitro evaluations. Steroids, 2017, 118, 119-127. | 1.8 | 8 |
| 9 | Degradative Behavior and Toxicity of Alkylated Imidazoles. Industrial & Engineering Chemistry Research, 2020, 59, 587-595. | 3.7 | 6 |
| 10 | Development of new LXR modulators that regulate LXR target genes and reduce lipogenesis in human cell models. European Journal of Medicinal Chemistry, 2014, 74, 258-263. | 5.5 | 5 |
| 11 | Synthesis, in vitro and in vivo biological evaluation of new oxysterols as modulators of the liver X receptors. Journal of Steroid Biochemistry and Molecular Biology, 2017, 165, 323-330. | 2.5 | 5 |
| 12 | Zinc-Chelating Compounds as Inhibitors of Human and Bacterial Zinc Metalloproteases. Molecules, 2022, 27, 56. | 3.8 | 3 |
| 13 | Crystal structure of (S)-2-[(3S,8S,9S,10R,13S,14S,17R)-3-hydroxy-10,13-dimethyl-2,3,4,7,8,9,10,11,12,13,14,15,16,17-tetradecahyd (Fernholz Weinreb amide). Acta Crystallographica Section E: Crystallographic Communications, 2015, 71, 275-277. | dro-1H-cyc 0.5 | :lopenta[a]pho |
| 14 | (2S,3S)-2,6-Dimethylheptane-1,3-diol, the oxygenated side chain of 22(S)-hydroxycholestrol, and its synthetic precursor (R)-4-benzyl-3-[(2R,3S)-3-hydroxy-2,6-dimethylheptanoyl]-1,3-oxazolidin-2-one. Acta Crystallographica Section C: Crystal Structure Communications, 2013, 69, 647-650. | 0.4 | 0 |
| 15 | (Acetonitrile){2-[bis(pyridin-2-ylmethyl-κ2N)amino-κN]-N-(2,6-dimethylphenyl)acetamide-κO}(perchlorato-κO)z (acetonitrile){2-[bis(pyridin-2-ylmethyl-κ2N)amino-κN]-N-(2,6-dimethylphenyl)acetamide-κO}zinc tris(perchlorate). Acta Crystallographica Section E: Structure Reports Online, 2013, 69, m112-m113. | inc 0.2 | 0 |