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List of Publications by Year in descending order

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papers

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