## Ivo Frydrych

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Substituted dienes prepared from betulinic acid – Synthesis, cytotoxicity, mechanism of action, and pharmacological parameters. European Journal of Medicinal Chemistry, 2021, 224, 113706.	2.6	6
2	A Novel Biological Role for Peptidyl-Arginine Deiminases: Citrullination of Cathelicidin LL-37 Controls the Immunostimulatory Potential of Cell-Free DNA. Journal of Immunology, 2018, 200, 2327-2340.	0.4	27
3	Peloruside A-Induced Cell Death in Hypoxia Is p53 Dependent in HCT116 Colorectal Cancer Cells. Journal of Natural Products, 2018, 81, 634-640.	1.5	5
4	Synthesis and Cytotoxic and Antiviral Profiling of Pyrrolo- and Furo-Fused 7-Deazapurine Ribonucleosides. Journal of Medicinal Chemistry, 2018, 61, 9347-9359.	2.9	24
5	Cellular effects of the microtubule-targeting agent peloruside A in hypoxia-conditioned colorectal carcinoma cells. Biochimica Et Biophysica Acta - General Subjects, 2017, 1861, 1833-1843.	1.1	5
6	Trilobolide-steroid hybrids: Synthesis, cytotoxic and antimycobacterial activity. Steroids, 2017, 117, 97-104.	0.8	15
7	Looking for ugly ducklings: The role of the stability of BrdU-antibody complex and the improved method of the detection of DNA replication. PLoS ONE, 2017, 12, e0174893.	1.1	9
8	Cell cycle profiling by image and flow cytometry: The optimised protocol for the detection of replicational activity using 5-Bromo-2′-deoxyuridine, low concentration of hydrochloric acid and exonuclease III. PLoS ONE, 2017, 12, e0175880.	1.1	8
9	Cells and Stripes: A novel quantitative photo-manipulation technique. Scientific Reports, 2016, 6, 19567.	1.6	13
10	Synthesis of cytotoxic 2,2-difluoroderivatives of dihydrobetulinic acid and allobetulin and study of their impact on cancer cells. European Journal of Medicinal Chemistry, 2015, 96, 482-490.	2.6	27
11	Caffeine–hydrazones as anticancer agents with pronounced selectivity toward T-lymphoblastic leukaemia cells. Bioorganic Chemistry, 2015, 60, 19-29.	2.0	42
12	Effects of synthetic A3 adenosine receptor agonists on cell proliferation and viability are receptor independent at micromolar concentrations. Journal of Physiology and Biochemistry, 2013, 69, 405-417.	1.3	12
13	Cyclosporin A sensitises Bcr-Abl positive cells to imatinib mesylate independently of P-glycoprotein expression. Toxicology in Vitro, 2009, 23, 1482-1490.	1.1	10
14	Serine protease inhibitors <i>N</i> â€Î±â€Tosylâ€ <scp>L</scp> â€Lysinylâ€Chloromethylketone (TLCK) and <i>N</i> â€Tosylâ€ <scp>L</scp> â€Phenylalaninylâ€Chloromethylketone (TPCK) are potent inhibitors of activated caspase proteases. Journal of Cellular Biochemistry, 2008, 103, 1646-1656.	1.2	30
15	Serine protease inhibitors <i>N</i> â€Î±â€tosylâ€ <scp>L</scp> â€lysinylâ€chloromethylketone (TLCK) and <i>N</i> â€tosylâ€ <scp>L</scp> â€phenylalaninylâ€chloromethylketone (TPCK) do not inhibit caspaseâ€3 and caspaseâ€7 processing in cells exposed to proâ€apoptotic inducing stimuli. Journal of Cellular Biochemistry. 2008. 105. 1501-1506.	1.2	5
16	The broad-spectrum caspase inhibitor Boc-Asp-CMK induces cell death in human leukaemia cells. Toxicology in Vitro, 2008, 22, 1356-1360.	1.1	5