

Agnieszka PotÄga

List of Publications by Year in descending order

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12
papers

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1478505

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#	ARTICLE	IF	CITATIONS
1	The Imidazoacridinone Antitumor Drug, C-1311, Is Metabolized by Flavin Monooxygenases but Not by Cytochrome P450s. <i>Drug Metabolism and Disposition</i> , 2011, 39, 1423-1432.	3.3	22
2	Modulation of CYP3A4 activity and induction of apoptosis, necrosis and senescence by the anti-tumour imidazoacridinone C-1311 in human hepatoma cells. <i>Cell Biology International</i> , 2013, 37, 109-120.	3.0	13
3	Diminished toxicity of C-1748, 4-methyl-9-hydroxyethylamino-1-nitroacridine, compared with its demethyl analog, C-857, corresponds to its resistance to metabolism in HepG2 cells. <i>Biochemical Pharmacology</i> , 2012, 84, 30-42.	4.4	10
4	Electrochemical simulation of metabolism for antitumor-active imidazoacridinone C-1311 and in silico prediction of drug metabolic reactions. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2019, 169, 269-278.	2.8	10
5	Phase I and phase II metabolism simulation of antitumor-active 2-hydroxyacridinone with electrochemistry coupled on-line with mass spectrometry. <i>Xenobiotica</i> , 2019, 49, 922-934.	1.1	9
6	Novel insights into conjugation of antitumor-active unsymmetrical bisacridine C-2028 with glutathione: Characteristics of non-enzymatic and glutathione S-transferase-mediated reactions. <i>Journal of Pharmaceutical Analysis</i> , 2021, 11, 791-798.	5.3	7
7	Mechanism-based inactivation of human cytochrome P450 1A2 and 3A4 isoenzymes by anti-tumor triazoloacridinone C-1305. <i>Xenobiotica</i> , 2016, 46, 1056-1065.	1.1	6
8	Imidazoacridinone antitumor agent C-1311 as a selective mechanism-based inactivator of human cytochrome P450 1A2 and 3A4 isoenzymes. <i>Pharmacological Reports</i> , 2016, 68, 663-670.	3.3	6
9	Electrochemical and in silico approaches for liver metabolic oxidation of antitumor-active triazoloacridinone C-1305. <i>Journal of Pharmaceutical Analysis</i> , 2020, 10, 376-384.	5.3	6
10	Metabolic Profiles of New Unsymmetrical Bisacridine Antitumor Agents in Electrochemical and Enzymatic Noncellular Systems and in Tumor Cells. <i>Pharmaceuticals</i> , 2021, 14, 317.	3.8	6
11	Acid-Base Equilibrium and Self-Association in Relation to High Antitumor Activity of Selected Unsymmetrical Bisacridines Established by Extensive Chemometric Analysis. <i>Molecules</i> , 2022, 27, 3995.	3.8	5
12	Electrochemical simulation of metabolic reduction and conjugation reactions of unsymmetrical bisacridine antitumor agents, C-2028 and C-2053. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2021, 197, 113970.	2.8	3