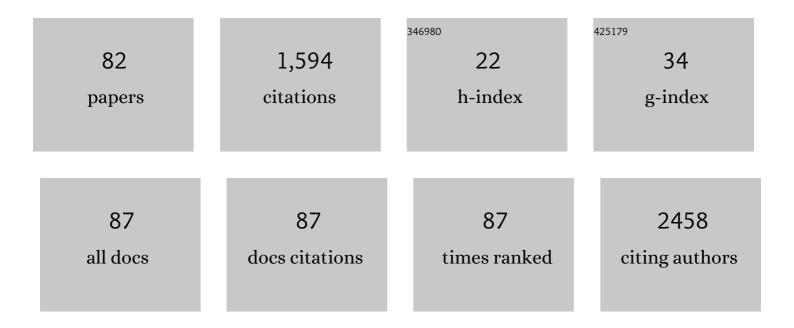
Krzysztof Bielawski

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	2-{5-[(Z,2Z)-2-Chloro-3-(4-nitrophenyl)-2-propenylidene]-4-oxo-2-thioxothiazolidin-3-yl}-3-methylbutanoic Acid as a Potential Anti-Breast Cancer Molecule. International Journal of Molecular Sciences, 2022, 23, 4091.	1.8	6
2	Exploration of novel heterofused 1,2,4-triazine derivative in colorectal cancer. Journal of Enzyme Inhibition and Medicinal Chemistry, 2021, 36, 535-548.	2.5	18
3	[1,2,4]triazyny – potencjalne leki w chemioterapii nowotworów. Postepy Higieny I Medycyny Doswiadczalnej, 2021, 75, 64-84.	0.1	0
4	In Vitro Anticancer Potential of Jasione montana and Its Main Components against Human Amelanotic Melanoma Cells. International Journal of Molecular Sciences, 2021, 22, 3345.	1.8	12
5	The Anticancer Action of a Novel 1,2,4-Triazine Sulfonamide Derivative in Colon Cancer Cells. Molecules, 2021, 26, 2045.	1.7	14
6	Synthesis and Anticancer Activity Evaluation of 5-[2-Chloro-3-(4-nitrophenyl)-2-propenylidene]-4-thiazolidinones. Molecules, 2021, 26, 3057.	1.7	14
7	Mechanism of Anticancer Action of Novel Imidazole Platinum(II) Complex Conjugated with G2 PAMAM-OH Dendrimer in Breast Cancer Cells. International Journal of Molecular Sciences, 2021, 22, 5581.	1.8	8
8	Autophagy Modulators in Cancer Therapy. International Journal of Molecular Sciences, 2021, 22, 5804.	1.8	37
9	Anti-HER2 monoclonal antibodies intensify the susceptibility of human gastric cancer cells to etoposide by promoting apoptosis, but not autophagy. PLoS ONE, 2021, 16, e0255585.	1.1	4
10	Mucin 1 as a Molecular Target of a Novel Diisoquinoline Derivative Combined with Anti-MUC1 Antibody in AGS Gastric Cancer Cells. Molecules, 2021, 26, 6504.	1.7	2
11	DNA topoisomerases as molecular targets for anticancer drugs. Journal of Enzyme Inhibition and Medicinal Chemistry, 2020, 35, 1781-1799.	2.5	58
12	The Effect of Novel 7-methyl-5-phenyl-pyrazolo[4,3-e]tetrazolo[4,5-b][1,2,4]triazine Sulfonamide Derivatives on Apoptosis and Autophagy in DLD-1 and HT-29 Colon Cancer Cells. International Journal of Molecular Sciences, 2020, 21, 5221.	1.8	18
13	The intensification of anticancer activity of LFM-A13 by erythropoietin as a possible option for inhibition of breast cancer. Journal of Enzyme Inhibition and Medicinal Chemistry, 2020, 35, 1697-1711.	2.5	4
14	1,2,4-Triazine Sulfonamides: Synthesis by Sulfenamide Intermediates, In Vitro Anticancer Screening, Structural Characterization, and Molecular Docking Study. Molecules, 2020, 25, 2324.	1.7	11
15	Which salivary components can differentiate metabolic obesity?. PLoS ONE, 2020, 15, e0235358.	1.1	10
16	Evaluation of the Anticancer Activities of Novel Transition Metal Complexes with Berenil and Nitroimidazole. Molecules, 2020, 25, 2860.	1.7	18
17	Monoclonal anti‑MUC1 antibody with novel octahydropyrazino[2,1‑a:5,4‑a']diisoquinoline derivative as a potential multi‑targeted strategy in MCF‑7 breast cancer cells. Oncology Reports, 2019, 42, 1391-1403.	1.2	8
18	<effect 2nd="" 3rd="" and="" cytokines="" dendrimers="" differentiation,="" fibroblasts<="" generation="" human="" in="" keratinocytes="" of="" on="" p="" pamam="" pro-inflammatory="" proliferation,="">. International Journal of Nanomedicine, 2019, Volume 14, 7123-7139.</effect>	3.3	20

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19	Antioxidant and cytotoxic activity of new di- and polyamine caffeine analogues. Free Radical Research, 2018, 52, 724-736.	1.5	10
20	The molecular mechanism of anticancer action of novel octahydropyrazino[2,1-a:5,4-a′]diisoquinoline derivatives in human gastric cancer cells. Investigational New Drugs, 2018, 36, 970-984.	1.2	14
21	Synthesis of unsymmetrical disulfanes bearing 1,2,4-triazine scaffold and their in vitro screening towards anti-breast cancer activity. Monatshefte Für Chemie, 2018, 149, 1409-1420.	0.9	24
22	Erythropoietin Intensifies the Proapoptotic Activity of LFM-A13 in Cells and in a Mouse Model of Colorectal Cancer. International Journal of Molecular Sciences, 2018, 19, 1262.	1.8	5
23	A novel series of pyrazole-platinum(II) complexes as potential anti-cancer agents that induce cell cycle arrest and apoptosis in breast cancer cells. Journal of Enzyme Inhibition and Medicinal Chemistry, 2018, 33, 1006-1023.	2.5	50
24	Dual Antibacterial and Anticancer Activity of 4-Benzoyl-1-dichlorobenzoylthiosemicarbazide Derivatives. Anti-Cancer Agents in Medicinal Chemistry, 2018, 18, 529-540.	0.9	8
25	Synergistic action of cisplatin and echistatin in MDA-MB-231 breast cancer cells. Molecular and Cellular Biochemistry, 2017, 427, 13-22.	1.4	20
26	Biological evaluation of octahydropyrazin[2,1-a:5,4-a′]diisoquinoline derivatives as potent anticancer agents. Tumor Biology, 2017, 39, 101042831770164.	0.8	7
27	Mechanism of anticancer action of novel berenil complex of platinum(II) combined with anti-MUC1 in MCF-7 breast cancer cells. Oncology Letters, 2017, 15, 2340-2348.	0.8	9
28	Anticancer Effect of a Novel Octahydropyrazino[2,1-a:5,4-aâ€2]diisoquinoline Derivative and Its Synergistic Action with <i> Nigella sativa</i> in Human Gastric Cancer Cells. BioMed Research International, 2017, 2017, 1-13.	0.9	9
29	Synthesis and antimicrobial activity of chiral quaternary N -spiro ammonium bromides with 3',4'-dihydro-1'H-spiro[isoindoline-2,2'-isoquinoline] skeleton. Drug Design, Development and Therapy, 2017, Volume 11, 2015-2028.	2.0	3
30	Biological evaluation of dimethylpyridine–platinum complexes with potent antiproliferative activity. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 150-165.	2.5	20
31	Effect of dentine bondings in combination with peroxide bleaching agents on the biosynthesis of DNA in human gingival fibroblasts. Journal of Stomatology, 2016, 69, 153-161.	0.1	0
32	Synthetic Approaches for Sulfur Derivatives Containing 1,2,4-Triazine Moiety: Their Activity for <i>in Vitro</i> Screening towards Two Human Cancer Cell Lines. Chemical and Pharmaceutical Bulletin, 2015, 63, 531-537.	0.6	16
33	New pyrazolo[4,3-e][1,2,4]triazine sulfonamides as carbonic anhydrase inhibitors. Bioorganic and Medicinal Chemistry, 2015, 23, 3674-3680.	1.4	36
34	The combined treatment with novel platinum(II) complex and anti-MUC1 increases apoptotic response in MDA-MB-231 breast cancer cells. Molecular and Cellular Biochemistry, 2015, 408, 103-113.	1.4	20
35	Search for human DNA topoisomerase II poisons in the group of 2,5-disubstituted-1,3,4-thiadiazoles. Journal of Enzyme Inhibition and Medicinal Chemistry, 2015, 30, 1021-1026.	2.5	13
36	Cytotoxic activity of octahydropyrazin[2,1-a:5,4-a′]diisoquinoline derivatives in human breast cancer cells. Archives of Pharmacal Research, 2015, 38, 628-641.	2.7	16

KRZYSZTOF BIELAWSKI

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37	Effects of Novel Alkyl Pyridine Platinum Complexes on Apoptosis in Ishikawa Endometrial Cancer Cells. Medicinal Chemistry, 2015, 11, 540-550.	0.7	11
38	Mucin levels in saliva of adolescents with dental caries. Medical Science Monitor, 2014, 20, 72-77.	0.5	36
39	Pyrazolo[4,3-e][1,2,4]triazine sulfonamides as carbonic anhydrase inhibitors with antitumor activity. Bioorganic and Medicinal Chemistry, 2014, 22, 2643-2647.	1.4	36
40	Synthesis and kinase inhibitory activity of new sulfonamide derivatives of pyrazolo[4,3-e][1,2,4]triazines. European Journal of Medicinal Chemistry, 2014, 78, 217-224.	2.6	27
41	Cytotoxic efficacy of a novel dinuclear platinum(II) complex used with anti-MUC1 in human breast cancer cells. Molecular and Cellular Biochemistry, 2014, 392, 161-174.	1.4	20
42	Cytotoxicity and topoisomerase I/II inhibition activity of novel 4-aryl/alkyl-1-(piperidin-4-yl)-carbonylthiosemicarbazides and 4-benzoylthiosemicarbazides. Journal of Enzyme Inhibition and Medicinal Chemistry, 2014, 29, 243-248.	2.5	7
43	The assessment of sIgA, histatin-5, and lactoperoxidase levels in saliva of adolescents with dental caries. Medical Science Monitor, 2014, 20, 1095-1100.	0.5	29
44	Cytotoxic effect and molecular docking of 4-ethoxycarbonylmethyl-1-(piperidin-4-ylcarbonyl)-thiosemicarbazide—a novel topoisomerase II inhibitor. Journal of Molecular Modeling, 2013, 19, 1319-1324.	0.8	13
45	C2-Symmetric hemiaminal ethers and diamines: new ligands for copper-catalyzed desymmetrization of meso-1,2-diols and asymmetric Henry reactions. Tetrahedron: Asymmetry, 2013, 24, 1435-1442.	1.8	23
46	Cytotoxicity and induction of apoptosis of human breast cancer cells by novel platinum(II) complexes. Environmental Toxicology and Pharmacology, 2013, 35, 254-264.	2.0	22
47	Effect of novel dinuclear platinum(II) complexes on redox status of MOLT-4 leukemic cells. Toxicology Mechanisms and Methods, 2013, 23, 641-649.	1.3	5
48	Pro-inflammatory cytokines in saliva of adolescents with dental caries disease. Annals of Agricultural and Environmental Medicine, 2012, 19, 711-6.	0.5	50
49	Cytotoxic activity of G3 PAMAM-NH2 dendrimer-chlorambucil conjugate in human breast cancer cells. Environmental Toxicology and Pharmacology, 2011, 32, 364-372.	2.0	42
50	Cytotoxic efficacy of a novel dinuclear platinum(II) complex in human breast cancer cells. European Journal of Pharmacology, 2010, 643, 34-41.	1.7	20
51	Dual effects of ouabain, digoxin and proscillaridin A on the regulation of apoptosis in human fibroblasts. Natural Product Research, 2010, 24, 274-285.	1.0	44
52	Synthesis and cytotoxic activity of G3 PAMAM-NH2 dendrimer-modified digoxin and proscillaridin A conjugates in breast cancer cells. Pharmacological Reports, 2010, 62, 414-423.	1.5	23
53	Novel dinuclear platinum(II) complexes targets NFkappaB signaling pathway to induce apoptosis and inhibit metabolism of MCF-7 breast cancer cells Folia Histochemica Et Cytobiologica, 2010, 47, S141-6.	0.6	6
54	The effect of a novel dinuclear platinum complex with berenil and 2-picoline ligands on growth of human breast cancer cells. Acta Poloniae Pharmaceutica, 2010, 67, 609-14.	0.3	8

KRZYSZTOF BIELAWSKI

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55	Synthesis and Cytotoxic Activity of Novel Amidine Analogues of Bis(2â€chloroethyl)amine. Archiv Der Pharmazie, 2009, 342, 484-490.	2.1	9
56	Proline Analogue of Nitrosourea as a New Cytotoxic Prodrug. Archiv Der Pharmazie, 2009, 342, 632-639.	2.1	1
57	The Effect of Generation 2 and 3 Poly(amidoamine) Dendrimers on Viability of Human Breast Cancer Cells. Journal of Health Science, 2009, 55, 169-177.	0.9	33
58	Smallâ€Molecule based Delivery Systems for Alkylating Antineoplastic Compounds. ChemMedChem, 2008, 3, 536-542.	1.6	22
59	Antiproliferative Activity of Derivatives of Ouabain, Digoxin and Proscillaridin A in Human MCF-7 and MDA-MB-231 Breast Cancer Cells. Biological and Pharmaceutical Bulletin, 2008, 31, 1131-1140.	0.6	72
60	Proline-linked nitrosoureas as prolidase-convertible prodrugs in human breast cancer cells. Pharmacological Reports, 2008, 60, 171-82.	1.5	11
61	Synthesis, DNA-binding affinity and cytotoxicity of the dinuclear platinum(II) complexes with berenil and amines ligands. Acta Poloniae Pharmaceutica, 2008, 65, 363-70.	0.3	5
62	Amidine Analogues of Melphalan: Synthesis, Cytotoxic Activity, and DNA Binding Properties. Archiv Der Pharmazie, 2007, 340, 251-257.	2.1	12
63	Apoptosis-mediated cytotoxicity of ouabain, digoxin and proscillaridin A in the estrogen independent MDA-MB-231 breast cancer cells. Archives of Pharmacal Research, 2007, 30, 1216-1224.	2.7	46
64	Inhibition of DNA Topoisomerases I and II, and Growth Inhibition of Breast Cancer MCF-7 Cells by Ouabain, Digoxin and Proscillaridin A. Biological and Pharmaceutical Bulletin, 2006, 29, 1493-1497.	0.6	109
65	Novel amidine analogue of melphalan as a specific multifunctional inhibitor of growth and metabolism of human breast cancer cells. Biochemical Pharmacology, 2006, 72, 320-331.	2.0	20
66	Cardiac glycosides in cancer research and cancer therapy. Acta Poloniae Pharmaceutica, 2006, 63, 109-15.	0.3	46
67	Synthesis, DNA Binding, Topoisomerase Inhibition and Cytotoxic Properties of 2-Chloroethylnitrosourea Derivatives of Hoechst 33258. Biological and Pharmaceutical Bulletin, 2005, 28, 1004-1009.	0.6	26
68	Inhibition of collagen and DNA biosynthesis by a novel amidine analogue of chlorambucil is accompanied by deregulation of β1-integrin and IGF-I receptor signaling in MDA-MB 231 cells. Environmental Toxicology and Pharmacology, 2005, 20, 118-124.	2.0	52
69	Amidine analogue of chlorambucil is a stronger inhibitor of protein and DNA synthesis in breast cancer MCF-7 cells than is the parent drug. European Journal of Pharmacology, 2004, 492, 95-101.	1.7	14
70	Synthesis and biological evaluation of new cyclic amidine analogs of chlorambucil. Il Farmaco, 2004, 59, 111-117.	0.9	28
71	Acetylsalicylic acid as a potential regulator of prolidase-convertible pro-drugs in control and neoplastic cells. Il Farmaco, 2004, 59, 679-684.	0.9	3
72	Synthesis and Biological Evaluation of New Cyclic Amidine Analogues of Chlorambucil ChemInform, 2004, 35, no.	0.1	0

KRZYSZTOF BIELAWSKI

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73	Aromatic analogues of DNA minor groove binders—synthesis and biological evaluation. European Journal of Medicinal Chemistry, 2004, 39, 99-105.	2.6	16
74	Structure-Activity Studies of Novel Amidine Analogues of Chlorambucil: Correlation of Cytotoxic Activity with DNA-Binding Affinity and Topoisomerase II Inhibition. Archiv Der Pharmazie, 2003, 336, 293-299.	2.1	14
75	Synthesis, DNA-Binding Activity and Cytotoxicity of Carbamate Derivatives of Hoechst 33258 in Breast Cancer MCF-7 Cells Biological and Pharmaceutical Bulletin, 2002, 25, 916-919.	0.6	11
76	Carbocyclic Analogues of Netropsin and Distamycin: DNA-Binding Properties and Inhibition of DNA Topoisomerases. Archiv Der Pharmazie, 2002, 335, 422-426.	2.1	12
77	Elongation factor 2 as a target for selective inhibition of protein synthesis in vitro by the novel aromatic bisamidine. Molecular and Cellular Biochemistry, 2002, 233, 159-164.	1.4	4
78	DNA-Binding Activity and Cytotoxicity of the Extended Diphenylfuran Bisamidines in Breast Cancer MCF-7 Cells Biological and Pharmaceutical Bulletin, 2001, 24, 704-706.	0.6	16
79	Synthesis, molecular modelling, and antiproliferative and cytotoxic effects of carbocyclic derivatives of distamycin with chlorambucil moiety. European Journal of Medicinal Chemistry, 2001, 36, 461-467.	2.6	23
80	Aromatic Extended Bisamidines: Synthesis, Inhibition of Topoisomerases, and Anticancer Cytotoxicity in Vitro. Archiv Der Pharmazie, 2001, 334, 235-240.	2.1	9
81	Cytotoxicity and effect on collagen biosynthesis of proline analogue of melphalan as a prolidase-convertible prodrug in cultured human skin fibroblasts. Il Farmaco, 2001, 56, 701-706.	0.9	7
82	Prolidase-activated prodrug for cancer chemotherapy. Il Farmaco, 2000, 55, 736-741.	0.9	9