

Dan Gibson

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3176465/publications.pdf>

Version: 2024-02-01

65
papers

4,356
citations

101543

36
h-index

106344

65
g-index

67
all docs

67
docs citations

67
times ranked

3298
citing authors

#	ARTICLE	IF	CITATIONS
1	Study of the DNA binding mechanism and <i>in vitro</i> activity against cancer cells of iron(III) and aluminium(III) kojic acid derivative complexes. Dalton Transactions, 2022, , .	3.3	2
2	Platinum(IV) anticancer agents; are we en route to the holy grail or to a dead end?. Journal of Inorganic Biochemistry, 2021, 217, 111353.	3.5	70
3	Pt(IV) Anticancer Prodrugs – A Tale of Mice and Men. ChemMedChem, 2021, 16, 2188-2191.	3.2	19
4	Are Pt(IV) Prodrugs That Release Combretastatin A4 True Multi-action Prodrugs?. Journal of Medicinal Chemistry, 2021, 64, 11364-11378.	6.4	30
5	Dibenzofuran annulated 1-azepines: Synthesis and cytotoxicity. Synthetic Communications, 2020, 50, 438-445.	2.1	2
6	Platinum(IV)-Estramustine Multiaction Prodrugs Are Effective Antiproliferative Agents against Prostate Cancer Cells. Journal of Medicinal Chemistry, 2020, 63, 13861-13877.	6.4	30
7	<i>trans</i> -Platinum(IV) pro-drugs that exhibit unusual resistance to reduction by endogenous reductants and blood serum but are rapidly activated inside cells: ¹ H NMR and XANES spectroscopy study. Dalton Transactions, 2020, 49, 7722-7736.	3.3	21
8	Oxidation of <i>cis</i> -Diamminediacetato Pt(II) with Hydrogen Peroxide Can Give Rise to Two Isomeric Pt(IV) Products. Chemistry - A European Journal, 2020, 26, 9475-9480.	3.3	5
9	Multiaction Pt(IV) Carbamate Complexes Can Codeliver Pt(II) Drugs and Amine Containing Bioactive Molecules. Inorganic Chemistry, 2020, 59, 5182-5193.	4.0	37
10	A Multi-action and Multi-target Ru(II)-Pt(IV) Conjugate Combining Cancer-Activated Chemotherapy and Photodynamic Therapy to Overcome Drug Resistant Cancers. Angewandte Chemie - International Edition, 2020, 59, 7069-7075.	13.8	172
11	A Multi-action and Multi-target Ru(II)-Pt(IV) Conjugate Combining Cancer-Activated Chemotherapy and Photodynamic Therapy to Overcome Drug Resistant Cancers. Angewandte Chemie, 2020, 132, 7135-7141.	2.0	25
12	Expanding the Arsenal of Pt(IV) Anticancer Agents: Multi-action Pt(IV) Anticancer Agents with Bioactive Ligands Possessing a Hydroxy Functional Group. Angewandte Chemie, 2019, 131, 18386-18391.	2.0	11
13	Expanding the Arsenal of Pt(IV) Anticancer Agents: Multi-action Pt(IV) Anticancer Agents with Bioactive Ligands Possessing a Hydroxy Functional Group. Angewandte Chemie - International Edition, 2019, 58, 18218-18223.	13.8	47
14	A Subset of New Platinum Antitumor Agents Kills Cells by a Multimodal Mechanism of Action Also Involving Changes in the Organization of the Microtubule Cytoskeleton. Journal of Medicinal Chemistry, 2019, 62, 5176-5190.	6.4	48
15	Dual-Targeting Dual-Action Platinum(IV) Platform for Enhanced Anticancer Activity and Reduced Nephrotoxicity. Angewandte Chemie, 2019, 131, 8193-8198.	2.0	24
16	Dual-Targeting Dual-Action Platinum(IV) Platform for Enhanced Anticancer Activity and Reduced Nephrotoxicity. Angewandte Chemie - International Edition, 2019, 58, 8109-8114.	13.8	81
17	An Anticancer Pt(IV) Prodrug That Acts by Mechanisms Involving DNA Damage and Different Epigenetic Effects. Chemistry - A European Journal, 2019, 25, 5235-5245.	3.3	31
18	Synthesis and Cytotoxicity of Water-Soluble Dual- and Triple-Action Satraplatin Derivatives: Replacement of Equatorial Chlorides of Satraplatin by Acetates. Inorganic Chemistry, 2019, 58, 16676-16688.	4.0	13

#	ARTICLE	IF	CITATIONS
19	Multi-action Pt(IV) anticancer agents; do we understand how they work?. <i>Journal of Inorganic Biochemistry</i> , 2019, 191, 77-84.	3.5	124
20	Triple action Pt(IV) derivatives of cisplatin: a new class of potent anticancer agents that overcome resistance. <i>Chemical Science</i> , 2018, 9, 4299-4307.	7.4	121
21	Probing the Interactions of Cytotoxic [Pt(1 <i>S</i> ,2 <i>S</i> -DACH)(5,6-dimethyl-1,10-phenanthroline)] and Its Pt ^{IV} Derivatives with Human Serum. <i>ChemMedChem</i> , 2017, 12, 510-519.	3.2	8
22	Synthesis, characterization and in vitro and in vivo anticancer activity of Pt(IV) derivatives of [Pt(1 <i>S</i> ,2 <i>S</i> -DACH)(5,6-dimethyl-1,10-phenanthroline)]. <i>Dalton Transactions</i> , 2017, 46, 7005-7019.	3.3	43
23	Epigenetic and antitumor effects of platinum(IV)-octanoato conjugates. <i>Scientific Reports</i> , 2017, 7, 3751.	3.3	38
24	The timing of caffeic acid treatment with cisplatin determines sensitization or resistance of ovarian carcinoma cell lines. <i>Redox Biology</i> , 2017, 11, 170-175.	9.0	34
25	A Quadruple-Action Platinum(IV) Prodrug with Anticancer Activity Against KRAS Mutated Cancer Cell Lines. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 11539-11544.	13.8	100
26	A Quadruple-Action Platinum(IV) Prodrug with Anticancer Activity Against KRAS Mutated Cancer Cell Lines. <i>Angewandte Chemie</i> , 2017, 129, 11697-11702.	2.0	22
27	Platinum(IV) anticancer prodrugs – hypotheses and facts. <i>Dalton Transactions</i> , 2016, 45, 12983-12991.	3.3	230
28	A Lipophilic Pt(IV) Oxaliplatin Derivative Enhances Antitumor Activity. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 9035-9046.	6.4	59
29	Potential of mitochondrial dysfunction in tumor cells by conjugates of metabolic modulator dichloroacetate with a Pt(IV) derivative of oxaliplatin. <i>Journal of Inorganic Biochemistry</i> , 2016, 156, 89-97.	3.5	60
30	Pt(IV) derivatives of cisplatin and oxaliplatin with phenylbutyrate axial ligands are potent cytotoxic agents that act by several mechanisms of action. <i>Chemical Science</i> , 2016, 7, 2381-2391.	7.4	155
31	On the Stability of Pt ^{IV} Pro-Drugs with Haloacetato Ligands in the Axial Positions. <i>Chemistry - A European Journal</i> , 2015, 21, 3108-3114.	3.3	45
32	The role of the catecholic and the electrophilic moieties of caffeic acid in Nrf2/Keap1 pathway activation in ovarian carcinoma cell lines. <i>Redox Biology</i> , 2015, 4, 48-59.	9.0	55
33	New insights into the molecular and epigenetic effects of antitumor Pt(IV)-valproic acid conjugates in human ovarian cancer cells. <i>Biochemical Pharmacology</i> , 2015, 95, 133-144.	4.4	78
34	Antitumor platinum(IV) derivatives of oxaliplatin with axial valproato ligands. <i>Journal of Inorganic Biochemistry</i> , 2014, 140, 72-79.	3.5	69
35	In vivo biodistribution of platinum-based drugs encapsulated into multi-walled carbon nanotubes. <i>Nanomedicine: Nanotechnology, Biology, and Medicine</i> , 2014, 10, 1465-1475.	3.3	56
36	Activation of trans geometry in bifunctional mononuclear platinum complexes by a non-bulky methylamine ligand. <i>Journal of Inorganic Biochemistry</i> , 2013, 126, 46-54.	3.5	6

#	ARTICLE	IF	CITATIONS
37	Facile Preparation of Mono-, Di- and Mixed-Carboxylato Platinum(IV) Complexes for Versatile Anticancer Prodrug Design. <i>Chemistry - A European Journal</i> , 2013, 19, 1672-1676.	3.3	108
38	Platinum(IV) Prodrugs with Haloacetato Ligands in the Axial Positions can Undergo Hydrolysis under Biologically Relevant Conditions. <i>Angewandte Chemie - International Edition</i> , 2013, 52, 6059-6062.	13.8	80
39	Toxicity in tumor cells, DNA binding mode, and resistance to decomposition by sulfur nucleophiles of new dinuclear bifunctional trans-Pt(II) complexes containing long alkane linkers. <i>Pure and Applied Chemistry</i> , 2012, 85, 343-354.	1.9	5
40	Cellular interactions of platinum drugs. <i>Inorganica Chimica Acta</i> , 2012, 393, 75-83.	2.4	60
41	What do we know about the reduction of Pt(IV) pro-drugs?. <i>Journal of Inorganic Biochemistry</i> , 2012, 117, 220-229.	3.5	307
42	Pt(IV) analogs of oxaliplatin that do not follow the expected correlation between electrochemical reduction potential and rate of reduction by ascorbate. <i>Chemical Communications</i> , 2012, 48, 847-849.	4.1	174
43	New reduction pathways for $\text{cis-}[\text{PtCl}_2(\text{CH}_3\text{CO})_2(\text{NH}_3)_2](\text{Am})$ anticancer prodrugs. <i>Chemical Communications</i> , 2010, 46, 1842-1844.	4.1	76
44	Studies on Cellular Accumulation of Satraplatin and Its Major Metabolite JM118 and Their Interactions with Glutathione. <i>Molecular Pharmaceutics</i> , 2010, 7, 2093-2102.	4.6	27
45	Trans labilization of am(m)ine ligands from platinum(II) complexes by cancer cell extracts. <i>Journal of Biological Inorganic Chemistry</i> , 2009, 14, 387-399.	2.6	28
46	Is Glutathione the Major Cellular Target of Cisplatin? A Study of the Interactions of Cisplatin with Cancer Cell Extracts. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 4319-4328.	6.4	125
47	The mechanism of action of platinum anticancer agents—what do we really know about it?. <i>Dalton Transactions</i> , 2009, , 10681.	3.3	169
48	Peculiar mechanistic and structural features of the carboplatin—cytochrome c system revealed by ESI-MS analysis. <i>Journal of Biological Inorganic Chemistry</i> , 2008, 13, 755-764.	2.6	35
49	Reduction of $\text{cis,trans-}[\text{PtCl}_2(\text{OCOCH}_3)_2(\text{NH}_3)_2]$ by Aqueous Extracts of Cancer Cells. <i>Journal of Medicinal Chemistry</i> , 2007, 50, 5554-5556.	6.4	100
50	Cationic Nonsymmetric Transplatinum Complexes with Piperidinopiperidine Ligands. Preparation, Characterization, in Vitro Cytotoxicity, in Vivo Toxicity, and Anticancer Efficacy Studies. <i>Journal of Medicinal Chemistry</i> , 2006, 49, 4665-4673.	6.4	32
51	The trans labilization of $\text{cis-}[\text{PtCl}_2(13\text{CH}_3\text{NH}_2)_2]$ by glutathione can be monitored at physiological pH by $[1\text{H},13\text{C}]$ HSQC NMR. <i>Journal of Biological Inorganic Chemistry</i> , 2006, 11, 179-188.	2.6	20
52	Mass spectrometric studies of the formation and reactivity of $\text{trans-}[\text{PtCl}_2(\text{Am})(\text{piperidinopiperidine})]\text{H}^+\text{Cl}^-$ complexes with ubiquitin. <i>Rapid Communications in Mass Spectrometry</i> , 2005, 19, 3666-3672.	1.5	9
53	Ligand effects on the binding of cis- and $\text{trans-}[\text{PtCl}_2\text{Am}_1\text{Am}_2]$ to proteins. <i>Journal of Biological Inorganic Chemistry</i> , 2003, 8, 167-175.	2.6	35
54	Drug—DNA interactions and novel drug design. <i>Pharmacogenomics Journal</i> , 2002, 2, 275-276.	2.0	44

#	ARTICLE	IF	CITATIONS
55	Novel Soluble Cationic trans-Diaminedichloroplatinum(II) Complexes that Are Active against Cisplatin Resistant Ovarian Cancer Cell Lines. <i>Journal of Medicinal Chemistry</i> , 2002, 45, 5189-5195.	6.4	104
56	Novel Apoptosis-Inducing trans-Platinum Piperidine Derivatives: Synthesis and Biological Characterization. <i>Journal of Medicinal Chemistry</i> , 2002, 45, 5196-5204.	6.4	64
57	Interactions of cisplatin and transplatin with proteins. <i>Journal of Inorganic Biochemistry</i> , 2002, 91, 306-311.	3.5	118
58	Cisplatin-Protein Adducts Are Efficiently Removed by Glutathione but Not by 5'-Guanosine Monophosphate. <i>Journal of the American Chemical Society</i> , 2001, 123, 3171-3172.	13.7	69
59	A mass spectral study of the binding of the anticancer drug cisplatin to ubiquitin. <i>European Journal of Mass Spectrometry</i> , 1999, 5, 501.	0.7	77
60	Monofunctional platinum amine complexes destabilize DNA significantly. <i>FEBS Journal</i> , 1998, 256, 253-260.	0.2	35
61	Anthraquinone intercalators as carrier molecules for second-generation platinum anticancer drugs. <i>European Journal of Medicinal Chemistry</i> , 1997, 32, 823-831.	5.5	27
62	Acylphosphonamidates and α -hydroxyiminophosphonamidates. Synthesis of N-acylphosphordiamidates by Beckmann rearrangement. Crystal structure of (E)- α -hydroxyiminobenzyl-1-pyrrolidinylphosphinate. <i>Heteroatom Chemistry</i> , 1996, 7, 515-520.	0.7	14
63	Rearrangement and Fragmentation Reactions of α -Hydroxyiminophosphinates. On the Nature of the Metaphosphonate Intermediate Involved in Phosphonylations by α -Hydroxyiminophosphinate. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 1990, 49-50, 81-84.	1.6	6
64	Structure and Reactivity of 2-Hydroxyiminobenzyl-2-oxo-4,4,5,5-tetramethyl[1,3,2]dioxaphospholanes. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 1989, 41, 433-437.	1.6	4
65	X-ray structure of the major adduct of the anticancer drug cisplatin with DNA: cis-[Pt(NH ₃) ₂ (d(pGpG))]. <i>Science</i> , 1985, 230, 412-417.	12.6	410