

# Patric J Jansson

## List of Publications by Year in descending order

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Version: 2024-02-01

81  
papers

11,100  
citations

76196

40  
h-index

62479

80  
g-index

82  
all docs

82  
docs citations

82  
times ranked

20664  
citing authors

#	ARTICLE	IF	CITATIONS
1	Guidelines for the use and interpretation of assays for monitoring autophagy (3rd edition). <i>Autophagy</i> , 2016, 12, 1-222.	4.3	4,701
2	Guidelines for the use and interpretation of assays for monitoring autophagy (4th) <i>Autophagy</i> , 2016, 12, 1-222.	4.3	1,430
3	Thiosemicarbazones from the Old to New: Iron Chelators That Are More Than Just Ribonucleotide Reductase Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 5271-5294.	2.9	338
4	Cellular iron uptake, trafficking and metabolism: Key molecules and mechanisms and their roles in disease. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2015, 1853, 1130-1144.	1.9	275
5	Antitumor Activity of Metal-Chelating Compound Dp44mT Is Mediated by Formation of a Redox-Active Copper Complex That Accumulates in Lysosomes. <i>Cancer Research</i> , 2011, 71, 5871-5880.	0.4	258
6	Cancer cell iron metabolism and the development of potent iron chelators as anti-tumour agents. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2009, 1790, 702-717.	1.1	214
7	Novel Thiosemicarbazones of the ApT and DpT Series and Their Copper Complexes: Identification of Pronounced Redox Activity and Characterization of Their Antitumor Activity. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 5759-5769.	2.9	205
8	Molecular Pharmacology of ABCG2 and Its Role in Chemoresistance. <i>Molecular Pharmacology</i> , 2013, 84, 655-669.	1.0	180
9	Novel Second-Generation Di-2-Pyridylketone Thiosemicarbazones Show Synergism with Standard Chemotherapeutics and Demonstrate Potent Activity against Lung Cancer Xenografts after Oral and Intravenous Administration in Vivo. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 7230-7244.	2.9	165
10	P-glycoprotein Mediates Drug Resistance via a Novel Mechanism Involving Lysosomal Sequestration. <i>Journal of Biological Chemistry</i> , 2013, 288, 31761-31771.	1.6	164
11	Zinc(II) Thiosemicarbazone Complexes Are Localized to the Lysosomal Compartment Where They Transmetallate with Copper Ions to Induce Cytotoxicity. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 4965-4984.	2.9	148
12	Roads to melanoma: Key pathways and emerging players in melanoma progression and oncogenic signaling. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2016, 1863, 770-784.	1.9	148
13	The old and new biochemistry of polyamines. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2018, 1862, 2053-2068.	1.1	145
14	The renaissance of polypharmacology in the development of anti-cancer therapeutics: Inhibition of the "Triad of Death" in cancer by Di-2-pyridylketone thiosemicarbazones. <i>Pharmacological Research</i> , 2015, 100, 255-260.	3.1	127
15	Metastasis suppressor, NDRG1, mediates its activity through signaling pathways and molecular motors. <i>Carcinogenesis</i> , 2013, 34, 1943-1954.	1.3	117
16	Redox cycling metals: Pedaling their roles in metabolism and their use in the development of novel therapeutics. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2016, 1863, 727-748.	1.9	111
17	The Iron Chelator, Deferasirox, as a Novel Strategy for Cancer Treatment: Oral Activity Against Human Lung Tumor Xenografts and Molecular Mechanism of Action. <i>Molecular Pharmacology</i> , 2013, 83, 179-190.	1.0	106
18	Copper and conquer: copper complexes of di-2-pyridylketone thiosemicarbazones as novel anti-cancer therapeutics. <i>Metallomics</i> , 2016, 8, 874-886.	1.0	105

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19	Di-2-pyridylketone 4,4-Dimethyl-3-thiosemicarbazone (Dp44mT) Overcomes Multidrug Resistance by a Novel Mechanism Involving the Hijacking of Lysosomal P-Glycoprotein (Pgp). <i>Journal of Biological Chemistry</i> , 2015, 290, 9588-9603.	1.6	103
20	Molecular functions of the iron-regulated metastasis suppressor, NDRG1, and its potential as a molecular target for cancer therapy. <i>Biochimica Et Biophysica Acta: Reviews on Cancer</i> , 2014, 1845, 1-19.	3.3	88
21	The Metastasis Suppressor, N-myc Downstream-regulated Gene 1 (NDRG1), Inhibits Stress-induced Autophagy in Cancer Cells. <i>Journal of Biological Chemistry</i> , 2014, 289, 9692-9709.	1.6	83
22	Structure-Activity Relationships of Di-2-pyridylketone, 2-Benzoylpyridine, and 2-Acetylpyridine Thiosemicarbazones for Overcoming Pgp-Mediated Drug Resistance. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 8601-8620.	2.9	82
23	The role of NDRG1 in the pathology and potential treatment of human cancers. <i>Journal of Clinical Pathology</i> , 2013, 66, 911-917.	1.0	72
24	A mechanism for overcoming P-glycoprotein-mediated drug resistance: novel combination therapy that releases stored doxorubicin from lysosomes via lysosomal permeabilization using Dp44mT or DpC. <i>Cell Death and Disease</i> , 2016, 7, e2510-e2510.	2.7	72
25	Deferasirox (ICL670A) effectively inhibits oesophageal cancer growth <i>in vitro</i> and <i>in vivo</i> . <i>British Journal of Pharmacology</i> , 2013, 168, 1316-1328.	2.7	68
26	The proto-oncogene c-Src and its downstream signaling pathways are inhibited by the metastasis suppressor, NDRG1. <i>Oncotarget</i> , 2015, 6, 8851-8874.	0.8	64
27	Identification of differential anti-neoplastic activity of copper bis(thiosemicarbazones) that is mediated by intracellular reactive oxygen species generation and lysosomal membrane permeabilization. <i>Journal of Inorganic Biochemistry</i> , 2015, 152, 20-37.	1.5	64
28	Glucose modulation induces reactive oxygen species and increases P-glycoprotein-mediated multidrug resistance to chemotherapeutics. <i>British Journal of Pharmacology</i> , 2015, 172, 2557-2572.	2.7	63
29	Molecular and Functional Alterations in a Mouse Cardiac Model of Friedreich Ataxia. <i>American Journal of Pathology</i> , 2013, 183, 745-757.	1.9	62
30	The iron complex of Dp44mT is redox-active and induces hydroxyl radical formation: An EPR study. <i>Journal of Inorganic Biochemistry</i> , 2010, 104, 1224-1228.	1.5	59
31	The Anticancer Agent Di-2-pyridylketone 4,4-Dimethyl-3-thiosemicarbazone (Dp44mT) Overcomes Prosurvival Autophagy by Two Mechanisms. <i>Journal of Biological Chemistry</i> , 2014, 289, 33568-33589.	1.6	59
32	Gene of the month: BECN1. <i>Journal of Clinical Pathology</i> , 2014, 67, 656-660.	1.0	57
33	Alkyl Substituted 2-Benzoylpyridine Thiosemicarbazone Chelators with Potent and Selective Anti-Neoplastic Activity: Novel Ligands that Limit Methemoglobin Formation. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 357-370.	2.9	56
34	Pharmacological targeting of mitochondria in cancer stem cells: An ancient organelle at the crossroad of novel anti-cancer therapies. <i>Pharmacological Research</i> , 2019, 139, 298-313.	3.1	55
35	Methemoglobin Formation by Triapine, Di-2-pyridylketone-4,4-dimethyl-3-thiosemicarbazone (Dp44mT), and Other Anticancer Thiosemicarbazones: Identification of Novel Thiosemicarbazones and Therapeutics That Prevent This Effect. <i>Molecular Pharmacology</i> , 2012, 82, 105-114.	1.0	54
36	Turning the gun on cancer: Utilizing lysosomal P-glycoprotein as a new strategy to overcome multi-drug resistance. <i>Free Radical Biology and Medicine</i> , 2016, 96, 432-445.	1.3	52

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37	Halogenated 2- <sup>2</sup> -Benzoylpyridine Thiosemicarbazone (XBpT) Chelators with Potent and Selective Anti-Neoplastic Activity: Relationship to Intracellular Redox Activity. <i>Journal of Medicinal Chemistry</i> , 2011, 54, 6936-6948.	2.9	51
38	Glucose Modulation Induces Lysosome Formation and Increases Lysosomotropic Drug Sequestration via the P-Glycoprotein Drug Transporter. <i>Journal of Biological Chemistry</i> , 2016, 291, 3796-3820.	1.6	51
39	The molecular effect of metastasis suppressors on Src signaling and tumorigenesis: new therapeutic targets. <i>Oncotarget</i> , 2015, 6, 35522-35541.	0.8	43
40	Coupling of the polyamine and iron metabolism pathways in the regulation of proliferation: Mechanistic links to alterations in key polyamine biosynthetic and catabolic enzymes. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2018, 1864, 2793-2813.	1.8	41
41	Novel Mechanism of Cytotoxicity for the Selective Selenosemicarbazone, 2-Acetylpyridine 4,4-Dimethyl-3-selenosemicarbazone (Ap44mSe): Lysosomal Membrane Permeabilization. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 294-312.	2.9	39
42	Mitochondrial dysfunction in the neuro-degenerative and cardio-degenerative disease, Friedreich's ataxia. <i>Neurochemistry International</i> , 2018, 117, 35-48.	1.9	38
43	Mechanism of the induction of endoplasmic reticulum stress by the anti-cancer agent, di-2-pyridylketone 4,4-dimethyl-3-thiosemicarbazone (Dp44mT): Activation of PERK/eIF2 <sup>±</sup> , IRE1 <sup>±</sup> , ATF6 and calmodulin kinase. <i>Biochemical Pharmacology</i> , 2016, 109, 27-47.	2.0	36
44	Identification of differential phosphorylation and sub-cellular localization of the metastasis suppressor, NDRG1. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2018, 1864, 2644-2663.	1.8	36
45	Tumor stressors induce two mechanisms of intracellular P-glycoprotein-mediated resistance that are overcome by lysosomal-targeted thiosemicarbazones. <i>Journal of Biological Chemistry</i> , 2018, 293, 3562-3587.	1.6	36
46	Lysosomal membrane stability plays a major role in the cytotoxic activity of the anti-proliferative agent, di-2-pyridylketone 4,4-dimethyl-3-thiosemicarbazone (Dp44mT). <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2016, 1863, 1665-1681.	1.9	34
47	A Nitric Oxide Storage and Transport System That Protects Activated Macrophages from Endogenous Nitric Oxide Cytotoxicity. <i>Journal of Biological Chemistry</i> , 2016, 291, 27042-27061.	1.6	32
48	Novel chelators based on adamantane-derived semicarbazones and hydrazones that target multiple hallmarks of Alzheimer's disease. <i>Dalton Transactions</i> , 2018, 47, 7190-7205.	1.6	30
49	Tumour Microenvironment Stress Promotes the Development of Drug Resistance. <i>Antioxidants</i> , 2021, 10, 1801.	2.2	29
50	The Novel Iron Chelator, 2-Pyridylcarboxaldehyde 2-Thiophenecarboxyl Hydrazone, Reduces Catecholamine-Mediated Myocardial Toxicity. <i>Chemical Research in Toxicology</i> , 2009, 22, 208-217.	1.7	27
51	The biochemical and molecular mechanisms involved in the role of tumor micro-environment stress in development of drug resistance. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2019, 1863, 1390-1397.	1.1	26
52	Oxidative Decomposition of Vitamin C in Drinking Water. <i>Free Radical Research</i> , 2004, 38, 855-860.	1.5	25
53	Unique targeting of androgen-dependent and -independent AR signaling in prostate cancer to overcome androgen resistance. <i>FASEB Journal</i> , 2020, 34, 11511-11528.	0.2	25
54	The mechanistic role of chemically diverse metal ions in the induction of autophagy. <i>Pharmacological Research</i> , 2017, 119, 118-127.	3.1	24

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55	Measurement of Ascorbic Acid (Vitamin C) Induced Hydroxyl Radical Generation in Household Drinking Water. <i>Free Radical Research</i> , 2002, 36, 1271-1276.	1.5	23
56	Breaking the cycle: Targeting of NDRG1 to inhibit bidirectional oncogenic cross-talk between pancreatic cancer and stroma. <i>FASEB Journal</i> , 2021, 35, e21347.	0.2	23
57	Overcoming tamoxifen resistance in oestrogen receptor-positive breast cancer using the novel thiosemicarbazone anticancer agent, <sc>DpC</sc>. <i>British Journal of Pharmacology</i> , 2020, 177, 2365-2380.	2.7	21
58	Autophagy: A promising target for triple negative breast cancers. <i>Pharmacological Research</i> , 2022, 175, 106006.	3.1	20
59	Targeting Wnt/tenascin C-mediated cross talk between pancreatic cancer cells and stellate cells via activation of the metastasis suppressor NDRG1. <i>Journal of Biological Chemistry</i> , 2022, 298, 101608.	1.6	20
60	Exploiting Cancer Metal Metabolism using Anti-Cancer Metal-Binding Agents. <i>Current Medicinal Chemistry</i> , 2019, 26, 302-322.	1.2	19
61	Thiosemicarbazones suppress expression of the c-Met oncogene by mechanisms involving lysosomal degradation and intracellular shedding. <i>Journal of Biological Chemistry</i> , 2020, 295, 481-503.	1.6	18
62	Making a case for albumin – a highly promising drug-delivery system. <i>Future Medicinal Chemistry</i> , 2015, 7, 553-556.	1.1	17
63	IRON METABOLISM AND AUTOPHAGY: A POORLY EXPLORED RELATIONSHIP THAT HAS IMPORTANT CONSEQUENCES FOR HEALTH AND DISEASE. <i>Nagoya Journal of Medical Science</i> , 2015, 77, 1-6.	0.6	17
64	Bonnie and Clyde: Vitamin C and iron are partners in crime in iron deficiency anaemia and its potential role in the elderly. <i>Aging</i> , 2016, 8, 1150-1152.	1.4	16
65	Regulation of autophagy and apoptosis by Dp44mT-mediated activation of AMPK in pancreatic cancer cells. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2020, 1866, 165657.	1.8	16
66	The growing evidence for targeting P-glycoprotein in lysosomes to overcome resistance. <i>Future Medicinal Chemistry</i> , 2020, 12, 473-477.	1.1	16
67	NDRG1 suppresses basal and hypoxia-induced autophagy at both the initiation and degradation stages and sensitizes pancreatic cancer cells to lysosomal membrane permeabilization. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2020, 1864, 129625.	1.1	13
68	Role of ABCB1 in mediating chemoresistance of triple-negative breast cancers. <i>Bioscience Reports</i> , 2021, 41, .	1.1	13
69	Iron prevents ascorbic acid (vitamin C) induced hydrogen peroxide accumulation in copper contaminated drinking water. <i>Free Radical Research</i> , 2005, 39, 1233-1239.	1.5	11
70	Tumor-induced neoangiogenesis and receptor tyrosine kinases – Mechanisms and strategies for acquired resistance. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2019, 1863, 1217-1225.	1.1	9
71	Vitamin C (Ascorbic Acid) Induced Hydroxyl Radical Formation in Copper Contaminated Household Drinking Water: Role of Bicarbonate Concentration. <i>Free Radical Research</i> , 2003, 37, 901-905.	1.5	8
72	Acireductone dioxygenase 1 (ADI1) is regulated by cellular iron by a mechanism involving the iron chaperone, PCBP1, with PCBP2 acting as a potential co-chaperone. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2020, 1866, 165844.	1.8	8

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73	Vesicular ATP-binding cassette transporters in human disease: relevant aspects of their organization for future drug development. <i>Future Drug Discovery</i> , 2020, 2, .	0.8	8
74	The thiosemicarbazone, DpC, broadly synergizes with multiple anti-cancer therapeutics and demonstrates temperature- and energy-dependent uptake by tumor cells. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2022, 1866, 130152.	1.1	8
75	Effects of iron on Vitamin C/copper-induced hydroxyl radical generation in bicarbonate-rich water. <i>Free Radical Research</i> , 2005, 39, 565-570.	1.5	7
76	During mitosis ZEB1 “switches” from being a chromatin-bound epithelial gene repressor, to become a microtubule-associated protein. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2020, 1867, 118673.	1.9	6
77	The redox-active, anti-cancer drug Dp44mT inhibits T-cell activation and CD25 through a copper-dependent mechanism. <i>Redox Report</i> , 2013, 18, 48-50.	1.4	3
78	Ascorbate and Tumor Cell Iron Metabolism: The Evolving Story and Its Link to Pathology. <i>Antioxidants and Redox Signaling</i> , 2020, 33, 816-838.	2.5	3
79	Emerging Role of Autophagy in the Development and Progression of Oral Squamous Cell Carcinoma. <i>Cancers</i> , 2021, 13, 6152.	1.7	3
80	Targeting autophagy in antitumor agent design: furthering the “lysosomal love”™ strategy. <i>Future Medicinal Chemistry</i> , 2016, 8, 727-729.	1.1	0
81	NDRG1: A Novel Therapeutic Target against Metastatic Cancers. <i>International Clinical Pathology Journal</i> , 2015, 1, .	0.1	0