Sumit Sahni

List of Publications by Year
in descending order

64 papers	7,505 citations	136885 32 h-index	114418 63 g-index
65 all docs	65 docs citations	65 times ranked	17354 citing authors

#	Article	IF	Citations
1	Autophagy: A promising target for triple negative breast cancers. Pharmacological Research, 2022, 175, 106006.	3.1	20
2	Targeting Wnt/tenascin C-mediated cross talk between pancreatic cancer cells and stellate cells via activation ofÂtheÂmetastasis suppressor NDRG1. Journal of Biological Chemistry, 2022, 298, 101608.	1.6	20
3	Data independent acquisition of plasma biomarkers of response to neoadjuvant chemotherapy in pancreatic ductal adenocarcinoma. Journal of Proteomics, 2021, 231, 103998.	1.2	10
4	A unique urinary metabolomic signature for the detection of pancreatic ductal adenocarcinoma. International Journal of Cancer, 2021, 148, 1508-1518.	2.3	14
5	Breaking the cycle: Targeting of NDRG1 to inhibit biâ€directional oncogenic crossâ€talk between pancreatic cancer and stroma. FASEB Journal, 2021, 35, e21347.	0.2	23
6	Role of ABCB1 in mediating chemoresistance of triple-negative breast cancers. Bioscience Reports, 2021, 41, .	1.1	13
7	Optimal Upfront Treatment in Surgically Resectable Pancreatic Cancer Candidates: A High-Volume Center Retrospective Analysis. Journal of Clinical Medicine, 2021, 10, 2700.	1.0	5
8	Serum Biomarker Panel for Diagnosis and Prognosis of Pancreatic Ductal Adenocarcinomas. Frontiers in Oncology, 2021, 11, 708963.	1.3	9
9	Urinary metabolite prognostic biomarker panel for pancreatic ductal adenocarcinomas. Biochimica Et Biophysica Acta - General Subjects, 2021, 1865, 129966.	1.1	8
10	A Critical Assessment of Postneoadjuvant Therapy Pancreatic Cancer Regression Grading Schemes With a Proposal for a Novel Approach. American Journal of Surgical Pathology, 2021, 45, 394-404.	2.1	15
11	Emerging Role of Autophagy in the Development and Progression of Oral Squamous Cell Carcinoma. Cancers, 2021, 13, 6152.	1.7	3
12	Mechanically stressed cancer microenvironment: Role in pancreatic cancer progression. Biochimica Et Biophysica Acta: Reviews on Cancer, 2020, 1874, 188418.	3.3	21
13	PSMD11, PTPRM and PTPRB as novel biomarkers of pancreatic cancer progression. Biochimica Et Biophysica Acta - General Subjects, 2020, 1864, 129682.	1.1	15
14	Small Molecule KRAS Inhibitors: The Future for Targeted Pancreatic Cancer Therapy?. Cancers, 2020, 12, 1341.	1.7	34
15	Identification of Novel Biomarkers in Pancreatic Tumor Tissue to Predict Response to Neoadjuvant Chemotherapy. Frontiers in Oncology, 2020, 10, 237.	1.3	22
16	Tissue biomarker panel as a surrogate marker for squamous subtype of pancreatic cancer. European Journal of Surgical Oncology, 2020, 46, 1539-1542.	0.5	6
17	NDRG1 suppresses basal and hypoxia-induced autophagy at both the initiation and degradation stages and sensitizes pancreatic cancer cells to lysosomal membrane permeabilization. Biochimica Et Biophysica Acta - General Subjects, 2020, 1864, 129625.	1.1	13
18	The Role of the Antioxidant Response in Mitochondrial Dysfunction in Degenerative Diseases: Cross-Talk between Antioxidant Defense, Autophagy, and Apoptosis. Oxidative Medicine and Cellular Longevity, 2019, 2019, 1-26.	1.9	92

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19	Exploiting Cancer Metal Metabolism using Anti-Cancer Metal- Binding Agents. Current Medicinal Chemistry, 2019, 26, 302-322.	1.2	19
20	Two mechanisms involving the autophagic and proteasomal pathways process the metastasis suppressor protein, N-myc downstream regulated gene 1. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2019, 1865, 1361-1378.	1.8	12
21	Nitric oxide reduces oxidative stress in cancer cells by forming dinitrosyliron complexes. Nitric Oxide - Biology and Chemistry, 2018, 76, 37-44.	1.2	36
22	Identification of differential phosphorylation and sub-cellular localization of the metastasis suppressor, NDRG1. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2018, 1864, 2644-2663.	1.8	36
23	Tumor stressors induce two mechanisms of intracellular P-glycoproteinâ€“mediated resistance that are overcome by lysosomal-targeted thiosemicarbazones. Journal of Biological Chemistry, 2018, 293, 3562-3587.	1.6	36
24	The mechanistic role of chemically diverse metal ions in the induction of autophagy. Pharmacological Research, 2017, 119, 118-127.	3.1	24
25	Interplay of the iron-regulated metastasis suppressor NDRG1 with epidermal growth factor receptor (EGFR) and oncogenic signaling. Journal of Biological Chemistry, 2017, 292, 12772-12782.	1.6	48
26	A novel class of thiosemicarbazones show multi-functional activity for the treatment of Alzheimer's disease. European Journal of Medicinal Chemistry, 2017, 139, 612-632.	2.6	64
27	Letter to the Editor: â€œAnalysis of the Interaction of Dp44mT with Human Serum Albumin and Calf Thymus DNA Using Molecular Docking and Spectroscopic Techniquesâ€. International Journal of Molecular Sciences, 2016, 17, 1916.	1.8	3
28	Targeting autophagy in antitumor agent design: furthering the â€˜lysosomal loveâ€™ strategy. Future Medicinal Chemistry, 2016, 8, 727-729.	1.1	0
29	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/eIF2Î±, IRE1Î±, ATF6 and calmodulin kinase. Biochemical Pharmacology, 2016, 109, 27-47.	2.0	36
30	Targeting the Metastasis Suppressor, N-Myc Downstream Regulated Gene-1, with Novel Di-2-Pyridylketone Thiosemicarbazones: Suppression of Tumor Cell Migration and Cell-Collagen Adhesion by Inhibiting Focal Adhesion Kinase/Paxillin Signaling. Molecular Pharmacology, 2016, 89, 521-540.	1.0	45
31	Frataxin and the molecular mechanism of mitochondrial iron-loading in Friedreich's ataxia. Clinical Science, 2016, 130, 853-870.	1.8	45
32	A Nitric Oxide Storage and Transport System That Protects Activated Macrophages from Endogenous Nitric Oxide Cytotoxicity. Journal of Biological Chemistry, 2016, 291, 27042-27061.	1.6	32
33	The Anticancer Agent, Di-2-Pyridylketone 4,4-Dimethyl-3-Thiosemicarbazone (Dp44mT), Up-Regulates the AMPK-Dependent Energy Homeostasis Pathway in Cancer Cells. Biochimica Et Biophysica Acta - Molecular Cell Research, 2016, 1863, 2916-2933.	1.9	36
34	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). Biochimica Et Biophysica Acta - Molecular Cell Research, 2016, 1863, 1665-1681.	1.9	34
35	Copper and conquer: copper complexes of di-2-pyridylketone thiosemicarbazones as novel anti-cancer therapeutics. Metallomics, 2016, 8, 874-886.	1.0	105
36	Guidelines for the use and interpretation of assays for monitoring autophagy (3rd edition). Autophagy, 2016, 12, 1-222.	4.3	4,701

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37	The Metastasis Suppressor, N-MYC Downstream-regulated Gene-1 (NDRG1), Down-regulates the ErbB Family of Receptors to Inhibit Downstream Oncogenic Signaling Pathways. Journal of Biological Chemistry, 2016, 291, 1029-1052.	1.6	65
38	Novel Mechanism of Cytotoxicity for the Selective Selenosemicarbazone, 2-Acetylpyridine 4,4-Dimethyl-3-selenosemicarbazone (Ap44mSe): Lysosomal Membrane Permeabilization. Journal of Medicinal Chemistry, 2016, 59, 294-312.	2.9	39
39	Roads to melanoma: Key pathways and emerging players in melanoma progression and oncogenic signaling. Biochimica Et Biophysica Acta - Molecular Cell Research, 2016, 1863, 770-784.	1.9	148
40	Redox cycling metals: Pedaling their roles in metabolism and their use in the development of novel therapeutics. Biochimica Et Biophysica Acta - Molecular Cell Research, 2016, 1863, 727-748.	1.9	111
41	Copper that cancer with lysosomal love!. Aging, 2016, 8, 210-211.	1.4	10
42	Duodenal Cytochrome b (DCYTB) in Iron Metabolism: An Update on Function and Regulation. Nutrients, 2015, 7, 2274-2296.	1.7	103
43	The proto-oncogene c-Src and its downstream signaling pathways are inhibited by the metastasis suppressor, NDRG1. Oncotarget, 2015, 6, 8851-8874.	0.8	64
44	Making a case for albumin â€“ a highly promising drug-delivery system. Future Medicinal Chemistry, 2015, 7, 553-556.	1.1	17
45	Novel Thiosemicarbazones Regulate the Signal Transducer and Activator of Transcription 3 (STAT3) Pathway: Inhibition of Constitutive and Interleukin 6â€“Induced Activation by Iron Depletion. Molecular Pharmacology, 2015, 87, 543-560.	1.0	37
46	The use of iron chelators in biocidal compositions: evaluation of patent, WO2014059417A1. Expert Opinion on Therapeutic Patents, 2015, 25, 367-372.	2.4	1
47	Di-2-pyridylketone 4,4-Dimethyl-3-thiosemicarbazone (Dp44mT) Overcomes Multidrug Resistance by a Novel Mechanism Involving the Hijacking of Lysosomal P-Glycoprotein (Pgp). Journal of Biological Chemistry, 2015, 290, 9588-9603.	1.6	103
48	The renaissance of polypharmacology in the development of anti-cancer therapeutics: Inhibition of the â€œTriad of Deathâ€•in cancer by Di-2-pyridylketone thiosemicarbazones. Pharmacological Research, 2015, 100, 255-260.	3.1	127
49	Adenosine Monophosphateâ€“Activated Kinase and Its Key Role in Catabolism: Structure, Regulation, Biological Activity, and Pharmacological Activation. Molecular Pharmacology, 2015, 87, 363-377.	1.0	74
50	In Vitro Characterization of the Pharmacological Properties of the Anti-Cancer Chelator, Bp4eT, and Its Phase I Metabolites. PLoS ONE, 2015, 10, e0139929.	1.1	7
51	Potentiating the cellular targeting and anti-tumor activity of Dp44mTviabinding to human serum albumin: two saturable mechanisms of Dp44mT uptake by cells. Oncotarget, 2015, 6, 10374-10398.	0.8	28
52	The molecular effect of metastasis suppressors on Src signaling and tumorigenesis: new therapeutic targets. Oncotarget, 2015, 6, 35522-35541.	0.8	43
53	IRON METABOLISM AND AUTOPHAGY: A POORLY EXPLORED RELATIONSHIP THAT HAS IMPORTANT CONSEQUENCES FOR HEALTH AND DISEASE. Nagoya Journal of Medical Science, 2015, 77, 1-6.	0.6	17
54	NDRG1 as a molecular target to inhibit the epithelialâ€“mesenchymal transition: the case for developing inhibitors of metastasis. Future Medicinal Chemistry, 2014, 6, 1241-1244.	1.1	9

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55	The Metastasis Suppressor, N-myc Downstream-regulated Gene 1 (NDRG1), Inhibits Stress-induced Autophagy in Cancer Cells. Journal of Biological Chemistry, 2014, 289, 9692-9709.	1.6	83
56	Molecular functions of the iron-regulated metastasis suppressor, NDRG1, and its potential as a molecular target for cancer therapy. Biochimica Et Biophysica Acta: Reviews on Cancer, 2014, 1845, 1-19.	3.3	88
57	Gene of the month: BECN1. Journal of Clinical Pathology, 2014, 67, 656-660.	1.0	57
58	AMP kinase (PRKAA1). Journal of Clinical Pathology, 2014, 67, 758-763.	1.0	51
59	P-glycoprotein Mediates Drug Resistance via a Novel Mechanism Involving Lysosomal Sequestration. Journal of Biological Chemistry, 2013, 288, 31761-31771.	1.6	164
60	Metastasis suppressor, NDRG1, mediates its activity through signaling pathways and molecular motors. Carcinogenesis, 2013, 34, 1943-1954.	1.3	117
61	The role of NDRG1 in the pathology and potential treatment of human cancers. Journal of Clinical Pathology, 2013, 66, 911-917.	1.0	72
62	Cellular Uptake of the Antitumor Agent Dp44mT Occurs via a Carrier/Receptor-Mediated Mechanism. Molecular Pharmacology, 2013, 84, 911-924.	1.0	19
63	Dinitrosyliron complexes are the most abundant nitric oxide-derived cellular adduct: biological parameters of assembly and disappearance. Free Radical Biology and Medicine, 2011, 51, 1558-1566.	1.3	127
64	Nitric Oxide Suppresses Tumor Cell Migration through N-Myc Downstream-regulated Gene-1 (NDRG1) Expression. Journal of Biological Chemistry, 2011, 286, 41413-41424.	1.6	69