

Claudio Festuccia

List of Publications by Year in descending order

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86
papers

2,865
citations

147801

31
h-index

214800

47
g-index

90
all docs

90
docs citations

90
times ranked

4421
citing authors

#	ARTICLE	IF	CITATIONS
1	Nucleo-cytoplasmic transport as a therapeutic target of cancer. <i>Journal of Hematology and Oncology</i> , 2014, 7, 85.	17.0	202
2	Antitumor Effects of Saffron-Derived Carotenoids in Prostate Cancer Cell Models. <i>BioMed Research International</i> , 2014, 2014, 1-12.	1.9	95
3	From glioblastoma to endothelial cells through extracellular vesicles: messages for angiogenesis. <i>Tumor Biology</i> , 2016, 37, 12743-12753.	1.8	83
4	Plasminogen activator system modulates invasive capacity and proliferation in prostatic tumor cells. <i>Clinical and Experimental Metastasis</i> , 1998, 16, 513-528.	3.3	82
5	Suppression of EGF-R signaling reduces the incidence of prostate cancer metastasis in nude mice. <i>Endocrine-Related Cancer</i> , 2006, 13, 197-210.	3.1	79
6	Osteoblast conditioned media contain TGF- β 1 and modulate the migration of prostate tumor cells and their interactions with extracellular matrix components. , 1999, 81, 395-403.		78
7	Vesicle-associated urokinase plasminogen activator promotes invasion in prostate cancer cell lines. <i>Clinical and Experimental Metastasis</i> , 2000, 18, 163-170.	3.3	74
8	HDAC4 and HDAC6 sustain DNA double strand break repair and stem-like phenotype by promoting radioresistance in glioblastoma cells. <i>Cancer Letters</i> , 2017, 397, 1-11.	7.2	72
9	Prostate cancer cell proliferation is strongly reduced by the epidermal growth factor receptor tyrosine kinase inhibitor ZD1839 in vitro on human cell lines and primary cultures. <i>Journal of Cancer Research and Clinical Oncology</i> , 2003, 129, 165-174.	2.5	71
10	Hypoxia sustains glioblastoma radioresistance through ERKs/DNA-PKcs/HIF-1 α functional interplay. <i>International Journal of Oncology</i> , 2014, 44, 2121-2131.	3.3	64
11	Azacitidine improves antitumor effects of docetaxel and cisplatin in aggressive prostate cancer models. <i>Endocrine-Related Cancer</i> , 2009, 16, 401-413.	3.1	63
12	Pyrazolo[3,4-d]pyrimidines c-Src inhibitors reduce epidermal growth factor-induced migration in prostate cancer cells. <i>European Journal of Cancer</i> , 2006, 42, 2838-2845.	2.8	62
13	Crocetin and Crocin from Saffron in Cancer Chemotherapy and Chemoprevention. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2019, 19, 38-47.	1.7	60
14	Osteoblast-derived TGF- β 1 modulates matrix degrading protease expression and activity in prostate cancer cells. <i>International Journal of Cancer</i> , 2000, 85, 407-415.	5.1	59
15	XPO1/CRM1-Selective Inhibitors of Nuclear Export (SINE) reduce tumor spreading and improve overall survival in preclinical models of prostate cancer (PCa). <i>Journal of Hematology and Oncology</i> , 2014, 7, 46.	17.0	59
16	The brain-penetrating CXCR4 antagonist, PRX177561, increases the antitumor effects of bevacizumab and sunitinib in preclinical models of human glioblastoma. <i>Journal of Hematology and Oncology</i> , 2017, 10, 5.	17.0	56
17	Cyclin D1 silencing suppresses tumorigenicity, impairs DNA double strand break repair and thus radiosensitizes androgen-independent prostate cancer cells to DNA damage. <i>Oncotarget</i> , 2016, 7, 5383-5400.	1.8	53
18	Increased matrix metalloproteinase-9 secretion in short-term tissue cultures of prostatic tumor cells. <i>International Journal of Cancer</i> , 1996, 69, 386-393.	5.1	50

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19	KPT-330, a potent and selective exportin-1 (XPO-1) inhibitor, shows antitumor effects modulating the expression of cyclin D1 and survivin in prostate cancer models. <i>BMC Cancer</i> , 2015, 15, 941.	2.6	50
20	Hormonal Therapy Promotes Hormone-Resistant Phenotype by Increasing DNMT Activity and Expression in Prostate Cancer Models. <i>Endocrinology</i> , 2011, 152, 4550-4561.	2.8	48
21	CXCR4 pharmacological inhibition reduces bone and soft tissue metastatic burden by affecting tumor growth and tumorigenic potential in prostate cancer preclinical models. <i>Prostate</i> , 2015, 75, 1227-1246.	2.3	45
22	The novel CXCR4 antagonist, PRX177561, reduces tumor cell proliferation and accelerates cancer stem cell differentiation in glioblastoma preclinical models. <i>Tumor Biology</i> , 2017, 39, 101042831769552.	1.8	44
23	PXD101 potentiates hormonal therapy and prevents the onset of castration-resistant phenotype modulating androgen receptor, HSP90, and CRM1 in preclinical models of prostate cancer. <i>Endocrine-Related Cancer</i> , 2013, 20, 321-337.	3.1	43
24	Close correlation between MEK/ERK and Aurora-B signaling pathways in sustaining tumorigenic potential and radioresistance of gynecological cancer cell lines. <i>International Journal of Oncology</i> , 2014, 44, 285-294.	3.3	43
25	The Small Molecule Ephrin Receptor Inhibitor, GLPG1790, Reduces Renewal Capabilities of Cancer Stem Cells, Showing Anti-Tumour Efficacy on Preclinical Glioblastoma Models. <i>Cancers</i> , 2019, 11, 359.	3.7	42
26	Potent and selective aldo-keto reductase 1C3 (AKR1C3) inhibitors based on the benzoisoxazole moiety: application of a bioisosteric scaffold hopping approach to flufenamic acid. <i>European Journal of Medicinal Chemistry</i> , 2018, 150, 930-945.	5.5	39
27	Î ⁵ -Cholenoyl-amino acids as selective and orally available antagonists of the Ephâ€“ephrin system. <i>European Journal of Medicinal Chemistry</i> , 2015, 103, 312-324.	5.5	38
28	Crocetin Extracted from Saffron Shows Antitumor Effects in Models of Human Glioblastoma. <i>International Journal of Molecular Sciences</i> , 2020, 21, 423.	4.1	37
29	Surgical and Biologic Outcomes After Neoadjuvant Bicalutamide Treatment in Prostate Cancer. <i>Urology</i> , 2007, 70, 728-733.	1.0	35
30	The TORC1/TORC2 inhibitor, Palomid 529, reduces tumor growth and sensitizes to docetaxel and cisplatin in aggressive and hormone-refractory prostate cancer cells. <i>Endocrine-Related Cancer</i> , 2011, 18, 385-400.	3.1	35
31	Torc1/Torc2 inhibitor, Palomid 529, enhances radiation response modulating CRM1â€“mediated survivin function and delaying DNA repair in prostate cancer models. <i>Prostate</i> , 2014, 74, 852-868.	2.3	35
32	Phenotypic characterization of human prostatic stromal cells in primary cultures derived from human tissue samples. <i>International Journal of Oncology</i> , 2013, 42, 2116-2122.	3.3	33
33	Chronic azacitidine treatment results in differentiating effects, sensitizes against bicalutamide in androgen-independent prostate cancer cells. <i>Prostate</i> , 2008, 68, 793-801.	2.3	31
34	The Importance of Tumor Stem Cells in Glioblastoma Resistance to Therapy. <i>International Journal of Molecular Sciences</i> , 2021, 22, 3863.	4.1	31
35	Bicalutamide increases phospho-Akt levels through Her2 in patients with prostate cancer. <i>Endocrine-Related Cancer</i> , 2007, 14, 601-611.	3.1	29
36	Akt downâ€“modulation induces apoptosis of human prostate cancer cells and synergizes with EGFR tyrosine kinase inhibitors. <i>Prostate</i> , 2008, 68, 965-974.	2.3	29

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37	Pharmacological targeting of the ephrin receptor kinase signalling by GLPG1790 in vitro and in vivo reverts oncophenotype, induces myogenic differentiation and radiosensitizes embryonal rhabdomyosarcoma cells. <i>Journal of Hematology and Oncology</i> , 2017, 10, 161.	17.0	29
38	NRF2 orchestrates the redox regulation induced by radiation therapy, sustaining embryonal and alveolar rhabdomyosarcoma cells radioresistance. <i>Journal of Cancer Research and Clinical Oncology</i> , 2019, 145, 881-893.	2.5	28
39	At the Bench: Pre-clinical evidence for multiple functions of CXCR4 in cancer. <i>Journal of Leukocyte Biology</i> , 2021, 109, 969-989.	3.3	28
40	UniPR1331, a small molecule targeting Eph/ephrin interaction, prolongs survival in glioblastoma and potentiates the effect of antiangiogenic therapy in mice. <i>Oncotarget</i> , 2018, 9, 24347-24363.	1.8	28
41	Tyrosine kinase inhibitor CEP-701 blocks the NTRK1/NGF receptor and limits the invasive capability of prostate cancer cells in vitro. <i>International Journal of Oncology</i> , 2007, 30, 193-200.	3.3	28
42	Differential effects of PXD101 (belinostat) on androgen-dependent and androgen-independent prostate cancer models. <i>International Journal of Oncology</i> , 2011, 40, 711-20.	3.3	27
43	Tebuconazole and Econazole Act Synergistically in Mediating Mitochondrial Stress, Energy Imbalance, and Sequential Activation of Autophagy and Apoptosis in Mouse Sertoli TM4 Cells: Possible Role of AMPK/ULK1 Axis. <i>Toxicological Sciences</i> , 2019, 169, 209-223.	3.1	25
44	Histone deacetylase inhibitor ITF2357 (givinostat) reverts transformed phenotype and counteracts stemness in in vitro and in vivo models of human glioblastoma. <i>Journal of Cancer Research and Clinical Oncology</i> , 2019, 145, 393-409.	2.5	25
45	Characterization of Prostate Cancer DU145 Cells Expressing the Recombinant Androgen Receptor. <i>Oncology Research</i> , 2003, 14, 101-112.	1.5	24
46	Effects of EGFR tyrosine kinase inhibitor erlotinib in prostate cancer cells in vitro. <i>Prostate</i> , 2009, 69, 1529-1537.	2.3	24
47	The possible prognostic role of histone deacetylase and transforming growth factor β /Smad signaling in high grade gliomas treated by radio-chemotherapy: a preliminary immunohistochemical study. <i>European Journal of Histochemistry</i> , 2017, 61, 2732.	1.5	24
48	The first-in-class alkylating deacetylase inhibitor molecule tinostamustine shows antitumor effects and is synergistic with radiotherapy in preclinical models of glioblastoma. <i>Journal of Hematology and Oncology</i> , 2018, 11, 32.	17.0	24
49	Bombesin-Dependent Pro-MMP-9 Activation in Prostatic Cancer Cells Requires β 1 Integrin Engagement. <i>Experimental Cell Research</i> , 2002, 280, 1-11.	2.6	22
50	Dual PI3K/mTOR inhibitor, XL765 (SAR245409), shows superior effects to sole PI3K [XL147 (SAR245408)] or mTOR [rapamycin] inhibition in prostate cancer cell models. <i>Tumor Biology</i> , 2016, 37, 341-351.	1.8	22
51	Pro-differentiating and radiosensitizing effects of inhibiting HDACs by PXD-101 (Belinostat) in in vitro and in vivo models of human rhabdomyosarcoma cell lines. <i>Cancer Letters</i> , 2019, 461, 90-101.	7.2	22
52	Ozarelix, a fourth generation GnRH antagonist, induces apoptosis in hormone refractory androgen receptor negative prostate cancer cells modulating expression and activity of death receptors. <i>Prostate</i> , 2010, 70, 1340-1349.	2.3	21
53	Trifluoroibuprofen Inhibits β -Methylacyl Coenzyme A Racemase (AMACR/P504S), Reduces Cancer Cell Proliferation and Inhibits in vivo Tumor Growth in Aggressive Prostate Cancer Models. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2014, 14, 1031-1041.	1.7	21
54	In vitro and in vivo effects of bicalutamide on the expression of TrkA and P75 neurotrophin receptors in prostate carcinoma. <i>Prostate</i> , 2007, 67, 1255-1264.	2.3	20

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55	Dual CXCR4 and E-Selectin Inhibitor, GMI-1359, Shows Anti-Bone Metastatic Effects and Synergizes with Docetaxel in Prostate Cancer Cell Intraosseous Growth. <i>Cells</i> , 2020, 9, 32.	4.1	19
56	Effects of Dutasteride on Prostate Carcinoma Primary Cultures: A Comparative Study With Finasteride and MK386. <i>Journal of Urology</i> , 2008, 180, 367-372.	0.4	18
57	Impregnation of Curcumin into a Biodegradable (Poly-lactic-co-glycolic acid, PLGA) Support, to Transfer Its Well Known In Vitro Effect to an In Vivo Prostate Cancer Model. <i>Nutrients</i> , 2019, 11, 2312.	4.1	18
58	Clinically relevant radioresistant rhabdomyosarcoma cell lines: functional, molecular and immune-related characterization. <i>Journal of Biomedical Science</i> , 2020, 27, 90.	7.0	18
59	The Botanical Drug PBI-05204, a Supercritical CO ₂ Extract of Nerium Oleander, Inhibits Growth of Human Glioblastoma, Reduces Akt/mTOR Activities, and Modulates GSC Cell-Renewal Properties. <i>Frontiers in Pharmacology</i> , 2020, 11, 552428.	3.5	17
60	Pharmacological treatment with inhibitors of nuclear export enhances the antitumor activity of docetaxel in human prostate cancer. <i>Oncotarget</i> , 2017, 8, 111225-111245.	1.8	16
61	MS-275 (Entinostat) Promotes Radio-Sensitivity in PAX3-FOXO1 Rhabdomyosarcoma Cells. <i>International Journal of Molecular Sciences</i> , 2021, 22, 10671.	4.1	14
62	Can the AGE/RAGE/ERK signalling pathway and the epithelial-to-mesenchymal transition interact in the pathogenesis of chronic rhinosinusitis with nasal polyps?. <i>European Journal of Histochemistry</i> , 2020, 64, .	1.5	13
63	Romidepsin (FK228) fails in counteracting the transformed phenotype of rhabdomyosarcoma cells but efficiently radiosensitizes, in <i>À</i> vitro and in <i>À</i> vivo, the alveolar phenotype subtype. <i>International Journal of Radiation Biology</i> , 2021, 97, 943-957.	1.8	13
64	Enhancement of radiosensitivity by the novel anticancer quinolone derivative vosaroxin in preclinical glioblastoma models. <i>Oncotarget</i> , 2017, 8, 29865-29886.	1.8	12
65	Evaluation of metastatic potential in prostate carcinoma: an in vivo model. <i>International Journal of Oncology</i> , 2004, 25, 1713-20.	3.3	12
66	The Brain Penetrating and Dual TORC1/TORC2 Inhibitor, RES529, Elicits Anti-Glioma Activity and Enhances the Therapeutic Effects of Anti-Angiogenetic Compounds in Preclinical Murine Models. <i>Cancers</i> , 2019, 11, 1604.	3.7	11
67	Uncoupling of the epidermal growth factor receptor from downstream signal transduction molecules guides the acquired resistance to gefitinib in prostate cancer cells. <i>Oncology Reports</i> , 2007, 18, 503-11.	2.6	11
68	Targeted Molecular Therapy in Glioblastoma. <i>Journal of Oncology</i> , 2020, 2020, 1-3.	1.3	10
69	ATX-101, a Peptide Targeting PCNA, Has Antitumor Efficacy Alone or in Combination with Radiotherapy in Murine Models of Human Glioblastoma. <i>Cancers</i> , 2022, 14, 289.	3.7	10
70	Synergistic Activity of Ketoconazole and Miconazole with Prochloraz in Inducing Oxidative Stress, GSH Depletion, Mitochondrial Dysfunction, and Apoptosis in Mouse Sertoli TM4 Cells. <i>International Journal of Molecular Sciences</i> , 2022, 23, 5429.	4.1	10
71	Her2 crosstalks with TrkA in a subset of prostate cancer cells: Rationale for a guided dual treatment. <i>Prostate</i> , 2009, 69, 337-345.	2.3	9
72	Antitumorigenic Effects of Inhibiting Ephrin Receptor Kinase Signaling by GLPG1790 against Colorectal Cancer Cell Lines <i><i>In Vitro</i></i> and <i><i>In Vivo</i></i> . <i>Journal of Oncology</i> , 2020, 2020, 1-16.	1.3	9

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73	<i>Lactobacillus sakei</i> Pro-Bio65 Reduces TNF- α Expression and Upregulates GSH Content and Antioxidant Enzymatic Activities in Human Conjunctival Cells. <i>Translational Vision Science and Technology</i> , 2021, 10, 8.	2.2	9
74	Evaluation of metastatic potential in prostate carcinoma: An in vivo model. <i>International Journal of Oncology</i> , 2004, 25, 1713.	3.3	8
75	Effects of 5 alpha reductase inhibitors on androgen-dependent human prostatic carcinoma cells. <i>Journal of Cancer Research and Clinical Oncology</i> , 2005, 131, 243-254.	2.5	8
76	Antitumor effects of carnitinib in castration resistant prostate cancer models: A comparative study with erlotinib. <i>Prostate</i> , 2011, 71, 1481-1491.	2.3	8
77	Dual PI3K/mTOR inhibition reduces prostate cancer bone engraftment altering tumor-induced bone remodeling. <i>Tumor Biology</i> , 2018, 40, 101042831877177.	1.8	7
78	The Botanical Drug PBI-05204, a Supercritical CO2 Extract of Nerium Oleander, Is Synergistic With Radiotherapy in Models of Human Glioblastoma. <i>Frontiers in Pharmacology</i> , 2022, 13, 852941.	3.5	7
79	Targeting DDX3X Helicase Activity with BA103 Shows Promising Therapeutic Effects in Preclinical Glioblastoma Models. <i>Cancers</i> , 2021, 13, 5569.	3.7	6
80	Gefitinib and bicalutamide show synergistic effects in primary cultures of prostate cancer derived from androgen-dependent naive patients. <i>Oncology Reports</i> , 2007, 18, 1321-7.	2.6	6
81	Downmodulation of dimethyl transferase activity enhances tumor necrosis factor-related apoptosis-inducing ligand-induced apoptosis in prostate cancer cells. <i>International Journal of Oncology</i> , 2008, 33, 381-8.	3.3	6
82	Increased matrix metalloproteinase-9 secretion in short-term tissue cultures of prostatic tumor cells. <i>International Journal of Cancer</i> , 1996, 69, 386-393.	5.1	4
83	Multiple Antitumor Molecular Mechanisms Are Activated by a Fully Synthetic and Stabilized Pharmaceutical Product Delivering the Active Compound Sulforaphane (SFX-01) in Preclinical Model of Human Glioblastoma. <i>Pharmaceuticals</i> , 2021, 14, 1082.	3.8	4
84	Long-term presence of androgens and anti-androgens modulate EGF-receptor expression and MAP-kinase phosphorylation in androgen receptor-prostate positive cancer cells. <i>International Journal of Oncology</i> , 2004, 25, 97-104.	3.3	4
85	Investigational serine/threonine kinase inhibitors against prostate cancer metastases. <i>Expert Opinion on Investigational Drugs</i> , 2017, 26, 25-34.	4.1	1
86	Editorial (Thematic Issue: New Drug Targets for Treatment of Recurrent/Metastatic Prostate Cancer). <i>Current Drug Targets</i> , 2016, 17, 254-256.	2.1	0