

Patrizia Limonta

List of Publications by Year in descending order

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Version: 2024-02-01

83
papers

3,129
citations

136950
32
h-index

168389
53
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84
all docs

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docs citations

84
times ranked

3397
citing authors

#	ARTICLE	IF	CITATIONS
1	Molecular Mechanisms of Cancer Drug Resistance: Emerging Biomarkers and Promising Targets to Overcome Tumor Progression. <i>Cancers</i> , 2022, 14, 1614.	3.7	15
2	Molecular mechanisms and genetic alterations in prostate cancer: From diagnosis to targeted therapy. <i>Cancer Letters</i> , 2022, 534, 215619.	7.2	18
3	Exploiting the Metabolic Consequences of PTEN Loss and Akt/Hexokinase 2 Hyperactivation in Prostate Cancer: A New Role for Î-Tocotrienol. <i>International Journal of Molecular Sciences</i> , 2022, 23, 5269.	4.1	10
4	Melanoma Stem Cells Educate Neutrophils to Support Cancer Progression. <i>Cancers</i> , 2022, 14, 3391.	3.7	15
5	Cancer Stem Cellsâ€”Key Players in Tumor Relapse. <i>Cancers</i> , 2021, 13, 376.	3.7	74
6	Ca ²⁺ overload- and ROS-associated mitochondrial dysfunction contributes to Î-tocotrienol-mediated paraptosis in melanoma cells. <i>Apoptosis: an International Journal on Programmed Cell Death</i> , 2021, 26, 277-292.	4.9	39
7	Dissecting the Hormonal Signaling Landscape in Castration-Resistant Prostate Cancer. <i>Cells</i> , 2021, 10, 1133.	4.1	13
8	In Vitro 3D Cultures to Model the Tumor Microenvironment. <i>Cancers</i> , 2021, 13, 2970.	3.7	40
9	The multifaceted roles of mitochondria at the crossroads of cell life and death in cancer. <i>Free Radical Biology and Medicine</i> , 2021, 176, 203-221.	2.9	20
10	Î-Tocotrienol sensitizes and re-sensitizes ovarian cancer cells to cisplatin via induction of G1 phase cell cycle arrest and ROS/MAPK-mediated apoptosis. <i>Cell Proliferation</i> , 2021, 54, e13111.	5.3	24
11	Beneficial effects of Î-tocotrienol against oxidative stress in osteoblastic cells: studies on the mechanisms of action. <i>European Journal of Nutrition</i> , 2020, 59, 1975-1987.	3.9	24
12	The emerging role of paraptosis in tumor cell biology: Perspectives for cancer prevention and therapy with natural compounds. <i>Biochimica Et Biophysica Acta: Reviews on Cancer</i> , 2020, 1873, 188338.	7.4	79
13	Mitochondrial functional and structural impairment is involved in the antitumor activity of Î-tocotrienol in prostate cancer cells. <i>Free Radical Biology and Medicine</i> , 2020, 160, 376-390.	2.9	17
14	Three-Dimensional Cell Cultures as an In Vitro Tool for Prostate Cancer Modeling and Drug Discovery. <i>International Journal of Molecular Sciences</i> , 2020, 21, 6806.	4.1	34
15	Gonadotropin-Releasing Hormone Receptors in Prostate Cancer: Molecular Aspects and Biological Functions. <i>International Journal of Molecular Sciences</i> , 2020, 21, 9511.	4.1	23
16	Natural Compounds in Prostate Cancer Prevention and Treatment: Mechanisms of Action and Molecular Targets. <i>Cells</i> , 2020, 9, 460.	4.1	60
17	Anticancer properties of tocotrienols: A review of cellular mechanisms and molecular targets. <i>Journal of Cellular Physiology</i> , 2019, 234, 1147-1164.	4.1	45
18	Cellular and molecular biology of cancer stem cells in melanoma: Possible therapeutic implications. <i>Seminars in Cancer Biology</i> , 2019, 59, 221-235.	9.6	39

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19	New insights in melanoma biology: Running fast towards precision medicine. <i>Seminars in Cancer Biology</i> , 2019, 59, 161-164.	9.6	2
20	Unraveling the molecular mechanisms and the potential chemopreventive/therapeutic properties of natural compounds in melanoma. <i>Seminars in Cancer Biology</i> , 2019, 59, 266-282.	9.6	23
21	Role of Endoplasmic Reticulum Stress in the Anticancer Activity of Natural Compounds. <i>International Journal of Molecular Sciences</i> , 2019, 20, 961.	4.1	93
22	Tocotrienols and Cancer: From the State of the Art to Promising Novel Patents. <i>Recent Patents on Anti-Cancer Drug Discovery</i> , 2019, 14, 5-18.	1.6	19
23	Î-tocotrienol induces apoptosis, involving endoplasmic reticulum stress and autophagy, and paraptosis in prostate cancer cells. <i>Cell Proliferation</i> , 2019, 52, e12576.	5.3	69
24	Epithelial-To-Mesenchymal Transition Markers and CD44 Isoforms Are Differently Expressed in 2D and 3D Cell Cultures of Prostate Cancer Cells. <i>Cells</i> , 2019, 8, 143.	4.1	46
25	Targeting melanoma stem cells with the Vitamin E derivative Î-tocotrienol. <i>Scientific Reports</i> , 2018, 8, 587.	3.3	46
26	Semi-preparative HPLC purification of Î-tocotrienol (Î-T3) from <i>Elaeis guineensis</i> Jacq. and <i>Bixa orellana</i> L. and evaluation of its <i>in vitro</i> anticancer activity in human A375 melanoma cells. <i>Natural Product Research</i> , 2018, 32, 1130-1135.	1.8	24
27	GnRH in the Human Female Reproductive Axis. <i>Vitamins and Hormones</i> , 2018, 107, 27-66.	1.7	39
28	Editorial (Thematic Issue: Novel Therapeutic Strategies for Castration-resistant Prostate Cancer:) <i>TJ ETQq0 0 0 rgBT /Overlock 10 Tf 50 3</i>	0.3	0
29	Vitamin E Î-tocotrienol triggers endoplasmic reticulum stress-mediated apoptosis in human melanoma cells. <i>Scientific Reports</i> , 2016, 6, 30502.	3.3	56
30	GnRH and GnRH receptors in the pathophysiology of the human female reproductive system. <i>Human Reproduction Update</i> , 2016, 22, 358-381.	10.8	156
31	Oxime bond-linked daunorubicin-GnRH-III bioconjugates exert antitumor activity in castration-resistant prostate cancer cells via the type I GnRH receptor. <i>International Journal of Oncology</i> , 2015, 46, 243-253.	3.3	16
32	Estrogen Receptor Î ² Agonists Differentially Affect the Growth of Human Melanoma Cell Lines. <i>PLoS ONE</i> , 2015, 10, e0134396.	2.5	38
33	FROM EMERGING BIOLOGICAL INSIGHTS TO NOVEL TREATMENT STRATEGIES IN PROSTATE CANCER. Istituto Lombardo - Accademia Di Scienze E Lettere - Rendiconti Di Scienze, 2014, , .	0.0	0
34	Gonadotropin-Releasing Hormone Agonists Sensitize, and Resensitize, Prostate Cancer Cells to Docetaxel in a p53-Dependent Manner. <i>PLoS ONE</i> , 2014, 9, e93713.	2.5	14
35	Targeting Hormonal Signaling Pathways in Castration Resistant Prostate Cancer. <i>Recent Patents on Anti-Cancer Drug Discovery</i> , 2014, 9, 267-285.	1.6	10
36	Gonadotropin-releasing hormone receptors as molecular therapeutic targets in prostate cancer: Current options and emerging strategies. <i>Cancer Treatment Reviews</i> , 2013, 39, 647-663.	7.7	56

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37	Castration Resistant Prostate Cancer: From Emerging Molecular Pathways to Targeted Therapeutic Approaches. <i>Clinical Cancer Drugs</i> , 2013, 1, 11-27.	0.3	1
38	GnRH Receptors in Cancer: From Cell Biology to Novel Targeted Therapeutic Strategies. <i>Endocrine Reviews</i> , 2012, 33, 784-811.	20.1	137
39	Molecular mechanisms of the antimetastatic activity of nuclear clusterin in prostate cancer cells. <i>International Journal of Oncology</i> , 2011, 39, 225-34.	3.3	8
40	Evaluation of a Stable Gonadotropin-Releasing Hormone Analog in Mice for the Treatment of Endocrine Disorders and Prostate Cancer. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2011, 336, 613-623.	2.5	17
41	Dual Targeting of Tumor and Endothelial Cells by Gonadotropin-Releasing Hormone Agonists to Reduce Melanoma Angiogenesis. <i>Endocrinology</i> , 2010, 151, 4643-4653.	2.8	15
42	Type I Gonadotropin-Releasing Hormone Receptor Mediates the Antiproliferative Effects of GnRH-II on Prostate Cancer Cells. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2009, 94, 1761-1767.	3.6	36
43	miR-205 Exerts Tumor-Suppressive Functions in Human Prostate through Down-regulation of Protein Kinase C δ . <i>Cancer Research</i> , 2009, 69, 2287-2295.	0.9	334
44	Novel insights into GnRH receptor activity: Role in the control of human glioblastoma cell proliferation. <i>Oncology Reports</i> , 2009, 21, 1277-82.	2.6	18
45	Clusterin Isoforms Differentially Affect Growth and Motility of Prostate Cells: Possible Implications in Prostate Tumorigenesis. <i>Cancer Research</i> , 2007, 67, 10325-10333.	0.9	53
46	Gonadotropin-releasing hormone agonists reduce the migratory and the invasive behavior of androgen-independent prostate cancer cells by interfering with the activity of IGF-I. <i>International Journal of Oncology</i> , 2007, 30, 261.	3.3	6
47	Gonadotropin-releasing hormone agonists reduce the migratory and the invasive behavior of androgen-independent prostate cancer cells by interfering with the activity of IGF-I. <i>International Journal of Oncology</i> , 2007, 30, 261-71.	3.3	4
48	Gonadotropin-Releasing Hormone (GnRH) Receptors in Tumors: a New Rationale for the Therapeutical Application of GnRH Analogs in Cancer Patients?. <i>Current Cancer Drug Targets</i> , 2006, 6, 257-269.	1.6	54
49	Activation of the orphan nuclear receptor ROR α counteracts the proliferative effect of fatty acids on prostate cancer cells: Crucial role of 5-lipoxygenase. <i>International Journal of Cancer</i> , 2004, 112, 87-93.	5.1	45
50	The biology of gonadotropin hormone-releasing hormone: role in the control of tumor growth and progression in humans. <i>Frontiers in Neuroendocrinology</i> , 2003, 24, 279-295.	5.2	114
51	Inhibitory activity of luteinizing hormone-releasing hormone on tumor growth and progression.. <i>Endocrine-Related Cancer</i> , 2003, 10, 161-167.	3.1	35
52	Locally Expressed LHRH Receptors Mediate the Oncostatic and Antimetastatic Activity of LHRH Agonists on Melanoma Cells. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2002, 87, 3791-3797.	3.6	53
53	Locally Expressed LHRH Receptors Mediate the Oncostatic and Antimetastatic Activity of LHRH Agonists on Melanoma Cells. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2002, 87, 3791-3797.	3.6	14
54	Oncostatic activity of a thiazolidinedione derivative on human androgen-dependent prostate cancer cells. <i>International Journal of Cancer</i> , 2001, 92, 733-737.	5.1	20

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55	Activation of the orphan nuclear receptor ROR γ induces growth arrest in androgen-independent DU 145 prostate cancer cells. <i>Prostate</i> , 2001, 46, 327-335.	2.3	25
56	LHRH analogues as anticancer agents: pituitary and extrapituitary sites of action. <i>Expert Opinion on Investigational Drugs</i> , 2001, 10, 709-720.	4.1	90
57	The Luteinizing Hormone-Releasing Hormone Receptor in Human Prostate Cancer Cells: Messenger Ribonucleic Acid Expression, Molecular Size, and Signal Transduction Pathway ¹ . <i>Endocrinology</i> , 1999, 140, 5250-5256.	2.8	123
58	The Luteinizing Hormone-Releasing Hormone Receptor in Human Prostate Cancer Cells: Messenger Ribonucleic Acid Expression, Molecular Size, and Signal Transduction Pathway. <i>Endocrinology</i> , 1999, 140, 5250-5256.	2.8	30
59	Growth-inhibitory effects of luteinizing hormone-releasing hormone (LHRH) agonists on xenografts of the DU 145 human androgen-independent prostate cancer cell line in nude mice. <i>International Journal of Cancer</i> , 1998, 76, 506-511.	5.1	42
60	Growth factors in steroid-responsive prostatic tumor cells. <i>Steroids</i> , 1996, 61, 222-225.	1.8	7
61	LH-RH and Somatostatin: Examples of Peptidergic Control of Prostate Cancer Growth. <i>Contributions To Oncology / Beitrage Zur Onkologie</i> , 1995, 50, 332-344.	0.1	0
62	Growth of the androgen-dependent tumor of the prostate: Role of androgens and of locally expressed growth modulatory factors. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 1995, 53, 401-405.	2.5	20
63	Effects of steroids on the brain opioid system. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 1995, 53, 343-348.	2.5	71
64	Effect of aging on opioid and LHRH receptors in the brain, pituitary, and testis of the male rat. <i>Neurobiology of Aging</i> , 1994, 15, 553-557.	3.1	6
65	Androgen-dependent prostatic tumors: biosynthesis and possible actions of LHRH. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 1994, 49, 347-350.	2.5	15
66	Characterization of a soluble LHRH-degrading activity in the rat ventral prostate. <i>Prostate</i> , 1993, 23, 315-328.	2.3	9
67	Binding Characteristics of Hypothalamic Mu Opioid Receptors throughout the Estrous Cycle in the Rat. <i>Neuroendocrinology</i> , 1993, 58, 366-372.	2.5	79
68	Gonadotropin-releasing hormone agonists suppress melanoma cell motility and invasiveness through the inhibition of α_3 integrin and MMP-2 expression and activity. <i>International Journal of Oncology</i> , 1992, 33, 405.	3.3	7
69	Modulation of the binding characteristics of hypothalamic mu opioid receptors in rats by gonadal steroids. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 1991, 40, 113-121.	2.5	22
70	Testosterone and postnatal ontogenesis of hypothalamic μ ([³ H]dihydromorphine) opioid receptors in the rat. <i>Developmental Brain Research</i> , 1991, 62, 131-136.	1.7	20
71	Hypothalamic Opiatergic Tone During Pregnancy, Parturition and Lactation in the Rat. <i>Neuroendocrinology</i> , 1991, 53, 460-466.	2.5	65
72	Effect of ovarian steroids on the concentration of μ opiate receptors in different regions of the brain of the female rat. <i>Pharmacological Research</i> , 1989, 21, 91-92.	7.1	13

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73	Distribution of kappa opioid receptors in the brain of young and old male rats. Life Sciences, 1989, 45, 2085-2092.	4.3	22
74	Effects of aging on pituitary and testicular luteinizing hormone-releasing hormone receptors in the rat. Life Sciences, 1988, 42, 335-342.	4.3	13
75	Further Evidence that Gonadal Steroids do not Modulate Brain Opiate Receptors in Male Rats.. Endocrinologia Japonica, 1987, 34, 521-529.	0.5	11
76	Decrease of mu opioid receptors in the brain and in the hypothalamus of the aged male rat. Life Sciences, 1987, 40, 391-398.	4.3	44
77	Stimulatory and Inhibitory Effects of the Opioids on Gonadotropin Secretion. Neuroendocrinology, 1986, 42, 504-512.	2.5	60
78	Species differences in the sensitivity to GnRH analogs. The Journal of Steroid Biochemistry, 1985, 23, 811-817.	1.1	13
79	Unexpected effects of nalmefene, a new opiate antagonist, on the hypothalamic-pituitary-gonadal axis in the male rat. Steroids, 1985, 46, 955-965.	1.8	14
80	Species differences in the sensitivity to a GnRH antagonist. Contraception, 1985, 32, 75-85.	1.5	5
81	Role of the subfornical organ (SFO) in the control of gonadotropin secretion. Brain Research, 1981, 229, 75-84.	2.2	23
82	Cholinergic inputs to the amygdala and the control of gonadotrophin release. European Journal of Endocrinology, 1980, 93, 1-6.	3.7	20
83	REPRODUCTIVE FUNCTION AND ANTITUMOR ACTIVITY: DIFFERENT ROLES FOR THE HYPOTHALAMIC HORMONE GnRH. Istituto Lombardo - Accademia Di Scienze E Lettere - Incontri Di Studio, 0, , .	0.0	0