## Cinzia Lanzi

List of Publications by Year in descending order

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| #  | Article   | IF  | CITATIONS |
|----|---|-----|-----------|
| 1  | Upregulation of ERK-EGR1-heparanase axis by HDAC inhibitors provides targets for rational therapeutic intervention in synovial sarcoma. Journal of Experimental and Clinical Cancer Research, 2021, 40, 381.                      | 3.5 | 9         |
| 2  | Receptor tyrosine kinases and heparan sulfate proteoglycans: Interplay providing anticancer targeting strategies and new therapeutic opportunities. Biochemical Pharmacology, 2020, 178, 114084.                                  | 2.0 | 20        |
| 3  | Editorial: Heparan Sulfate Proteoglycans and Their Endogenous Modifying Enzymes: Cancer Players,<br>Biomarkers and Therapeutic Targets. Frontiers in Oncology, 2020, 10, 195.   | 1.3 | 6         |
| 4  | Heparanase: A Potential Therapeutic Target in Sarcomas. Advances in Experimental Medicine and<br>Biology, 2020, 1221, 405-431.  | 0.8 | 3         |
| 5  | Overactive IGF1/Insulin Receptors and NRASQ61R Mutation Drive Mechanisms of Resistance to<br>Pazopanib and Define Rational Combination Strategies to Treat Synovial Sarcoma. Cancers, 2019, 11,<br>408.                           | 1.7 | 10        |
| 6  | Microenvironment modulation and enhancement of antilymphoma therapy by the heparanase inhibitor roneparstat. Hematological Oncology, 2018, 36, 360-362.   | 0.8 | 15        |
| 7  | Supersulfated low-molecular weight heparin synergizes with IGF1R/IR inhibitor to suppress synovial sarcoma growth and metastases. Cancer Letters, 2018, 415, 187-197.   | 3.2 | 24        |
| 8  | Heparan Sulfate Mimetics in Cancer Therapy: The Challenge to Define Structural Determinants and the<br>Relevance of Targets for Optimal Activity. Molecules, 2018, 23, 2915.  | 1.7 | 46        |
| 9  | Axl molecular targeting counteracts aggressiveness but not platinum-resistance of ovarian carcinoma cells. Biochemical Pharmacology, 2017, 136, 40-50.  | 2.0 | 16        |
| 10 | Targeting Heparan Sulfate Proteoglycans and their Modifying Enzymes to Enhance Anticancer<br>Chemotherapy Efficacy and Overcome Drug Resistance. Current Medicinal Chemistry, 2017, 24,<br>2860-2886.                             | 1.2 | 42        |
| 11 | Targeting ErbB3 activation in drug-resistant ovarian carcinoma cells over-expressing the receptor tyrosine kinase Axl. European Journal of Cancer, 2016, 69, S71-S72.   | 1.3 | 0         |
| 12 | Synthetic sulfoglycolipids targeting the serine–threonine protein kinase Akt. Bioorganic and<br>Medicinal Chemistry, 2016, 24, 3396-3405.   | 1.4 | 9         |
| 13 | The heparanase/heparan sulfate proteoglycan axis: A potential new therapeutic target in sarcomas.<br>Cancer Letters, 2016, 382, 245-254.  | 3.2 | 25        |
| 14 | Guidelines for the use and interpretation of assays for monitoring autophagy (3rd edition).<br>Autophagy, 2016, 12, 1-222.  | 4.3 | 4,701     |
| 15 | Targeting of <i>RET</i> oncogene by naphthalene diimide-mediated gene promoter G-quadruplex<br>stabilization exerts anti-tumor activity in oncogene-addicted human medullary thyroid cancer.<br>Oncotarget, 2016, 7, 49649-49663. | 0.8 | 22        |
| 16 | Antitumor efficacy of the heparan sulfate mimic roneparstat (SST0001) against sarcoma models involves multi-target inhibition of receptor tyrosine kinases. Oncotarget, 2016, 7, 47848-47863.                                     | 0.8 | 43        |
| 17 | Role of the Receptor Tyrosine Kinase Axl and its Targeting in Cancer Cells. Current Medicinal Chemistry, 2016, 23, 1496-1512.   | 1.2 | 31        |
| 18 | Abstract 3289: Microenvironment modulation and enhancement of cytotoxic therapy by the  |     | 0         |

heparanase inhibitor Roneparstat against human B-non Hodgkin lymphomas. , 2016, , .

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|----|---|-----|-----------|
| 19 | New mechanisms for old drugs: Insights into DNA-unrelated effects of platinum compounds and drug resistance determinants. Drug Resistance Updates, 2015, 20, 1-11.  | 6.5 | 47        |
| 20 | Targeting the invasive phenotype of cisplatin-resistant Non-Small Cell Lung Cancer cells by a novel histone deacetylase inhibitor. Biochemical Pharmacology, 2015, 94, 79-90.   | 2.0 | 22        |
| 21 | PLK1 is a critical determinant of tumor cell sensitivity to CPT11 and its inhibition enhances the drug antitumor efficacy in squamous cell carcinoma models sensitive and resistant to camptothecins. Oncotarget, 2015, 6, 8736-8749. | 0.8 | 17        |
| 22 | Synergistic Cooperation Between Sunitinib and Cisplatin Promotes Apoptotic Cell Death in Human<br>Medullary Thyroid Cancer. Journal of Clinical Endocrinology and Metabolism, 2014, 99, 498-509.                                      | 1.8 | 23        |
| 23 | Stratification of clear cell renal cell carcinoma by signaling pathway analysis. Expert Review of Proteomics, 2014, 11, 237-249.  | 1.3 | 9         |
| 24 | 141: RET/PTC1 in vitro models unveil a novel tumor suppressor miRNA in papillary thyroid carcinoma.<br>European Journal of Cancer, 2014, 50, S31.   | 1.3 | 0         |
| 25 | Differential outcome of MEK1/2 inhibitor-platinum combinations in platinum-sensitive and -resistant ovarian carcinoma cells. Cancer Letters, 2014, 347, 212-224.  | 3.2 | 26        |
| 26 | miR-199a-3p displays tumor suppressor functions in papillary thyroid carcinoma. Oncotarget, 2014, 5, 2513-2528.   | 0.8 | 98        |
| 27 | Medullary Thyroid Cancer Targeted Therapy. , 2014, , 1-4.   |     | Ο         |
| 28 | Medullary Thyroid Cancer Targeted Therapy. , 2014, , 2699-2702.   |     | 0         |
| 29 | Antitumor efficacy of the heparanase inhibitor SST0001 alone and in combination with antiangiogenic agents in the treatment of human pediatric sarcoma models. Biochemical Pharmacology, 2013, 85, 1424-1432.                         | 2.0 | 75        |
| 30 | DUSP6/MKP3 is overexpressed in papillary and poorly differentiated thyroid carcinoma and contributes to neoplastic properties of thyroid cancer cells. Endocrine-Related Cancer, 2013, 20, 23-37.                                     | 1.6 | 41        |
| 31 | Modulation of Sensitivity to Antitumor Agents by Targeting the MAPK Survival Pathway. Current<br>Pharmaceutical Design, 2013, 19, 883-894.  | 0.9 | 47        |
| 32 | Targeting the Akt Kinase to Modulate Survival, Invasiveness and Drug Resistance of Cancer Cells.<br>Current Medicinal Chemistry, 2013, 20, 1923-1945.   | 1.2 | 86        |
| 33 | Abstract A93: Targeting the increased invasive capability of non-small cell lung cancer platinum-resistant cells by histone deacetylase inhibitors , 2013, , .  |     | Ο         |
| 34 | Abstract A64: Synergistic cooperation between sunitinib and cisplatin promotes apoptotic cell death in human medullary thyroid cancer , 2013, , .   |     | 0         |
| 35 | The curative efficacy of namitecan (ST1968) in preclinical models of pediatric sarcoma is associated with antiangiogenic effects. Biochemical Pharmacology, 2012, 84, 163-171.  | 2.0 | 29        |
| 36 | Modulation of Sensitivity to Antitumor Agents by Targeting the MAPK Survival Pathway. Current<br>Pharmaceutical Design, 2012, 19, 883-894.  | 0.9 | 23        |

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|----|--|-----|-----------|
| 37 | Signaling pathway-based stratification of clear cell renal cell carcinoma Journal of Clinical<br>Oncology, 2012, 30, 434-434.  | 0.8 | 0         |
| 38 | Identification of MET and SRC Activation in Melanoma Cell Lines Showing Primary Resistance to PLX4032. Neoplasia, 2011, 13, 1132-IN17.   | 2.3 | 89        |
| 39 | Pre-clinical and clinical significance of heparanase in Ewing's sarcoma. Journal of Cellular and<br>Molecular Medicine, 2011, 15, 1857-1864.   | 1.6 | 53        |
| 40 | Interplay between Ret and Fap-1 regulates CD95-mediated apoptosis in medullary thyroid cancer cells.<br>Biochemical Pharmacology, 2011, 82, 778-788.   | 2.0 | 13        |
| 41 | Abstract C206: High antitumor complete response rate to Namitecan (ST1968) in preclinical model of pediatric sarcomas , 2011, , .  |     | 0         |
| 42 | Targeting RET for thyroid cancer therapy. Biochemical Pharmacology, 2009, 77, 297-309.   | 2.0 | 62        |
| 43 | Concomitant downregulation of proliferation/survival pathways dependent on FGF-R3, JAK2 and BCMA<br>in human multiple myeloma cells by multi-kinase targeting. Biochemical Pharmacology, 2009, 78,<br>1139-1147.   | 2.0 | 9         |
| 44 | Proteomics study of medullary thyroid carcinomas expressing RET germâ€line mutations: Identification of new signaling elements. Molecular Carcinogenesis, 2009, 48, 220-231.   | 1.3 | 26        |
| 45 | RET/PTC1-Driven Neoplastic Transformation and Proinvasive Phenotype of Human Thyrocytes Involve Met Induction and β-Catenin Nuclear Translocation. Neoplasia, 2009, 11, 10-21.   | 2.3 | 55        |
| 46 | Modulation of Survival Pathways in Ovarian Carcinoma Cells Resistant to Platinum Compounds. , 2009, , 195-200.   |     | 0         |
| 47 | Abstract B224: Multiâ€ŧyrosine kinase targeting resulting in concomitant downregulation of proliferation/survival pathways dependent on FGFâ€R3, Jak2 and BCMA in human multiple myeloma cells. , 2009, , .  |     | 0         |
| 48 | Synthesis, Modeling, and RET Protein Kinase Inhibitory Activity of 3- and 4-Substituted β-Carbolin-1-ones.<br>Journal of Medicinal Chemistry, 2008, 51, 7777-7787.   | 2.9 | 36        |
| 49 | Modulation of survival pathways in ovarian carcinoma cell lines resistant to platinum compounds.<br>Molecular Cancer Therapeutics, 2008, 7, 679-687.   | 1.9 | 52        |
| 50 | Synthesis and RET protein kinase inhibitory activity of 3-arylureidobenzylidene-indolin-2-ones.<br>Bioorganic and Medicinal Chemistry Letters, 2007, 17, 3962-3968.  | 1.0 | 21        |
| 51 | Apoptotic cell death induction and angiogenesis inhibition in large established medullary thyroid carcinoma xenografts by Ret inhibitor RPI-1. Biochemical Pharmacology, 2006, 72, 405-414.  | 2.0 | 30        |
| 52 | Inhibition of c-Met and prevention of spontaneous metastatic spreading by the 2-indolinone RPI-1.<br>Molecular Cancer Therapeutics, 2006, 5, 2388-2397.  | 1.9 | 42        |
| 53 | Modulation of Survival Signaling Pathways and Persistence of the Genotoxic Stress as a Basis for the Synergistic Interaction between the Atypical Retinoid ST1926 and the Epidermal Growth Factor Receptor Inhibitor ZD1839. Cancer Research, 2005, 65, 2364-2372. | 0.4 | 26        |
| 54 | Development of Resistance to the Atypical Retinoid, ST1926, in the Lung Carcinoma Cell Line H460 Is<br>Associated with Reduced Formation of DNA Strand Breaks and a Defective DNA Damage Response.<br>Neoplasia, 2005, 7, 667-677.                                 | 2.3 | 27        |

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|----|--|-----|-----------|
| 55 | Cellular Effects and Antitumor Activity of RET Inhibitor RPI-1 on MEN2A-Associated Medullary Thyroid<br>Carcinoma. Journal of the National Cancer Institute, 2004, 96, 1006-1014.  | 3.0 | 106       |
| 56 | Induction of apoptosis and stress response in ovarian carcinoma cell lines treated with ST1926, an atypical retinoid. Cell Death and Differentiation, 2004, 11, 280-289.   | 5.0 | 54        |
| 57 | Role of c-myc protein in hormone refractory prostate carcinoma: cellular response to paclitaxel.<br>Biochemical Pharmacology, 2004, 68, 923-931.   | 2.0 | 14        |
| 58 | Antitumour and antiangiogenic effects of IDN 5390, a novel C-seco taxane, in a paclitaxel-resistant human ovarian tumour xenograft. British Journal of Cancer, 2004, 90, 1464-1468.  | 2.9 | 18        |
| 59 | Inactivation of Ret/Ptc1 oncoprotein and inhibition of papillary thyroid carcinoma cell proliferation by indolinone RPI-1. Cellular and Molecular Life Sciences, 2003, 60, 1449-1459.                                      | 2.4 | 45        |
| 60 | IDN 5390: an oral taxane candidate for protracted treatment schedules. British Journal of Cancer, 2003, 88, 965-972.   | 2.9 | 18        |
| 61 | Antiangiogenic effects of the novel camptothecin ST1481 (gimatecan) in human tumor xenografts.<br>Molecular Cancer Research, 2003, 1, 863-70.  | 1.5 | 35        |
| 62 | Cell cycle checkpoint efficiency and cellular response to paclitaxel in prostate cancer cells. Prostate, 2001, 48, 254-264.  | 1.2 | 68        |
| 63 | A role for loss of p53 function in sensitivity of ovarian carcinoma cells to taxanes. International<br>Journal of Cancer, 2001, 92, 738-747.   | 2.3 | 61        |
| 64 | Inhibition of transforming activity of the ret/ptc1 oncoprotein by a 2-indolinone derivative.<br>International Journal of Cancer, 2000, 85, 384-390.   | 2.3 | 57        |
| 65 | Structure elucidation of clavilactone D: an inhibitor of protein tyrosine kinases. Phytochemistry, 2000, 53, 1039-1041.  | 1.4 | 44        |
| 66 | Clavilactones, a novel class of tyrosine kinase inhibitors of fungal origin. Biochemical Pharmacology, 2000, 59, 1539-1547.  | 2.0 | 59        |
| 67 | Decreased Drug Accumulation and Increased Tolerance to DNA Damage in Tumor Cells with a Low<br>Level of Cisplatin Resistance. Biochemical Pharmacology, 1998, 55, 1247-1254.   | 2.0 | 55        |
| 68 | Lipid peroxidation, phosphoinositide turnover and protein kinase C activation in human platelets<br>treated with anthracyclines and their complexes with Fe(III). Biochemical Pharmacology, 1992, 43,<br>1521-1527.        | 2.0 | 8         |
| 69 | Protein kinase C activation by anthracyclines in swiss 3T3 cells. International Journal of Cancer, 1991, 47, 136-142.  | 2.3 | 12        |
| 70 | Selection of monoclonal antibodies which induce internalization and phosphorylation of P185HER2<br>and growth inhibition of cells with HER2/neu gene amplification. International Journal of Cancer,<br>1991, 47, 933-937. | 2.3 | 99        |
| 71 | Protein Kinase C Activation and Lipid Peroxidation by Doxorubicin Analogues. Tumori, 1989, 75, 358-361.  | 0.6 | 4         |
| 72 | Diversity of effects of two antitumor anthracycline analogs on the pathway of activation of PKC in intact human platelets. Biochemical Pharmacology, 1988, 37, 3497-3504.  | 2.0 | 9         |

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|----|--|-----|-----------|
| 73 | Role of Daunosamine and Hydroxyacetyl Side Chain in Reaction With Iron and Lipid Peroxidation by<br>Anthracyclines. Journal of the National Cancer Institute, 1988, 80, 1104-1111. | 3.0 | 42        |