## Atanasio Pandiella

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Membrane-Anchored Growth Factors. Annual Review of Biochemistry, 1993, 62, 515-541.	5.0	641
2	Sox2 expression in breast tumours and activation in breast cancer stem cells. Oncogene, 2012, 31, 1354-1365.	2.6	447
3	Antitumor Effects of Doxorubicin in Combination With Anti-epidermal Growth Factor Receptor Monoclonal Antibodies. Journal of the National Cancer Institute, 1993, 85, 1327-1333.	3.0	372
4	Neutrophils in cancer: prognostic role and therapeutic strategies. Molecular Cancer, 2017, 16, 137.	7.9	295
5	Extracellular Signal-regulated Kinase Phosphorylates Tumor Necrosis Factor α-converting Enzyme at Threonine 735: A Potential Role in Regulated Shedding. Molecular Biology of the Cell, 2002, 13, 2031-2044.	0.9	273
6	Inhibition of Src Family Kinases and Receptor Tyrosine Kinases by Dasatinib: Possible Combinations in Solid Tumors. Clinical Cancer Research, 2011, 17, 5546-5552.	3.2	247
7	The Histone Deacetylase Inhibitor LBH589 Is a Potent Antimyeloma Agent that Overcomes Drug Resistance. Cancer Research, 2006, 66, 5781-5789.	0.4	233
8	Cleavage of the membrane precursor for transforming growth factor alpha is a regulated process Proceedings of the National Academy of Sciences of the United States of America, 1991, 88, 1726-1730.	3.3	219
9	Bortezomib induces selective depletion of alloreactive T lymphocytes and decreases the production of Th1 cytokines. Blood, 2006, 107, 3575-3583.	0.6	188
10	HER3 Overexpression and Survival in Solid Tumors: A Meta-analysis. Journal of the National Cancer Institute, 2013, 105, 266-273.	3.0	168
11	Erk5 Participates in Neuregulin Signal Transduction and Is Constitutively Active in Breast Cancer Cells Overexpressing ErbB2. Molecular and Cellular Biology, 2002, 22, 270-285.	1.1	163
12	Transforming Growth Factor Î <sup>2</sup> Engages TACE and ErbB3 To Activate Phosphatidylinositol-3 Kinase/Akt in ErbB2-Overexpressing Breast Cancer and Desensitizes Cells to Trastuzumab. Molecular and Cellular Biology, 2008, 28, 5605-5620.	1.1	153
13	Differential Shedding of Transmembrane Neuregulin Isoforms by the Tumor Necrosis Factor-α-Converting Enzyme. Molecular and Cellular Neurosciences, 2000, 16, 631-648.	1.0	152
14	Preclinical development of molecular-targeted agents for cancer. Nature Reviews Clinical Oncology, 2011, 8, 200-209.	12.5	145
15	In vitro and in vivo rationale for the triple combination of panobinostat (LBH589) and dexamethasone with either bortezomib or lenalidomide in multiple myeloma. Haematologica, 2010, 95, 794-803.	1.7	144
16	Activation of the PI3K/mTOR/AKT Pathway and Survival in Solid Tumors: Systematic Review and Meta-Analysis. PLoS ONE, 2014, 9, e95219.	1.1	140
17	The cytoplasmic carboxy-terminal amino acid specifies cleavage of membrane TGFα into soluble growth factor. Cell, 1992, 71, 1157-1165.	13.5	136
18	Resistance to Antibody–Drug Conjugates. Cancer Research, 2018, 78, 2159-2165.	0.4	136

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19	Mesenchymal stem cells from multiple myeloma patients display distinct genomic profile as compared with those from normal donors. Leukemia, 2009, 23, 1515-1527.	3.3	122
20	New drugs in multiple myeloma: mechanisms of action and phase I/II clinical findings. Lancet Oncology, The, 2008, 9, 1157-1165.	5.1	116
21	The epoxyketone-based proteasome inhibitors carfilzomib and orally bioavailable oprozomib have anti-resorptive and bone-anabolic activity in addition to anti-myeloma effects. Leukemia, 2013, 27, 430-440.	3.3	112
22	Genetic Abnormalities and Patterns of Antigenic Expression in Multiple Myeloma. Clinical Cancer Research, 2005, 11, 3661-3667.	3.2	109
23	TrkA receptor ectodomain cleavage generates a tyrosine-phosphorylated cell-associated fragment Journal of Cell Biology, 1996, 132, 427-436.	2.3	104
24	Resistance to the Antibody–Drug Conjugate T-DM1 Is Based in a Reduction in Lysosomal Proteolytic Activity. Cancer Research, 2017, 77, 4639-4651.	0.4	103
25	Expression of Erk5 in Early Stage Breast Cancer and Association with Disease Free Survival Identifies this Kinase as a Potential Therapeutic Target. PLoS ONE, 2009, 4, e5565.	1.1	99
26	Aplidin, a Marine Organism–Derived Compound with Potent Antimyeloma Activity <i>In vitro</i> and <i>In vivo</i> . Cancer Research, 2008, 68, 5216-5225.	0.4	98
27	Neuregulins and Cancer. Clinical Cancer Research, 2008, 14, 3237-3241.	3.2	95
28	P-Rex1 participates in Neuregulin-ErbB signal transduction and its expression correlates with patient outcome in breast cancer. Oncogene, 2011, 30, 1059-1071.	2.6	92
29	EGF raises cytosolic Ca2+ in A431 and Swiss 3T3 cells by a dual mechanism. Experimental Cell Research, 1987, 170, 175-185.	1.2	89
30	Early rise of cytosolic Ca2+induced by NGF in PC12 and chromaffin cells. FEBS Letters, 1986, 208, 48-51.	1.3	88
31	Cellular Plasticity Confers Migratory and Invasive Advantages to a Population of Glioblastoma-Initiating Cells that Infiltrate Peritumoral Tissue. Stem Cells, 2013, 31, 1075-1085.	1.4	83
32	Multifunctional role of Erk5 in multiple myeloma. Blood, 2005, 105, 4492-4499.	0.6	82
33	ω-Conotoxin binding and effects on calcium channel function in human neuroblastoma and rat pheochromocytoma cell lines. FEBS Letters, 1988, 235, 178-182.	1.3	78
34	Zalypsis: a novel marine-derived compound with potent antimyeloma activity that reveals high sensitivity of malignant plasma cells to DNA double-strand breaks. Blood, 2009, 113, 3781-3791.	0.6	78
35	Active kinase profiling, genetic and pharmacological data define mTOR as an important common target in triple-negative breast cancer. Oncogene, 2014, 33, 148-156.	2.6	78
36	TGF-β1 induces COX-2 expression and PGE2 synthesis through MAPK and PI3K pathways in human mesangial cells. Kidney International, 2006, 70, 901-909.	2.6	75

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37	Clinical significance of CD81 expression by clonal plasma cells in high-risk smoldering and symptomatic multiple myeloma patients. Leukemia, 2012, 26, 1862-1869.	3.3	73
38	In vivo murine model of acquired resistance in myeloma reveals differential mechanisms for lenalidomide and pomalidomide in combination with dexamethasone. Leukemia, 2015, 29, 705-714.	3.3	72
39	Autocrine Regulation of Membrane Transforming Growth Factor-Î $\pm$ Cleavage. Journal of Biological Chemistry, 1996, 271, 3279-3284.	1.6	69
40	Endoglin Modulation of TGF-ß1-Induced Collagen Synthesis is Dependent on ERK1/2 MAPK Activation. Cellular Physiology and Biochemistry, 2006, 18, 135-142.	1.1	65
41	Oncogenic Targets, Magnitude of Benefit, and Market Pricing of Antineoplastic Drugs. Journal of Clinical Oncology, 2011, 29, 2543-2549.	0.8	64
42	Activation of ErbB2 by Overexpression or by Transmembrane Neuregulin Results in Differential Signaling and Sensitivity to Herceptin. Cancer Research, 2005, 65, 6801-6810.	0.4	63
43	Activated release of membrane-anchored TGF-alpha in the absence of cytosol. Journal of Cell Biology, 1993, 122, 95-101.	2.3	62
44	ERK5 Activates NF-κB in Leukemic T Cells and Is Essential for Their Growth In Vivo. Journal of Immunology, 2006, 177, 7607-7617.	0.4	62
45	Activity of BET-proteolysis targeting chimeric (PROTAC) compounds in triple negative breast cancer. Journal of Experimental and Clinical Cancer Research, 2019, 38, 383.	3.5	62
46	An Overview of Antibody Conjugated Polymeric Nanoparticles for Breast Cancer Therapy. Pharmaceutics, 2020, 12, 802.	2.0	62
47	Dasatinib as a Bone-Modifying Agent: Anabolic and Anti-Resorptive Effects. PLoS ONE, 2012, 7, e34914.	1.1	61
48	Defective Cyclin B1 Induction in Trastuzumab-emtansine (T-DM1) Acquired Resistance in HER2-positive Breast Cancer. Clinical Cancer Research, 2017, 23, 7006-7019.	3.2	61
49	Stimulation of cleavage of membrane proteins by calmodulin inhibitors. Biochemical Journal, 2000, 346, 359-367.	1.7	59
50	Transforming growth factor-β1 induces collagen synthesis and accumulation via p38 mitogen-activated protein kinase (MAPK) pathway in cultured L6E9myoblasts. FEBS Letters, 2002, 513, 282-288.	1.3	59
51	Bortezomib is an efficient agent in plasma cell leukemias. International Journal of Cancer, 2005, 114, 665-667.	2.3	59
52	Role of metalloproteinases MMP-9 and MT1-MMP in CXCL12-promoted myeloma cell invasion across basement membranes. Journal of Pathology, 2006, 208, 108-118.	2.1	59
53	The synergy of panobinostat plus doxorubicin in acute myeloid leukemia suggests a role for HDAC inhibitors in the control of DNA repair. Leukemia, 2009, 23, 2265-2274.	3.3	58
54	The mitogen-activated protein kinase ERK5 regulates the development and growth of hepatocellular carcinoma. Gut, 2015, 64, 1454-1465.	6.1	58

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55	Trastuzumab Emtansine: Mechanisms of Action and Resistance, Clinical Progress, and Beyond. Trends in Cancer, 2020, 6, 130-146.	3.8	58
56	Synergic antitumoral effect of an IGF-IR inhibitor and trastuzumab on HER2-overexpressing breast cancer cells. Annals of Oncology, 2008, 19, 1860-1869.	0.6	57
57	The effect of the proteasome inhibitor bortezomib on acute myeloid leukemia cells and drug resistance associated with the CD34+ immature phenotype. Haematologica, 2008, 93, 57-66.	1.7	56
58	Colorectal cancer and medicinal plants: Principle findings from recent studies. Biomedicine and Pharmacotherapy, 2018, 107, 408-423.	2.5	56
59	Imatinib mesylate (STI571) inhibits multiple myeloma cell proliferation and potentiates the effect of common antimyeloma agents. British Journal of Haematology, 2003, 123, 858-868.	1.2	53
60	Neuregulin Expression Modulates Clinical Response to Trastuzumab in Patients With Metastatic Breast Cancer. Journal of Clinical Oncology, 2007, 25, 2656-2663.	0.8	53
61	ERK5/BMK1 Is a Novel Target of the Tumor Suppressor VHL: Implication in Clear Cell Renal Carcinoma. Neoplasia, 2013, 15, 649-IN17.	2.3	53
62	Ubiquitin-conjugating enzyme E2T (UBE2T) and denticleless protein homolog (DTL) are linked to poor outcome in breast and lung cancers. Scientific Reports, 2017, 7, 17530.	1.6	53
63	HER2 heterogeneity and resistance to anti-HER2 antibody-drug conjugates. Breast Cancer Research, 2020, 22, 15.	2.2	53
64	A new simple whole blood flow cytometry-based method for simultaneous identification of activated cells and quantitative evaluation of cytokines released during activation. Laboratory Investigation, 2004, 84, 1387-1398.	1.7	52
65	Personalized therapies in the cancer "omics" era. Molecular Cancer, 2010, 9, 202.	7.9	52
66	Mitogen-activated protein kinase-dependent and -independent routes control shedding of transmembrane growth factors through multiple secretases. Biochemical Journal, 2002, 363, 211-221.	1.7	51
67	Targeting the EGF/HER Ligand-Receptor System in Cancer. Current Pharmaceutical Design, 2016, 22, 5887-5898.	0.9	51
68	Androgen-independent prostate cancer cells circumvent EGFR inhibition by overexpression of alternative HER receptors and ligands. International Journal of Oncology, 2012, 41, 1128-1138.	1.4	50
69	Cleavage of the TrkA neurotrophin receptor by multiple metalloproteases generates signalling-competent truncated forms. European Journal of Neuroscience, 1999, 11, 1421-1430.	1.2	49
70	Induction of B-Chronic Lymphocytic Leukemia Cell Apoptosis by Arsenic Trioxide Involves Suppression of the Phosphoinositide 3-Kinase/Akt Survival Pathway via <i>c-jun</i> -NH2 Terminal Kinase Activation and PTEN Upregulation. Clinical Cancer Research, 2010, 16, 4382-4391.	3.2	49
71	Phospho-kinase profile of triple negative breast cancer and androgen receptor signaling. BMC Cancer, 2014, 14, 302.	1.1	49
72	PDGF-induced receptor phosphorylation and phosphoinositide hydrolysis are unaffected by protein kinase C activation in mouse Swiss 3T3 and human skin fibroblasts. Biochemical and Biophysical Research Communications, 1986, 137, 343-350.	1.0	48

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73	ODZ1 allows glioblastoma to sustain invasiveness through a Myc-dependent transcriptional upregulation of RhoA. Oncogene, 2017, 36, 1733-1744.	2.6	48
74	Predominance of mTORC1 over mTORC2 in the Regulation of Proliferation of Ovarian Cancer Cells: Therapeutic Implications. Molecular Cancer Therapeutics, 2012, 11, 1342-1352.	1.9	47
75	Endoglin Expression Regulates Basal and TGF-β1-induced Extracellular Matrix Synthesis in Cultured L <sub>6</sub> E <sub>9</sub> Myoblasts. Cellular Physiology and Biochemistry, 2004, 14, 301-310.	1.1	46
76	Targeting oncogenic vulnerabilities in triple negative breast cancer: biological bases and ongoing clinical studies. Oncotarget, 2017, 8, 22218-22234.	0.8	46
77	Potent Antimyeloma Activity of a Novel ERK5/CDK Inhibitor. Clinical Cancer Research, 2013, 19, 2677-2687.	3.2	45
78	Impaired Trafficking and Activation of Tumor Necrosis Factor-α-converting Enzyme in Cell Mutants Defective in Protein Ectodomain Shedding. Journal of Biological Chemistry, 2003, 278, 25933-25939.	1.6	44
79	Multisite phosphorylation of Erk5 in mitosis. Journal of Cell Science, 2010, 123, 3146-3156.	1.2	44
80	Synthetic Lethality Interaction Between Aurora Kinases and CHEK1 Inhibitors in Ovarian Cancer. Molecular Cancer Therapeutics, 2017, 16, 2552-2562.	1.9	44
81	Mitogen-activated protein kinase-dependent and -independent routes control shedding of transmembrane growth factors through multiple secretases. Biochemical Journal, 2002, 363, 211.	1.7	43
82	Effect of Multikinase Inhibitors on Caspase-Independent Cell Death and DNA Damage in HER2-Overexpressing Breast Cancer Cells. Journal of the National Cancer Institute, 2010, 102, 1432-1446.	3.0	43
83	Transcriptomic profile induced in bone marrow mesenchymal stromal cells after interaction with multiple myeloma cells: implications in myeloma progression and myeloma bone disease. Oncotarget, 2014, 5, 8284-8305.	0.8	43
84	Identifying Breast Cancer Druggable Oncogenic Alterations: Lessons Learned and Future Targeted Options. Clinical Cancer Research, 2008, 14, 961-970.	3.2	42
85	Differential action of small molecule HER kinase inhibitors on receptor heterodimerization: Therapeutic implications. International Journal of Cancer, 2012, 131, 244-252.	2.3	42
86	NADPH Oxidases as Therapeutic Targets in Chronic Myelogenous Leukemia. Clinical Cancer Research, 2014, 20, 4014-4025.	3.2	42
87	ERK2, but Not ERK1, Mediates Acquired and "De novo―Resistance to Imatinib Mesylate: Implication for CML Therapy. PLoS ONE, 2009, 4, e6124.	1.1	41
88	Targeting receptor tyrosine kinases and their signal transduction routes in head and neck cancer. Annals of Oncology, 2007, 18, 421-430.	0.6	40
89	Breast Cancer Heterogeneity and Response to Novel Therapeutics. Cancers, 2020, 12, 3271.	1.7	40
90	Therapeutic potential of ERK5 targeting in triple negative breast cancer. Oncotarget, 2014, 5, 11308-11318.	0.8	40

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91	Targeting HER Receptors in Cancer. Current Pharmaceutical Design, 2013, 19, 808-817.	0.9	39
92	Mechanism of apoptosis induced by IFN-α in human myeloma cells: Role of Jak1 and Bim and potentiation by rapamycin. Cellular Signalling, 2007, 19, 844-854.	1.7	38
93	Prognostic Value of Lymphocyte-Activation Gene 3 (LAG3) in Cancer: A Meta-Analysis. Frontiers in Oncology, 2019, 9, 1040.	1.3	38
94	Effect of p95HER2/611CTF on the Response to Trastuzumab and Chemotherapy. Journal of the National Cancer Institute, 2014, 106, .	3.0	36
95	Proteolysis targeting chimeras (PROTACs) in cancer therapy. Journal of Experimental and Clinical Cancer Research, 2020, 39, 189.	3.5	36
96	Plasma membrane hyperpolarization and [Ca2+]i increase induced by fibroblast growth factor in NIH-3T3 fibroblasts: Resemblance to early signals generated by platelet-derived growth factor. Biochemical and Biophysical Research Communications, 1989, 163, 1325-1331.	1.0	35
97	Erk5 is activated and acts as a survival factor in mitosis. Cellular Signalling, 2007, 19, 1964-1972.	1.7	35
98	The insulin-like growth factor-I receptor inhibitor NVP-AEW541 provokes cell cycle arrest and apoptosis in multiple myeloma cells. British Journal of Haematology, 2008, 141, 470-482.	1.2	35
99	CD20 positive cells are undetectable in the majority of multiple myeloma cell lines and are not associated with a cancer stem cell phenotype. Haematologica, 2012, 97, 1110-1114.	1.7	34
100	β-Lapachone analogs with enhanced antiproliferative activity. European Journal of Medicinal Chemistry, 2012, 53, 264-274.	2.6	34
101	Prognostic relevance of receptor tyrosine kinase expression in breast cancer: A meta-analysis. Cancer Treatment Reviews, 2014, 40, 1048-1055.	3.4	34
102	Expression of MHC class I, HLA-A and HLA-B identifies immune-activated breast tumors with favorable outcome. Oncolmmunology, 2019, 8, e1629780.	2.1	34
103	Efficacy and safety of dasatinib with trastuzumab and paclitaxel in first line HER2-positive metastatic breast cancer: results from the phase II GEICAM/2010-04 study. Breast Cancer Research and Treatment, 2019, 174, 693-701.	1.1	34
104	Deficient Spindle Assembly Checkpoint in Multiple Myeloma. PLoS ONE, 2011, 6, e27583.	1.1	33
105	Overexpression of HER2 signaling to WAVE2–Arp2/3 complex activates MMP-independent migration in breast cancer. Breast Cancer Research and Treatment, 2011, 126, 311-318.	1.1	33
106	BET inhibitors as novel therapeutic agents in breast cancer. Oncotarget, 2017, 8, 71285-71291.	0.8	33
107	Tumor-Infiltrating Lymphocytes in Breast Cancer: Ready for Prime Time?. Journal of Clinical Oncology, 2015, 33, 1298-1299.	0.8	32
108	Trastuzumab and Antiestrogen Therapy. American Journal of Clinical Oncology: Cancer Clinical Trials, 2006, 29, 90-95.	0.6	31

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109	Expression of c-Kit isoforms in multiple myeloma: differences in signaling and drug sensitivity. Haematologica, 2008, 93, 851-859.	1.7	31
110	Interaction between Hormonal Receptor Status, Age and Survival in Patients with BRCA1/2 Germline Mutations: A Systematic Review and Meta-Regression. PLoS ONE, 2016, 11, e0154789.	1.1	31
111	A new method for detecting TNF-α-secreting cells using direct-immunofluorescence surface membrane stainings. Journal of Immunological Methods, 2002, 264, 77-87.	0.6	30
112	Circulating DNA and Survival in Solid Tumors. Cancer Epidemiology Biomarkers and Prevention, 2016, 25, 399-406.	1.1	30
113	HER3 targeting with an antibodyâ€drug conjugate bypasses resistance to antiâ€HER2 therapies. EMBO Molecular Medicine, 2020, 12, e11498.	3.3	30
114	Novel ADCs and Strategies to Overcome Resistance to Anti-HER2 ADCs. Cancers, 2022, 14, 154.	1.7	30
115	Erk5 nuclear location is independent on dual phosphorylation, and favours resistance to TRAIL-induced apoptosis. Cellular Signalling, 2007, 19, 1473-1487.	1.7	29
116	Transcriptomic immunologic signature associated with favorable clinical outcome in basal-like breast tumors. PLoS ONE, 2017, 12, e0175128.	1.1	28
117	Influence of companion diagnostics on efficacy and safety of targeted anti-cancer drugs: systematic review and meta-analyses. Oncotarget, 2015, 6, 39538-39549.	0.8	27
118	Pemetrexed acts as an antimyeloma agent by provoking cell cycle blockade and apoptosis. Leukemia, 2007, 21, 797-804.	3.3	26
119	Intracellular Calcium Homeostasis in a Human Neuroblastoma Cell Line: Modulation by Depolarization, Cholinergic Receptors, and ?-Latrotoxin. Journal of Neurochemistry, 1988, 50, 1708-1713.	2.1	25
120	Oleic Acid Blocks Epidermal Growth Factor-Activated Early Intracellular Signals without Altering the Ensuing Mitogenic Response. Experimental Cell Research, 1993, 205, 365-373.	1.2	24
121	Signalling-competent truncated forms of ErbB2 in breast cancer cells: differential regulation by protein kinase C and phosphatidylinositol 3-kinase. Biochemical Journal, 1999, 344, 339-348.	1.7	24
122	Molecular Pathways: P-Rex in Cancer. Clinical Cancer Research, 2013, 19, 4564-4569.	3.2	24
123	A phase I study of the SRC kinase inhibitor dasatinib with trastuzumab and paclitaxel as first line therapy for patients with HER2-overexpressing advanced breast cancer. CEICAM/2010-04 study. Oncotarget, 2017, 8, 73144-73153.	0.8	24
124	A Transcriptomic Immunologic Signature Predicts Favorable Outcome in Neoadjuvant Chemotherapy Treated Triple Negative Breast Tumors. Frontiers in Immunology, 2019, 10, 2802.	2.2	24
125	Antitumor activity of the novel multi-kinase inhibitor EC-70124 in triple negative breast cancer. Oncotarget, 2015, 6, 27923-27937.	0.8	24
126	Protein kinase C-mediated feed back inhibition of the Ca2+ response at the EGF receptor. Biochemical and Biophysical Research Communications, 1987, 149, 145-151.	1.0	23

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127	The Extracellular Linker of pro-Neuregulin-α2c Is Required for Efficient Sorting and Juxtacrine Function. Molecular Biology of the Cell, 2007, 18, 380-393.	0.9	23
128	The mitogen-activated protein kinase Erk5 mediates human mesangial cell activation. Nephrology Dialysis Transplantation, 2008, 23, 3403-3411.	0.4	23
129	<i>In Silico</i> Analysis Guides Selection of BET Inhibitors for Triple-Negative Breast Cancer Treatment. Molecular Cancer Therapeutics, 2016, 15, 1823-1833.	1.9	23
130	Targeting basal-like breast tumors with bromodomain and extraterminal domain (BET) and polo-like kinase inhibitors. Oncotarget, 2017, 8, 19478-19490.	0.8	23
131	α1-Adrenergic Stimulation ofin VitroGrowth Hormone Release and Cytosolic Free Ca2+in Rat Somatotrophs*. Endocrinology, 1988, 122, 1419-1425.	1.4	22
132	Autophagy inhibition sensitizes multiple myeloma cells to 17-dimethylaminoethylamino-17-demethoxygeldanamycin-induced apoptosis. Leukemia Research, 2010, 34, 1533-1538.	0.4	22
133	Genomic Mapping Identifies Mutations in RYR2 and AHNAK as Associated with Favorable Outcome in Basal-Like Breast Tumors Expressing PD1/PD-L1. Cancers, 2020, 12, 2243.	1.7	22
134	<i>In silico</i> analyses identify gene-sets, associated with clinical outcome in ovarian cancer: role of mitotic kinases. Oncotarget, 2016, 7, 22865-22872.	0.8	21
135	Neuregulin expression in solid tumors: Prognostic value and predictive role to anti-HER3 therapies. Oncotarget, 2016, 7, 45042-45051.	0.8	21
136	Enhancement of antiproliferative activity by molecular simplification of catalpol. Bioorganic and Medicinal Chemistry, 2010, 18, 2515-2523.	1.4	20
137	The evolving landscape of protein kinases in breast cancer: Clinical implications. Cancer Treatment Reviews, 2013, 39, 68-76.	3.4	20
138	Genomic Signatures of Immune Activation Predict Outcome in Advanced Stages of Ovarian Cancer and Basal-Like Breast Tumors. Frontiers in Oncology, 2019, 9, 1486.	1.3	20
139	Stimulation of cleavage of membrane proteins by calmodulin inhibitors. Biochemical Journal, 2000, 346, 359.	1.7	19
140	N-terminal cleavage of proTGFα occurs at the cell surface by a TACE-independent activity. Biochemical Journal, 2005, 389, 161-172.	1.7	19
141	Mitotic Arrest Induced by a Novel Family of DNA Topoisomerase II Inhibitors. Journal of Medicinal Chemistry, 2010, 53, 3835-3839.	2.9	18
142	Effect of Oncoxin Oral Solution in HER2-Overexpressing Breast Cancer. Nutrition and Cancer, 2015, 67, 1159-1169.	0.9	18
143	Breast cancer dissemination promoted by a neuregulin-collagenase 3 signalling node. Oncogene, 2016, 35, 2756-2765.	2.6	18
144	Mitotic read-out genes confer poor outcome in luminal A breast cancer tumors. Oncotarget, 2017, 8, 21733-21740.	0.8	18

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145	Novel Tyrosine Kinase Inhibitors in the Treatment of Cancer. Current Drug Targets, 2009, 10, 575-576.	1.0	18
146	The Activation of the Sox2 RR2 Pluripotency Transcriptional Reporter in Human Breast Cancer Cell Lines is Dynamic and Labels Cells with Higher Tumorigenic Potential. Frontiers in Oncology, 2014, 4, 308.	1.3	17
147	Antitumoral effect of Ocoxin on acute myeloid leukemia. Oncotarget, 2016, 7, 6231-6242.	0.8	17
148	Refining Early Antitumoral Drug Development. Trends in Pharmacological Sciences, 2018, 39, 922-925.	4.0	17
149	Antitumoral activity of the mithralog EC-8042 in triple negative breast cancer linked to cell cycle arrest in G2. Oncotarget, 2015, 6, 32856-32867.	0.8	17
150	ErbBs inhibition by lapatinib blocks tumor growth in an orthotopic model of human testicular germ cell tumor. International Journal of Cancer, 2013, 133, 235-246.	2.3	16
151	Phosphorylation of P-Rex1 at serine 1169 participates in IGF-1R signaling in breast cancer cells. Cellular Signalling, 2013, 25, 2281-2289.	1.7	16
152	Genetic mutational status of genes regulating epigenetics: Role of the histone methyltransferase KMT2D in triple negative breast tumors. PLoS ONE, 2019, 14, e0209134.	1.1	16
153	In silico transcriptomic mapping of integrins and immune activation in Basal-like and HER2+ breast cancer. Cellular Oncology (Dordrecht), 2021, 44, 569-580.	2.1	16
154	Clinical, genetic and pharmacological data support targeting the MEK5/ERK5 module in lung cancer. Npj Precision Oncology, 2021, 5, 78.	2.3	16
155	Identification of therapeutic targets in ovarian cancer through active tyrosine kinase profiling. Oncotarget, 2015, 6, 30057-30071.	0.8	15
156	Functional transcriptomic annotation and protein–protein interaction analysis identify <scp>EZH</scp> 2 and <scp>UBE</scp> 2C as key upregulated proteins in ovarian cancer. Cancer Medicine, 2018, 7, 1896-1907.	1.3	14
157	Dual targeting of HER2-positive breast cancer with trastuzumab emtansine and pertuzumab: understanding clinical trial results. Oncotarget, 2018, 9, 31915-31919.	0.8	14
158	Prognostic value of receptor tyrosine kinase-like orphan receptor (ROR) family in cancer: A meta-analysis. Cancer Treatment Reviews, 2019, 77, 11-19.	3.4	14
159	Antimyeloma Efficacy of Plitidepsin (Aplidin®): From Bench to the Bedside Blood, 2007, 110, 1178-1178.	0.6	14
160	Zalypsis has in vitro activity in acute myeloid blasts and leukemic progenitor cells through the induction of a DNA damage response. Haematologica, 2011, 96, 687-695.	1.7	13
161	Inhibition of the mitotic kinase PLK1 overcomes therapeutic resistance to BET inhibitors in triple negative breast cancer. Cancer Letters, 2020, 491, 50-59.	3.2	13
162	DNA-damage related genes and clinical outcome in hormone receptor positive breast cancer. Oncotarget, 2017, 8, 62834-62841.	0.8	13

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163	Do We Have to Change the Way Targeted Drugs Are Developed?. Journal of Clinical Oncology, 2010, 28, e420-e421.	0.8	12
164	Paclitaxel-Trastuzumab Mixed Nanovehicle to Target HER2-Overexpressing Tumors. Nanomaterials, 2019, 9, 948.	1.9	12
165	TRAIL receptor activation overcomes resistance to trastuzumab in HER2 positive breast cancer cells. Cancer Letters, 2019, 453, 34-44.	3.2	12
166	Pharmacological screening and transcriptomic functional analyses identify a synergistic interaction between dasatinib and olaparib in tripleâ€negative breast cancer. Journal of Cellular and Molecular Medicine, 2020, 24, 3117-3127.	1.6	12
167	Preclinical and Clinical Characterization of Fibroblast-derived Neuregulin-1 on Trastuzumab and Pertuzumab Activity in HER2-positive Breast Cancer. Clinical Cancer Research, 2021, 27, 5096-5108.	3.2	12
168	Transforming growth factor- $\hat{l}\pm$ . Biochemical Society Transactions, 1991, 19, 259-262.	1.6	11
169	RAF265, a dual BRAF and VEGFR2 inhibitor, prevents osteoclast formation and resorption. Therapeutic implications. Investigational New Drugs, 2013, 31, 200-205.	1.2	11
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