

Juan Carlos Montero

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

54
papers

2,067
citations

26
h-index

45
g-index

55
ext. papers

2,288
ext. citations

6
avg, IF

4.4
L-index

#	Paper	IF	Citations
54	Surfaceome analyses uncover CD98hc as an antibody drug-conjugate target in triple negative breast cancer.. <i>Journal of Experimental and Clinical Cancer Research</i> , 2022 , 41, 106	12.8	0
53	JKST6, a novel multikinase modulator of the BCR-ABL1/STAT5 signaling pathway that potentiates direct BCR-ABL1 inhibition and overcomes imatinib resistance in chronic myelogenous leukemia. <i>Biomedicine and Pharmacotherapy</i> , 2021 , 144, 112330	7.5	0
52	MZ1 co-operates with trastuzumab in HER2 positive breast cancer. <i>Journal of Experimental and Clinical Cancer Research</i> , 2021 , 40, 106	12.8	4
51	PDCD4 limits prooncogenic neuregulin-ErbB signaling. <i>Cellular and Molecular Life Sciences</i> , 2021 , 78, 1799-1815	10.3	2
50	Altered proTGF β cleaved TGF β ratios offer new therapeutic strategies in renal carcinoma. <i>Journal of Experimental and Clinical Cancer Research</i> , 2021 , 40, 256	12.8	0
49	Preclinical and Clinical Characterization of Fibroblast-derived Neuregulin-1 on Trastuzumab and Pertuzumab Activity in HER2-positive Breast Cancer. <i>Clinical Cancer Research</i> , 2021 , 27, 5096-5108	12.9	2
48	Pharmacological screening and transcriptomic functional analyses identify a synergistic interaction between dasatinib and olaparib in triple-negative breast cancer. <i>Journal of Cellular and Molecular Medicine</i> , 2020 , 24, 3117-3127	5.6	6
47	Inhibition of the mitotic kinase PLK1 overcomes therapeutic resistance to BET inhibitors in triple negative breast cancer. <i>Cancer Letters</i> , 2020 , 491, 50-59	9.9	5
46	Activity of BET-proteolysis targeting chimeric (PROTAC) compounds in triple negative breast cancer. <i>Journal of Experimental and Clinical Cancer Research</i> , 2019 , 38, 383	12.8	32
45	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. <i>Breast Cancer Research and Treatment</i> , 2019 , 174, 693-701	4.4	18
44	The immunoglobulin-like domain of neuregulins potentiates ErbB3/HER3 activation and cellular proliferation. <i>Molecular Oncology</i> , 2018 , 12, 1061-1076	7.9	4
43	Regulation of the prometastatic neuregulin-MMP13 axis by SRC family kinases: therapeutic implications. <i>Molecular Oncology</i> , 2017 , 11, 1788-1805	7.9	5
42	Synthetic Lethality Interaction Between Aurora Kinases and CHEK1 Inhibitors in Ovarian Cancer. <i>Molecular Cancer Therapeutics</i> , 2017 , 16, 2552-2562	6.1	32
41	CM363, a novel naphthoquinone derivative which acts as multikinase modulator and overcomes imatinib resistance in chronic myelogenous leukemia. <i>Oncotarget</i> , 2017 , 8, 29679-29698	3.3	8
40	Targeting basal-like breast tumors with bromodomain and extraterminal domain (BET) and polo-like kinase inhibitors. <i>Oncotarget</i> , 2017 , 8, 19478-19490	3.3	22
39	In Silico Analysis Guides Selection of BET Inhibitors for Triple-Negative Breast Cancer Treatment. <i>Molecular Cancer Therapeutics</i> , 2016 , 15, 1823-33	6.1	20
38	Multisite phosphorylation of P-Rex1 by protein kinase C. <i>Oncotarget</i> , 2016 , 7, 77937-77949	3.3	6

37	Neuregulin expression in solid tumors: prognostic value and predictive role to anti-HER3 therapies. <i>Oncotarget</i> , 2016 , 7, 45042-45051	3.3	14
36	Targeting the EGF/HER Ligand-Receptor System in Cancer. <i>Current Pharmaceutical Design</i> , 2016 , 22, 5887-5898	3.5	35
35	Antitumor activity of the novel multi-kinase inhibitor EC-70124 in triple negative breast cancer. <i>Oncotarget</i> , 2015 , 6, 27923-37	3.3	19
34	Identification of therapeutic targets in ovarian cancer through active tyrosine kinase profiling. <i>Oncotarget</i> , 2015 , 6, 30057-71	3.3	13
33	Phospho-kinase profile of colorectal tumors guides in the selection of multi-kinase inhibitors. <i>Oncotarget</i> , 2015 , 6, 31272-83	3.3	6
32	Antitumoral activity of the mithralog EC-8042 in triple negative breast cancer linked to cell cycle arrest in G2. <i>Oncotarget</i> , 2015 , 6, 32856-67	3.3	15
31	Phospho-kinase profile of triple negative breast cancer and androgen receptor signaling. <i>BMC Cancer</i> , 2014 , 14, 302	4.8	37
30	AchillesHeel of triple negative cancer. <i>Oncoscience</i> , 2014 , 1, 763-4	0.8	2
29	Phosphorylation of P-Rex1 at serine 1169 participates in IGF-1R signaling in breast cancer cells. <i>Cellular Signalling</i> , 2013 , 25, 2281-9	4.9	14
28	Molecular pathways: P-Rex in cancer. <i>Clinical Cancer Research</i> , 2013 , 19, 4564-9	12.9	20
27	Elapachone analogs with enhanced antiproliferative activity. <i>European Journal of Medicinal Chemistry</i> , 2012 , 53, 264-74	6.8	29
26	Predominance of mTORC1 over mTORC2 in the regulation of proliferation of ovarian cancer cells: therapeutic implications. <i>Molecular Cancer Therapeutics</i> , 2012 , 11, 1342-52	6.1	38
25	Inhibition of SRC family kinases and receptor tyrosine kinases by dasatinib: possible combinations in solid tumors. <i>Clinical Cancer Research</i> , 2011 , 17, 5546-52	12.9	204
24	Transautocrine signaling by membrane neuregulins requires cell surface targeting, which is controlled by multiple domains. <i>Journal of Biological Chemistry</i> , 2011 , 286, 24350-63	5.4	3
23	Effect of multikinase inhibitors on caspase-independent cell death and DNA damage in HER2-overexpressing breast cancer cells. <i>Journal of the National Cancer Institute</i> , 2010 , 102, 1432-46	9.7	39
22	Mitotic arrest induced by a novel family of DNA topoisomerase II inhibitors. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 3835-9	8.3	17
21	Enhancement of antiproliferative activity by molecular simplification of catalpol. <i>Bioorganic and Medicinal Chemistry</i> , 2010 , 18, 2515-23	3.4	16
20	Expression of Erk5 in early stage breast cancer and association with disease free survival identifies this kinase as a potential therapeutic target. <i>PLoS ONE</i> , 2009 , 4, e5565	3.7	76

19	Zalypsis: a novel marine-derived compound with potent antimyeloma activity that reveals high sensitivity of malignant plasma cells to DNA double-strand breaks. <i>Blood</i> , 2009 , 113, 3781-91	2.2	66
18	The insulin-like growth factor-I receptor inhibitor NVP-AEW541 provokes cell cycle arrest and apoptosis in multiple myeloma cells. <i>British Journal of Haematology</i> , 2008 , 141, 470-82	4.5	33
17	Neuregulins and cancer. <i>Clinical Cancer Research</i> , 2008 , 14, 3237-41	12.9	81
16	The effect of the proteasome inhibitor bortezomib on acute myeloid leukemia cells and drug resistance associated with the CD34+ immature phenotype. <i>Haematologica</i> , 2008 , 93, 57-66	6.6	50
15	Aplidin, a marine organism-derived compound with potent antimyeloma activity in vitro and in vivo. <i>Cancer Research</i> , 2008 , 68, 5216-25	10.1	79
14	Expression of c-Kit isoforms in multiple myeloma: differences in signaling and drug sensitivity. <i>Haematologica</i> , 2008 , 93, 851-9	6.6	26
13	Erk5 is activated and acts as a survival factor in mitosis. <i>Cellular Signalling</i> , 2007 , 19, 1964-72	4.9	26
12	Neuregulin expression modulates clinical response to trastuzumab in patients with metastatic breast cancer. <i>Journal of Clinical Oncology</i> , 2007 , 25, 2656-63	2.2	51
11	The extracellular linker of pro-neuregulin-alpha2c is required for efficient sorting and juxtacrine function. <i>Molecular Biology of the Cell</i> , 2007 , 18, 380-93	3.5	22
10	Multifunctional role of Erk5 in multiple myeloma. <i>Blood</i> , 2005 , 105, 4492-9	2.2	70
9	N-terminal cleavage of proTGFalpha occurs at the cell surface by a TACE-independent activity. <i>Biochemical Journal</i> , 2005 , 389, 161-72	3.8	17
8	Activation of ErbB2 by overexpression or by transmembrane neuregulin results in differential signaling and sensitivity to herceptin. <i>Cancer Research</i> , 2005 , 65, 6801-10	10.1	59
7	Erk5 participates in neuregulin signal transduction and is constitutively active in breast cancer cells overexpressing ErbB2. <i>Molecular and Cellular Biology</i> , 2002 , 22, 270-85	4.8	144
6	Extracellular signal-regulated kinase phosphorylates tumor necrosis factor alpha-converting enzyme at threonine 735: a potential role in regulated shedding. <i>Molecular Biology of the Cell</i> , 2002 , 13, 2031-44	3.5	251
5	Mitogen-activated protein kinase-dependent and -independent routes control shedding of transmembrane growth factors through multiple secretases. <i>Biochemical Journal</i> , 2002 , 363, 211-21	3.8	37
4	Mitogen-activated protein kinase-dependent and -independent routes control shedding of transmembrane growth factors through multiple secretases. <i>Biochemical Journal</i> , 2002 , 363, 211-221	3.8	47
3	Stimulation of cleavage of membrane proteins by calmodulin inhibitors. <i>Biochemical Journal</i> , 2000 , 346, 359-367	3.8	57
2	Differential shedding of transmembrane neuregulin isoforms by the tumor necrosis factor-alpha-converting enzyme. <i>Molecular and Cellular Neurosciences</i> , 2000 , 16, 631-48	4.8	147

- 1 Cleavage of the TrkA neurotrophin receptor by multiple metalloproteases generates signalling-competent truncated forms. *European Journal of Neuroscience*, **1999**, 11, 1421-30 3·5 4¹