Juan Carlos Montero

List of Publications by Citations

Source: https://exaly.com/author-pdf/9466393/juan-carlos-montero-publications-by-citations.pdf

Version: 2024-04-23

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

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

54
papers2,067
citations26
h-index45
g-index55
ext. papers2,288
ext. citations6
avg, IF4.4
L-index

#	Paper	IF	Citations
54	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
53	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
52	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
51	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
50	Neuregulins and cancer. Clinical Cancer Research, 2008, 14, 3237-41	12.9	81
49	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
48	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
47	Multifunctional role of Erk5 in multiple myeloma. <i>Blood</i> , 2005 , 105, 4492-9	2.2	70
46	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
45	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
44	Stimulation of cleavage of membrane proteins by calmodulin inhibitors. <i>Biochemical Journal</i> , 2000 , 346, 359-367	3.8	57
43	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
42	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
41	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
40	Cleavage of the TrkA neurotrophin receptor by multiple metalloproteases generates signalling-competent truncated forms. <i>European Journal of Neuroscience</i> , 1999 , 11, 1421-30	3.5	41
39	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
38	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

(2010-2014)

Phospho-kinase profile of triple negative breast cancer and androgen receptor signaling. <i>BMC Cancer</i> , 2014 , 14, 302	4.8	37
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
Targeting the EGF/HER Ligand-Receptor System in Cancer. Current Pharmaceutical Design, 2016, 22, 58	8 3. 5 89	835
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
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
Synthetic Lethality Interaction Between Aurora Kinases and CHEK1 Inhibitors in Ovarian Cancer. <i>Molecular Cancer Therapeutics</i> , 2017 , 16, 2552-2562	6.1	32
Lapachone analogs with enhanced antiproliferative activity. <i>European Journal of Medicinal Chemistry</i> , 2012 , 53, 264-74	6.8	29
Expression of c-Kit isoforms in multiple myeloma: differences in signaling and drug sensitivity. Haematologica, 2008 , 93, 851-9	6.6	26
Erk5 is activated and acts as a survival factor in mitosis. Cellular Signalling, 2007, 19, 1964-72	4.9	26
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
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
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
Molecular pathways: P-Rex in cancer. Clinical Cancer Research, 2013, 19, 4564-9	12.9	20
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
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
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
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
Enhancement of antiproliferative activity by molecular simplification of catalpol. <i>Bioorganic and Medicinal Chemistry</i> , 2010 , 18, 2515-23	3.4	16
	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 Targeting the EGF/HER Ligand-Receptor System in Cancer. <i>Current Pharmaceutical Design</i> , 2016, 22, 58 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 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 Synthetic Lethality interaction Between Aurora Kinases and CHEK1 Inhibitors in Ovarian Cancer. <i>Molecular Cancer Therapeutics</i> , 2017, 16, 2552-2562 Lapachone analogs with enhanced antiproliferative activity. <i>European Journal of Medicinal Chemistry</i> , 2012, 53, 264-74 Expression of c-Kit isoforms in multiple myeloma: differences in signaling and drug sensitivity. <i>Haematologica</i> , 2008, 93, 851-9 ErkS is activated and acts as a survival factor in mitosis. <i>Cellular Signalling</i> , 2007, 19, 1964-72 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 Targeting basal-like breast tumors with bromodomain and extraterminal domain (BET) and polo-like kinase inhibitors. <i>Oncotarget</i> , 2017, 8, 19478-19490 In Silico Analysis Guides Selection of BET Inhibitors for Triple-Negative Breast Cancer Treatment. <i>Molecular Cancer Therapeutics</i> , 2016, 15, 1823-33 Molecular pathways: P-Rex in cancer. <i>Clinical Cancer Research</i> , 2013, 19, 4564-9 Antitumor activity of the novel multi-kinase inhibitor EC-70124 in triple negative breast cancer. <i>Oncotarget</i> , 2015, 6, 27923-37 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, 69	Mitogen-activated protein kinase-dependent and -independent routes control shedding of transmembrane growth factors through multiple secretases. Biochemical Journal, 2002, 363, 211-21 Targeting the ECF/HER Ligand-Receptor System in Cancer. Current Pharmaceutical Design, 2016, 22, 5883-589. 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, 470-82 Activity of BET-proteolysis targeting chimeric (PROTAC) compounds in triple negative breast cancer. Journal of Experimental and Clinical Cancer Research, 2019, 38, 383 Synthetic Lethality interaction Between Aurora Kinases and CHEK1 Inhibitors in Ovarian Cancer. Molecular Cancer Therapeutics, 2017, 16, 2552-2562 Blapachone analogs with enhanced antiproliferative activity. European Journal of Medicinal Chemistry, 2012, 53, 264-74 Expression of c-Kit isoforms in multiple myeloma: differences in signaling and drug sensitivity. Haematologica, 2008, 93, 851-9 ErkS is activated and acts as a survival factor in mitosis. Cellular Signalling, 2007, 19, 1964-72 The extracellular linker of pro-neuregulin-alpha2c is required for efficient sorting and juxtacrine function. Molecular Biology of the Cell, 2007, 18, 380-93 Targeting basal-like breast tumors with bromodomain and extraterminal domain (BET) and polo-like kinase inhibitors. Oncotarget, 2017, 8, 19478-19490 In Silico Analysis Guides Selection of BET Inhibitors for Triple-Negative Breast Cancer Treatment. Molecular Cancer Therapeutics, 2016, 15, 1823-33 Molecular pathways: P-Rex in cancer. Clinical Cancer Research, 2013, 19, 4564-9 12-9 Antitumor activity of the novel multi-kinase inhibitor EC-70124 in triple negative breast cancer. Oncotarget, 2015, 6, 27923-37 Mitotic arrest induced by a novel family of DNA topoisomerase II inhibitors. Journal of Medicinal Chemistry, 2010, 53, 3833-9 N-terminal cleavage of proTGFalpha occurs at the cell surface by a TACE-independent activity. Biochemical

19	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
18	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
17	Neuregulin expression in solid tumors: prognostic value and predictive role to anti-HER3 therapies. <i>Oncotarget</i> , 2016 , 7, 45042-45051	3.3	14
16	Identification of therapeutic targets in ovarian cancer through active tyrosine kinase profiling. <i>Oncotarget</i> , 2015 , 6, 30057-71	3.3	13
15	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
14	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
13	Multisite phosphorylation of P-Rex1 by protein kinase C. <i>Oncotarget</i> , 2016 , 7, 77937-77949	3.3	6
12	Phospho-kinase profile of colorectal tumors guides in the selection of multi-kinase inhibitors. <i>Oncotarget</i> , 2015 , 6, 31272-83	3.3	6
11	Regulation of the prometastatic neuregulin-MMP13 axis by SRC family kinases: therapeutic implications. <i>Molecular Oncology</i> , 2017 , 11, 1788-1805	7.9	5
10	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
9	The immunoglobulin-like domain of neuregulins potentiates ErbB3/HER3 activation and cellular proliferation. <i>Molecular Oncology</i> , 2018 , 12, 1061-1076	7.9	4
8	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
7	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
6	AchillesTheel of triple negative cancer. <i>Oncoscience</i> , 2014 , 1, 763-4	0.8	2
5	PDCD4 limits prooncogenic neuregulin-ErbB signaling. <i>Cellular and Molecular Life Sciences</i> , 2021 , 78, 1799-1815	10.3	2
4	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
3	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	O
2	Altered proTGFIcleaved TGFIratios offer new therapeutic strategies in renal carcinoma. <i>Journal of Experimental and Clinical Cancer Research</i> , 2021 , 40, 256	12.8	0

Surfaceome analyses uncover CD98hc as an antibody drug-conjugate target in triple negative breast cancer.. *Journal of Experimental and Clinical Cancer Research*, **2022**, 41, 106

12.8 0