

# Silvia La Monica

## List of Publications by Year in descending order

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Version: 2024-02-01

47  
papers

1,914  
citations

186209

28  
h-index

254106

43  
g-index

47  
all docs

47  
docs citations

47  
times ranked

3080  
citing authors

#	ARTICLE	IF	CITATIONS
1	Human myeloma cells stimulate the receptor activator of nuclear factor- $\kappa$ B ligand (RANKL) in T lymphocytes: a potential role in multiple myeloma bone disease. <i>Blood</i> , 2002, 100, 4615-4621.	0.6	241
2	Combined Inhibition of CDK4/6 and PI3K/AKT/mTOR Pathways Induces a Synergistic Anti-Tumor Effect in Malignant Pleural Mesothelioma Cells. <i>Neoplasia</i> , 2017, 19, 637-648.	2.3	81
3	Overcoming acquired resistance to letrozole by targeting the PI3K/AKT/mTOR pathway in breast cancer cell clones. <i>Cancer Letters</i> , 2012, 323, 77-87.	3.2	78
4	Everolimus restores gefitinib sensitivity in resistant non-small cell lung cancer cell lines. <i>Biochemical Pharmacology</i> , 2009, 78, 460-468.	2.0	71
5	Trastuzumab emtansine delays and overcomes resistance to the third-generation EGFR-TKI osimertinib in NSCLC EGFR mutated cell lines. <i>Journal of Experimental and Clinical Cancer Research</i> , 2017, 36, 174.	3.5	70
6	The anti-tumor efficacy of CDK4/6 inhibition is enhanced by the combination with PI3K/AKT/mTOR inhibitors through impairment of glucose metabolism in TNBC cells. <i>Journal of Experimental and Clinical Cancer Research</i> , 2018, 37, 72.	3.5	68
7	Effects of sorafenib on energy metabolism in breast cancer cells: role of AMPK and mTORC1 signaling. <i>Breast Cancer Research and Treatment</i> , 2013, 141, 67-78.	1.1	65
8	Multiple effects of CDK4/6 inhibition in cancer: From cell cycle arrest to immunomodulation. <i>Biochemical Pharmacology</i> , 2019, 170, 113676.	2.0	64
9	Pre-treatment with the CDK4/6 inhibitor palbociclib improves the efficacy of paclitaxel in TNBC cells. <i>Scientific Reports</i> , 2019, 9, 13014.	1.6	62
10	Amino acid signaling through the mammalian target of rapamycin (mTOR) pathway: Role of glutamine and of cell shrinkage. <i>Journal of Cellular Physiology</i> , 2005, 204, 155-165.	2.0	61
11	Combination of Gefitinib and Pemetrexed Prevents the Acquisition of TKI Resistance in NSCLC Cell Lines Carrying EGFR- Activating Mutation. <i>Journal of Thoracic Oncology</i> , 2016, 11, 1051-1063.	0.5	58
12	Trastuzumab emtansine is active on HER-2 overexpressing NSCLC cell lines and overcomes gefitinib resistance. <i>Molecular Cancer</i> , 2014, 13, 143.	7.9	55
13	Combination of EGFR-TKIs and chemotherapy in advanced EGFR mutated NSCLC: Review of the literature and future perspectives. <i>Critical Reviews in Oncology/Hematology</i> , 2020, 146, 102820.	2.0	53
14	Effect of ABCG2/BCRP Expression on Efflux and Uptake of Gefitinib in NSCLC Cell Lines. <i>PLoS ONE</i> , 2015, 10, e0141795.	1.1	51
15	Novel Irreversible Epidermal Growth Factor Receptor Inhibitors by Chemical Modulation of the Cysteine-Trap Portion. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 2038-2050.	2.9	49
16	Third generation EGFR inhibitor osimertinib combined with pemetrexed or cisplatin exerts long-lasting anti-tumor effect in EGFR-mutated pre-clinical models of NSCLC. <i>Journal of Experimental and Clinical Cancer Research</i> , 2019, 38, 222.	3.5	45
17	Inhibition of PI3K Pathway Reduces Invasiveness and Epithelial-to-Mesenchymal Transition in Squamous Lung Cancer Cell Lines Harboring PIK3CA Gene Alterations. <i>Molecular Cancer Therapeutics</i> , 2015, 14, 1916-1927.	1.9	43
18	Gefitinib Inhibits Invasive Phenotype and Epithelial-Mesenchymal Transition in Drug-Resistant NSCLC Cells with MET Amplification. <i>PLoS ONE</i> , 2013, 8, e78656.	1.1	39

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19	New therapeutic strategies for malignant pleural mesothelioma. <i>Biochemical Pharmacology</i> , 2017, 123, 8-18.	2.0	38
20	MYC Amplification as a Potential Mechanism of Primary Resistance to Crizotinib in ALK-Rearranged Non-Small Cell Lung Cancer: A Brief Report. <i>Translational Oncology</i> , 2019, 12, 116-121.	1.7	37
21	Metabolism of the EGFR tyrosin kinase inhibitor gefitinib by cytochrome P450 1A1 enzyme in EGFR-wild type non small cell lung cancer cell lines. <i>Molecular Cancer</i> , 2011, 10, 143.	7.9	36
22	Synergistic activity of letrozole and sorafenib on breast cancer cells. <i>Breast Cancer Research and Treatment</i> , 2010, 124, 79-88.	1.1	35
23	Combined use of anti-ErbB monoclonal antibodies and erlotinib enhances antibody-dependent cellular cytotoxicity of wild-type erlotinib-sensitive NSCLC cell lines. <i>Molecular Cancer</i> , 2012, 11, 91.	7.9	35
24	Expanding the Arsenal of FGFR Inhibitors: A Novel Chloroacetamide Derivative as a New Irreversible Agent With Anti-proliferative Activity Against FGFR1-Amplified Lung Cancer Cell Lines. <i>Frontiers in Oncology</i> , 2019, 9, 179.	1.3	34
25	Acquired BRAF G469A Mutation as a Resistance Mechanism to First-Line Osimertinib Treatment in NSCLC Cell Lines Harboring an EGFR Exon 19 Deletion. <i>Targeted Oncology</i> , 2019, 14, 619-626.	1.7	33
26	Functional characterization of gefitinib uptake in non-small cell lung cancer cell lines. <i>Biochemical Pharmacology</i> , 2010, 80, 179-187.	2.0	31
27	Concurrent Acquired BRAF V600E Mutation and MET Amplification as Resistance Mechanism of First-Line Osimertinib Treatment in a Patient with EGFR-Mutated NSCLC. <i>Journal of Thoracic Oncology</i> , 2018, 13, e89-e91.	0.5	31
28	Efficacy of the CDK4/6 Dual Inhibitor Abemaciclib in EGFR-Mutated NSCLC Cell Lines with Different Resistance Mechanisms to Osimertinib. <i>Cancers</i> , 2021, 13, 6.	1.7	30
29	Antitumor Efficacy of Dual Blockade of EGFR Signaling by Osimertinib in Combination With Selumetinib or Cetuximab in Activated EGFR Human NCLC Tumor Models. <i>Journal of Thoracic Oncology</i> , 2018, 13, 810-820.	0.5	29
30	A sulfonyl fluoride derivative inhibits EGFR L858R/T790M/C797S by covalent modification of the catalytic lysine. <i>European Journal of Medicinal Chemistry</i> , 2021, 225, 113786.	2.6	28
31	Enhancement of the anti-tumor activity of FGFR1 inhibition in squamous cell lung cancer by targeting downstream signaling involved in glucose metabolism. <i>Oncotarget</i> , 2017, 8, 91841-91859.	0.8	28
32	Zoledronic acid determines S-phase arrest but fails to induce apoptosis in cholangiocarcinoma cells. <i>Biochemical Pharmacology</i> , 2009, 78, 133-141.	2.0	27
33	Small Cell Lung Cancer Transformation as a Resistance Mechanism to Osimertinib in Epidermal Growth Factor Receptor-Mutated Lung Adenocarcinoma: Case Report and Literature Review. <i>Frontiers in Oncology</i> , 2021, 11, 642190.	1.3	26
34	Epidermal Growth Factor Receptor Tyrosine Kinase Inhibitors: Current Status and Future Perspectives in the Development of Novel Irreversible Inhibitors for the Treatment of Mutant Non-small Cell Lung Cancer. <i>Current Pharmaceutical Design</i> , 2013, 19, 818-832.	0.9	24
35	Pemetrexed Enhances Membrane PD-L1 Expression and Potentiates T Cell-Mediated Cytotoxicity by Anti-PD-L1 Antibody Therapy in Non-Small-Cell Lung Cancer. <i>Cancers</i> , 2020, 12, 666.	1.7	24
36	Dual Inhibition of CDK4/6 and PI3K/AKT/mTOR Signaling Impairs Energy Metabolism in MPM Cancer Cells. <i>International Journal of Molecular Sciences</i> , 2020, 21, 5165.	1.8	21

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37	Reprogramming of Lipid Metabolism in Lung Cancer: An Overview with Focus on EGFR-Mutated Non-Small Cell Lung Cancer. <i>Cells</i> , 2022, 11, 413.	1.8	21
38	Enhanced efficacy of AKT and FAK kinase combined inhibition in squamous cell lung carcinomas with stable reduction in PTEN. <i>Oncotarget</i> , 2017, 8, 53068-53083.	0.8	19
39	Simultaneous Combination of the CDK4/6 Inhibitor Palbociclib With Regorafenib Induces Enhanced Anti-tumor Effects in Hepatocarcinoma Cell Lines. <i>Frontiers in Oncology</i> , 2020, 10, 563249.	1.3	18
40	Fighting tertiary mutations in EGFR-driven lung-cancers: Current advances and future perspectives in medicinal chemistry. <i>Biochemical Pharmacology</i> , 2021, 190, 114643.	2.0	11
41	Hematologic Malignancies With Extramedullary Spread of Disease. <i>Journal of Clinical Oncology</i> , 2003, 21, 1887-1888.	0.8	10
42	Afatinib therapy in case of EGFR G724S emergence as resistance mechanism to osimertinib. <i>Anti-Cancer Drugs</i> , 2021, 32, 758-762.	0.7	9
43	Inhibition of Human Malignant Pleural Mesothelioma Growth by Mesenchymal Stromal Cells. <i>Cells</i> , 2021, 10, 1427.	1.8	9
44	Generation and Characterization of a New Preclinical Mouse Model of EGFR-Driven Lung Cancer with MET-Induced Osimertinib Resistance. <i>Cancers</i> , 2021, 13, 3441.	1.7	8
45	Physico-chemical characterization and biological evaluation of two fibroin materials. <i>Journal of Tissue Engineering and Regenerative Medicine</i> , 2014, 8, 874-885.	1.3	4
46	YES1 and MYC Amplifications as Synergistic Resistance Mechanisms to Different Generation ALK Tyrosine Kinase Inhibitors in Advanced NSCLC: Brief Report of Clinical and Preclinical Proofs. <i>JTO Clinical and Research Reports</i> , 2022, 3, 100278.	0.6	3
47	EGFR Signaling in Non-Small Cell Lung Cancer: From Molecular Mechanisms to Therapeutic Opportunities. <i>Cells</i> , 2022, 11, 1344.	1.8	1