

# Martin L Sos

## List of Publications by Year in descending order

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Version: 2024-02-01

55  
papers

12,134  
citations

94433

37  
h-index

155660

55  
g-index

56  
all docs

56  
docs citations

56  
times ranked

18577  
citing authors

#	ARTICLE	IF	CITATIONS
1	Report of the First International Symposium on NUT Carcinoma. <i>Clinical Cancer Research</i> , 2022, 28, 2493-2505.	7.0	23
2	CD74-NRG1 Fusions Are Oncogenic <i>In Vivo</i> and Induce Therapeutically Tractable ERBB2:ERBB3 Heterodimerization. <i>Molecular Cancer Therapeutics</i> , 2022, 21, 821-830.	4.1	4
3	Insight into Targeting Exon20 Insertion Mutations of the Epidermal Growth Factor Receptor with Wild Type-Sparing Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 6643-6655.	6.4	12
4	The next tier of EGFR resistance mutations in lung cancer. <i>Oncogene</i> , 2021, 40, 1-11.	5.9	77
5	Ferroptosis response segregates small cell lung cancer (SCLC) neuroendocrine subtypes. <i>Nature Communications</i> , 2021, 12, 2048.	12.8	66
6	MAPK-pathway inhibition mediates inflammatory reprogramming and sensitizes tumors to targeted activation of innate immunity sensor RIG-I. <i>Nature Communications</i> , 2021, 12, 5505.	12.8	30
7	Clonal dynamics of BRAF-driven drug resistance in EGFR-mutant lung cancer. <i>Npj Precision Oncology</i> , 2021, 5, 102.	5.4	11
8	Sorafenib and everolimus in patients with advanced solid tumors and KRAS-mutated NSCLC: A phase I trial with early pharmacodynamic FDG-PET assessment. <i>Cancer Medicine</i> , 2020, 9, 4991-5007.	2.8	14
9	New Approaches to SCLC Therapy: From the Laboratory to the Clinic. <i>Journal of Thoracic Oncology</i> , 2020, 15, 520-540.	1.1	119
10	MYC paralog-dependent apoptotic priming orchestrates a spectrum of vulnerabilities in small cell lung cancer. <i>Nature Communications</i> , 2019, 10, 3485.	12.8	54
11	Genomic Profiling Identifies Outcome-Relevant Mechanisms of Innate and Acquired Resistance to Third-Generation Epidermal Growth Factor Receptor Tyrosine Kinase Inhibitor Therapy in Lung Cancer. <i>JCO Precision Oncology</i> , 2019, 3, 1-14.	3.0	17
12	K-ras Mutation Subtypes in NSCLC and Associated Co-occurring Mutations in Other Oncogenic Pathways. <i>Journal of Thoracic Oncology</i> , 2019, 14, 606-616.	1.1	178
13	Structural Alterations of MET Trigger Response to MET Kinase Inhibition in Lung Adenocarcinoma Patients. <i>Clinical Cancer Research</i> , 2018, 24, 1337-1343.	7.0	71
14	Loss of G2032R Resistance Mutation Upon Chemotherapy Treatment Enables Successful Crizotinib Rechallenge in a Patient With ROS1-Rearranged NSCLC. <i>JCO Precision Oncology</i> , 2018, 2, 1-6.	3.0	2
15	Overcoming EGFRG724S-mediated osimertinib resistance through unique binding characteristics of second-generation EGFR inhibitors. <i>Nature Communications</i> , 2018, 9, 4655.	12.8	107
16	MYC Drives Progression of Small Cell Lung Cancer to a Variant Neuroendocrine Subtype with Vulnerability to Aurora Kinase Inhibition. <i>Cancer Cell</i> , 2017, 31, 270-285.	16.8	406
17	Mechanisms of Primary Drug Resistance in FGFR1-Amplified Lung Cancer. <i>Clinical Cancer Research</i> , 2017, 23, 5527-5536.	7.0	44
18	Drugging the catalytically inactive state of RET kinase in RET-rearranged tumors. <i>Science Translational Medicine</i> , 2017, 9, .	12.4	55

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19	Systematic Kinase Inhibitor Profiling Identifies CDK9 as a Synthetic Lethal Target in NUT Midline Carcinoma. <i>Cell Reports</i> , 2017, 20, 2833-2845.	6.4	40
20	Family matters: How MYC family oncogenes impact small cell lung cancer. <i>Cell Cycle</i> , 2017, 16, 1489-1498.	2.6	75
21	Resistance Mechanisms to AZD9291 and Rociletinib Response. <i>Clinical Cancer Research</i> , 2017, 23, 3967-3968.	7.0	3
22	Heterogeneous Mechanisms of Primary and Acquired Resistance to Third-Generation EGFR Inhibitors. <i>Clinical Cancer Research</i> , 2016, 22, 4837-4847.	7.0	223
23	Implementation of Amplicon Parallel Sequencing Leads to Improvement of Diagnosis and Therapy of Lung Cancer Patients. <i>Journal of Thoracic Oncology</i> , 2015, 10, 1049-1057.	1.1	85
24	Spatial Tumor Heterogeneity in Lung Cancer with Acquired Epidermal Growth Factor Receptor-Tyrosine Kinase Inhibitor Resistance: Targeting High-Level MET-Amplification and EGFR T790M Mutation Occurring at Different Sites in the Same Patient. <i>Journal of Thoracic Oncology</i> , 2015, 10, e40-e43.	1.1	33
25	Squamous Cell Lung Cancer: From Tumor Genomics to Cancer Therapeutics. <i>Clinical Cancer Research</i> , 2015, 21, 2236-2243.	7.0	147
26	Discovery and functional characterization of a neomorphic PTEN mutation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, 13976-13981.	7.1	38
27	Targeting Drug Resistance in EGFR with Covalent Inhibitors: A Structure-Based Design Approach. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 6844-6863.	6.4	92
28	Oncogene Mimicry as a Mechanism of Primary Resistance to BRAF Inhibitors. <i>Cell Reports</i> , 2014, 8, 1037-1048.	6.4	69
29	Targeting Gain of Function and Resistance Mutations in Abl and KIT by Hybrid Compound Design. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 5757-5772.	6.4	17
30	A Neo-Substrate that Amplifies Catalytic Activity of Parkinson's-Disease-Related Kinase PINK1. <i>Cell</i> , 2013, 154, 737-747.	28.9	229
31	K-Ras(G12C) inhibitors allosterically control GTP affinity and effector interactions. <i>Nature</i> , 2013, 503, 548-551.	27.8	1,713
32	Characteristics of Lung Cancers Harboring <i>NRAS</i> Mutations. <i>Clinical Cancer Research</i> , 2013, 19, 2584-2591.	7.0	134
33	Staurosporine-Derived Inhibitors Broaden the Scope of Analog-Sensitive Kinase Technology. <i>Journal of the American Chemical Society</i> , 2013, 135, 18153-18159.	13.7	31
34	Differential Protein Stability and ALK Inhibitor Sensitivity of EML4-ALK Fusion Variants. <i>Clinical Cancer Research</i> , 2012, 18, 4682-4690.	7.0	252
35	Definition of a fluorescence in-situ hybridization score identifies high- and low-level FGFR1 amplification types in squamous cell lung cancer. <i>Modern Pathology</i> , 2012, 25, 1473-1480.	5.5	118
36	A framework for identification of actionable cancer genome dependencies in small cell lung cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 17034-17039.	7.1	167

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37	AATF/Che-1 acts as a phosphorylation-dependent molecular modulator to repress p53-driven apoptosis. <i>EMBO Journal</i> , 2012, 31, 3961-3975.	7.8	53
38	Integrative genome analyses identify key somatic driver mutations of small-cell lung cancer. <i>Nature Genetics</i> , 2012, 44, 1104-1110.	21.4	1,186
39	Optimization of Dosing for EGFR-Mutant Non-Small Cell Lung Cancer with Evolutionary Cancer Modeling. <i>Science Translational Medicine</i> , 2011, 3, 90ra59.	12.4	457
40	Mutations in the <i>DDR2</i> Kinase Gene Identify a Novel Therapeutic Target in Squamous Cell Lung Cancer. <i>Cancer Discovery</i> , 2011, 1, 78-89.	9.4	455
41	Synthesis and biological evaluation of 7-substituted-1-(3-bromophenylamino)isoquinoline-4-carbonitriles as inhibitors of myosin light chain kinase and epidermal growth factor receptor. <i>Bioorganic and Medicinal Chemistry</i> , 2011, 19, 429-439.	3.0	10
42	ALK Mutations Conferring Differential Resistance to Structurally Diverse ALK Inhibitors. <i>Clinical Cancer Research</i> , 2011, 17, 7394-7401.	7.0	179
43	A crucial requirement for Hedgehog signaling in small cell lung cancer. <i>Nature Medicine</i> , 2011, 17, 1504-1508.	30.7	224
44	Frequent and Focal <i>FGFR1</i> Amplification Associates with Therapeutically Tractable <i>FGFR1</i> Dependency in Squamous Cell Lung Cancer. <i>Science Translational Medicine</i> , 2010, 2, 62ra93.	12.4	761
45	Chemogenomic Profiling Provides Insights into the Limited Activity of Irreversible EGFR Inhibitors in Tumor Cells Expressing the T790M EGFR Resistance Mutation. <i>Cancer Research</i> , 2010, 70, 868-874.	0.9	191
46	Synthesis and Biological Evaluation of 4-Anilinoquinolines as Potent Inhibitors of Epidermal Growth Factor Receptor. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 2892-2901.	6.4	54
47	Analysis of Compound Synergy in High-Throughput Cellular Screens by Population-Based Lifetime Modeling. <i>PLoS ONE</i> , 2010, 5, e8919.	2.5	24
48	Identifying genotype-dependent efficacy of single and combined PI3K- and MAPK-pathway inhibition in cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 18351-18356.	7.1	251
49	Systematically linking drug susceptibility to cancer genome aberrations. <i>Cell Cycle</i> , 2009, 8, 3652-3656.	2.6	3
50	<i>PTEN</i> Loss Contributes to Erlotinib Resistance in EGFR-Mutant Lung Cancer by Activation of Akt and EGFR. <i>Cancer Research</i> , 2009, 69, 3256-3261.	0.9	480
51	Systematic RNA interference reveals that oncogenic KRAS-driven cancers require TBK1. <i>Nature</i> , 2009, 462, 108-112.	27.8	2,707
52	Predicting drug susceptibility of non-small cell lung cancers based on genetic lesions. <i>Journal of Clinical Investigation</i> , 2009, 119, 1727-1740.	8.2	230
53	Expression of Signaling Mediators Downstream of EGF-Receptor Predict Sensitivity to Small Molecule Inhibitors Directed Against the EGF-Receptor Pathway. <i>Journal of Thoracic Oncology</i> , 2008, 3, 170-173.	1.1	13
54	Early Detection of Erlotinib Treatment Response in NSCLC by $^{18}\text{F}$ -Deoxy- $^{18}\text{F}$ -[ $^{18}\text{F}$ ]-Fluoro-L-Thymidine ([ $^{18}\text{F}$ ]FLT) Positron Emission Tomography (PET). <i>PLoS ONE</i> , 2008, 3, e3908.	2.5	80

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55	Inhibition of Nuclear Translocation of Nuclear Factor- $\kappa$ B Despite Lack of Functional I $\kappa$ B $\beta$ Protein Overcomes Multiple Defects in Apoptosis Signaling in Human B-Cell Malignancies. <i>Clinical Cancer Research</i> , 2005, 11, 8186-8194.	7.0	19