

Erik S Knudsen

List of Publications by Year in descending order

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Version: 2024-02-01

62
papers

7,025
citations

87888

38
h-index

123424

61
g-index

64
all docs

64
docs citations

64
times ranked

11746
citing authors

#	ARTICLE	IF	CITATIONS
1	The history and future of targeting cyclin-dependent kinases in cancer therapy. <i>Nature Reviews Drug Discovery</i> , 2015, 14, 130-146.	46.4	1,316
2	Whole-exome sequencing of pancreatic cancer defines genetic diversity and therapeutic targets. <i>Nature Communications</i> , 2015, 6, 6744.	12.8	879
3	Tailoring to RB: tumour suppressor status and therapeutic response. <i>Nature Reviews Cancer</i> , 2008, 8, 714-724.	28.4	311
4	Differential Regulation of Retinoblastoma Protein Function by Specific Cdk Phosphorylation Sites. <i>Journal of Biological Chemistry</i> , 1996, 271, 8313-8320.	3.4	264
5	The meaning of p16 ^{ink4a} expression in tumors. <i>Cell Cycle</i> , 2011, 10, 2497-2503.	2.6	240
6	The Strange Case of CDK4/6 Inhibitors: Mechanisms, Resistance, and Combination Strategies. <i>Trends in Cancer</i> , 2017, 3, 39-55.	7.4	206
7	MCT4 Defines a Glycolytic Subtype of Pancreatic Cancer with Poor Prognosis and Unique Metabolic Dependencies. <i>Cell Reports</i> , 2014, 9, 2233-2249.	6.4	182
8	CDK4/6 inhibitors have potent activity in combination with pathway selective therapeutic agents in models of pancreatic cancer. <i>Oncotarget</i> , 2014, 5, 6512-6525.	1.8	180
9	Kinome-Wide RNA Interference Screen Reveals a Role for PDK1 in Acquired Resistance to CDK4/6 Inhibition in ER-Positive Breast Cancer. <i>Cancer Research</i> , 2017, 77, 2488-2499.	0.9	178
10	The retinoblastoma tumor suppressor modifies the therapeutic response of breast cancer. <i>Journal of Clinical Investigation</i> , 2007, 117, 218-228.	8.2	178
11	Targeting the RB-pathway in Cancer Therapy. <i>Clinical Cancer Research</i> , 2010, 16, 1094-1099.	7.0	177
12	RB-pathway disruption in breast cancer. <i>Cell Cycle</i> , 2010, 9, 4153-4163.	2.6	163
13	Metabolic Reprogramming of Pancreatic Cancer Mediated by CDK4/6 Inhibition Elicits Unique Vulnerabilities. <i>Cell Reports</i> , 2016, 14, 979-990.	6.4	160
14	CDK4/6 inhibition antagonizes the cytotoxic response to anthracycline therapy. <i>Cell Cycle</i> , 2012, 11, 2747-2755.	2.6	147
15	Stratification of Pancreatic Ductal Adenocarcinoma: Combinatorial Genetic, Stromal, and Immunologic Markers. <i>Clinical Cancer Research</i> , 2017, 23, 4429-4440.	7.0	142
16	p27 allosterically activates cyclin-dependent kinase 4 and antagonizes palbociclib inhibition. <i>Science</i> , 2019, 366, .	12.6	132
17	Modification of the DNA Damage Response by Therapeutic CDK4/6 Inhibition. <i>Journal of Biological Chemistry</i> , 2012, 287, 29075-29087.	3.4	128
18	Cell Cycle and Beyond: Exploiting New RB1 Controlled Mechanisms for Cancer Therapy. <i>Trends in Cancer</i> , 2019, 5, 308-324.	7.4	113

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19	Unbiased analysis of RB-mediated transcriptional repression identifies novel targets and distinctions from E2F action. <i>Cancer Research</i> , 2002, 62, 6587-97.	0.9	106
20	Retinoblastoma tumor suppressor pathway in breast cancer: prognosis, precision medicine, and therapeutic interventions. <i>Breast Cancer Research</i> , 2014, 16, 207.	5.0	101
21	CDK4/6 inhibition provides a potent adjunct to Her2-targeted therapies in preclinical breast cancer models. <i>Genes and Cancer</i> , 2014, 5, 261-272.	1.9	101
22	Genetic Diversity of Pancreatic Ductal Adenocarcinoma and Opportunities for Precision Medicine. <i>Gastroenterology</i> , 2016, 150, 48-63.	1.3	90
23	Binary pan-cancer classes with distinct vulnerabilities defined by pro- or anti-cancer YAP/TEAD activity. <i>Cancer Cell</i> , 2021, 39, 1115-1134.e12.	16.8	86
24	Integrated Patient-Derived Models Delineate Individualized Therapeutic Vulnerabilities of Pancreatic Cancer. <i>Cell Reports</i> , 2016, 16, 2017-2031.	6.4	84
25	The Changing Landscape of Hepatocellular Carcinoma. <i>American Journal of Pathology</i> , 2014, 184, 574-583.	3.8	82
26	Pancreatic cancer cell lines as patient-derived avatars: genetic characterisation and functional utility. <i>Gut</i> , 2018, 67, 508-520.	12.1	81
27	Immunologic and Metabolic Features of Pancreatic Ductal Adenocarcinoma Define Prognostic Subtypes of Disease. <i>Clinical Cancer Research</i> , 2016, 22, 3606-3617.	7.0	73
28	Phase I Clinical Trial of Combination Propranolol and Pembrolizumab in Locally Advanced and Metastatic Melanoma: Safety, Tolerability, and Preliminary Evidence of Antitumor Activity. <i>Clinical Cancer Research</i> , 2021, 27, 87-95.	7.0	72
29	Targeting the Vulnerability of RB Tumor Suppressor Loss in Triple-Negative Breast Cancer. <i>Cell Reports</i> , 2018, 22, 1185-1199.	6.4	71
30	Cyclin A Is a Functional Target of Retinoblastoma Tumor Suppressor Protein-mediated Cell Cycle Arrest. <i>Journal of Biological Chemistry</i> , 1999, 274, 27632-27641.	3.4	66
31	RB-Pathway Disruption Is Associated with Improved Response to Neoadjuvant Chemotherapy in Breast Cancer. <i>Clinical Cancer Research</i> , 2012, 18, 5110-5122.	7.0	64
32	Selective impact of CDK4/6 suppression on patient-derived models of pancreatic cancer. <i>Oncotarget</i> , 2015, 6, 15788-15801.	1.8	51
33	Pan-cancer molecular analysis of the RB tumor suppressor pathway. <i>Communications Biology</i> , 2020, 3, 158.	4.4	50
34	Targeting dual signalling pathways in concert with immune checkpoints for the treatment of pancreatic cancer. <i>Gut</i> , 2021, 70, 127-138.	12.1	49
35	CDK/cyclin dependencies define extreme cancer cell-cycle heterogeneity and collateral vulnerabilities. <i>Cell Reports</i> , 2022, 38, 110448.	6.4	48
36	Cell cycle plasticity driven by MTOR signaling: integral resistance to CDK4/6 inhibition in patient-derived models of pancreatic cancer. <i>Oncogene</i> , 2019, 38, 3355-3370.	5.9	46

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37	Unique metabolic features of pancreatic cancer stroma: relevance to the tumor compartment, prognosis, and invasive potential. <i>Oncotarget</i> , 2016, 7, 78396-78411.	1.8	45
38	Kinase-independent role of cyclin D1 in chromosomal instability and mammary tumorigenesis. <i>Oncotarget</i> , 2015, 6, 8525-8538.	1.8	43
39	The RB tumor suppressor at the intersection of proliferation and immunity: relevance to disease immune evasion and immunotherapy. <i>Cell Cycle</i> , 2015, 14, 3812-3819.	2.6	42
40	Loss of RB compromises specific heterochromatin modifications and modulates HP1 \pm dynamics. <i>Journal of Cellular Physiology</i> , 2007, 211, 131-137.	4.1	41
41	Functional Determinants of Cell Cycle Plasticity and Sensitivity to CDK4/6 Inhibition. <i>Cancer Research</i> , 2021, 81, 1347-1360.	0.9	40
42	Composite analysis of immunological and metabolic markers defines novel subtypes of triple negative breast cancer. <i>Modern Pathology</i> , 2018, 31, 288-298.	5.5	38
43	Chemotherapy and CDK4/6 Inhibitors: Unexpected Bedfellows. <i>Molecular Cancer Therapeutics</i> , 2020, 19, 1575-1588.	4.1	35
44	EZH2 and ALDH1 expression in ductal carcinoma in situ: Complex association with recurrence and progression to invasive breast cancer. <i>Cell Cycle</i> , 2013, 12, 2042-2050.	2.6	31
45	Retinoblastoma protein potentiates the innate immune response in hepatocytes: Significance for hepatocellular carcinoma. <i>Hepatology</i> , 2014, 60, 1231-1240.	7.3	28
46	Coordinately Targeting Cell-Cycle Checkpoint Functions in Integrated Models of Pancreatic Cancer. <i>Clinical Cancer Research</i> , 2019, 25, 2290-2304.	7.0	26
47	Defining the transcriptional and biological response to CDK4/6 inhibition in relation to ER+/HER2- breast cancer. <i>Oncotarget</i> , 2016, 7, 69111-69123.	1.8	26
48	Chemotherapy impacts on the cellular response to CDK4/6 inhibition: distinct mechanisms of interaction and efficacy in models of pancreatic cancer. <i>Oncogene</i> , 2020, 39, 1831-1845.	5.9	25
49	Retinoblastoma and Phosphate and Tensin Homolog Tumor Suppressors: Impact on Ductal Carcinoma In Situ Progression. <i>Journal of the National Cancer Institute</i> , 2012, 104, 1825-1836.	6.3	24
50	Identification of highly penetrant Rb-related synthetic lethal interactions in triple negative breast cancer. <i>Oncogene</i> , 2018, 37, 5701-5718.	5.9	24
51	Systematically Defining Single-Gene Determinants of Response to Neoadjuvant Chemotherapy Reveals Specific Biomarkers. <i>Clinical Cancer Research</i> , 2014, 20, 4837-4848.	7.0	19
52	Selective CDK4/6 Inhibitors: Biologic Outcomes, Determinants of Sensitivity, Mechanisms of Resistance, Combinatorial Approaches, and Pharmacodynamic Biomarkers. <i>American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting</i> , 2020, 40, 115-126.	3.8	16
53	A Phase I Study of Ribociclib Plus Everolimus in Patients with Metastatic Pancreatic Adenocarcinoma Refractory to Chemotherapy. <i>Journal of Pancreatic Cancer</i> , 2020, 6, 45-54.	0.9	15
54	Cancer cell cycle dystopia: heterogeneity, plasticity, and therapy. <i>Trends in Cancer</i> , 2022, 8, 711-725.	7.4	12

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55	Phase Ib/II Study of Cetuximab plus Pembrolizumab in Patients with Advanced RAS Wild-Type Colorectal Cancer. <i>Clinical Cancer Research</i> , 2021, 27, 6726-6736.	7.0	8
56	Real-World Experience with CDK4/6 Inhibitors for Metastatic HR+/HER2 ⁺ Breast Cancer at a Single Cancer Center. <i>Oncologist</i> , 2022, 27, 646-654.	3.7	7
57	TP53, CDKN2A/P16, and NFE2L2/NRF2 regulate the incidence of pure- and combined-small cell lung cancer in mice. <i>Oncogene</i> , 2022, 41, 3423-3432.	5.9	7
58	RB Tumor Suppressive Function in Response to Xenobiotic Hepatocarcinogens. <i>American Journal of Pathology</i> , 2014, 184, 1853-1859.	3.8	6
59	RB loss determines selective resistance and novel vulnerabilities in ER-positive breast cancer models. <i>Oncogene</i> , 0, , .	5.9	6
60	Interrogating Mutant Allele Expression via Customized Reference Genomes to Define Influential Cancer Mutations. <i>Scientific Reports</i> , 2019, 9, 12766.	3.3	5
61	The transcriptome of CDK4/6 inhibition. <i>Aging</i> , 2017, 9, 1859-1860.	3.1	2
62	Cell cycle: mechanisms of control and dysregulation in cancer. , 0, , 452-464.		0