

Roderick L Beijersbergen

List of Publications by Year in descending order

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Version: 2024-02-01

88
papers

18,148
citations

53660

45
h-index

51492

86
g-index

97
all docs

97
docs citations

97
times ranked

23873
citing authors

#	ARTICLE	IF	CITATIONS
1	Creation of human tumour cells with defined genetic elements. <i>Nature</i> , 1999, 400, 464-468.	13.7	2,148
2	Unresponsiveness of colon cancer to BRAF(V600E) inhibition through feedback activation of EGFR. <i>Nature</i> , 2012, 483, 100-103.	13.7	1,769
3	hEST2, the Putative Human Telomerase Catalytic Subunit Gene, Is Up-Regulated in Tumor Cells and during Immortalization. <i>Cell</i> , 1997, 90, 785-795.	13.5	1,689
4	A Functional Genetic Approach Identifies the PI3K Pathway as a Major Determinant of Trastuzumab Resistance in Breast Cancer. <i>Cancer Cell</i> , 2007, 12, 395-402.	7.7	1,471
5	Inhibition of telomerase limits the growth of human cancer cells. <i>Nature Medicine</i> , 1999, 5, 1164-1170.	15.2	983
6	A large-scale RNAi screen in human cells identifies new components of the p53 pathway. <i>Nature</i> , 2004, 428, 431-437.	13.7	955
7	Reversible and adaptive resistance to BRAF(V600E) inhibition in melanoma. <i>Nature</i> , 2014, 508, 118-122.	13.7	702
8	Dissociation among in vitro telomerase activity, telomere maintenance, and cellular immortalization. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1998, 95, 14723-14728.	3.3	582
9	Statistical methods for analysis of high-throughput RNA interference screens. <i>Nature Methods</i> , 2009, 6, 569-575.	9.0	532
10	Phosphatidylinositol 3-Kinase Hyperactivation Results in Lapatinib Resistance that Is Reversed by the mTOR/Phosphatidylinositol 3-Kinase Inhibitor NVP-BE235. <i>Cancer Research</i> , 2008, 68, 9221-9230.	0.4	474
11	Stabilization of N-Myc Is a Critical Function of Aurora A in Human Neuroblastoma. <i>Cancer Cell</i> , 2009, 15, 67-78.	7.7	464
12	The ubiquitin-specific protease USP28 is required for MYC stability. <i>Nature Cell Biology</i> , 2007, 9, 765-774.	4.6	391
13	MED12 Controls the Response to Multiple Cancer Drugs through Regulation of TGF- β 2 Receptor Signaling. <i>Cell</i> , 2012, 151, 937-950.	13.5	371
14	CRISPR knockout screening outperforms shRNA and CRISPRi in identifying essential genes. <i>Nature Biotechnology</i> , 2016, 34, 631-633.	9.4	344
15	E2F-4, a new member of the E2F gene family, has oncogenic activity and associates with p107 in vivo.. <i>Genes and Development</i> , 1994, 8, 2680-2690.	2.7	312
16	Intracellular bacterial growth is controlled by a kinase network around PKB/AKT1. <i>Nature</i> , 2007, 450, 725-730.	13.7	310
17	Intrinsic Resistance to MEK Inhibition in KRAS Mutant Lung and Colon Cancer through Transcriptional Induction of ERBB3. <i>Cell Reports</i> , 2014, 7, 86-93.	2.9	266
18	A Genetic Screen Identifies PITX1 as a Suppressor of RAS Activity and Tumorigenicity. <i>Cell</i> , 2005, 121, 849-858.	13.5	257

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19	Inducing and exploiting vulnerabilities for the treatment of liver cancer. <i>Nature</i> , 2019, 574, 268-272.	13.7	249
20	Neutrophils Kill Antibody-Opsonized Cancer Cells by Trogoptosis. <i>Cell Reports</i> , 2018, 23, 3946-3959.e6.	2.9	245
21	E2F-5, a New E2F Family Member That Interacts with p130 In Vivo. <i>Molecular and Cellular Biology</i> , 1995, 15, 3082-3089.	1.1	228
22	An shRNA barcode screen provides insight into cancer cell vulnerability to MDM2 inhibitors. <i>Nature Chemical Biology</i> , 2006, 2, 202-206.	3.9	196
23	EGFR activation limits the response of liver cancer to lenvatinib. <i>Nature</i> , 2021, 595, 730-734.	13.7	183
24	PTPN11 Is a Central Node in Intrinsic and Acquired Resistance to Targeted Cancer Drugs. <i>Cell Reports</i> , 2015, 12, 1978-1985.	2.9	163
25	A Genome-wide Multidimensional RNAi Screen Reveals Pathways Controlling MHC Class II Antigen Presentation. <i>Cell</i> , 2011, 145, 268-283.	13.5	151
26	High-Throughput Functional Genetic and Compound Screens Identify Targets for Senescence Induction in Cancer. <i>Cell Reports</i> , 2017, 21, 773-783.	2.9	136
27	ZNF423 Is Critically Required for Retinoic Acid-Induced Differentiation and Is a Marker of Neuroblastoma Outcome. <i>Cancer Cell</i> , 2009, 15, 328-340.	7.7	132
28	Cloning, expression and chromosomal localization of a new putative receptor-like protein tyrosine phosphatase. <i>FEBS Letters</i> , 1991, 290, 123-130.	1.3	125
29	Advances in CRISPR-Cas9 genome engineering: lessons learned from RNA interference. <i>Nucleic Acids Research</i> , 2015, 43, 3407-3419.	6.5	124
30	A Large Scale shRNA Barcode Screen Identifies the Circadian Clock Component ARNTL as Putative Regulator of the p53 Tumor Suppressor Pathway. <i>PLoS ONE</i> , 2009, 4, e4798.	1.1	118
31	shRNA libraries and their use in cancer genetics. <i>Nature Methods</i> , 2006, 3, 701-706.	9.0	116
32	Impact of chromatin context on Cas9-induced DNA double-strand break repair pathway balance. <i>Molecular Cell</i> , 2021, 81, 2216-2230.e10.	4.5	106
33	Involvement of MINK, a Ste20 Family Kinase, in Ras Oncogene-Induced Growth Arrest in Human Ovarian Surface Epithelial Cells. <i>Molecular Cell</i> , 2005, 20, 673-685.	4.5	96
34	Cell cycle regulation by the retinoblastoma family of growth inhibitory proteins. <i>Biochimica Et Biophysica Acta: Reviews on Cancer</i> , 1996, 1287, 103-120.	3.3	94
35	The Cancer SENESCopedia: A delineation of cancer cell senescence. <i>Cell Reports</i> , 2021, 36, 109441.	2.9	84
36	Loss of p53 induces cell proliferation via Ras-independent activation of the Raf/Mek/Erk signaling pathway. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, 15155-15160.	3.3	80

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37	Phospho-ERK is a biomarker of response to a synthetic lethal drug combination of sorafenib and MEK inhibition in liver cancer. <i>Journal of Hepatology</i> , 2018, 69, 1057-1065.	1.8	74
38	CDK12 inhibition mediates DNA damage and is synergistic with sorafenib treatment in hepatocellular carcinoma. <i>Gut</i> , 2020, 69, 727-736.	6.1	74
39	ARID1A mutation sensitizes most ovarian clear cell carcinomas to BET inhibitors. <i>Oncogene</i> , 2018, 37, 4611-4625.	2.6	72
40	Miz1 and HectH9 regulate the stability of the checkpoint protein, TopBP1. <i>EMBO Journal</i> , 2008, 27, 2851-2861.	3.5	70
41	A Vulnerability of a Subset of Colon Cancers with Potential Clinical Utility. <i>Cell</i> , 2016, 165, 317-330.	13.5	70
42	Telomerase-independent Regulation of ATR by Human Telomerase RNA. <i>Journal of Biological Chemistry</i> , 2006, 281, 40503-40514.	1.6	66
43	Effects of Monomethylfumarate on Human Granulocytes. <i>Journal of Investigative Dermatology</i> , 1993, 101, 37-42.	0.3	59
44	A Functional Genetic Screen Identifies the Phosphoinositide 3-kinase Pathway as a Determinant of Resistance to Fibroblast Growth Factor Receptor Inhibitors in FGFR Mutant Urothelial Cell Carcinoma. <i>European Urology</i> , 2017, 71, 858-862.	0.9	59
45	An integrated genomic approach identifies that the PI3K/AKT/FOXO pathway is involved in breast cancer tumor initiation. <i>Oncotarget</i> , 2016, 7, 2596-2610.	0.8	52
46	A CRISPR screen identifies CDK7 as a therapeutic target in hepatocellular carcinoma. <i>Cell Research</i> , 2018, 28, 690-692.	5.7	46
47	Inhibition of Ataxia-Telangiectasia Mutated and RAD3-Related (<i>ATR</i>) Overcomes Oxaliplatin Resistance and Promotes Antitumor Immunity in Colorectal Cancer. <i>Cancer Research</i> , 2019, 79, 2933-2946.	0.4	46
48	The Histone Demethylase Jarid1b (Kdm5b) Is a Novel Component of the Rb Pathway and Associates with E2f-Target Genes in MEFs during Senescence. <i>PLoS ONE</i> , 2011, 6, e25235.	1.1	44
49	Loss of <i>ARID1A</i> Activates <i>ANXA1</i> , which Serves as a Predictive Biomarker for Trastuzumab Resistance. <i>Clinical Cancer Research</i> , 2016, 22, 5238-5248.	3.2	43
50	EGFR overexpression induces activation of telomerase via PI3K/AKT-mediated phosphorylation and transcriptional regulation through Hif1 α in a cellular model of oral esophageal carcinogenesis. <i>Cancer Science</i> , 2011, 102, 351-360.	1.7	42
51	Functional Analysis of Burkitt's Lymphoma Mutant c-Myc Proteins. <i>Journal of Biological Chemistry</i> , 1996, 271, 5513-5518.	1.6	39
52	Identification of Fox only protein 7 as a negative regulator of NF κ B signalling. <i>Journal of Cellular and Molecular Medicine</i> , 2012, 16, 2140-2149.	1.6	38
53	Identification of signalling cascades involved in red blood cell shrinkage and vesiculation. <i>Bioscience Reports</i> , 2015, 35, .	1.1	37
54	SMARCE1 suppresses EGFR expression and controls responses to MET and ALK inhibitors in lung cancer. <i>Cell Research</i> , 2015, 25, 445-458.	5.7	36

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55	Synthetic Lethality in Cancer Therapeutics. <i>Annual Review of Cancer Biology</i> , 2017, 1, 141-161.	2.3	36
56	Glucocorticoid receptor triggers a reversible drug-tolerant dormancy state with acquired therapeutic vulnerabilities in lung cancer. <i>Nature Communications</i> , 2021, 12, 4360.	5.8	35
57	Parallel InÂVivo and InÂVitro Melanoma RNAi Dropout Screens Reveal Synthetic Lethality between Hypoxia and DNA Damage Response Inhibition. <i>Cell Reports</i> , 2014, 9, 1375-1386.	2.9	34
58	Sensitizing Triple-Negative Breast Cancer to PI3K Inhibition by Cotargeting IGF1R. <i>Molecular Cancer Therapeutics</i> , 2016, 15, 1545-1556.	1.9	30
59	Ezh2 inhibition in Kras-driven lung cancer amplifies inflammation and associated vulnerabilities. <i>Journal of Experimental Medicine</i> , 2018, 215, 3115-3135.	4.2	29
60	VAV3 mediates resistance to breast cancer endocrine therapy. <i>Breast Cancer Research</i> , 2014, 16, R53.	2.2	28
61	PIM Kinases Are a Potential Prognostic Biomarker and Therapeutic Target in Neuroblastoma. <i>Molecular Cancer Therapeutics</i> , 2018, 17, 849-857.	1.9	28
62	A role for the unfolded protein response stress sensor ERN1 in regulating the response to MEK inhibitors in KRAS mutant colon cancers. <i>Genome Medicine</i> , 2018, 10, 90.	3.6	28
63	The Corepressor CTBP2 Is a Coactivator of Retinoic Acid Receptor/Retinoid X Receptor in Retinoic Acid Signaling. <i>Molecular and Cellular Biology</i> , 2013, 33, 3343-3353.	1.1	25
64	The lethal response to Cdk1 inhibition depends on sister chromatid alignment errors generated by KIF4 and isoform 1 of PRC1. <i>Scientific Reports</i> , 2015, 5, 14798.	1.6	25
65	TLE3 loss confers AR inhibitor resistance by facilitating GR-mediated human prostate cancer cell growth. <i>ELife</i> , 2019, 8, .	2.8	25
66	A genome-wide RNAi screen in mouse embryonic stem cells identifies Mp1 as a key mediator of differentiation. <i>Journal of Experimental Medicine</i> , 2011, 208, 2675-2689.	4.2	24
67	Targeting CDC7 potentiates ATR-Chk1 signaling inhibition through induction of DNA replication stress in liver cancer. <i>Genome Medicine</i> , 2021, 13, 166.	3.6	19
68	Differential transcriptional regulation of human telomerase in a cellular model representing important genetic alterations in esophageal squamous carcinogenesis. <i>Carcinogenesis</i> , 2005, 26, 1879-1889.	1.3	16
69	Intrinsic resistance to PIM kinase inhibition in AML through p38Î±-mediated feedback activation of mTOR signaling. <i>Oncotarget</i> , 2016, 7, 37407-37419.	0.8	16
70	Screening for modulators of cisplatin sensitivity: Unbiased screens reveal common themes. <i>Cell Cycle</i> , 2011, 10, 380-386.	1.3	15
71	Exploration of synthetic lethal interactions as cancer drug targets. <i>Future Oncology</i> , 2010, 6, 1789-1802.	1.1	14
72	Integrative Modeling Identifies Key Determinants of Inhibitor Sensitivity in Breast Cancer Cell Lines. <i>Cancer Research</i> , 2018, 78, 4396-4410.	0.4	14

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73	High-throughput compound screen reveals mTOR inhibitors as potential therapeutics to reduce (auto)antibody production by human plasma cells. <i>European Journal of Immunology</i> , 2020, 50, 73-85.	1.6	12
74	Old drugs with new tricks. <i>Nature Cancer</i> , 2020, 1, 153-155.	5.7	10
75	Integrated <i>in vivo</i> genetic and pharmacologic screening identifies co-inhibition of EGFR and ROCK as a potential treatment regimen for triple-negative breast cancer. <i>Oncotarget</i> , 2016, 7, 42859-42872.	0.8	10
76	Candidate Biomarkers of Response to an Experimental Cancer Drug Identified through a Large-scale RNA Interference Genetic Screen. <i>Clinical Cancer Research</i> , 2009, 15, 5811-5819.	3.2	9
77	A kinome-centered CRISPR-Cas9 screen identifies activated BRAF to modulate enzalutamide resistance with potential therapeutic implications in BRAF-mutated prostate cancer. <i>Scientific Reports</i> , 2021, 11, 13683.	1.6	8
78	Pooled shRNA Screening in Mammalian Cells as a Functional Genomic Discovery Platform. <i>Methods in Molecular Biology</i> , 2016, 1470, 49-73.	0.4	6
79	Identification of Autophagy-Related Genes as Targets for Senescence Induction Using a Customizable CRISPR-Based Suicide Switch Screen. <i>Molecular Cancer Research</i> , 2021, 19, 1613-1621.	1.5	6
80	Genetic and compound screens uncover factors modulating cancer cell response to indisulam. <i>Life Science Alliance</i> , 2022, 5, e202101348.	1.3	6
81	Using large-scale RNAi screens to identify novel drug targets for cancer. <i>IDrugs: the Investigational Drugs Journal</i> , 2010, 13, 772-7.	0.7	5
82	A loss-of-adhesion CRISPR-Cas9 screening platform to identify cell adhesion-regulatory proteins and signaling pathways. <i>Nature Communications</i> , 2022, 13, 2136.	5.8	4
83	Interleukin-1 α -Associated Kinase 2 Is a Novel Modulator of the Transforming Growth Factor β Signaling Cascade. <i>Molecular Cancer Research</i> , 2010, 8, 592-603.	1.5	3
84	Ribociclib Induces Broad Chemotherapy Resistance and EGFR Dependency in ESR1 Wildtype and Mutant Breast Cancer. <i>Cancers</i> , 2021, 13, 6314.	1.7	3
85	The Good, the Bad, and the Ugly: in search of gold standards for assessing functional genetic screen quality. <i>Molecular Systems Biology</i> , 2014, 10, 738.	3.2	2
86	It takes two to tango, and the right music: Synergistic drug combinations with cell-cycle phase-dependent sensitivities. <i>EBioMedicine</i> , 2021, 69, 103448.	2.7	1
87	Bioassay development. , 0, , 67-84.		0
88	Functional Subtyping of Breast Cancer. <i>Cancer Discovery</i> , 2011, 1, 205-206.	7.7	0