List of Publications by Year in descending order

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		57631	56606
133	7,854	44	83
papers	citations	h-index	g-index
135	135	135	9793
all docs	docs citations	times ranked	citing authors

ANDERSON L RVAN

#	Article	IF	CITATIONS
1	CHK1 inhibition exacerbates replication stress induced by IGF blockade. Oncogene, 2022, 41, 476-488.	2.6	4
2	Antiâ€ŧumoural activity of the Gâ€quadruplex ligand pyridostatin against BRCA1/2â€deficient tumours. EMBO Molecular Medicine, 2022, 14, e14501.	3.3	13
3	Targeting TOPK sensitises tumour cells to radiation-induced damage by enhancing replication stress. Cell Death and Differentiation, 2021, 28, 1333-1346.	5.0	13
4	Targeting IGF Perturbs Global Replication through Ribonucleotide Reductase Dysfunction. Cancer Research, 2021, 81, 2128-2141.	0.4	10
5	Germline and Somatic Genetic Variants in the p53 Pathway Interact to Affect Cancer Risk, Progression, and Drug Response. Cancer Research, 2021, 81, 1667-1680.	0.4	32
6	Olaparib increases the therapeutic index of hemithoracic irradiation compared with hemithoracic irradiation alone in a mouse lung cancer model. British Journal of Cancer, 2021, 124, 1809-1819.	2.9	5
7	Beyond cancer cells: Targeting the tumor microenvironment with gene therapy and armed oncolytic virus. Molecular Therapy, 2021, 29, 1668-1682.	3.7	33
8	DNAPK Inhibition Preferentially Compromises the Repair of Radiation-induced DNA Double-strand Breaks in Chronically Hypoxic Tumor Cells in Xenograft Models. Molecular Cancer Therapeutics, 2021, 20, 1663-1671.	1.9	5
9	CONCORDE: A phase I platform study of novel agents in combination with conventional radiotherapy in non-small-cell lung cancer. Clinical and Translational Radiation Oncology, 2020, 25, 61-66.	0.9	15
10	Adefovir dipivoxil induces DNA replication stress and augments ATR inhibitorâ€related cytotoxicity. International Journal of Cancer, 2020, 147, 1474-1484.	2.3	7
11	Identification of anticancer drugs to radiosensitise BRAF-wild-type and mutant colorectal cancer. Cancer Biology and Medicine, 2019, 16, 234.	1.4	4
12	Overcoming acquired resistance to HSP90 inhibition by targeting JAK-STAT signalling in triple-negative breast cancer. BMC Cancer, 2019, 19, 102.	1.1	29
13	Chlorambucil targets <scp>BRCA</scp> 1/2â€deficient tumours and counteracts <scp>PARP</scp> inhibitor resistance. EMBO Molecular Medicine, 2019, 11, e9982.	3.3	26
14	<scp>RASSF</scp> 1A controls tissue stiffness and cancer stemâ€like cells in lung adenocarcinoma. EMBO Journal, 2019, 38, e100532.	3.5	83
15	Cardio-Respiratory synchronized bSSFP MRI for high throughput in vivo lung tumour quantification. PLoS ONE, 2019, 14, e0212172.	1.1	7
16	Selective DNA-PKcs inhibition extends the therapeutic index of localized radiotherapy and chemotherapy. Journal of Clinical Investigation, 2019, 130, 258-271.	3.9	45
17	PARP Inhibition Combined With Thoracic Irradiation Exacerbates Esophageal and Skin Toxicity in C57BL6 Mice. International Journal of Radiation Oncology Biology Physics, 2018, 100, 767-775.	0.4	22
18	Pre-clinical Profile and Expectations for Pharmacological ATM Inhibition. Cancer Drug Discovery and Development, 2018, , 155-183.	0.2	0

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19	Persistent DNA strand breaks induce a CAF-like phenotype in normal fibroblasts. Oncotarget, 2018, 9, 13666-13681.	0.8	20
20	The HGF/c-MET Pathway Is a Driver and Biomarker of VEGFR-inhibitor Resistance and Vascular Remodeling in Non–Small Cell Lung Cancer. Clinical Cancer Research, 2017, 23, 5489-5501.	3.2	55
21	<scp>BRCA</scp> 1 and <scp>BRCA</scp> 2 tumor suppressors protect against endogenous acetaldehyde toxicity. EMBO Molecular Medicine, 2017, 9, 1398-1414.	3.3	57
22	TOPK modulates tumour-specific radiosensitivity and correlates with recurrence after prostate radiotherapy. British Journal of Cancer, 2017, 117, 503-512.	2.9	20
23	Abstract B30: Novel targets for combination therapy in EGFR mutated NSCLC. , 2017, , .		0
24	Phenotypic consequences of somatic mutations in the ataxia-telangiectasia mutated gene in non-small cell lung cancer. Oncotarget, 2016, 7, 60807-60822.	0.8	23
25	Hypoxia Potentiates the Radiation-Sensitizing Effect of Olaparib in Human Non-Small Cell Lung Cancer Xenografts by Contextual Synthetic Lethality. International Journal of Radiation Oncology Biology Physics, 2016, 95, 772-781.	0.4	39
26	Targeting BRCA1 and BRCA2 Deficiencies with G-Quadruplex-Interacting Compounds. Molecular Cell, 2016, 61, 449-460.	4.5	185
27	Abstract B40: WEE1 inhibition selectively kills histone H3K36me3-deficient cancers by dNTP starvation. , 2016, , .		0
28	Vascular endothelial growth factor directly stimulates tumour cell proliferation in non-small cell lung cancer. International Journal of Oncology, 2015, 47, 849-856.	1.4	29
29	Inhibiting WEE1 Selectively Kills Histone H3K36me3-Deficient Cancers by dNTP Starvation. Cancer Cell, 2015, 28, 557-568.	7.7	244
30	Acute vascular response to cediranib treatment in human non-small-cell lung cancer xenografts with different tumour stromal architecture. Lung Cancer, 2015, 90, 191-198.	0.9	14
31	ATM and ATR as therapeutic targets in cancer. , 2015, 149, 124-138.		487
32	EGFR biomarkers predict benefit from vandetanib in combination with docetaxel in a randomized phase III study of second-line treatment of patients with advanced non-small cell lung cancer. Annals of Oncology, 2014, 25, 1941-1948.	0.6	20
33	Changes in Signaling Pathways Induced by Vandetanib in a Human Medullary Thyroid Carcinoma Model, as Analyzed by Reverse Phase Protein Array. Thyroid, 2014, 24, 43-51.	2.4	8
34	Erlotinib, Gefitinib, and Vandetanib Inhibit Human Nucleoside Transporters and Protect Cancer Cells from Gemcitabine Cytotoxicity. Clinical Cancer Research, 2014, 20, 176-186.	3.2	37
35	Short-Course Treatment With Gefitinib Enhances Curative Potential of Radiation Therapy in a Mouse Model of Human Non-Small Cell Lung Cancer. International Journal of Radiation Oncology Biology Physics, 2014, 88, 947-954.	0.4	26
36	Combining AKT inhibition with chloroquine and gefitinib prevents compensatory autophagy and induces cell death in EGFR mutated NSCLC cells. Oncotarget, 2014, 5, 4765-4778.	0.8	42

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37	Abstract 3785: Olaparib increases the effectiveness of radiation in hypoxic tumor cells in xenograft models of human non-small-cell lung cancer. , 2014, , .		0
38	Replication Stress and Chromatin Context Link ATM Activation to a Role in DNA Replication. Molecular Cell, 2013, 52, 758-766.	4.5	102
39	An Integrated Inspection of the Somatic Mutations in a Lung Squamous Cell Carcinoma Using Next-Generation Sequencing. PLoS ONE, 2013, 8, e78823.	1.1	5
40	Abstract B91: Combined ATM and ATR kinase inhibition selectively kills p53-mutated non-small cell lung cancer (NSCLC) cells Molecular Cancer Therapeutics, 2013, 12, B91-B91.	1.9	3
41	Changes in signaling pathways induced by vandetanib in a human medullary thyroid carcinoma model, as analyzed by Reverse Phase Protein Array Thyroid, 2013, , 130703231537001.	2.4	Ο
42	Abstract B276: A live-cell apoptosis assay identifies novel combination approaches that selectively kill EGFR mutant NSCLC cells , 2013, , .		0
43	Reply to JF. Chatal et al. Journal of Clinical Oncology, 2012, 30, 2166-2167.	0.8	Ο
44	Vandetanib in Patients With Locally Advanced or Metastatic Medullary Thyroid Cancer: A Randomized, Double-Blind Phase III Trial. Journal of Clinical Oncology, 2012, 30, 134-141.	0.8	1,295
45	False-negative MRI biomarkers of tumour response to targeted cancer therapeutics. British Journal of Cancer, 2012, 106, 1960-1966.	2.9	10
46	Combined MEK and VEGFR Inhibition in Orthotopic Human Lung Cancer Models Results in Enhanced Inhibition of Tumor Angiogenesis, Growth, and Metastasis. Clinical Cancer Research, 2012, 18, 1641-1654.	3.2	51
47	Inhibition of Aurora-B kinase activity confers antitumor efficacy in preclinical mouse models of early and advanced gastrointestinal neoplasia. International Journal of Oncology, 2012, 41, 1475-1485.	1.4	10
48	Evaluation of novel combined carbogen USPIO (CUSPIO) imaging biomarkers in assessing the antiangiogenic effects of cediranib (AZD2171) in rat C6 gliomas. International Journal of Cancer, 2012, 131, 1854-1862.	2.3	9
49	Antitumor effect of the vascular-disrupting agent ZD6126 in a murine renal cell carcinoma model. International Journal of Oncology, 2011, 38, 455-64.	1.4	3
50	Vandetanib inhibits both VEGFR-2 and EGFR signalling at clinically relevant drug levels in preclinical models of human cancer. International Journal of Oncology, 2011, 39, 271-8.	1.4	13
51	Investigating temporal fluctuations in tumor vasculature with combined carbogen and ultrasmall superparamagnetic iron oxide particle (CUSPIO) imaging. Magnetic Resonance in Medicine, 2011, 66, 227-234.	1.9	11
52	Abstract 5471: Pharmacokinetic investigation of vandetanib in patients with medullary thyroid carcinoma using liquid chromatography-tandem mass spectrometry. , 2011, , .		0
53	Abstract 3569: Changes in signaling pathways induced by vandetanib in a human medullary thyroid carcinoma model as detected by reverse phase protein arrays. , 2011, , .		0
54	Abstract 629: MEK inhibition by selumetinib (AZD6244) inhibits progression with a multicentric antiangiogenic effect and enhances the efficacy of cediranib in orthotopic human lung cancer models. , 2011, , .		0

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55	Abstract 3279: Effects of the isoforms of the angiogenic growth factor VEGF on neo-vascularization and tumor response to the tyrosine kinase inhibitor cediranib. , 2011, , .		0
56	Effects of vandetanib on adenoma formation in a dextran sodium sulphate enhanced ApcMIN/+ mouse model. International Journal of Oncology, 2010, 37, 767-72.	1.4	3
57	Distinct Patterns of Cytokine and Angiogenic Factor Modulation and Markers of Benefit for Vandetanib and/or Chemotherapy in Patients With Non–Small-Cell Lung Cancer. Journal of Clinical Oncology, 2010, 28, 193-201.	0.8	131
58	Vascular Endothelial Growth Factor Receptors VEGFR-2 and VEGFR-3 Are Localized Primarily to the Vasculature in Human Primary Solid Cancers. Clinical Cancer Research, 2010, 16, 3548-3561.	3.2	202
59	The tyrosine kinase inhibitor ZD6474 blocks proliferation of RET mutant medullary thyroid carcinoma cells. Endocrine-Related Cancer, 2010, 18, 1-11.	1.6	58
60	Assessment of Acute Antivascular Effects of Vandetanib with High-Resolution Dynamic Contrast-Enhanced Computed Tomographic Imaging in a Human Colon Tumor Xenograft Model in the Nude Rat. Neoplasia, 2010, 12, 697-707.	2.3	20
61	Circulating free DNA as a surrogate for tumor material for EGFR and KRAS analysis. , 2010, , .		1
62	The antiangiogenic agent ZD4190 prevents tumour outgrowth in a model of minimal residual carcinoma in deep tissues. British Journal of Cancer, 2009, 101, 418-423.	2.9	1
63	Targeting vascular endothelial growth factor receptor-1 and -3 with cediranib (AZD2171): effects on migration and invasion of gastrointestinal cancer cell lines. Molecular Cancer Therapeutics, 2009, 8, 2546-2558.	1.9	40
64	Baseline Vascular Endothelial Growth Factor Concentration as a Potential Predictive Marker of Benefit from Vandetanib in Non–Small Cell Lung Cancer. Clinical Cancer Research, 2009, 15, 3600-3609.	3.2	90
65	Identification of tyrosine 806 as a molecular determinant of RET kinase sensitivity to ZD6474. Endocrine-Related Cancer, 2009, 16, 233-241.	1.6	37
66	Combination of Vandetanib, Radiotherapy, and Irinotecan in the LoVo Human Colorectal Cancer Xenograft Model. International Journal of Radiation Oncology Biology Physics, 2009, 75, 854-861.	0.4	13
67	DCE-MRI assessment of the effect of vandetanib on tumor vasculature in patients with advanced colorectal cancer and liver metastases: a randomized phase I study. Journal of Angiogenesis Research, 2009, 1, 5.	2.9	47
68	The Effects of Vandetanib on Paclitaxel Tumor Distribution and Antitumor Activity in a Xenograft Model of Human Ovarian Carcinoma. Neoplasia, 2009, 11, 1155-IN7.	2.3	31
69	Antitumor and antiangiogenic activity of cediranib in a preclinical model of renal cell carcinoma. Anticancer Research, 2009, 29, 5065-76.	0.5	15
70	Longitudinal in vivo susceptibility contrast MRI measurements of LS174T colorectal liver metastasis in nude mice. Journal of Magnetic Resonance Imaging, 2008, 28, 1451-1458.	1.9	19
71	Liposomal encapsulation enhances the antitumour efficacy of the vascular disrupting agent ZD6126 in murine B16.F10 melanoma. British Journal of Cancer, 2008, 99, 1256-1264.	2.9	28
72	Novel dual targeting strategy with vandetanib induces tumor cell apoptosis and inhibits angiogenesis in malignant pleural mesothelioma cells expressing RET oncogenic rearrangement. Cancer Letters, 2008, 265, 55-66.	3.2	26

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73	Dual inhibition of VEGFR and EGFR signaling reduces the incidence and size of intestinal adenomas in ApcMin/+ mice. Molecular Cancer Therapeutics, 2008, 7, 590-598.	1.9	36
74	Vascular Endothelial Growth Factor Receptor-1 Contributes to Resistance to Anti–Epidermal Growth Factor Receptor Drugs in Human Cancer Cells. Clinical Cancer Research, 2008, 14, 5069-5080.	3.2	139
75	Vandetanib Inhibits Growth of Adenoid Cystic Carcinoma in an Orthotopic Nude Mouse Model. Clinical Cancer Research, 2008, 14, 5081-5089.	3.2	23
76	Potent antitumor effects of ZD6474 on neuroblastoma via dual targeting of tumor cells and tumor endothelium. Molecular Cancer Therapeutics, 2008, 7, 418-424.	1.9	47
77	Vandetanib (ZACTIMAâ"¢; ZD6474): Preclinical and Clinical Development. , 2008, , 741-759.		0
78	Targeted therapy of orthotopic human lung cancer by combined vascular endothelial growth factor and epidermal growth factor receptor signaling blockade. Molecular Cancer Therapeutics, 2007, 6, 471-483.	1.9	84
79	Distribution of radioactivity and metabolite profiling in tumour and plasma following intravenous administration of a colchicine derivative (14C-ZD6126) to tumour-bearing mice. Xenobiotica, 2007, 37, 328-340.	0.5	10
80	Effect of Pretreatment With Atenolol and Nifedipine on ZD6126-Induced Cardiac Toxicity in Rats. Journal of the National Cancer Institute, 2007, 99, 1724-1728.	3.0	26
81	Antiangiogenic and antitumor activity of a novel vascular endothelial growth factor receptor-2 tyrosine kinase inhibitor ZD6474 in a metastatic human pancreatic tumor model. Anti-Cancer Drugs, 2007, 18, 569-579.	0.7	28
82	Vandetanib (ZD6474): an orally available receptor tyrosine kinase inhibitor that selectively targets pathways critical for tumor growth and angiogenesis. Expert Opinion on Investigational Drugs, 2007, 16, 239-249.	1.9	131
83	Correlation of MRI Biomarkers with Tumor Necrosis in Hras5 Tumor Xenograft in Athymic Rats. Neoplasia, 2007, 9, 382-391.	2.3	32
84	A5-02: Targeted therapy against VEGFR and/or EGFR signaling with AZD2171, vandetanib, and gefitinib as part of a combined modality approach for the treatment of non-small-cell lung cancer. Journal of Thoracic Oncology, 2007, 2, S323.	0.5	0
85	Susceptibility Contrast Magnetic Resonance Imaging Determination of Fractional Tumor Blood Volume: A Noninvasive Imaging Biomarker of Response to the Vascular Disrupting Agent ZD6126. International Journal of Radiation Oncology Biology Physics, 2007, 69, 872-879.	0.4	26
86	Targeted Therapy Against VEGFR and EGFR With ZD6474 Enhances the Therapeutic Efficacy of Irradiation in an Orthotopic Model of Human Non–Small-Cell Lung Cancer. International Journal of Radiation Oncology Biology Physics, 2007, 69, 1534-1543.	0.4	58
87	Sequence dependent antitumour efficacy of the vascular disrupting agent ZD6126 in combination with paclitaxel. British Journal of Cancer, 2007, 97, 888-894.	2.9	49
88	The Response of RIF-1 Fibrosarcomas to the Vascular-Disrupting Agent ZD6126 Assessed by In Vivo and Ex Vivo1H Magnetic Resonance Spectroscopy. Neoplasia, 2006, 8, 560-567.	2.3	36
89	Tumour overexpression of inducible nitric oxide synthase (iNOS) increases angiogenesis and may modulate the anti-tumour effects of the vascular disrupting agent ZD6126. Microvascular Research, 2006, 71, 76-84.	1.1	32
90	ZD6474, an Inhibitor of Vascular Endothelial Growth Factor Receptor Tyrosine Kinase, Inhibits Growth of Experimental Lung Metastasis and Production of Malignant Pleural Effusions in a Non-Small Cell Lung Cancer Model. Oncology Research, 2006, 16, 15-26.	0.6	45

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91	Inhibiting vascular endothelial growth factor receptor-2 signaling reduces tumor burden in the ApcMin/+ mouse model of early intestinal cancer. Carcinogenesis, 2006, 27, 2133-2139.	1.3	56
92	A Longitudinal Study of R2* and R2 Magnetic Resonance Imaging Relaxation Rate Measurements in Murine Liver After a Single Administration of 3 Different Iron Oxide-Based Contrast Agents. Investigative Radiology, 2005, 40, 784-791.	3.5	32
93	ZD6474 – a novel inhibitor of VEGFR and EGFR tyrosine kinase activity. British Journal of Cancer, 2005, 92, S6-S13.	2.9	160
94	ZD6474, a Novel Tyrosine Kinase Inhibitor of Vascular Endothelial Growth Factor Receptor and Epidermal Growth Factor Receptor, Inhibits Tumor Growth of Multiple Nervous System Tumors. Clinical Cancer Research, 2005, 11, 8145-8157.	3.2	94
95	Differential Effects of Vascular Endothelial Growth Factor Receptor-2 Inhibitor ZD6474 on Circulating Endothelial Progenitors and Mature Circulating Endothelial Cells: Implications for Use as a Surrogate Marker of Antiangiogenic Activity. Clinical Cancer Research, 2005, 11, 3514-3522.	3.2	145
96	Potential Antagonism of Tubulin-Binding Anticancer Agents in Combination Therapies. Clinical Cancer Research, 2005, 11, 2720-2726.	3.2	23
97	Antitumor Vascular Strategy for Controlling Experimental Metastatic Spread of Human Small-Cell Lung Cancer Cells with ZD6474 in Natural Killer Cell–Depleted Severe Combined Immunodeficient Mice. Clinical Cancer Research, 2005, 11, 8789-8798.	3.2	45
98	ZD6474 Suppresses Oncogenic RET Isoforms in a Drosophila Model for Type 2 Multiple Endocrine Neoplasia Syndromes and Papillary Thyroid Carcinoma. Cancer Research, 2005, 65, 3538-3541.	0.4	133
99	Cooperative Antitumor Effect of Multitargeted Kinase Inhibitor ZD6474 and Ionizing Radiation in Glioblastoma. Clinical Cancer Research, 2005, 11, 5639-5644.	3.2	83
100	Acute Tumor Response to ZD6126 Assessed by Intrinsic Susceptibility Magnetic Resonance Imaging. Neoplasia, 2005, 7, 466-474.	2.3	32
101	Vascular-Targeting Agents and Radiation Therapy in Lung Cancer: Where Do We Stand in 2005?. Clinical Lung Cancer, 2005, 7, 175-179.	1.1	6
102	Vascular Targeting in Pancreatic Cancer: The Novel Tubulin-Binding Agent ZD6126 Reveals Antitumor Activity in Primary and Metastatic Tumor Models. Neoplasia, 2005, 7, 957-966.	2.3	17
103	Effect of the tumor vascular-damaging agent, ZD6126, on the radioresponse of U87 glioblastoma. Clinical Cancer Research, 2005, 11, 835-42.	3.2	84
104	Regulation of p27Kip1 Protein Levels Contributes to Mitogenic Effects of the RET/PTC Kinase in Thyroid Carcinoma Cells. Cancer Research, 2004, 64, 3823-3829.	0.4	45
105	Magnetic Resonance Imaging Measurements of the Response of Murine and Human Tumors to the Vascular-Targeting Agent ZD6126. Clinical Cancer Research, 2004, 10, 3650-3657.	3.2	162
106	Inhibition of Src Tyrosine Kinase as Treatment for Human Pancreatic Cancer Growing Orthotopically in Nude Mice. Clinical Cancer Research, 2004, 10, 8028-8036.	3.2	128
107	ZD6126 inhibits orthotopic growth and peritoneal carcinomatosis in a mouse model of human gastric cancer. British Journal of Cancer, 2004, 90, 705-711.	2.9	21
108	Disease associated mutations at valine 804 in the RET receptor tyrosine kinase confer resistance to selective kinase inhibitors. Oncogene, 2004, 23, 6056-6063.	2.6	227

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109	In vivo videomicroscopy reveals differential effects of the vascular-targeting agent ZD6126 and the anti-angiogenic agent ZD6474 on vascular function in a liver metastasis model. Angiogenesis, 2004, 7, 157-164.	3.7	22
110	The VEGF receptor tyrosine kinase inhibitor, ZD6474, inhibits angiogenesis and affects microvascular architecture within an orthotopically implanted renal cell carcinoma. Angiogenesis, 2004, 7, 347-354.	3.7	36
111	Single Dose of the Antivascular Agent, ZD6126 (N-Acetylcoichinol-O-Phosphate), Reduces Perfusion for at Least 96 Hours in the GH3 Prolactinoma Rat Tumor Model. Neoplasia, 2004, 6, 150-157.	2.3	34
112	ZD6474, a vascular endothelial growth factor receptor tyrosine kinase inhibitor with additional activity against epidermal growth factor receptor tyrosine kinase, inhibits orthotopic growth and angiogenesis of gastric cancer. Molecular Cancer Therapeutics, 2004, 3, 1041-8.	1.9	75
113	Tumour dose response to the antivascular agent ZD6126 assessed by magnetic resonance imaging. British Journal of Cancer, 2003, 88, 1592-1597.	2.9	114
114	ZD6126: A novel small molecule vascular targeting agent. International Journal of Radiation Oncology Biology Physics, 2002, 54, 1497-1502.	0.4	66
115	Antitumor activity of the novel vascular targeting agent ZD6126 in a panel of tumor models. Clinical Cancer Research, 2002, 8, 1974-83.	3.2	135
116	ZD6474, an orally available inhibitor of KDR tyrosine kinase activity, efficiently blocks oncogenic RET kinases. Cancer Research, 2002, 62, 7284-90.	0.4	463
117	Lack of correlation between residual radiation-induced DNA damage, in keratinocytes assayed directly from skin, and late radiotherapy reactions in breast cancer patients. International Journal of Radiation Oncology Biology Physics, 1999, 43, 481-487.	0.4	16
118	A correlation between residual radiation-induced DNA double-strand breaks in cultured fibroblasts and late radiotherapy reactions in breast cancer patients. Radiotherapy and Oncology, 1999, 51, 55-65.	0.3	55
119	Targeting Double-Strand Breaks to Replicating DNA Identifies a Subpathway of DSB Repair That Is Defective in Ataxia-Telangiectasia Cells. Biochemical and Biophysical Research Communications, 1999, 261, 317-325.	1.0	53
120	SYBR Green I staining of pulsed field agarose gels is a sensitive and inexpensive way of quantitating DNA double-strand breaks in mammalian cells. Nucleic Acids Research, 1997, 25, 2945-2946.	6.5	47
121	A correlation between residual DNA double-strand breaks and clonogenic measurements of radiosensitivity in fibroblasts from preradiotherapy cervix cancer patients. International Journal of Radiation Oncology Biology Physics, 1997, 39, 1137-1144.	0.4	36
122	Dominant genetic instability and sensitivity to DNA damaging agents in a mammalian cell line. Somatic Cell and Molecular Genetics, 1996, 22, 177-189.	0.7	1
123	Different fates of camptothecin-induced replication fork-associated double-strand DNA breaks in mammalian cells. Carcinogenesis, 1994, 15, 823-828.	1.3	47
124	Problems and paradigms: Fine tuning of DNA repair in transcribed genes: Mechanisms, prevalence and consequences. BioEssays, 1993, 15, 209-216.	1.2	26
125	Characterisation and correction of a mammalian cell mutant defective in late step of base excision repair. Somatic Cell and Molecular Genetics, 1992, 18, 529-541.	0.7	1
126	A class of amphipathic proteins associated with lipid storage bodies in plants. Possible similarities with animal serum apolipoproteins. Biochimica Et Biophysica Acta Gene Regulatory Mechanisms, 1991, 1088, 86-94.	2.4	70

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127	Camptothecin cytotoxicity in mammalian cells is associated with the induction of persistent double strand breaks in replicating DNA. Nucleic Acids Research, 1991, 19, 3295-3300.	6.5	211
128	Genomic sequence of a 12S seed storage protein from oilseed rape (Brassica napus c.v.jet neuf). Nucleic Acids Research, 1989, 17, 3584-3584.	6.5	23
129	Increased oncogene expression in livers of rats after exposure to dimethylnitrosamine. Biochemical Society Transactions, 1988, 16, 1058-1059.	1.6	4
130	Micrococcal nuclease digests intranucleosomal DNA of inactive genes more rapidly than active genes. Biochemical Society Transactions, 1988, 16, 1060-1061.	1.6	0
131	Highly selective binding of the carcinogen benzo[<i>a</i>]pyrene diol epoxide to nuclear matrix/scaffold DNA. Biochemical Society Transactions, 1986, 14, 1164-1165.	1.6	1
132	Selective repair of methylated purines in regions of chromatin DNA. Carcinogenesis, 1986, 7, 1497-1503.	1.3	43
133	Preferential binding of the carcinogen benzo[a]pyrene to DNA in active chromatin and the nuclear matrix. Carcinogenesis, 1986, 7, 907-913.	1.3	47