

# Scott H Kaufmann

## List of Publications by Year in descending order

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336  
papers

28,461  
citations

5569

82  
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6128

159  
g-index

434  
all docs

434  
docs citations

434  
times ranked

29223  
citing authors

#	ARTICLE	IF	CITATIONS
1	Mammalian Caspases: Structure, Activation, Substrates, and Functions During Apoptosis. Annual Review of Biochemistry, 1999, 68, 383-424.	5.0	2,499
2	PARP inhibition: PARP1 and beyond. Nature Reviews Cancer, 2010, 10, 293-301.	12.8	1,166
3	Induction of Apoptosis by Cancer Chemotherapy. Experimental Cell Research, 2000, 256, 42-49.	1.2	1,101
4	Rucaparib in relapsed, platinum-sensitive high-grade ovarian carcinoma (ARIEL2 Part 1): an international, multicentre, open-label, phase 2 trial. Lancet Oncology, The, 2017, 18, 75-87.	5.1	975
5	Cathepsin B contributes to TNF- $\alpha$ -mediated hepatocyte apoptosis by promoting mitochondrial release of cytochrome c. Journal of Clinical Investigation, 2000, 106, 1127-1137.	3.9	635
6	Programmed cell death: alive and well in the new millennium. Trends in Cell Biology, 2001, 11, 526-534.	3.6	603
7	The Current Status of Camptothecin Analogues as Antitumor Agents. Journal of the National Cancer Institute, 1993, 85, 271-291.	3.0	574
8	The role of proteases during apoptosis. FASEB Journal, 1996, 10, 587-597.	0.2	538
9	Caspases and caspase inhibitors. Trends in Biochemical Sciences, 1997, 22, 388-393.	3.7	517
10	Phase II Trial of Single-Agent Temsirolimus (CCI-779) for Relapsed Mantle Cell Lymphoma. Journal of Clinical Oncology, 2005, 23, 5347-5356.	0.8	509
11	Toxic bile salts induce rodent hepatocyte apoptosis via direct activation of Fas. Journal of Clinical Investigation, 1999, 103, 137-145.	3.9	485
12	Nonhomologous end joining drives poly(ADP-ribose) polymerase (PARP) inhibitor lethality in homologous recombination-deficient cells. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 3406-3411.	3.3	475
13	Clinical and biologic activity of the farnesyltransferase inhibitor R115777 in adults with refractory and relapsed acute leukemias: a phase 1 clinical-laboratory correlative trial. Blood, 2001, 97, 3361-3369.	0.6	445
14	Secondary Somatic Mutations Restoring <i>RAD51C</i> and <i>RAD51D</i> Associated with Acquired Resistance to the PARP Inhibitor Rucaparib in High-Grade Ovarian Carcinoma. Cancer Discovery, 2017, 7, 984-998.	7.7	310
15	Poly (ADP-Ribose) Polymerase Inhibitors: Recent Advances and Future Development. Journal of Clinical Oncology, 2015, 33, 1397-1406.	0.8	295
16	<i>BRCA</i> Reversion Mutations in Circulating Tumor DNA Predict Primary and Acquired Resistance to the PARP Inhibitor Rucaparib in High-Grade Ovarian Carcinoma. Cancer Discovery, 2019, 9, 210-219.	7.7	278
17	Considerations in the isolation of rat liver nuclear matrix, nuclear envelope, and pore complex lamina. Experimental Cell Research, 1981, 132, 105-123.	1.2	270
18	Elevated Expression of the Apoptotic Regulator Mcl-1 at the Time of Leukemic Relapse. Blood, 1998, 91, 991-1000.	0.6	265

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19	Elevated Expression of the Apoptotic Regulator Mcl-1 at the Time of Leukemic Relapse. <i>Blood</i> , 1998, 91, 991-1000.	0.6	263
20	The role of Mcl-1 downregulation in the proapoptotic activity of the multikinase inhibitor BAY 43-9006. <i>Oncogene</i> , 2005, 24, 6861-6869.	2.6	254
21	Alterations in the apoptotic machinery and their potential role in anticancer drug resistance. <i>Oncogene</i> , 2003, 22, 7414-7430.	2.6	253
22	Comparison of Apoptosis in Wild-Type and Fas-Resistant Cells: Chemotherapy-Induced Apoptosis Is Not Dependent on Fas/Fas Ligand Interactions. <i>Blood</i> , 1997, 90, 935-943.	0.6	247
23	Cell death induced by topoisomerase-targeted drugs: more questions than answers. <i>Biochimica Et Biophysica Acta Gene Regulatory Mechanisms</i> , 1998, 1400, 195-211.	2.4	239
24	Synthetic Smac/DIABLO Peptides Enhance the Effects of Chemotherapeutic Agents by Binding XIAP and cIAP1 in Situ. <i>Journal of Biological Chemistry</i> , 2002, 277, 44236-44243.	1.6	239
25	Caspase-6 gene disruption reveals a requirement for lamin A cleavage in apoptotic chromatin condensation. <i>EMBO Journal</i> , 2002, 21, 1967-1977.	3.5	233
26	Low-dose, single-agent temsirolimus for relapsed mantle cell lymphoma. <i>Cancer</i> , 2008, 113, 508-514.	2.0	220
27	Loss of HSulf-1 Up-regulates Heparin-binding Growth Factor Signaling in Cancer. <i>Journal of Biological Chemistry</i> , 2003, 278, 23107-23117.	1.6	215
28	Mcl-1 Mediates Tumor Necrosis Factor-Related Apoptosis-Inducing Ligand Resistance in Human Cholangiocarcinoma Cells. <i>Cancer Research</i> , 2004, 64, 3517-3524.	0.4	204
29	Interleukin-6 Contributes to Mcl-1 Up-regulation and TRAIL Resistance via an Akt-Signaling Pathway in Cholangiocarcinoma Cells. <i>Gastroenterology</i> , 2005, 128, 2054-2065.	0.6	204
30	Failure of Iniparib to Inhibit Poly(ADP-Ribose) Polymerase <i>In Vitro</i> . <i>Clinical Cancer Research</i> , 2012, 18, 1655-1662.	3.2	204
31	Phase II Study of the Farnesyl Transferase Inhibitor R115777 in Patients With Advanced Non-Small-Cell Lung Cancer. <i>Journal of Clinical Oncology</i> , 2003, 21, 1760-1766.	0.8	200
32	COMMD1 is linked to the WASH complex and regulates endosomal trafficking of the copper transporter ATP7A. <i>Molecular Biology of the Cell</i> , 2015, 26, 91-103.	0.9	200
33	Activation of Multiple Interleukin-1 <sup>2</sup> Converting Enzyme Homologues in Cytosol and Nuclei of HL-60 Cells during Etoposide-induced Apoptosis. <i>Journal of Biological Chemistry</i> , 1997, 272, 7421-7430.	1.6	197
34	Methylation of all BRCA1 copies predicts response to the PARP inhibitor rucaparib in ovarian carcinoma. <i>Nature Communications</i> , 2018, 9, 3970.	5.8	192
35	Olaparib and $\pm$ -specific PI3K inhibitor alpelisib for patients with epithelial ovarian cancer: a dose-escalation and dose-expansion phase 1b trial. <i>Lancet Oncology</i> , The, 2019, 20, 570-580.	5.1	191
36	Apoptosis in cancer: cause and cure. <i>BioEssays</i> , 2000, 22, 1007-1017.	1.2	181

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37	A phase 2 study of the farnesyltransferase inhibitor tipifarnib in poor-risk and elderly patients with previously untreated acute myelogenous leukemia. <i>Blood</i> , 2007, 109, 1387-1394.	0.6	180
38	S Phase and G2 Arrests Induced by Topoisomerase I Poisons Are Dependent on ATR Kinase Function. <i>Journal of Biological Chemistry</i> , 2002, 277, 1599-1606.	1.6	179
39	Severe Graft-versus-Host Disease in a Liver-Transplant Recipient. <i>New England Journal of Medicine</i> , 1988, 318, 689-691.	13.9	174
40	G1 and G2 cell-cycle arrest following microtubule depolymerization in human breast cancer cells. <i>Journal of Clinical Investigation</i> , 2002, 110, 91-99.	3.9	173
41	The erasable Western blot. <i>Analytical Biochemistry</i> , 1987, 161, 89-95.	1.1	172
42	Tumorgrafts as <i>In Vivo</i> Surrogates for Women with Ovarian Cancer. <i>Clinical Cancer Research</i> , 2014, 20, 1288-1297.	3.2	168
43	Bile acids induce cyclooxygenase-2 expression via the epidermal growth factor receptor in a human cholangiocarcinoma cell line. <i>Gastroenterology</i> , 2002, 122, 985-993.	0.6	166
44	Farnesyltransferase inhibitor tipifarnib is well tolerated, induces stabilization of disease, and inhibits farnesylation and oncogenic/tumor survival pathways in patients with advanced multiple myeloma. <i>Blood</i> , 2004, 103, 3271-3277.	0.6	163
45	ATR Inhibition Broadly Sensitizes Ovarian Cancer Cells to Chemotherapy Independent of BRCA Status. <i>Cancer Research</i> , 2013, 73, 3683-3691.	0.4	160
46	A candidate tumor suppressor HtrA1 is downregulated in ovarian cancer. <i>Oncogene</i> , 2004, 23, 1636-1644.	2.6	157
47	Cytotoxic Effects of Topotecan Combined With Various Anticancer Agents in Human Cancer Cell Lines. <i>Journal of the National Cancer Institute</i> , 1996, 88, 734-741.	3.0	153
48	APOBEC3B Upregulation and Genomic Mutation Patterns in Serous Ovarian Carcinoma. <i>Cancer Research</i> , 2013, 73, 7222-7231.	0.4	153
49	A subset of non-histone nuclear proteins reversibly stabilized by the sulfhydryl cross-linking reagent tetrathionate. <i>Experimental Cell Research</i> , 1984, 155, 477-495.	1.2	152
50	Inhibition of histone deacetylase overcomes rapamycin-mediated resistance in diffuse large B-cell lymphoma by inhibiting Akt signaling through mTORC2. <i>Blood</i> , 2009, 114, 2926-2935.	0.6	152
51	Transient binding of an activator BH3 domain to the Bak BH3-binding groove initiates Bak oligomerization. <i>Journal of Cell Biology</i> , 2011, 194, 39-48.	2.3	139
52	Successful Virtual Screening of a Chemical Database for Farnesyltransferase Inhibitor Leads. <i>Journal of Medicinal Chemistry</i> , 2000, 43, 401-408.	2.9	130
53	Serine protease HtrA1 modulates chemotherapy-induced cytotoxicity. <i>Journal of Clinical Investigation</i> , 2006, 116, 1994-2004.	3.9	130
54	Chemotherapy-Induced Apoptosis. <i>Advances in Pharmacology</i> , 1997, 41, 461-499.	1.2	126

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55	Transition from Caspase-dependent to Caspase-independent Mechanisms at the Onset of Apoptotic Execution. <i>Journal of Cell Biology</i> , 1998, 143, 225-239.	2.3	122
56	Emerging understanding of Bcl-2 biology: Implications for neoplastic progression and treatment. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2015, 1853, 1658-1671.	1.9	122
57	Comparison of Caspase Activation and Subcellular Localization in HL-60 and K562 Cells Undergoing Etoposide-Induced Apoptosis. <i>Blood</i> , 1997, 90, 4283-4296.	0.6	119
58	Gemcitabine-Induced Activation of Checkpoint Signaling Pathways That Affect Tumor Cell Survival. <i>Molecular Pharmacology</i> , 2005, 68, 1636-1644.	1.0	119
59	Heat shock protein 90 inhibition sensitizes acute myelogenous leukemia cells to cytarabine. <i>Blood</i> , 2005, 106, 318-327.	0.6	118
60	Molecular correlates of platinum response in human high-grade serous ovarian cancer patient-derived xenografts. <i>Molecular Oncology</i> , 2014, 8, 656-668.	2.1	117
61	Molecular and clinical determinants of response and resistance to rucaparib for recurrent ovarian cancer treatment in ARIEL2 (Parts 1 and 2). <i>Nature Communications</i> , 2021, 12, 2487.	5.8	116
62	Caspases 3 and 9 Send a Pro-Apoptotic Signal from Synapse to Cell Body in Olfactory Receptor Neurons. <i>Journal of Neuroscience</i> , 2001, 21, 7099-7109.	1.7	114
63	Tumor Necrosis Factor-related Apoptosis-inducing Ligand Activates a Lysosomal Pathway of Apoptosis That Is Regulated by Bcl-2 Proteins. <i>Journal of Biological Chemistry</i> , 2007, 282, 28960-28970.	1.6	113
64	The molecular origin and taxonomy of mucinous ovarian carcinoma. <i>Nature Communications</i> , 2019, 10, 3935.	5.8	110
65	BCL2 mutations are associated with increased risk of transformation and shortened survival in follicular lymphoma. <i>Blood</i> , 2015, 125, 658-667.	0.6	108
66	Effects of the Bcr/abl kinase inhibitors STI571 and adaphostin (NSC 680410) on chronic myelogenous leukemia cells in vitro. <i>Blood</i> , 2002, 99, 664-671.	0.6	107
67	Death Receptor 5 Signaling Promotes Hepatocyte Lipoapoptosis. <i>Journal of Biological Chemistry</i> , 2011, 286, 39336-39348.	1.6	106
68	Human INCENP colocalizes with the Aurora-B/AIRK2 kinase on chromosomes and is overexpressed in tumour cells. <i>Chromosoma</i> , 2001, 110, 65-74.	1.0	104
69	MCL-1 as a Buffer for Proapoptotic BCL-2 Family Members during TRAIL-induced Apoptosis. <i>Journal of Biological Chemistry</i> , 2007, 282, 29831-29846.	1.6	104
70	The relationship of the nuclear matrix to cellular structure and function. <i>Advances in Enzyme Regulation</i> , 1979, 17, 213-248.	2.9	103
71	Cytotoxic synergy between the multikinase inhibitor sorafenib and the proteasome inhibitor bortezomib in vitro: induction of apoptosis through Akt and c-Jun NH2-terminal kinase pathways. <i>Molecular Cancer Therapeutics</i> , 2006, 5, 2378-2387.	1.9	102
72	Phase I and Pharmacologic Trial of Cytosine Arabinoside with the Selective Checkpoint 1 Inhibitor Sch 900776 in Refractory Acute Leukemias. <i>Clinical Cancer Research</i> , 2012, 18, 6723-6731.	3.2	100

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73	Retention of the Human Rad9 Checkpoint Complex in Extraction-resistant Nuclear Complexes after DNA Damage. <i>Journal of Biological Chemistry</i> , 2000, 275, 26343-26348.	1.6	99
74	Calpain-mediated X-linked Inhibitor of Apoptosis Degradation in Neutrophil Apoptosis and Its Impairment in Chronic Neutrophilic Leukemia. <i>Journal of Biological Chemistry</i> , 2002, 277, 33968-33977.	1.6	96
75	The Elephant and the Blind Men: Making Sense of PARP Inhibitors in Homologous Recombination Deficient Tumor Cells. <i>Frontiers in Oncology</i> , 2013, 3, 228.	1.3	95
76	A cell cycle-dependent BRCA1â€UHRF1 cascade regulates DNA double-strand break repair pathway choice. <i>Nature Communications</i> , 2016, 7, 10201.	5.8	95
77	Caspase-mediated Cleavage of DNA Topoisomerase I at Unconventional Sites during Apoptosis. <i>Journal of Biological Chemistry</i> , 1999, 274, 4335-4340.	1.6	94
78	Serine 64 Phosphorylation Enhances the Antiapoptotic Function of Mcl-1. <i>Journal of Biological Chemistry</i> , 2007, 282, 18407-18417.	1.6	94
79	Current status of clinical trials of farnesyltransferase inhibitors. <i>Current Opinion in Oncology</i> , 2001, 13, 470-476.	1.1	93
80	The Role of Checkpoint Kinase 1 in Sensitivity to Topoisomerase I Poisons. <i>Journal of Biological Chemistry</i> , 2005, 280, 14349-14355.	1.6	92
81	Dual mTORC1/mTORC2 inhibition diminishes Akt activation and induces Puma-dependent apoptosis in lymphoid malignancies. <i>Blood</i> , 2012, 119, 476-487.	0.6	91
82	Spartan deficiency causes accumulation of Topoisomerase 1 cleavage complexes and tumorigenesis. <i>Nucleic Acids Research</i> , 2017, 45, 4564-4576.	6.5	91
83	Enhanced Killing of Cancer Cells by Poly(ADP-ribose) Polymerase Inhibitors and Topoisomerase I Inhibitors Reflects Poisoning of Both Enzymes. <i>Journal of Biological Chemistry</i> , 2012, 287, 4198-4210.	1.6	89
84	Somatic Mosaic Mutations in <i>PPM1D</i> and <i>TP53</i> in the Blood of Women With Ovarian Carcinoma. <i>JAMA Oncology</i> , 2016, 2, 370.	3.4	88
85	Prime, Shock, and Kill: Priming CD4 T Cells from HIV Patients with a BCL-2 Antagonist before HIV Reactivation Reduces HIV Reservoir Size. <i>Journal of Virology</i> , 2016, 90, 4032-4048.	1.5	85
86	Phase I and Pharmacokinetic Study of Flavopiridol followed by 1- $\beta$ -d-Arabinofuranosylcytosine and Mitoxantrone in Relapsed and Refractory Adult Acute Leukemias. <i>Clinical Cancer Research</i> , 2005, 11, 8403-8412.	3.2	84
87	A Multistep Model for Paclitaxel-Induced Apoptosis in Human Breast Cancer Cell Lines. <i>Experimental Cell Research</i> , 2001, 270, 277-288.	1.2	81
88	Contribution of Bcl-2 Phosphorylation to Bak Binding and Drug Resistance. <i>Cancer Research</i> , 2013, 73, 6998-7008.	0.4	81
89	Inhibition of epidermal growth factor receptor kinase induces protease-dependent apoptosis in human colon cancer cells. <i>Gastroenterology</i> , 1998, 114, 930-939.	0.6	80
90	Noxa/Bcl-2 Protein Interactions Contribute to Bortezomib Resistance in Human Lymphoid Cells. <i>Journal of Biological Chemistry</i> , 2011, 286, 17682-17692.	1.6	80

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91	Pooled Clustering of High-Grade Serous Ovarian Cancer Gene Expression Leads to Novel Consensus Subtypes Associated with Survival and Surgical Outcomes. <i>Clinical Cancer Research</i> , 2017, 23, 4077-4085.	3.2	80
92	Comparison of Paclitaxel-, 5-Fluoro-2â€²-deoxyuridine-, and Epidermal Growth Factor (EGF)-induced Apoptosis. <i>Journal of Biological Chemistry</i> , 1999, 274, 15927-15936.	1.6	79
93	Apoptosis and the response to anticancer therapy. <i>Current Opinion in Oncology</i> , 2001, 13, 453-462.	1.1	79
94	Is TRAIL hepatotoxic?. <i>Hepatology</i> , 2001, 34, 3-6.	3.6	77
95	Selectively targeting Mcl-1 for the treatment of acute myelogenous leukemia and solid tumors: Figure 1.. <i>Genes and Development</i> , 2012, 26, 305-311.	2.7	77
96	Adaphostin-induced oxidative stress overcomes BCR/ABL mutation-dependent and -independent imatinib resistance. <i>Blood</i> , 2006, 107, 2501-2506.	0.6	76
97	Apoptosis-associated caspase activation assays. <i>Methods</i> , 2008, 44, 262-272.	1.9	76
98	Phorbol 12-myristate 13-Acetate Inhibits Death Receptor-mediated Apoptosis in Jurkat Cells by Disrupting Recruitment of Fas-associated Polypeptide with Death Domain. <i>Journal of Biological Chemistry</i> , 2002, 277, 3776-3783.	1.6	72
99	Comparison of Apoptosis in Wild-Type and Fas-Resistant Cells: Chemotherapy-Induced Apoptosis Is Not Dependent on Fas/Fas Ligand Interactions. <i>Blood</i> , 1997, 90, 935-943.	0.6	72
100	Bile acids inhibit Mcl-1 protein turnover via an epidermal growth factor receptor/Raf-1-dependent mechanism. <i>Cancer Research</i> , 2002, 62, 6500-5.	0.4	72
101	Inhibition of the phosphatidylinositol 3-kinase/mammalian target of rapamycin pathway in hematologic malignancies. <i>Current Treatment Options in Oncology</i> , 2006, 7, 285-294.	1.3	70
102	Death Receptor 5 Internalization Is Required for Lysosomal Permeabilization by TRAIL in Malignant Liver Cell Lines. <i>Gastroenterology</i> , 2009, 136, 2365-2376.e7.	0.6	68
103	FAM111A protects replication forks from protein obstacles via its trypsin-like domain. <i>Nature Communications</i> , 2020, 11, 1318.	5.8	67
104	Effect of adding the topoisomerase I poison 7-ethyl-10-hydroxycamptothecin (SN-38) to 5-fluorouracil and folinic acid in HCT-8 cells: elevated dTTP pools and enhanced cytotoxicity. <i>Cancer Chemotherapy and Pharmacology</i> , 1998, 42, 391-399.	1.1	66
105	Involvement of reactive oxygen species in adaphostin-induced cytotoxicity in human leukemia cells. <i>Blood</i> , 2003, 102, 4512-4519.	0.6	66
106	Phase 1 Trial of Flavopiridol Combined with Cisplatin or Carboplatin in Patients with Advanced Malignancies with the Assessment of Pharmacokinetic and Pharmacodynamic End Points. <i>Clinical Cancer Research</i> , 2005, 11, 5935-5941.	3.2	65
107	Apoptosis in the treatment of cancer: a promise kept?. <i>Current Opinion in Cell Biology</i> , 2006, 18, 668-676.	2.6	65
108	Evaluation of the BH3-only Protein Puma as a Direct Bak Activator. <i>Journal of Biological Chemistry</i> , 2014, 289, 89-99.	1.6	65



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109	Lack of Correlation between Caspase Activation and Caspase Activity Assays in Paclitaxel-treated MCF-7 Breast Cancer Cells. <i>Journal of Biological Chemistry</i> , 2002, 277, 804-815.	1.6	64
110	How does doxorubicin work?. <i>ELife</i> , 2012, 1, e00387.	2.8	64
111	Characterization of Caspase Processing and Activation in HL-60 Cell Cytosol Under Cell-free Conditions. <i>Journal of Biological Chemistry</i> , 1999, 274, 22635-22645.	1.6	63
112	Reutilization of Immunoblots after Chemiluminescent Detection. <i>Analytical Biochemistry</i> , 2001, 296, 283-286.	1.1	62
113	Analysis of the internal nuclear matrix. <i>Experimental Cell Research</i> , 1986, 164, 139-153.	1.2	61
114	Alteration of the Nucleolar Localization of Poly(ADP-ribose) Polymerase upon Treatment with Transcription Inhibitors. <i>Experimental Cell Research</i> , 1996, 227, 146-153.	1.2	61
115	Components of the Cell Death Machine and Drug Sensitivity of the National Cancer Institute Cell Line Panel. <i>Clinical Cancer Research</i> , 2004, 10, 6807-6820.	3.2	61
116	CXCR4 Chemokine Receptor Signaling Induces Apoptosis in Acute Myeloid Leukemia Cells via Regulation of the Bcl-2 Family Members Bcl-XL, Noxa, and Bak. <i>Journal of Biological Chemistry</i> , 2013, 288, 22899-22914.	1.6	59
117	APOBEC3G Expression Correlates with T-Cell Infiltration and Improved Clinical Outcomes in High-grade Serous Ovarian Carcinoma. <i>Clinical Cancer Research</i> , 2016, 22, 4746-4755.	3.2	59
118	Association of topoisomerase II with the hepatoma cell nuclear matrix: The role of intermolecular disulfide bond formation. <i>Experimental Cell Research</i> , 1991, 192, 511-523.	1.2	57
119	Thromboembolism in Adults with Acute Lymphoblastic Leukemia During Induction with L-Asparaginase-containing Multi-agent Regimens: Incidence, Risk Factors, and Possible Role of Antithrombin. <i>Leukemia and Lymphoma</i> , 2004, 45, 1545-1551.	0.6	57
120	Epigenetic silencing of TCEAL7 (Bex4) in ovarian cancer. <i>Oncogene</i> , 2005, 24, 5089-5100.	2.6	57
121	In vivo anti-tumor activity of the PARP inhibitor niraparib in homologous recombination deficient and proficient ovarian carcinoma. <i>Gynecologic Oncology</i> , 2016, 143, 379-388.	0.6	57
122	Association of poly(ADP-ribose) polymerase with the nuclear matrix: The role intermolecular disulfide bond formation, RNA retention, and cell type. <i>Experimental Cell Research</i> , 1991, 192, 524-535.	1.2	56
123	A Phase 1 Study of the PARP Inhibitor Veliparib in Combination with Temozolomide in Acute Myeloid Leukemia. <i>Clinical Cancer Research</i> , 2017, 23, 697-706.	3.2	56
124	53BP1 as a potential predictor of response in PARP inhibitor-treated homologous recombination-deficient ovarian cancer. <i>Gynecologic Oncology</i> , 2019, 153, 127-134.	0.6	56
125	Evaluation of Apaf-1 and procaspases-2, -3, -7, -8, and -9 as potential prognostic markers in acute leukemia. <i>Blood</i> , 2000, 96, 3922-3931.	0.6	54
126	Active oral regimen for elderly adults with newly diagnosed acute myelogenous leukemia: a preclinical and phase 1 trial of the farnesyltransferase inhibitor tipifarnib (R115777, Zarnestra) combined with etoposide. <i>Blood</i> , 2009, 113, 4841-4852.	0.6	54



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127	Maintenance of the HIV Reservoir Is Antagonized by Selective BCL2 Inhibition. <i>Journal of Virology</i> , 2017, 91, .	1.5	54
128	Expression of Insulin Receptor Isoform A and Insulin-like Growth Factor-1 Receptor in Human Acute Myelogenous Leukemia: Effect of the Dual-Receptor Inhibitor BMS-536924 <i>In vitro</i> . <i>Cancer Research</i> , 2009, 69, 7635-7643.	0.4	53
129	Effects of Selective Checkpoint Kinase 1 Inhibition on Cytarabine Cytotoxicity in Acute Myelogenous Leukemia Cells <i>In Vitro</i> . <i>Clinical Cancer Research</i> , 2012, 18, 5364-5373.	3.2	53
130	Central Role of Fas-associated Death Domain Protein in Apoptosis Induction by the Mitogen-activated Protein Kinase Kinase Inhibitor CI-1040 (PD184352) in Acute Lymphocytic Leukemia Cells <i>In Vitro</i> . <i>Journal of Biological Chemistry</i> , 2003, 278, 47326-47339.	1.6	52
131	Altered Formation of Topotecan-Stabilized Topoisomerase I-DNA Adducts in Human Leukemia Cells. <i>Blood</i> , 1997, 89, 2098-2104.	0.6	51
132	Decreased drug accumulation in a mitoxantrone-resistant gastric carcinoma cell line in the absence of P-glycoprotein. , 1997, 71, 817-824.		50
133	Detection of DNA Cleavage in Apoptotic Cells. <i>Methods in Enzymology</i> , 2000, 322, 3-15.	0.4	50
134	Prospects for the Use of ATR Inhibitors to Treat Cancer. <i>Pharmaceuticals</i> , 2010, 3, 1311-1334.	1.7	50
135	4EBP1/c-MYC/PLUMA and NF- $\kappa$ B/EGR1/BIM pathways underlie cytotoxicity of mTOR dual inhibitors in malignant lymphoid cells. <i>Blood</i> , 2016, 127, 2711-2722.	0.6	49
136	Therapeutic options for mucinous ovarian carcinoma. <i>Gynecologic Oncology</i> , 2020, 156, 552-560.	0.6	49
137	Phosphorylated Forms of Activated Caspases Are Present in Cytosol From HL-60 Cells During Etoposide-Induced Apoptosis. <i>Blood</i> , 1998, 92, 3042-3049.	0.6	48
138	Detection of Poly(ADP-Ribose) Polymerase and Its Apoptosis-Specific Fragment by a Nonisotopic Activityâ€“Western Blot Technique. <i>Analytical Biochemistry</i> , 1995, 232, 251-254.	1.1	47
139	Genomic Mechanisms of p210BCR-ABL Signaling. <i>Journal of Biological Chemistry</i> , 2004, 279, 35604-35615.	1.6	47
140	Context-dependent Bcl-2/Bak Interactions Regulate Lymphoid Cell Apoptosis. <i>Journal of Biological Chemistry</i> , 2009, 284, 18311-18322.	1.6	47
141	Poly(ADP-ribose) Polymerase Inhibitors Sensitize Cancer Cells to Death Receptor-mediated Apoptosis by Enhancing Death Receptor Expression. <i>Journal of Biological Chemistry</i> , 2014, 289, 20543-20558.	1.6	47
142	Platelet-derived Growth Factor Primes Cancer-associated Fibroblasts for Apoptosis. <i>Journal of Biological Chemistry</i> , 2014, 289, 22835-22849.	1.6	47
143	<i>TP53</i> mutations, tetraploidy and homologous recombination repair defects in early stage high-grade serous ovarian cancer. <i>Nucleic Acids Research</i> , 2015, 43, 6945-6958.	6.5	46
144	Tyrosine Phosphorylation of Mitochondrial Creatine Kinase 1 Enhances a Druggable Tumor Energy Shuttle Pathway. <i>Cell Metabolism</i> , 2018, 28, 833-847.e8.	7.2	46

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145	The DNA Cytosine Deaminase APOBEC3B is a Molecular Determinant of Platinum Responsiveness in Clear Cell Ovarian Cancer. <i>Clinical Cancer Research</i> , 2020, 26, 3397-3407.	3.2	45
146	RAS mutations drive proliferative chronic myelomonocytic leukemia via a KMT2A-PLK1 axis. <i>Nature Communications</i> , 2021, 12, 2901.	5.8	44
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