

# Khandan Keyomarsi

## List of Publications by Year in descending order

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Version: 2024-02-01

153  
papers

14,303  
citations

53660

45  
h-index

19136

118  
g-index

156  
all docs

156  
docs citations

156  
times ranked

15666  
citing authors

#	ARTICLE	IF	CITATIONS
1	Palbociclib plus endocrine therapy significantly enhances overall survival of <sc>HR</sc>+<sc>HER2</sc> metastatic breast cancer patients compared to endocrine therapy alone in the secondâ€line setting: A large institutional study. International Journal of Cancer, 2022, 150, 2025-2037.	2.3	16
2	Abstract P2-05-02: Low molecular weight cyclin E facilitate replication stress tolerance in breast cancer development. Cancer Research, 2022, 82, P2-05-02-P2-05-02.	0.4	0
3	PARP inhibitors as single agents and in combination therapy: the most promising treatment strategies in clinical trials for BRCA-mutant ovarian and triple-negative breast cancers. Expert Opinion on Investigational Drugs, 2022, 31, 607-631.	1.9	20
4	CHD1 Promotes Sensitivity to Aurora Kinase Inhibitors by Suppressing Interaction of AURKA with Its Coactivator TPX2. Cancer Research, 2022, 82, 3088-3101.	0.4	2
5	Cytoplasmic Cyclin E Expression Predicts for Response to Neoadjuvant Chemotherapy in Breast Cancer. Annals of Surgery, 2021, 274, e150-e159.	2.1	5
6	Targeting Replicative Stress and DNA Repair by Combining PARP and Wee1 Kinase Inhibitors Is Synergistic in Triple Negative Breast Cancers with Cyclin E or BRCA1 Alteration. Cancers, 2021, 13, 1656.	1.7	16
7	LMW cyclin E and its novel catalytic partner CDK5 are therapeutic targets and prognostic biomarkers in salivary gland cancers. Oncogenesis, 2021, 10, 40.	2.1	2
8	Phase I safety and efficacy study of autophagy inhibition with hydroxychloroquine to augment the antiproliferative and biological effects of preoperative palbociclib plus letrozole for estrogen receptor-positive, HER2-negative metastatic breast cancer (MBC).. Journal of Clinical Oncology, 2021, 39, 1067-1067.	0.8	1
9	Selective CDK4/6 Inhibitors: Biologic Outcomes, Determinants of Sensitivity, Mechanisms of Resistance, Combinatorial Approaches, and Pharmacodynamic Biomarkers. American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting, 2020, 40, 115-126.	1.8	16
10	Cytoplasmic Cyclin E Is an Independent Marker of Aggressive Tumor Biology and Breast Cancer-Specific Mortality in Women over 70 Years of Age. Cancers, 2020, 12, 712.	1.7	3
11	Specific, reversible G1 arrest by UCN-01 in vivo provides cytostatic protection of normal cells against cytotoxic chemotherapy in breast cancer. British Journal of Cancer, 2020, 122, 812-822.	2.9	11
12	Abstract PD2-05: Differential mechanisms of acquired resistance to abemaciclib versus palbociclib reveal novel therapeutic strategies for CDK4/6 therapy-resistant breast cancers. Cancer Research, 2020, 80, PD2-05-PD2-05.	0.4	4
13	Abstract P6-04-12: STAT3 as a therapeutic target in estrogen receptor positive breast cancer patients refractory to CDK4/6 inhibition. , 2020, , .		0
14	Abstract P3-10-02: Neutrophil elastase as a therapeutic target to inhibit metastasis in breast cancer. , 2020, , .		0
15	Abstract P2-05-04: Low molecular weight cyclin E facilitates replication stress tolerance in breast cancer development. , 2020, , .		1
16	Abstract P4-06-03: Assessment of intratumoral heterogeneity in early stage estrogen receptor (ER) positive breast cancer. , 2020, , .		0
17	Abstract P6-03-09: Role of IL-6 in promoting endocrine therapy and palbociclib resistance estrogen receptor positive breast cancer cells. , 2020, , .		0
18	Toxicity of Radiation Therapy Given Concomitantly with Palbociclib for Metastatic Breast Carcinoma. International Journal of Radiation Oncology Biology Physics, 2019, 105, E60.	0.4	1

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19	Arthur B. Pardee: In Memoriam (1921â€“2019). <i>Cancer Research</i> , 2019, 79, 2089-2090.	0.4	0
20	Combined Inhibition of STAT3 and DNA Repair in Palbociclib-Resistant ER-Positive Breast Cancer. <i>Clinical Cancer Research</i> , 2019, 25, 3996-4013.	3.2	77
21	Cytoplasmic cyclin E independently predicts recurrence in older patients with primary breast cancer.. <i>Journal of Clinical Oncology</i> , 2019, 37, 3128-3128.	0.8	0
22	Abstract 323: Combined inhibition of STAT-3 & DNA repair in palbociclib resistant breast cancer. , 2019, , .		0
23	Enhancer transcription reveals subtype-specific gene expression programs controlling breast cancer pathogenesis. <i>Genome Research</i> , 2018, 28, 159-170.	2.4	137
24	Synthetic Lethality of PARP Inhibitors in Combination with MYC Blockade Is Independent of BRCA Status in Triple-Negative Breast Cancer. <i>Cancer Research</i> , 2018, 78, 742-757.	0.4	98
25	Inhibiting CDK in Cancer Therapy: Current Evidence and Future Directions. <i>Targeted Oncology</i> , 2018, 13, 21-38.	1.7	78
26	Low-Molecular-Weight Cyclin E in Human Cancer: Cellular Consequences and Opportunities for Targeted Therapies. <i>Cancer Research</i> , 2018, 78, 5481-5491.	0.4	39
27	Cyclin E Overexpression Sensitizes Triple-Negative Breast Cancer to Wee1 Kinase Inhibition. <i>Clinical Cancer Research</i> , 2018, 24, 6594-6610.	3.2	70
28	Histone modification profiling in breast cancer cell lines highlights commonalities and differences among subtypes. <i>BMC Genomics</i> , 2018, 19, 150.	1.2	62
29	Strategic development of AZD1775, a Wee1 kinase inhibitor, for cancer therapy. <i>Expert Opinion on Investigational Drugs</i> , 2018, 27, 741-751.	1.9	43
30	Leveraging MYC as a therapeutic treatment option for TNBC. <i>Oncoscience</i> , 2018, 5, 137-139.	0.9	2
31	Abstract 1843: Combination therapy targeting Rb/Wee1 kinase pathways for rhabdomyosarcoma treatment. , 2018, , .		0
32	CDK4/6 Inhibitors Sensitize Rb-positive Sarcoma Cells to Wee1 Kinase Inhibition through Reversible Cell-Cycle Arrest. <i>Molecular Cancer Therapeutics</i> , 2017, 16, 1751-1764.	1.9	39
33	Cytoplasmic Cyclin E Mediates Resistance to Aromatase Inhibitors in Breast Cancer. <i>Clinical Cancer Research</i> , 2017, 23, 7288-7300.	3.2	29
34	CDK4/6 and autophagy inhibitors synergistically induce senescence in Rb positive cytoplasmic cyclin E negative cancers. <i>Nature Communications</i> , 2017, 8, 15916.	5.8	214
35	AXL Inhibition Suppresses the DNA Damage Response and Sensitizes Cells to PARP Inhibition in Multiple Cancers. <i>Molecular Cancer Research</i> , 2017, 15, 45-58.	1.5	73
36	Cytoplasmic Cyclin E Predicts Recurrence in Patients with Breast Cancer. <i>Clinical Cancer Research</i> , 2017, 23, 2991-3002.	3.2	46

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37	Rapid Breast Cancer Disease Progression Following Cyclin Dependent Kinase 4 and 6 Inhibitor Discontinuation. <i>Journal of Cancer</i> , 2017, 8, 2004-2009.	1.2	14
38	Abstract 2060: Characterizing acquired resistance to palbociclib in breast cancer. <i>Cancer Research</i> , 2017, 77, 2060-2060.	0.4	3
39	Cyclin E overexpression as a biomarker for combination treatment strategies in inflammatory breast cancer. <i>Oncotarget</i> , 2017, 8, 14897-14911.	0.8	35
40	Relationships of cyclin E with clinical outcome and biomarkers in older women with early operable primary breast cancer.. <i>Journal of Clinical Oncology</i> , 2017, 35, e12031-e12031.	0.8	1
41	Abstract 2338: CDK4/6 and autophagy inhibitors synergize to induce senescence in cancers with an intact G1/S checkpoint. , 2017, , .		0
42	Abstract A15: Downregulation of c-myc is synthetic lethal with PARP inhibitors in high MYC cancers independent of BRCA status. , 2017, , .		0
43	Targeting the Cell Cycle in Breast Cancer. <i>Breast Diseases</i> , 2016, 27, 256-260.	0.0	0
44	Estrogen receptor alpha is cell cycle-regulated and regulates the cell cycle in a ligand-dependent fashion. <i>Cell Cycle</i> , 2016, 15, 1579-1590.	1.3	31
45	Cyclin E Associates with the Lipogenic Enzyme ATP-Citrate Lyase to Enable Malignant Growth of Breast Cancer Cells. <i>Cancer Research</i> , 2016, 76, 2406-2418.	0.4	64
46	Cytoplasmic Cyclin E and Phospho- Cyclin-Dependent Kinase 2 Are Biomarkers of Aggressive Breast Cancer. <i>American Journal of Pathology</i> , 2016, 186, 1900-1912.	1.9	42
47	PAF-Wnt signaling-induced cell plasticity is required for maintenance of breast cancer cell stemness. <i>Nature Communications</i> , 2016, 7, 10633.	5.8	63
48	Sequential Combination Therapy of CDK Inhibition and Doxorubicin Is Synthetically Lethal in p53-Mutant Triple-Negative Breast Cancer. <i>Molecular Cancer Therapeutics</i> , 2016, 15, 593-607.	1.9	54
49	PKC $\alpha$ promotes ovarian tumor progression through deregulation of cyclin E. <i>Oncogene</i> , 2016, 35, 2428-2440.	2.6	17
50	Abstract PR06: Analysis of enhancer transcription reveals novel gene regulatory networks in breast cancer. , 2016, , .		0
51	Abstract 2989: An Intact G1/S checkpoint determines response to CDK4/6 inhibitor in breast cancer. , 2016, , .		0
52	A phase 1 study with dose expansion of the CDK inhibitor dinaciclib (SCH 727965) in combination with epirubicin in patients with metastatic triple negative breast cancer. <i>Investigational New Drugs</i> , 2015, 33, 890-894.	1.2	58
53	How will we recruit, train, and retain physicians and scientists to conduct translational cancer research?. <i>Cancer</i> , 2015, 121, 806-816.	2.0	13
54	The serine protease inhibitor elafin maintains normal growth control by opposing the mitogenic effects of neutrophil elastase. <i>Oncogene</i> , 2015, 34, 3556-3567.	2.6	29

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55	Abstract 1783: Pharmacological inhibition of CDK4/6 induces G1 arrest, autophagy and senescence in ER+ breast cancer. , 2015, , .		1
56	Abstract P6-03-09: Targeting the c-myc/E2F1 pathway in TNBC promotes a DNA damage dependent synthetic lethality. , 2015, , .		0
57	Abstract P5-05-05: Low molecular weight cyclin E regulates response to aromatase inhibitors in post-menopausal breast cancer patients. , 2015, , .		0
58	Abstract P5-08-02: Inhibition of CDK4/6 induces senescence and autophagy in ER positive breast cancers. , 2015, , .		0
59	Abstract 3772: Cyclin E as a prognostic marker and predictor of response to neoadjuvant chemotherapy and adjuvant hormonal therapy in patients with stage II-III breast cancer. , 2015, , .		0
60	Abstract 3091: Targeting low molecular weight (LMW) cyclin E-Cdk2 pathway for the prevention of breast cancer. , 2015, , .		0
61	Abstract 3579: Identification of biomarkers of AXL-mediated drug resistance in head and neck squamous cell carcinoma. , 2015, , .		0
62	Abstract 1422: Neutrophil Elastase plays a key role in epithelial-mesenchymal transition and metastasis in triple-negative breast cancers. , 2015, , .		0
63	Abstract B27: CDK inhibition impairs homologous recombination and induces PARP inhibitor sensitivity via loss of c-myc expression in TNBC. , 2015, , .		1
64	Indole-3-carbinol and its N-alkoxy derivatives preferentially target ER<math>\alpha</math>-positive breast cancer cells. Cell Cycle, 2014, 13, 2587-2599.	1.3	38
65	Elafin is downregulated during breast and ovarian tumorigenesis but its residual expression predicts recurrence. Breast Cancer Research, 2014, 16, 3417.	2.2	21
66	2-OMe-phosphorodithioate-modified siRNAs show increased loading into the RISC complex and enhanced anti-tumour activity. Nature Communications, 2014, 5, 3459.	5.8	103
67	Exploiting Cell Cycle Pathways in Cancer Therapy: New (and Old) Targets and Potential Strategies. Cancer Drug Discovery and Development, 2014, , 337-372.	0.2	1
68	A phase 1 study of dinaciclib (SCH 727965) in combination with epirubicin patients with metastatic triple-negative breast cancer.. Journal of Clinical Oncology, 2014, 32, 163-163.	0.8	2
69	Abstract 1327: Targeting the RB-pathway in sarcoma: Utility of CDK4/6 inhibitors. , 2014, , .		0
70	Elafin, an inhibitor of elastase, is a prognostic indicator in breast cancer. Breast Cancer Research, 2013, 15, R3.	2.2	40
71	EVI1 splice variants modulate functional responses in ovarian cancer cells. Molecular Oncology, 2013, 7, 647-668.	2.1	38
72	Staurosporine is chemoprotective by inducing G 1 arrest in a Chk1- and pRb-dependent manner. Carcinogenesis, 2013, 34, 2244-2252.	1.3	10

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73	Hbo1 Is a Cyclin E/CDK2 Substrate That Enriches Breast Cancer Stem-like Cells. <i>Cancer Research</i> , 2013, 73, 5556-5568.	0.4	46
74	LMW-E/CDK2 Deregulates Acinar Morphogenesis, Induces Tumorigenesis, and Associates with the Activated b-Raf-ERK1/2-mTOR Pathway in Breast Cancer Patients. <i>PLoS Genetics</i> , 2012, 8, e1002538.	1.5	35
75	Low molecular weight cyclin E is associated with p27-resistant, high-grade, high-stage and invasive bladder cancer. <i>Cell Cycle</i> , 2012, 11, 1468-1476.	1.3	18
76	Targeting low molecular weight cyclin E (LMW-E) in breast cancer. <i>Breast Cancer Research and Treatment</i> , 2012, 132, 575-588.	1.1	35
77	Abstract 2466: Elafin, a serine protease inhibitor, is deregulated during breast cancer progression. , 2012, , .		1
78	Abstract 3283: Conditional Low-molecular-weight (LMW) cyclin E-induced mammary tumorigenesis. , 2012, , .		0
79	Cyclin E amplification/overexpression is a mechanism of trastuzumab resistance in HER2+ breast cancer patients. <i>Breast Diseases</i> , 2011, 22, 266-267.	0.0	0
80	MDA-7 results in downregulation of AKT concomitant with apoptosis and cell cycle arrest in breast cancer cells. <i>Cancer Gene Therapy</i> , 2011, 18, 510-519.	2.2	8
81	Semi-high throughput method of measuring proteasome inhibition in vitro and in cultured cells. <i>Cell Biology and Toxicology</i> , 2011, 27, 123-131.	2.4	4
82	Cdk2 is Required for Breast Cancer Mediated by the Low-Molecular-Weight Isoform of Cyclin E. <i>Cancer Research</i> , 2011, 71, 3377-3386.	0.4	55
83	Breaking the cycle: An insight into the role of ER $\pm$ in eukaryotic cell cycles. <i>Journal of Carcinogenesis</i> , 2011, 10, 25.	2.5	12
84	A novel interaction between HER2/neu and cyclin E in breast cancer. <i>Oncogene</i> , 2010, 29, 3896-3907.	2.6	54
85	Cyclin E Deregulation Impairs Mitotic Progression through Premature Activation of Cdc25C. <i>Cancer Research</i> , 2010, 70, 5085-5095.	0.4	33
86	The Neutrophil Elastase Inhibitor Elafin Triggers Rb-Mediated Growth Arrest and Caspase-Dependent Apoptosis in Breast Cancer. <i>Cancer Research</i> , 2010, 70, 7125-7136.	0.4	26
87	Low Molecular Weight Cyclin E Overexpression Shortens Mitosis, Leading to Chromosome Missegregation and Centrosome Amplification. <i>Cancer Research</i> , 2010, 70, 5074-5084.	0.4	53
88	Low-Molecular-Weight Cyclin E Can Bypass Letrozole-Induced G1 Arrest in Human Breast Cancer Cells and Tumors. <i>Clinical Cancer Research</i> , 2010, 16, 1179-1190.	3.2	47
89	Absence of pRb facilitates E2F1-induced apoptosis in breast cancer cells. <i>Cell Cycle</i> , 2010, 9, 1122-1130.	1.3	19
90	Abstract 3057: The serine protease inhibitor, elafin, exhibits novel tumor suppressor functions in the context of breast cancer. , 2010, , .		0

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91	Altered Subcellular Localization of Tumor-Specific Cyclin E Isoforms Affects Cyclin-Dependent Kinase 2 Complex Formation and Proteasomal Regulation. <i>Cancer Research</i> , 2009, 69, 2817-2825.	0.4	39
92	Low molecular weight cyclin E is specific in breast cancer and is associated with mechanisms of tumor progression. <i>Cell Cycle</i> , 2009, 8, 1062-1068.	1.3	45
93	DEAR1 Is a Dominant Regulator of Acinar Morphogenesis and an Independent Predictor of Local Recurrence-Free Survival in Early-Onset Breast Cancer. <i>PLoS Medicine</i> , 2009, 6, e1000068.	3.9	37
94	Integrative Analysis of Cyclin Protein Levels Identifies Cyclin B1 as a Classifier and Predictor of Outcomes in Breast Cancer. <i>Clinical Cancer Research</i> , 2009, 15, 3654-3662.	3.2	121
95	Cyclin E deregulation is an early event in the development of breast cancer. <i>Breast Cancer Research and Treatment</i> , 2009, 115, 651-659.	1.1	32
96	Post-translational modification and stability of low molecular weight cyclin E. <i>Oncogene</i> , 2009, 28, 3167-3176.	2.6	10
97	Biomarkers in Neoadjuvant Trials. <i>Cancer Treatment and Research</i> , 2009, 147, 1-36.	0.2	0
98	A novel MCM-2 fragment with potential biological function in senescence. <i>Cell Cycle</i> , 2008, 7, 3479-3480.	1.3	0
99	Autophagy: A Novel Mechanism of Synergistic Cytotoxicity between Doxorubicin and Roscovitine in a Sarcoma Model. <i>Cancer Research</i> , 2008, 68, 7966-7974.	0.4	95
100	Synchronization of the cell cycle using Lovastatin. <i>Cell Cycle</i> , 2008, 7, 2434-2440.	1.3	68
101	Cyclin E-associated Kinase Activity Predicts Response to Platinum-Based Chemotherapy. <i>Clinical Cancer Research</i> , 2007, 13, 4800-4806.	3.2	15
102	Overexpression of the Low Molecular Weight Cyclin E in Transgenic Mice Induces Metastatic Mammary Carcinomas through the Disruption of the ARF-p53 Pathway. <i>Cancer Research</i> , 2007, 67, 7212-7222.	0.4	64
103	Differential Regulation of Elafin in Normal and Tumor-Derived Mammary Epithelial Cells Is Mediated by CCAAT/Enhancer Binding Protein $\beta$ . <i>Cancer Research</i> , 2007, 67, 11272-11283.	0.4	19
104	The Double-Stranded RNA-Activated Protein Kinase Mediates Radiation Resistance in Mouse Embryo Fibroblasts through Nuclear Factor $\kappa$ B and Akt Activation. <i>Clinical Cancer Research</i> , 2007, 13, 6032-6039.	3.2	26
105	Cell Cycle Deregulation in Breast Cancer: Insurmountable Chemoresistance or Achilles' Heel?. <i>Advances in Experimental Medicine and Biology</i> , 2007, 608, 52-69.	0.8	5
106	Deregulation of cyclin E meets dysfunction in p53: Closing the escape hatch on breast cancer. <i>Journal of Cellular Physiology</i> , 2006, 209, 686-694.	2.0	23
107	Anti-HER2 Antibody Trastuzumab Inhibits CDK2-Mediated NPAT and Histone H4 Expression via PI3K Pathway. <i>Cell Cycle</i> , 2006, 5, 1654-1661.	1.3	26
108	The Differential Staurosporine-Mediated G1 Arrest in Normal versus Tumor Cells Is Dependent on the Retinoblastoma Protein. <i>Cancer Research</i> , 2006, 66, 9744-9753.	0.4	15

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109	Farnesyl and Geranylgeranyl Transferase Inhibitors Induce G1 Arrest by Targeting the Proteasome. <i>Cancer Research</i> , 2006, 66, 1040-1051.	0.4	46
110	Cyclin E as a prognostic and predictive marker in breast cancer. <i>Seminars in Cancer Biology</i> , 2005, 15, 319-326.	4.3	56
111	Clinical outcome of patients with lymph node-negative breast carcinoma who have sentinel lymph node micrometastases detected by immunohistochemistry. <i>Cancer</i> , 2005, 104, 1779-1780.	2.0	0
112	Atypical PKC $\hat{A}$ contributes to poor prognosis through loss of apical-basal polarity and Cyclin E overexpression in ovarian cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005, 102, 12519-12524.	3.3	231
113	The Tumor-specific Hyperactive Forms of Cyclin E Are Resistant to Inhibition by p21 and p27. <i>Journal of Biological Chemistry</i> , 2005, 280, 15148-15157.	1.6	57
114	The low molecular weight cyclin E isoforms augment angiogenesis and metastasis of human melanoma cells in vivo. <i>Cancer Research</i> , 2005, 65, 692-7.	0.4	44
115	Activation of Cyclin-dependent Kinase 2 by Full Length and Low Molecular Weight Forms of Cyclin E in Breast Cancer Cells. <i>Journal of Biological Chemistry</i> , 2004, 279, 12695-12705.	1.6	31
116	Tumor-Specific Low Molecular Weight Forms of Cyclin E Induce Genomic Instability and Resistance to p21, p27, and Antiestrogens in Breast Cancer. <i>Cancer Research</i> , 2004, 64, 3198-3208.	0.4	134
117	Cyclin E deregulation alters the biologic properties of ovarian cancer cells. <i>Oncogene</i> , 2004, 23, 2648-2657.	2.6	58
118	Role of cell cycle in mediating sensitivity to radiotherapy. <i>International Journal of Radiation Oncology Biology Physics</i> , 2004, 59, 928-942.	0.4	870
119	Low-molecular-weight cyclin E: the missing link between biology and clinical outcome. <i>Breast Cancer Research</i> , 2004, 6, 188-91.	2.2	24
120	Cyclin E is a more powerful predictor of breast cancer outcome than proliferation. <i>Nature Medicine</i> , 2003, 9, 152-152.	15.2	25
121	The Low Molecular Weight Isoforms of Cyclin E Deregulate the Cell Cycle of Mammary Epithelial Cells. <i>Cell Cycle</i> , 2003, 2, 459-464.	1.3	36
122	Cyclin E and Its Low Molecular Weight Forms in Human Cancer and as Targets for Cancer Therapy. <i>Cancer Biology and Therapy</i> , 2003, 2, 37-46.	1.5	63
123	Cyclin E in Breast Cancer. <i>New England Journal of Medicine</i> , 2003, 348, 1063-1064.	13.9	3
124	The low molecular weight (LMW) isoforms of cyclin E deregulate the cell cycle of mammary epithelial cells. <i>Cell Cycle</i> , 2003, 2, 461-6.	1.3	19
125	Cyclin E and its low molecular weight forms in human cancer and as targets for cancer therapy. <i>Cancer Biology and Therapy</i> , 2003, 2, S38-47.	1.5	34
126	Selective protection of normal proliferating cells against the toxic effects of chemotherapeutic agents. <i>Progress in Cell Cycle Research</i> , 2003, 5, 527-32.	0.9	17

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127	Cyclin E and Survival in Patients with Breast Cancer. <i>New England Journal of Medicine</i> , 2002, 347, 1566-1575.	13.9	522
128	Phosphorylation-Dependent Ubiquitination of Cyclin E by the SCFFbw7 Ubiquitin Ligase. <i>Science</i> , 2001, 294, 173-177.	6.0	718
129	Taxol-induced apoptosis depends on MAP kinase pathways (ERK and p38) and is independent of p53. <i>Oncogene</i> , 2001, 20, 147-155.	2.6	332
130	Tumor-Specific Proteolytic Processing of Cyclin E Generates Hyperactive Lower-Molecular-Weight Forms. <i>Molecular and Cellular Biology</i> , 2001, 21, 6254-6269.	1.1	179
131	Activation of the Estrogen-Signaling Pathway by p21WAF1/CIP1 in Estrogen Receptor-Negative Breast Cancer Cells. <i>Journal of the National Cancer Institute</i> , 2000, 92, 1403-1413.	3.0	36
132	Differential mRNA expression of the human DNA methyltransferases (DNMTs) 1, 3a and 3b during the G0/G1 to S phase transition in normal and tumor cells. <i>Nucleic Acids Research</i> , 2000, 28, 2108-2113.	6.5	170
133	Novel splice variants of cyclin E with altered substrate specificity. <i>Nucleic Acids Research</i> , 2000, 28, 101e-101.	6.5	48
134	Expression of an estrogen receptor alpha variant protein in cell lines and tumors. <i>Molecular and Cellular Endocrinology</i> , 2000, 162, 167-180.	1.6	18
135	Lovastatin-mediated G1 arrest is through inhibition of the proteasome, independent of hydroxymethyl glutaryl-CoA reductase. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1999, 96, 7797-7802.	3.3	346
136	UCN-01-mediated G1 arrest in normal but not tumor breast cells is pRb-dependent and p53-independent. <i>Oncogene</i> , 1999, 18, 5691-5702.	2.6	62
137	The Biphasic Induction of p21 and p27 in Breast Cancer Cells by Modulators of cAMP Is Posttranscriptionally Regulated and Independent of the PKA Pathway. <i>Experimental Cell Research</i> , 1999, 252, 211-223.	1.2	19
138	Lovastatin mediated G1 arrest in normal and tumor breast cells is through inhibition of CDK2 activity and redistribution of p21 and p27, independent of p53. <i>Oncogene</i> , 1998, 17, 2393-2402.	2.6	177
139	Molecular Cloning, Characterization, and Regulation of the Human Mitochondrial Serine Hydroxymethyltransferase Gene. <i>Journal of Biological Chemistry</i> , 1997, 272, 1842-1848.	1.6	117
140	The role of cyclin E in cell proliferation, development and cancer. , 1997, 3, 171-191.		107
141	Cyclin E, a redundant cyclin in breast cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1996, 93, 15215-15220.	3.3	114
142	Synchronization of mammalian cells by Lovastatin. <i>Cytotechnology</i> , 1996, 18, 109-114.	0.7	4
143	Cyclin E " a better prognostic marker for breast cancer than cyclin D?. <i>Nature Medicine</i> , 1996, 2, 254-254.	15.2	29
144	Synthesis and Biological Activity of N.omega.-Hemiphthaloyl-.alpha.,.omega.-diaminoalkanoic Acid Analogs of Aminopterin and 3',5-Dichloroaminopterin. <i>Journal of Medicinal Chemistry</i> , 1994, 37, 2167-2174.	2.9	27

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145	The p21 Cdk-interacting protein Cip1 is a potent inhibitor of G1 cyclin-dependent kinases. <i>Cell</i> , 1993, 75, 805-816.	13.5	5,487
146	Redundant cyclin overexpression and gene amplification in breast cancer cells.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1993, 90, 1112-1116.	3.3	481
147	Transcriptional downregulation of gap-junction proteins blocks junctional communication in human mammary tumor cell lines.. <i>Journal of Cell Biology</i> , 1992, 118, 1213-1221.	2.3	226
148	Down-regulation of a member of the S100 gene family in mammary carcinoma cells and reexpression by azadeoxycytidine treatment.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1992, 89, 2504-2508.	3.3	142
149	Modification of cell proliferation with inhibitors. <i>Current Opinion in Cell Biology</i> , 1992, 4, 186-191.	2.6	24
150	Progression through the Cell Cycle: An Overview. <i>The American Review of Respiratory Disease</i> , 1990, 142, S3-S6.	2.9	6
151	An efficient deletion mutant packaging system for defective herpes simplex virus vectors: potential applications to human gene therapy and neuronal physiology.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1990, 87, 8950-8954.	3.3	263
152	[47] Preparation of (6S)-5-formyltetrahydrofolate labeled at high specific activity with 14C and 3H. <i>Methods in Enzymology</i> , 1986, 122, 309-312.	0.4	9
153	Understanding the Biology of Cancer. , 0, , 101-122.		0