

Vincent Cavailles

List of Publications by Year in descending order

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131
papers

6,723
citations

53660

45
h-index

66788

78
g-index

139
all docs

139
docs citations

139
times ranked

7652
citing authors

#	ARTICLE	IF	CITATIONS
1	Cytoplasmic Colocalization of RXR α and PPAR γ as an Independent Negative Prognosticator for Breast Cancer Patients. <i>Cells</i> , 2022, 11, 1244.	1.8	4
2	RIP140 regulates POLK gene expression and the response to alkylating drugs in colon cancer cells. <i>Cancer Drug Resistance (Alhambra, Calif)</i> , 2022, 5, 401-415.	0.9	2
3	RIP140 inhibits glycolysis-dependent proliferation of breast cancer cells by regulating GLUT3 expression through transcriptional crosstalk between hypoxia induced factor and p53. <i>Cellular and Molecular Life Sciences</i> , 2022, 79, 270.	2.4	10
4	The Transcription Coregulator RIP140 Inhibits Cancer Cell Proliferation by Targeting the Pentose Phosphate Pathway. <i>International Journal of Molecular Sciences</i> , 2022, 23, 7419.	1.8	3
5	Metastatic colorectal cancer cells maintain the TGF β 2 program and use TGFBI to fuel angiogenesis. <i>Theranostics</i> , 2021, 11, 1626-1640.	4.6	45
6	Fabrication of 3D printed antimicrobial polycaprolactone scaffolds for tissue engineering applications. <i>Materials Science and Engineering C</i> , 2021, 118, 111525.	3.8	90
7	An R package for generic modular response analysis and its application to estrogen and retinoic acid receptor crosstalk. <i>Scientific Reports</i> , 2021, 11, 7272.	1.6	5
8	Sacrificial mold-assisted 3D printing of stable biocompatible gelatin scaffolds. <i>Bioprinting</i> , 2021, 22, e00140.	2.9	17
9	Cytoplasmic LXR expression is an independent marker of poor prognosis for patients with early stage primary breast cancer. <i>Journal of Cancer Research and Clinical Oncology</i> , 2021, 147, 2535-2544.	1.2	8
10	RIP140 Represses Intestinal Paneth Cell Differentiation and Interplays with SOX9 Signaling in Colorectal Cancer. <i>Cancers</i> , 2021, 13, 3192.	1.7	4
11	Cytoplasmic Localization of RXR α Determines Outcome in Breast Cancer. <i>Cancers</i> , 2021, 13, 3756.	1.7	7
12	A Truncated NRIP1 Mutant Amplifies Microsatellite Instability of Colorectal Cancer by Regulating MSH2/MSH6 Expression, and Is a Prognostic Marker of Stage III Tumors. <i>Cancers</i> , 2021, 13, 4449.	1.7	5
13	Adsorption of proteins on TiO2 particles influences their aggregation and cell penetration. <i>Food Chemistry</i> , 2021, 360, 130003.	4.2	5
14	Development of new biocompatible 3D printed graphene oxide-based scaffolds. <i>Materials Science and Engineering C</i> , 2020, 110, 110595.	3.8	103
15	Long-Term Exposure of Early-Transformed Human Mammary Cells to Low Doses of Benzo[a]pyrene and/or Bisphenol A Enhances Their Cancerous Phenotype via an AhR/GPR30 Interplay. <i>Frontiers in Oncology</i> , 2020, 10, 712.	1.3	13
16	Boron Nitride Based Nanobiocomposites: Design by 3D Printing for Bone Tissue Engineering. <i>ACS Applied Bio Materials</i> , 2020, 3, 1865-1874.	2.3	42
17	Cytoplasmic PPAR γ is a marker of poor prognosis in patients with Cox-1 negative primary breast cancers. <i>Journal of Translational Medicine</i> , 2020, 18, 94.	1.8	19
18	Cytoplasmic and Nuclear Forms of Thyroid Hormone Receptor β 1 Are Inversely Associated with Survival in Primary Breast Cancer. <i>International Journal of Molecular Sciences</i> , 2020, 21, 330.	1.8	13

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19	Prognostic relevance of RIP140 and ER β expression in unifocal versus multifocal breast cancers: a preliminary report. <i>International Journal of Molecular Sciences</i> , 2019, 20, 418.	1.8	8
20	Increased expression of the HDAC9 gene is associated with antiestrogen resistance of breast cancers. <i>Molecular Oncology</i> , 2019, 13, 1534-1547.	2.1	36
21	The Prognostic Impact of the Aryl Hydrocarbon Receptor (AhR) in Primary Breast Cancer Depends on the Lymph Node Status. <i>International Journal of Molecular Sciences</i> , 2019, 20, 1016.	1.8	24
22	Importance of RIP140 and LCoR Sub-Cellular Localization for Their Association With Breast Cancer Aggressiveness and Patient Survival. <i>Translational Oncology</i> , 2018, 11, 1090-1096.	1.7	13
23	Complex regulation of LCoR signaling in breast cancer cells. <i>Oncogene</i> , 2017, 36, 4790-4801.	2.6	27
24	A Dominant Mutation in Nuclear Receptor Interacting Protein 1 Causes Urinary Tract Malformations via Dysregulation of Retinoic Acid Signaling. <i>Journal of the American Society of Nephrology: JASN</i> , 2017, 28, 2364-2376.	3.0	40
25	Identification of a tumor-promoter cholesterol metabolite in human breast cancers acting through the glucocorticoid receptor. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, E9346-E9355.	3.3	96
26	Design of Boron Nitride/Gelatin Electrospun Nanofibers for Bone Tissue Engineering. <i>ACS Applied Materials & Interfaces</i> , 2017, 9, 33695-33706.	4.0	135
27	Investigation of RIP140 and LCoR as independent markers for poor prognosis in cervical cancer. <i>Oncotarget</i> , 2017, 8, 105356-105371.	0.8	10
28	RIP140 and LCoR expression in gastrointestinal cancers. <i>Oncotarget</i> , 2017, 8, 111161-111175.	0.8	7
29	Expression and role of nuclear receptor coregulators in colorectal cancer. <i>World Journal of Gastroenterology</i> , 2017, 23, 4480.	1.4	16
30	Histone deacetylase 9 regulates breast cancer cell proliferation and the response to histone deacetylase inhibitors. <i>Oncotarget</i> , 2016, 7, 19693-19708.	0.8	49
31	Design of graphene oxide/gelatin electrospun nanocomposite fibers for tissue engineering applications. <i>RSC Advances</i> , 2016, 6, 109150-109156.	1.7	26
32	Novel biocompatible electrospun gelatin fiber mats with antibiotic drug delivery properties. <i>Journal of Materials Chemistry B</i> , 2016, 4, 1134-1141.	2.9	49
33	Effect of tamoxifen and fulvestrant long-term treatments on ROS production and (pro/anti)-oxidant enzymes mRNA levels in a MCF-7-derived breast cancer cell line. <i>Breast Cancer</i> , 2016, 23, 692-700.	1.3	6
34	The emerging role of the transcriptional coregulator RIP140 in solid tumors. <i>Biochimica Et Biophysica Acta: Reviews on Cancer</i> , 2015, 1856, 144-150.	3.3	17
35	Expression and role of RIP140/NRIP1 in chronic lymphocytic leukemia. <i>Journal of Hematology and Oncology</i> , 2015, 8, 20.	6.9	17
36	Synergistic activation of human pregnane X receptor by binary cocktails of pharmaceutical and environmental compounds. <i>Nature Communications</i> , 2015, 6, 8089.	5.8	125

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37	Transcriptional Repression of Estrogen Receptor $\hat{\pm}$ Signaling by SENP2 in Breast Cancer Cells. <i>Molecular Endocrinology</i> , 2014, 28, 183-196.	3.7	25
38	Regulation of intestinal homeostasis and tumorigenesis by the transcriptional coregulator RIP140. <i>Molecular and Cellular Oncology</i> , 2014, 1, e960761.	0.3	3
39	Research Resource: STR DNA Profile and Gene Expression Comparisons of Human BG-1 Cells and a BG-1/MCF-7 Clonal Variant. <i>Molecular Endocrinology</i> , 2014, 28, 2072-2081.	3.7	17
40	Structural and Functional Profiling of Environmental Ligands for Estrogen Receptors. <i>Environmental Health Perspectives</i> , 2014, 122, 1306-1313.	2.8	72
41	Affinity purification using recombinant PXR as a tool to characterize environmental ligands. <i>Environmental Toxicology</i> , 2014, 29, 207-215.	2.1	6
42	Selectivity of natural, synthetic and environmental estrogens for zebrafish estrogen receptors. <i>Toxicology and Applied Pharmacology</i> , 2014, 280, 60-69.	1.3	38
43	RIP140 increases APC expression and controls intestinal homeostasis and tumorigenesis. <i>Journal of Clinical Investigation</i> , 2014, 124, 1899-1913.	3.9	45
44	Efficient new constructs against triple negative breast cancer cells: synthesis and preliminary biological study of ferrocifenâ€™SAHA hybrids and related species. <i>Dalton Transactions</i> , 2013, 42, 15489.	1.6	34
45	Negative Regulation of Estrogen Signaling by ER $\hat{2}$ and RIP140 in Ovarian Cancer Cells. <i>Molecular Endocrinology</i> , 2013, 27, 1429-1441.	3.7	38
46	Dialogue between estrogen receptor and E2F signaling pathways: The transcriptional coregulator RIP140 at the crossroads. <i>Advances in Bioscience and Biotechnology (Print)</i> , 2013, 04, 45-54.	0.3	1
47	Structural and mechanistic insights into bisphenols action provide guidelines for risk assessment and discovery of bisphenol A substitutes. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 14930-14935.	3.3	313
48	Occurrence of androgens in sewage treatment plants influents is associated with antagonist activities on other steroid receptors. <i>Water Research</i> , 2012, 46, 1912-1922.	5.3	51
49	Long-term treatment with the pure anti-estrogen fulvestrant durably remodels estrogen signaling in BG-1 ovarian cancer cells. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2012, 132, 176-185.	1.2	8
50	The RIP140 Gene Is a Transcriptional Target of E2F1. <i>PLoS ONE</i> , 2012, 7, e35839.	1.1	26
51	Cognitive impairments in adult mice with constitutive inactivation of <i>RIP140</i> gene expression. <i>Genes, Brain and Behavior</i> , 2012, 11, 69-78.	1.1	36
52	Abstract 1056: Deregulated HDAC9 expression in breast cancer is associated with basal molecular subtype. , 2012, , .		0
53	Abstract 4704: Effects of the HDAC inhibitor S78454/PCI-24781 on ER signalling in ER $\hat{\pm}$ -positive antiestrogen-sensitive and -resistant breast cancer cells. , 2012, , .		0
54	Prognostic Significance of TRIM24/TIF-1 $\hat{\pm}$ Gene Expression in Breast Cancer. <i>American Journal of Pathology</i> , 2011, 178, 1461-1469.	1.9	73

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55	Manipulating Protein Acetylation in Breast Cancer: A Promising Approach in Combination with Hormonal Therapies?. Journal of Biomedicine and Biotechnology, 2011, 2011, 1-15.	3.0	28
56	Biological Analysis of Endocrine-Disrupting Compounds in Tunisian Sewage Treatment Plants. Archives of Environmental Contamination and Toxicology, 2010, 59, 1-12.	2.1	34
57	Estrogenic and AhR activities in dissolved phase and suspended solids from wastewater treatment plants. Science of the Total Environment, 2010, 408, 2608-2615.	3.9	36
58	The steroid receptor RNA activator protein is recruited to promoter regions and acts as a transcriptional repressor. FEBS Letters, 2010, 584, 2218-2224.	1.3	27
59	The Transcriptional Coregulator RIP140 Represses E2F1 Activity and Discriminates Breast Cancer Subtypes. Clinical Cancer Research, 2010, 16, 2959-2970.	3.2	52
60	Estrogen Receptor Interactions and Dynamics Monitored in Live Cells by Fluorescence Cross-Correlation Spectroscopy. Biochemistry, 2010, 49, 772-781.	1.2	36
61	Protein arginine methylation in estrogen signaling and estrogen-related cancers. Trends in Endocrinology and Metabolism, 2010, 21, 181-189.	3.1	41
62	Abstract 4970: Complex regulation of RIP140 gene expression by E2F transcription factors. , 2010, , .		0
63	A new mechanism of SOX9 action to regulate PKC ζ expression in the intestine epithelium. Journal of Cell Science, 2009, 122, 2191-2196.	1.2	19
64	New stably transfected bioluminescent cells expressing FLAG epitope-tagged estrogen receptors to study their chromatin recruitment. BMC Biotechnology, 2009, 9, 77.	1.7	2
65	Regulation of activities of steroid hormone receptors by tibolone and its primary metabolites. Journal of Steroid Biochemistry and Molecular Biology, 2009, 116, 8-14.	1.2	23
66	Profiling of benzophenone derivatives using fish and human estrogen receptor-specific in vitro bioassays. Toxicology and Applied Pharmacology, 2008, 232, 384-395.	1.3	127
67	Specific Activity of Class II Histone Deacetylases in Human Breast Cancer Cells. Molecular Cancer Research, 2008, 6, 1908-1919.	1.5	95
68	Regulation of Hormone Signaling by Nuclear Receptor Interacting Proteins. Advances in Experimental Medicine and Biology, 2008, 617, 121-127.	0.8	3
69	Tamoxifen Resistance and Epigenetic Modifications in Breast Cancer Cell Lines. Current Medicinal Chemistry, 2007, 14, 3035-3043.	1.2	45
70	Differential Regulation of Estrogen Receptor $\hat{\pm}$ Turnover and Transactivation by Mdm2 and Stress-Inducing Agents. Cancer Research, 2007, 67, 5513-5521.	0.4	92
71	Estrogens and antiestrogens activate hPXR. Toxicology Letters, 2007, 170, 19-29.	0.4	88
72	International Union of Pharmacology. LXIV. Estrogen Receptors. Pharmacological Reviews, 2006, 58, 773-781.	7.1	492

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73	Negative regulation of hormone signaling by RIP140. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2006, 102, 51-59.	1.2	46
74	The nuclear receptor transcriptional coregulator RIP140. <i>Nuclear Receptor Signaling</i> , 2006, 4, nrs.04024.	1.0	49
75	ER α and ER β expression and transcriptional activity are differentially regulated by HDAC inhibitors. <i>Oncogene</i> , 2006, 25, 1799-1806.	2.6	66
76	Evaluation of ligand selectivity using reporter cell lines stably expressing estrogen receptor alpha or beta. <i>Biochemical Pharmacology</i> , 2006, 71, 1459-1469.	2.0	171
77	Receptor-Interacting Protein 140 Is a Repressor of the Androgen Receptor Activity. <i>Molecular Endocrinology</i> , 2006, 20, 1506-1518.	3.7	40
78	Transcriptional Regulation of the Human NRIP1/RIP140 Gene by Estrogen Is Modulated by Dioxin Signalling. <i>Molecular Pharmacology</i> , 2006, 69, 1338-1346.	1.0	48
79	Identification of New Human Pregnane X Receptor Ligands among Pesticides Using a Stable Reporter Cell System. <i>Toxicological Sciences</i> , 2006, 91, 501-509.	1.4	162
80	Receptor-Interacting Protein 140 Differentially Regulates Estrogen Receptor-Related Receptor Transactivation Depending on Target Genes. <i>Molecular Endocrinology</i> , 2006, 20, 1035-1047.	3.7	98
81	The nuclear receptor liver receptor homolog-1 is an estrogen receptor target gene. <i>Oncogene</i> , 2005, 24, 8167-8175.	2.6	95
82	SHP Represses Transcriptional Activity via Recruitment of Histone Deacetylases. <i>Biochemistry</i> , 2005, 44, 6312-6320.	1.2	49
83	The Nuclear Receptor Coactivator PGC-1 α Exhibits Modes of Interaction with the Estrogen Receptor Distinct From those of SRC-1. <i>Journal of Molecular Biology</i> , 2005, 347, 921-934.	2.0	43
84	Involvement of HSP1 α protein in irreversible transcriptional inactivation by antiestrogens in breast cancer cells. <i>FEBS Letters</i> , 2005, 579, 4278-4286.	1.3	12
85	Histone deacetylase inhibition and estrogen receptor alpha levels modulate the transcriptional activity of partial antiestrogens. <i>Journal of Molecular Endocrinology</i> , 2004, 32, 583-594.	1.1	40
86	Transcriptional Regulation by the Repressor of Estrogen Receptor Activity via Recruitment of Histone Deacetylases. <i>Journal of Biological Chemistry</i> , 2004, 279, 24834-24843.	1.6	92
87	Multiple domains of the Receptor-Interacting Protein 140 contribute to transcription inhibition. <i>Nucleic Acids Research</i> , 2004, 32, 1957-1966.	6.5	67
88	Dimerization is required for transactivation by estrogen-receptor-related (ERR) orphan receptors: evidence from amphioxus ERR. <i>Journal of Molecular Endocrinology</i> , 2004, 33, 493-509.	1.1	28
89	Histone deacetylase inhibition and estrogen signalling in human breast cancer cells. <i>Biochemical Pharmacology</i> , 2004, 68, 1239-1246.	2.0	56
90	Ligands Differentially Modulate the Protein Interactions of the Human Estrogen Receptors α and β . <i>Journal of Molecular Biology</i> , 2003, 326, 77-92.	2.0	83

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91	Oestrogen receptor alpha increases p21(WAF1/CIP1) gene expression and the antiproliferative activity of histone deacetylase inhibitors in human breast cancer cells. <i>Journal of Endocrinology</i> , 2003, 179, 41-53.	1.2	60
92	Receptor-Interacting Protein 140 Binds c-Jun and Inhibits Estradiol-Induced Activator Protein-1 Activity by Reversing Glucocorticoid Receptor-Interacting Protein 1 Effect. <i>Molecular Endocrinology</i> , 2003, 17, 287-299.	3.7	37
93	Comparative activity of pulsed or continuous estradiol exposure on gene expression and proliferation of normal and tumoral human breast cells. <i>Journal of Molecular Endocrinology</i> , 2002, 28, 165-175.	1.1	23
94	The human estrogen receptor α dimer binds a single SRC-1 coactivator molecule with an affinity dictated by agonist structure ¹¹ Edited by K. Yamamoto. <i>Journal of Molecular Biology</i> , 2001, 306, 433-442.	2.0	70
95	Semiquantitative reverse transcription-polymerase chain reaction to evaluate the expression patterns of genes involved in the oestrogen pathway. <i>Journal of Molecular Endocrinology</i> , 2000, 24, 433-440.	1.1	37
96	Estrogen receptor cofactors expression in breast and endometrial human cancer cells. <i>Molecular and Cellular Endocrinology</i> , 1999, 156, 85-93.	1.6	87
97	Effect of Ligand and DNA Binding on the Interaction between Human Transcription Intermediary Factor 1 α and Estrogen Receptors. <i>Molecular Endocrinology</i> , 1999, 13, 2137-2150.	3.7	14
98	Transcriptional Activities of the Orphan Nuclear Receptor ERR α (Estrogen Receptor-Related Receptor- α). <i>Molecular Endocrinology</i> , 1999, 13, 764-773.	3.7	74
99	Les histone d α ac α tylases : de nouvelles cibles en chimioth α rapie ?. <i>Medecine/Sciences</i> , 1999, 15, 1318.	0.0	0
100	R α glage fin de la transcription par les co-facteurs des r α cepteurs hormonaux nucl α aires.. <i>Medecine/Sciences</i> , 1998, 14, 1127.	0.0	0
101	Differential Interaction of Nuclear Receptors with the Putative Human Transcriptional Coactivator hTIF1. <i>Journal of Biological Chemistry</i> , 1997, 272, 12062-12068.	1.6	91
102	A natural transactivation mutation in the thyroid hormone α receptor: Impaired interaction with putative transcriptional mediators. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1997, 94, 248-253.	3.3	112
103	LxxLL : une signature des coactivateurs de r α cepteurs hormonaux nucl α aires. <i>Medecine/Sciences</i> , 1997, 13, 1212.	0.0	0
104	Biochemical characterization and novel isolation of pure estrogen receptor hormone-binding domain. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 1996, 58, 467-477.	1.2	5
105	RIP-140 Interacts with Multiple Nuclear Receptors by Means of Two Distinct Sites. <i>Molecular and Cellular Biology</i> , 1996, 16, 6029-6036.	1.1	130
106	N-CoR et SMRT sont des cor α presseurs transcriptionnels des r α cepteurs des hormones thyro α idiennes et de l'acide r α tino α que.. <i>Medecine/Sciences</i> , 1996, 12, 234.	0.0	1
107	A la recherche des modulateurs de l'activit α transcriptionnelle des r α cepteurs nucl α aires. <i>Medecine/Sciences</i> , 1996, 12, 229.	0.0	2
108	Transcriptional Activators Differ in Their Responses to Overexpression of TATA-Box-Binding Protein. <i>Molecular and Cellular Biology</i> , 1995, 15, 1554-1563.	1.1	154

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109	Characterization of the proximal estrogen-responsive element of human cathepsin D gene. <i>Molecular Endocrinology</i> , 1994, 8, 693-703.	3.7	60
110	Insensitivity of Cathepsin D Gene to Estradiol in Endometrial Cells Is Determined by the Sequence of Its Estrogen Responsive Element. <i>Biochemical and Biophysical Research Communications</i> , 1994, 203, 711-718.	1.0	9
111	Interaction of proteins with transcriptionally active estrogen receptors.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1994, 91, 10009-10013.	3.3	342
112	Cathepsin D gene is controlled by a mixed promoter, and estrogens stimulate only TATA-dependent transcription in breast cancer cells.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1993, 90, 203-207.	3.3	144
113	Hormonal regulation of cathepsin D following transfection of the estrogen or progesterone receptor into three sex steroid hormone resistant cancer cell lines. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 1991, 40, 231-237.	1.2	30
114	Cathepsin D gene of human MCF7 cells contains estrogen-responsive sequences in its 5' proximal flanking region. <i>Biochemical and Biophysical Research Communications</i> , 1991, 174, 816-824.	1.0	42
115	Estradiol increases the secretion by MCF7 cells of several lysosomal pro-enzymes. <i>Biochemical and Biophysical Research Communications</i> , 1990, 171, 972-978.	1.0	14
116	Estrogen-induced Pro-cathepsin D and its Relationship to Breast Cancer Invasion and Metastasis. , 1990, , 100-110.		0
117	Oestrogen-induced pro-cathepsin D in breast cancer: from biology to clinical applications. <i>Proceedings of the Royal Society of Edinburgh Section B Biological Sciences</i> , 1989, 95, 107-118.	0.2	2
118	Regulation of Cathepsin-D and pS2 Gene Expression by Growth Factors in MCF7 Human Breast Cancer Cells. <i>Molecular Endocrinology</i> , 1989, 3, 552-558.	3.7	165
119	Differential regulation of cathepsin D by sex steroids in mammary cancer and uterine cells. <i>Molecular and Cellular Endocrinology</i> , 1989, 66, 231-238.	1.6	45
120	Overexpression and hormonal regulation of pro-cathepsin D in mammary and endometrial cancer. <i>The Journal of Steroid Biochemistry</i> , 1989, 34, 177-182.	1.3	71
121	Mapping on the calf estrogen receptor of the binding domain for an antibody interfering with receptor activation. <i>The Journal of Steroid Biochemistry</i> , 1989, 32, 769-780.	1.3	2
122	A hormone-regulated pro-cathepsin D secreted by human mammary cancer cells. <i>Biochemical Society Transactions</i> , 1989, 17, 31-33.	1.6	2
123	Estrogen Induced Cathepsin D in Breast Cancer: From Biology to Clinical Applications. , 1989, , 171-186.		2
124	Structure, function, regulation and clinical significance of the 52K pro-cathepsin D secreted by breast cancer cells. <i>Biochimie</i> , 1988, 70, 943-949.	1.3	56
125	Cloning and Sequencing of the 52K Cathepsin D Complementary Deoxyribonucleic Acid of MCF7 Breast Cancer Cells and Mapping on Chromosome 11. <i>Molecular Endocrinology</i> , 1988, 2, 186-192.	3.7	89
126	Estrogens and growth factors induce the mRNA of the 52K-pro-cathepsin-D secreted by breast cancer cells. <i>Nucleic Acids Research</i> , 1988, 16, 1903-1919.	6.5	121

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127	The 52K cathepsin-D of breast cancer: structure, regulation, function and clinical value. <i>Cancer Treatment and Research</i> , 1988, 40, 207-221.	0.2	5
128	An Estrogen Induced Protease in Breast Cancer: From Basic Research to Clinical Applications. , 1988, , 221-232.		0
129	The estrogen-regulated 52K-cathepsin-D in breast cancer: From biology to clinical applications. <i>International Journal of Radiation Applications and Instrumentation Part B, Nuclear Medicine and Biology</i> , 1987, 14, 377-384.	0.3	11
130	Immunohistochemical distribution of the 52-kDa protein in mammary tumors: A marker associated with cell proliferation rather than with hormone responsiveness. <i>The Journal of Steroid Biochemistry</i> , 1987, 27, 439-445.	1.3	38
131	Estrogen-induced lysosomal proteases secreted by breast cancer cells: A role in carcinogenesis?. <i>Journal of Cellular Biochemistry</i> , 1987, 35, 17-29.	1.2	184