

# Pamela M Pollock

## List of Publications by Year in Descending Order

**Source:** <https://exaly.com/author-pdf/1638668/pamela-m-pollock-publications-by-year.pdf>

**Version:** 2024-04-26

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

62  
papers

9,486  
citations

35  
h-index

66  
g-index

66  
ext. papers

11,040  
ext. citations

10.5  
avg, IF

5.13  
L-index

#	Paper	IF	Citations
62	Patient-derived xenograft models capture genomic heterogeneity in endometrial cancer.. <i>Genome Medicine</i> , <b>2022</b> , 14, 3	14.4	2
61	Fibroblast Growth Factor Receptor 2 Isoforms Detected via Novel RNA ISH as Predictive Biomarkers for Progestin Therapy in Atypical Hyperplasia and Low-Grade Endometrial Cancer. <i>Cancers</i> , <b>2021</b> , 13,	6.6	2
60	FGFR2c Mesenchymal Isoform Expression Is Associated with Poor Prognosis and Further Refines Risk Stratification within Endometrial Cancer Molecular Subtypes. <i>Clinical Cancer Research</i> , <b>2020</b> , 26, 4569-4580	12.9	3
59	Anti-CDCP1 immuno-conjugates for detection and inhibition of ovarian cancer. <i>Theranostics</i> , <b>2020</b> , 10, 2095-2114	12.1	10
58	Molecular Classification of the PORTEC-3 Trial for High-Risk Endometrial Cancer: Impact on Prognosis and Benefit From Adjuvant Therapy. <i>Journal of Clinical Oncology</i> , <b>2020</b> , 38, 3388-3397	2.2	116
57	Bcl-2 inhibitors enhance FGFR inhibitor-induced mitochondrial-dependent cell death in FGFR2-mutant endometrial cancer. <i>Molecular Oncology</i> , <b>2019</b> , 13, 738-756	7.9	7
56	Refinement of high-risk endometrial cancer classification using DNA damage response biomarkers: a TransPORTEC initiative. <i>Modern Pathology</i> , <b>2018</b> , 31, 1851-1861	9.8	24
55	FGFR2-activating mutations disrupt cell polarity to potentiate migration and invasion in endometrial cancer cell models. <i>Journal of Cell Science</i> , <b>2018</b> , 131,	5.3	11
54	PI3K Inhibitors Synergize with FGFR Inhibitors to Enhance Antitumor Responses in FGFR2 Endometrial Cancers. <i>Molecular Cancer Therapeutics</i> , <b>2017</b> , 16, 637-648	6.1	26
53	Markers of the p53 pathway further refine molecular profiling in high-risk endometrial cancer: A TransPORTEC initiative. <i>Gynecologic Oncology</i> , <b>2017</b> , 146, 327-333	4.9	16
52	FGFR2 mutations are associated with poor outcomes in endometrioid endometrial cancer: An NRG Oncology/Gynecologic Oncology Group study. <i>Gynecologic Oncology</i> , <b>2017</b> , 145, 366-373	4.9	26
51	Loss of Rearranged L-Myc Fusion (RLF) results in defects in heart development in the mouse. <i>Differentiation</i> , <b>2017</b> , 94, 8-20	3.5	6
50	Immunological profiling of molecularly classified high-risk endometrial cancers identifies -mutant and microsatellite unstable carcinomas as candidates for checkpoint inhibition. <i>Onc Immunology</i> , <b>2017</b> , 6, e1264565	7.2	79
49	hSSB1 phosphorylation is dynamically regulated by DNA-PK and PPP-family protein phosphatases. <i>DNA Repair</i> , <b>2017</b> , 54, 30-39	4.3	9
48	Endometrial cancer cells exhibit high expression of p110 $\beta$ and its selective inhibition induces variable responses on PI3K signaling, cell survival and proliferation. <i>Oncotarget</i> , <b>2017</b> , 8, 3881-3894	3.3	10
47	A Common Variant at the 14q32 Endometrial Cancer Risk Locus Activates AKT1 through YY1 Binding. <i>American Journal of Human Genetics</i> , <b>2016</b> , 98, 1159-1169	11	17
46	The "melanoma-enriched" microRNA miR-4731-5p acts as a tumour suppressor. <i>Oncotarget</i> , <b>2016</b> , 7, 49677-49687	3.3	18

45	The Prognostic and Predictive Value of Melanoma-related MicroRNAs Using Tissue and Serum: A MicroRNA Expression Analysis. <i>EBioMedicine</i> , <b>2015</b> , 2, 671-80	8.8	67
44	Refining prognosis and identifying targetable pathways for high-risk endometrial cancer; a TransPORTEC initiative. <i>Modern Pathology</i> , <b>2015</b> , 28, 836-44	9.8	222
43	Paralog-Specific Kinase Inhibition of FGFR4: Adding to the Arsenal of Anti-FGFR Agents. <i>Cancer Discovery</i> , <b>2015</b> , 5, 355-7	24.4	15
42	Fine-mapping of the HNF1B multicancer locus identifies candidate variants that mediate endometrial cancer risk. <i>Human Molecular Genetics</i> , <b>2015</b> , 24, 1478-92	5.6	46
41	miR-514a regulates the tumour suppressor NF1 and modulates BRAFi sensitivity in melanoma. <i>Oncotarget</i> , <b>2015</b> , 6, 17753-63	3.3	65
40	A phase II trial of brivanib in recurrent or persistent endometrial cancer: an NRG Oncology/Gynecologic Oncology Group Study. <i>Gynecologic Oncology</i> , <b>2014</b> , 135, 38-43	4.9	67
39	The N550K/H mutations in FGFR2 confer differential resistance to PD173074, dovitinib, and ponatinib ATP-competitive inhibitors. <i>Neoplasia</i> , <b>2013</b> , 15, 975-88	6.4	98
38	Integrated genomic characterization of endometrial carcinoma. <i>Nature</i> , <b>2013</b> , 497, 67-73	50.4	2800
37	XIAP downregulation accompanies mebendazole growth inhibition in melanoma xenografts. <i>Anti-Cancer Drugs</i> , <b>2013</b> , 24, 181-8	2.4	36
36	Sensitivity to the MEK inhibitor E6201 in melanoma cells is associated with mutant BRAF and wildtype PTEN status. <i>Molecular Cancer</i> , <b>2012</b> , 11, 75	42.1	30
35	Fibroblast growth factor receptor inhibition synergizes with Paclitaxel and Doxorubicin in endometrial cancer cells. <i>International Journal of Gynecological Cancer</i> , <b>2012</b> , 22, 1517-26	3.5	19
34	Lineage-specific biomarkers predict response to FGFR inhibition. <i>Cancer Discovery</i> , <b>2012</b> , 2, 1081-3	24.4	1
33	FGFR2 point mutations in 466 endometrioid endometrial tumors: relationship with MSI, KRAS, PIK3CA, CTNNB1 mutations and clinicopathological features. <i>PLoS ONE</i> , <b>2012</b> , 7, e30801	3.7	122
32	Targeting mutant fibroblast growth factor receptors in cancer. <i>Trends in Molecular Medicine</i> , <b>2011</b> , 17, 283-92	11.5	96
31	Cellular settings mediating Src Substrate switching between focal adhesion kinase tyrosine 861 and CUB-domain-containing protein 1 (CDCP1) tyrosine 734. <i>Journal of Biological Chemistry</i> , <b>2011</b> , 286, 42303-42315	5.4	159
30	p53 prevents progression of nevi to melanoma predominantly through cell cycle regulation. <i>Pigment Cell and Melanoma Research</i> , <b>2010</b> , 23, 781-94	4.5	50
29	FGFR2 mutations are rare across histologic subtypes of ovarian cancer. <i>Gynecologic Oncology</i> , <b>2010</b> , 117, 125-9	4.9	38
28	Active Notch1 confers a transformed phenotype to primary human melanocytes. <i>Cancer Research</i> , <b>2009</b> , 69, 5312-20	10.1	93

27	Homodimerization controls the fibroblast growth factor 9 subfamily B receptor binding and heparan sulfate-dependent diffusion in the extracellular matrix. <i>Molecular and Cellular Biology</i> , <b>2009</b> , 29, 4663-78	4.8	33
26	Loss-of-function fibroblast growth factor receptor-2 mutations in melanoma. <i>Molecular Cancer Research</i> , <b>2009</b> , 7, 41-54	6.6	100
25	FGFR2 as a molecular target in endometrial cancer. <i>Future Oncology</i> , <b>2009</b> , 5, 27-32	3.6	49
24	A crystallographic snapshot of tyrosine trans-phosphorylation in action. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2008</b> , 105, 19660-5	11.5	49
23	Common variation in the fibroblast growth factor receptor 2 gene is not associated with endometriosis risk. <i>Human Reproduction</i> , <b>2008</b> , 23, 1661-8	5.7	12
22	Inhibition of activated fibroblast growth factor receptor 2 in endometrial cancer cells induces cell death despite PTEN abrogation. <i>Cancer Research</i> , <b>2008</b> , 68, 6902-7	10.1	124
21	Brivanib alaninate, a dual inhibitor of vascular endothelial growth factor receptor and fibroblast growth factor receptor tyrosine kinases, induces growth inhibition in mouse models of human hepatocellular carcinoma. <i>Clinical Cancer Research</i> , <b>2008</b> , 14, 6146-53	12.9	195
20	Frequent activating FGFR2 mutations in endometrial carcinomas parallel germline mutations associated with craniosynostosis and skeletal dysplasia syndromes. <i>Oncogene</i> , <b>2007</b> , 26, 7158-62	9.2	254
19	Osteopontin is a downstream effector of the PI3-kinase pathway in melanomas that is inversely correlated with functional PTEN. <i>Carcinogenesis</i> , <b>2006</b> , 27, 1778-86	4.6	49
18	Proteasome inhibitors trigger NOXA-mediated apoptosis in melanoma and myeloma cells. <i>Cancer Research</i> , <b>2005</b> , 65, 6282-93	10.1	261
17	Microarray expression profiling in melanoma reveals a BRAF mutation signature. <i>Oncogene</i> , <b>2004</b> , 23, 4060-7	9.2	159
16	Deletion mapping suggests that the 1p22 melanoma susceptibility gene is a tumor suppressor localized to a 9-Mb interval. <i>Genes Chromosomes and Cancer</i> , <b>2004</b> , 41, 56-64	5	36
15	p53-independent NOXA induction overcomes apoptotic resistance of malignant melanomas. <i>Molecular Cancer Therapeutics</i> , <b>2004</b> , 3, 895-902	6.1	86
14	High frequency of BRAF mutations in nevi. <i>Nature Genetics</i> , <b>2003</b> , 33, 19-20	36.3	1355
13	Melanoma mouse model implicates metabotropic glutamate signaling in melanocytic neoplasia. <i>Nature Genetics</i> , <b>2003</b> , 34, 108-12	36.3	223
12	A genome-based strategy uncovers frequent BRAF mutations in melanoma. <i>Cancer Cell</i> , <b>2002</b> , 2, 5-7	24.3	128
11	PTEN inactivation is rare in melanoma tumours but occurs frequently in melanoma cell lines. <i>Melanoma Research</i> , <b>2002</b> , 12, 565-75	3.3	55
10	Mutations in exon 3 of the beta-catenin gene are rare in melanoma cell lines. <i>Melanoma Research</i> , <b>2002</b> , 12, 183-6	3.3	31

9	CDKN2A is not the principal target of deletions on the short arm of chromosome 9 in neuroendocrine (Merkel cell) carcinoma of the skin. <i>International Journal of Cancer</i> , <b>2001</b> , 93, 361-7	7.5	10
8	Mutation analysis of the CDKN2A promoter in Australian melanoma families. <i>Genes Chromosomes and Cancer</i> , <b>2001</b> , 32, 89-94	5	17
7	Molecular classification of cutaneous malignant melanoma by gene expression profiling. <i>Nature</i> , <b>2000</b> , 406, 536-40	50.4	1647
6	Haplotype analysis of two recurrent CDKN2A mutations in 10 melanoma families: evidence for common founders and independent mutations. <i>Human Mutation</i> , <b>1998</b> , 11, 424-31	4.7	49
5	Analysis of the CDKN2A, CDKN2B and CDK4 genes in 48 Australian melanoma kindreds. <i>Oncogene</i> , <b>1997</b> , 15, 2999-3005	9.2	74
4	CDKN2A mutation in a non-FAMMM kindred with cancers at multiple sites results in a functionally abnormal protein. <i>International Journal of Cancer</i> , <b>1997</b> , 73, 531-6	7.5	37
3	Compilation of somatic mutations of the CDKN2 gene in human cancers: non-random distribution of base substitutions. <i>Genes Chromosomes and Cancer</i> , <b>1996</b> , 15, 77-88	5	133
2	A homologue of the Drosophila Son of sevenless gene maps to mouse chromosome 17. <i>Genomics</i> , <b>1993</b> , 18, 733-4	4.3	1
1	Genomic analysis of patient-derived xenograft models reveals intra-tumor heterogeneity in endometrial cancer and can predict tumor growth inhibition with talazoparib		2