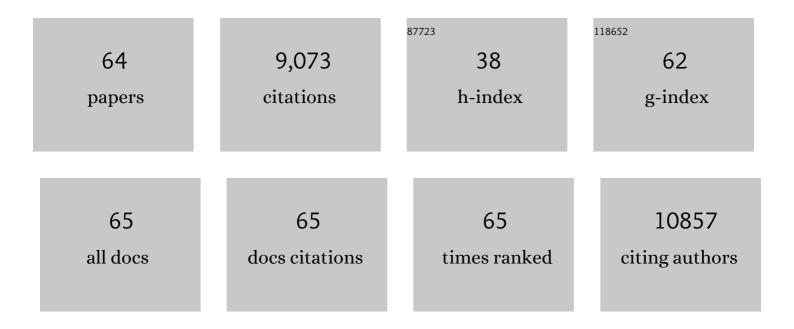
Colin Collins

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Modeling Androgen Deprivation Therapy–Induced Prostate Cancer Dormancy and Its Clinical Implications. Molecular Cancer Research, 2022, 20, 782-793.	1.5	10
2	ZRSR2 overexpression is a frequent and early event in castration-resistant prostate cancer development. Prostate Cancer and Prostatic Diseases, 2021, 24, 775-785.	2.0	0
3	Androgen receptor (AR) antagonism triggers acute succinateâ€mediated adaptive responses to reactivate AR signaling. EMBO Molecular Medicine, 2021, 13, e13427.	3.3	11
4	The long noncoding RNA H19 regulates tumor plasticity in neuroendocrine prostate cancer. Nature Communications, 2021, 12, 7349.	5.8	51
5	Immune-focused multi-omics analysis of prostate cancer: leukocyte Ig-Like receptors are associated with disease progression. Oncolmmunology, 2020, 9, 1851950.	2.1	8
6	RNA Splicing of the BHC80 Gene Contributes to Neuroendocrine Prostate Cancer Progression. European Urology, 2019, 76, 157-166.	0.9	19
7	Widespread and Functional RNA Circularization in Localized Prostate Cancer. Cell, 2019, 176, 831-843.e22.	13.5	317
8	Patient-derived Hormone-naive Prostate Cancer Xenograft Models Reveal Growth Factor Receptor Bound Protein 10 as an Androgen Receptor-repressed Gene Driving the Development of Castration-resistant Prostate Cancer. European Urology, 2018, 73, 949-960.	0.9	19
9	SRRM4 Drives Neuroendocrine Transdifferentiation of Prostate Adenocarcinoma Under Androgen Receptor Pathway Inhibition. European Urology, 2017, 71, 68-78.	0.9	136
10	Quantification of large scale DNA organization for predicting prostate cancer recurrence. Cytometry Part A: the Journal of the International Society for Analytical Cytology, 2017, 91, 1164-1174.	1.1	10
11	miR-100-5p inhibition induces apoptosis in dormant prostate cancer cells and prevents the emergence of castration-resistant prostate cancer. Scientific Reports, 2017, 7, 4079.	1.6	37
12	Therapy-induced developmental reprogramming of prostate cancer cells and acquired therapy resistance. Oncotarget, 2017, 8, 18949-18967.	0.8	47
13	Establishment of a neuroendocrine prostate cancer model driven by the RNA splicing factor SRRM4. Oncotarget, 2017, 8, 66878-66888.	0.8	21
14	The role of epigenetics and long noncoding RNA MIAT in neuroendocrine prostate cancer. Epigenomics, 2016, 8, 721-731.	1.0	94
15	Whole-Exome Sequencing of Metastatic Cancer and Biomarkers of Treatment Response. JAMA Oncology, 2015, 1, 466.	3.4	264
16	The long non-coding RNA PCGEM1 is regulated by androgen receptor activity in vivo. Molecular Cancer, 2015, 14, 46.	7.9	71
17	Expression Profiling of Primary and Metastatic Ovarian Tumors Reveals Differences Indicative of Aggressive Disease. PLoS ONE, 2014, 9, e94476.	1.1	66
18	REST mediates androgen receptor actions on gene repression and predicts early recurrence of prostate cancer. Nucleic Acids Research, 2014, 42, 999-1015.	6.5	125

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19	Identification of Ovarian Cancer Metastatic miRNAs. PLoS ONE, 2013, 8, e58226.	1.1	78
20	nFuse: Discovery of complex genomic rearrangements in cancer using high-throughput sequencing. Genome Research, 2012, 22, 2250-2261.	2.4	67
21	Isolation and genomic analysis of circulating tumor cells from castration resistant metastatic prostate cancer. BMC Cancer, 2012, 12, 78.	1.1	85
22	Detection of recurrent rearrangement breakpoints from copy number data. BMC Bioinformatics, 2011, 12, 114.	1.2	21
23	Comrad: detection of expressed rearrangements by integrated analysis of RNA-Seq and low coverage genome sequence data. Bioinformatics, 2011, 27, 1481-1488.	1.8	39
24	A Group of Genome-Based Biomarkers That Add to a Kattan Nomogram for Predicting Progression in Men with High-Risk Prostate Cancer. Clinical Cancer Research, 2010, 16, 195-202.	3.2	34
25	ZNF217 confers resistance to the pro-apoptotic signals of paclitaxel and aberrant expression of Aurora-A in breast cancer cells. Molecular Cancer, 2010, 9, 291.	7.9	42
26	Evaluation of Paired-End Sequencing Strategies for Detection of Genome Rearrangements in Cancer. PLoS Computational Biology, 2008, 4, e1000051.	1.5	72
27	Evaluation of Whole Genome Amplification Protocols for Array and Oligonucleotide CGH. Diagnostic Molecular Pathology, 2007, 16, 198-206.	2.1	11
28	TMPRSS2-ERG Fusion Prostate Cancer: An Early Molecular Event Associated With Invasion. American Journal of Surgical Pathology, 2007, 31, 882-888.	2.1	394
29	High resolution oligonucleotide CGH using DNA from archived prostate tissue. Prostate, 2007, 67, 1447-1455.	1.2	23
30	Genomic Profiling of Hormone-NaÃ ⁻ ve Lymph Node Metastases in Patients with Prostate Cancer. Neoplasia, 2006, 8, 1083-IN35.	2.3	21
31	Decoding the fine-scale structure of a breast cancer genome and transcriptome. Genome Research, 2006, 16, 394-404.	2.4	49
32	Integrating data on DNA copy number with gene expression levels and drug sensitivities in the NCI-60 cell line panel. Molecular Cancer Therapeutics, 2006, 5, 853-867.	1.9	157
33	High-resolution array comparative genomic hybridization of chromosome arm 8q: Evaluation of genetic progression markers for prostate cancer. Genes Chromosomes and Cancer, 2005, 44, 438-449.	1.5	85
34	Construction and application of a full-coverage, high-resolution, human chromosome 8q genomic microarray for comparative genomic hybridization. Cytometry Part A: the Journal of the International Society for Analytical Cytology, 2005, 63A, 10-19.	1.1	15
35	ZNF217 suppresses cell death associated with chemotherapy and telomere dysfunction. Human Molecular Genetics, 2005, 14, 3219-3225.	1.4	60
36	Overexpression, Amplification, and Androgen Regulation of TPD52 in Prostate Cancer. Cancer Research, 2004, 64, 3814-3822.	0.4	145

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37	Integration of high-resolution array comparative genomic hybridization analysis of chromosome 16q with expression array data refines common regions of loss at 16q23–qter and identifies underlying candidate tumor suppressor genes in prostate cancer. Oncogene, 2004, 23, 3487-3494.	2.6	78
38	Evolution of 8p loss in transformed human prostate epithelial cells. Cancer Genetics and Cytogenetics, 2004, 154, 36-43.	1.0	17
39	Evaluation of genetic patterns in different tumor areas of intermediate-grade prostatic adenocarcinomas by high-resolution genomic array analysis. Genes Chromosomes and Cancer, 2004, 39, 249-256.	1.5	46
40	Computational BAC clone contig assembly for comprehensive genome analysis. Genes Chromosomes and Cancer, 2004, 40, 66-71.	1.5	15
41	Whole genome scanning identifies genotypes associated with recurrence and metastasis in prostate tumors. Human Molecular Genetics, 2004, 13, 1303-1313.	1.4	171
42	Genome Amplification of Chromosome 20 in Breast Cancer. Breast Cancer Research and Treatment, 2003, 78, 337-345.	1.1	100
43	Chromosome aberrations in solid tumors. Nature Genetics, 2003, 34, 369-376.	9.4	702
44	High-Resolution Analysis of Paraffin-Embedded and Formalin-Fixed Prostate Tumors Using Comparative Genomic Hybridization to Genomic Microarrays. American Journal of Pathology, 2003, 162, 763-770.	1.9	76
45	Characterization of the novel amplified in breast cancer-1 (NABC1) gene product. Experimental Cell Research, 2003, 290, 402-413.	1.2	28
46	Reconstructing tumor genome architectures. Bioinformatics, 2003, 19, ii162-ii171.	1.8	58
47	End-sequence profiling: Sequence-based analysis of aberrant genomes. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 7696-7701.	3.3	119
48	Protein elongation factor EEF1A2 is a putative oncogene in ovarian cancer. Nature Genetics, 2002, 31, 301-305.	9.4	220
49	Genome scanning with array CGH delineates regional alterations in mouse islet carcinomas. Nature Genetics, 2001, 29, 459-464.	9.4	268
50	Comprehensive Genome Sequence Analysis of a Breast Cancer Amplicon. Genome Research, 2001, 11, 1034-1042.	2.4	27
51	Quantitative mapping of amplicon structure by array CGH identifies CYP24 as a candidate oncogene. Nature Genetics, 2000, 25, 144-146.	9.4	555
52	Genome changes and gene expression in human solid tumors. Carcinogenesis, 2000, 21, 443-452.	1.3	174
53	PIK3CA is implicated as an oncogene in ovarian cancer. Nature Genetics, 1999, 21, 99-102.	9.4	1,041
54	High resolution analysis of DNA copy number variation using comparative genomic hybridization to microarrays. Nature Genetics, 1998, 20, 207-211.	9.4	2,066

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#	Article	IF	CITATIONS
55	Zf9, a Kruppel-like transcription factor up-regulatedin vivoduring early hepatic fibrosis. Proceedings of the National Academy of Sciences of the United States of America, 1998, 95, 9500-9505.	3.3	234
56	Genome Scanning and Gene Discovery in Breast and Ovarian Cancer. , 1998, , 65-72.		0
57	Genomic Organization, Sequence, and Chromosomal Localization of the Human Helix–Loop–Helix Id1 Gene. Biochemical and Biophysical Research Communications, 1997, 231, 628-634.	1.0	33
58	ESX: a structurally unique Ets overexpressed early during human breast tumorigenesis. Oncogene, 1997, 14, 1617-1622.	2.6	159
59	A physical map of chromosome 20 established using fluorescence in situ hybridization and digital image analysis. Genomics, 1995, 26, 134-137.	1.3	50
60	The human β-subunit of rod photoreceptor cGMP phosphodiesterase: Complete retinal cDNA sequence and evidence for expression in brain. Genomics, 1992, 13, 698-704.	1.3	35
61	Exclusion of DNA changes in the β–subunit of the c–GMP phosphodiesterase gene as the cause for Huntington's disease. Nature Genetics, 1992, 1, 104-108.	9.4	12
62	Delineation of a 50 kilobase DNA segment containing the recombination site in a sporadic case of Huntington's disease. Nature Genetics, 1992, 2, 216-222.	9.4	7
63	Identification of multiple CpG islands and associated conserved sequences in a candidate region for the Huntington disease gene. Genomics, 1991, 11, 1113-1124.	1.3	21
64	Genomic organization and complete sequence of the human gene encoding the β-subunit of the cGMP phosphodiesterase and its localisation to 4p16.3. Nucleic Acids Research, 1991, 19, 6263-6268.	6.5	57