

William B Isaacs

List of Publications by Citations

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

340
papers

31,364
citations

89
h-index

168
g-index

359
ext. papers

35,490
ext. citations

8.6
avg, IF

6.43
L-index

#	Paper	IF	Citations
340	AR-V7 and resistance to enzalutamide and abiraterone in prostate cancer. <i>New England Journal of Medicine</i> , 2014 , 371, 1028-38	59.2	1753
339	Inflammation in prostate carcinogenesis. <i>Nature Reviews Cancer</i> , 2007 , 7, 256-69	31.3	1168
338	Prostate cancer. <i>New England Journal of Medicine</i> , 2003 , 349, 366-81	59.2	883
337	The evolutionary history of lethal metastatic prostate cancer. <i>Nature</i> , 2015 , 520, 353-357	50.4	857
336	Ligand-independent androgen receptor variants derived from splicing of cryptic exons signify hormone-refractory prostate cancer. <i>Cancer Research</i> , 2009 , 69, 16-22	10.1	779
335	Genome-wide association study identifies a second prostate cancer susceptibility variant at 8q24. <i>Nature Genetics</i> , 2007 , 39, 631-7	36.3	739
334	REVEL: An Ensemble Method for Predicting the Pathogenicity of Rare Missense Variants. <i>American Journal of Human Genetics</i> , 2016 , 99, 877-885	11	722
333	Evidence for a prostate cancer susceptibility locus on the X chromosome. <i>Nature Genetics</i> , 1998 , 20, 175-183	36.3	592
332	Frequency of homozygous deletion at p16/CDKN2 in primary human tumours. <i>Nature Genetics</i> , 1995 , 11, 210-2	36.3	554
331	Cumulative association of five genetic variants with prostate cancer. <i>New England Journal of Medicine</i> , 2008 , 358, 910-9	59.2	526
330	Copy number analysis indicates monoclonal origin of lethal metastatic prostate cancer. <i>Nature Medicine</i> , 2009 , 15, 559-65	50.5	513
329	Hereditary prostate cancer: epidemiologic and clinical features. <i>Journal of Urology</i> , 1993 , 150, 797-802	2.5	437
328	Androgen-induced TOP2B-mediated double-strand breaks and prostate cancer gene rearrangements. <i>Nature Genetics</i> , 2010 , 42, 668-75	36.3	436
327	Germline mutations in HOXB13 and prostate-cancer risk. <i>New England Journal of Medicine</i> , 2012 , 366, 141-9	59.2	424
326	Distinct transcriptional programs mediated by the ligand-dependent full-length androgen receptor and its splice variants in castration-resistant prostate cancer. <i>Cancer Research</i> , 2012 , 72, 3457-62	10.1	417
325	Hypermethylation of CpG islands in primary and metastatic human prostate cancer. <i>Cancer Research</i> , 2004 , 64, 1975-86	10.1	416
324	Pathological and molecular aspects of prostate cancer. <i>Lancet, The</i> , 2003 , 361, 955-64	40	361

323	Tracking the clonal origin of lethal prostate cancer. <i>Journal of Clinical Investigation</i> , 2013 , 123, 4918-22	15.9	355
322	Establishment and characterization of seven Dunning rat prostatic cancer cell lines and their use in developing methods for predicting metastatic abilities of prostatic cancers. <i>Prostate</i> , 1986 , 9, 261-81	4.2	350
321	Cyclooxygenases in cancer: progress and perspective. <i>Cancer Letters</i> , 2004 , 215, 1-20	9.9	341
320	Alpha-methylacyl-CoA racemase: a new molecular marker for prostate cancer. <i>Cancer Research</i> , 2002 , 62, 2220-6	10.1	339
319	A meta-analysis of 87,040 individuals identifies 23 new susceptibility loci for prostate cancer. <i>Nature Genetics</i> , 2014 , 46, 1103-9	36.3	331
318	Common sequence variants on 2p15 and Xp11.22 confer susceptibility to prostate cancer. <i>Nature Genetics</i> , 2008 , 40, 281-3	36.3	327
317	Phenotypic analysis of prostate-infiltrating lymphocytes reveals TH17 and Treg skewing. <i>Clinical Cancer Research</i> , 2008 , 14, 3254-61	12.9	314
316	Nuclear MYC protein overexpression is an early alteration in human prostate carcinogenesis. <i>Modern Pathology</i> , 2008 , 21, 1156-67	9.8	301
315	Prostate carcinogenesis and inflammation: emerging insights. <i>Carcinogenesis</i> , 2005 , 26, 1170-81	4.6	295
314	Germline mutations and sequence variants of the macrophage scavenger receptor 1 gene are associated with prostate cancer risk. <i>Nature Genetics</i> , 2002 , 32, 321-5	36.3	283
313	PTEN protein loss by immunostaining: analytic validation and prognostic indicator for a high risk surgical cohort of prostate cancer patients. <i>Clinical Cancer Research</i> , 2011 , 17, 6563-73	12.9	266
312	Mobile DNA in cancer. Extensive transduction of nonrepetitive DNA mediated by L1 retrotransposition in cancer genomes. <i>Science</i> , 2014 , 345, 1251343	33.3	250
311	The landscape of recombination in African Americans. <i>Nature</i> , 2011 , 476, 170-5	50.4	243
310	Two genome-wide association studies of aggressive prostate cancer implicate putative prostate tumor suppressor gene DAB2IP. <i>Journal of the National Cancer Institute</i> , 2007 , 99, 1836-44	9.7	218
309	Rb loss is characteristic of prostatic small cell neuroendocrine carcinoma. <i>Clinical Cancer Research</i> , 2014 , 20, 890-903	12.9	215
308	Human prostate cancer precursors and pathobiology. <i>Urology</i> , 2003 , 62, 55-62	1.6	212
307	DNA hypomethylation arises later in prostate cancer progression than CpG island hypermethylation and contributes to metastatic tumor heterogeneity. <i>Cancer Research</i> , 2008 , 68, 8954-67	10.1	209
306	Identification of a new prostate cancer susceptibility locus on chromosome 8q24. <i>Nature Genetics</i> , 2009 , 41, 1055-7	36.3	201

305	Sequence variants of toll-like receptor 4 are associated with prostate cancer risk: results from the CAncer Prostate in Sweden Study. <i>Cancer Research</i> , 2004 , 64, 2918-22	10.1	199
304	DNA methylation alterations exhibit intraindividual stability and interindividual heterogeneity in prostate cancer metastases. <i>Science Translational Medicine</i> , 2013 , 5, 169ra10	17.5	190
303	A germline DNA polymorphism enhances alternative splicing of the KLF6 tumor suppressor gene and is associated with increased prostate cancer risk. <i>Cancer Research</i> , 2005 , 65, 1213-22	10.1	182
302	Understanding the mechanisms of androgen deprivation resistance in prostate cancer at the molecular level. <i>European Urology</i> , 2015 , 67, 470-9	10.2	181
301	GSTP1 CpG island hypermethylation is responsible for the absence of GSTP1 expression in human prostate cancer cells. <i>American Journal of Pathology</i> , 2001 , 159, 1815-26	5.8	180
300	Human prostate-infiltrating CD8+ T lymphocytes are oligoclonal and PD-1+. <i>Prostate</i> , 2009 , 69, 1694-703	4.2	173
299	Germline Mutations in ATM and BRCA1/2 Distinguish Risk for Lethal and Indolent Prostate Cancer and are Associated with Early Age at Death. <i>European Urology</i> , 2017 , 71, 740-747	10.2	171
298	Genome-wide association study of prostate cancer in men of African ancestry identifies a susceptibility locus at 17q21. <i>Nature Genetics</i> , 2011 , 43, 570-3	36.3	171
297	A snapshot of the expression signature of androgen receptor splicing variants and their distinctive transcriptional activities. <i>Prostate</i> , 2011 , 71, 1656-67	4.2	159
296	Detection and analysis of beta-catenin mutations in prostate cancer. <i>Prostate</i> , 2000 , 45, 323-34	4.2	156
295	Deletional, mutational, and methylation analyses of CDKN2 (p16/MTS1) in primary and metastatic prostate cancer. <i>Genes Chromosomes and Cancer</i> , 1997 , 19, 90-96	5	146
294	In vitro evidence for complex modes of nuclear beta-catenin signaling during prostate growth and tumorigenesis. <i>Oncogene</i> , 2002 , 21, 2679-94	9.2	145
293	Genome-wide association study identifies new prostate cancer susceptibility loci. <i>Human Molecular Genetics</i> , 2011 , 20, 3867-75	5.6	143
292	Pathological and molecular mechanisms of prostate carcinogenesis: implications for diagnosis, detection, prevention, and treatment. <i>Journal of Cellular Biochemistry</i> , 2004 , 91, 459-77	4.7	143
291	Evidence for two independent prostate cancer risk-associated loci in the HNF1B gene at 17q12. <i>Nature Genetics</i> , 2008 , 40, 1153-5	36.3	139
290	Sequence variants in Toll-like receptor gene cluster (TLR6-TLR1-TLR10) and prostate cancer risk. <i>Journal of the National Cancer Institute</i> , 2005 , 97, 525-32	9.7	139
289	Global patterns of prostate cancer incidence, aggressiveness, and mortality in men of african descent. <i>Prostate Cancer</i> , 2013 , 2013, 560857	1.9	136
288	CYP3A4-V and prostate cancer in African Americans: causal or confounding association because of population stratification?. <i>Human Genetics</i> , 2002 , 110, 553-60	6.3	136

287	HOXB13 is a susceptibility gene for prostate cancer: results from the International Consortium for Prostate Cancer Genetics (ICPCG). <i>Human Genetics</i> , 2013 , 132, 5-14	6.3	134
286	Ligand-dependent inhibition of beta-catenin/TCF signaling by androgen receptor. <i>Oncogene</i> , 2002 , 21, 8453-69	9.2	133
285	A molecular analysis of prokaryotic and viral DNA sequences in prostate tissue from patients with prostate cancer indicates the presence of multiple and diverse microorganisms. <i>Prostate</i> , 2008 , 68, 306-20	4.2	131
284	DNA copy number alterations in prostate cancers: a combined analysis of published CGH studies. <i>Prostate</i> , 2007 , 67, 692-700	4.2	129
283	A combined genomewide linkage scan of 1,233 families for prostate cancer-susceptibility genes conducted by the international consortium for prostate cancer genetics. <i>American Journal of Human Genetics</i> , 2005 , 77, 219-29	11	129
282	Peroxisomal branched chain fatty acid beta-oxidation pathway is upregulated in prostate cancer. <i>Prostate</i> , 2005 , 63, 316-23	4.2	129
281	Linkage and association studies of prostate cancer susceptibility: evidence for linkage at 8p22-23. <i>American Journal of Human Genetics</i> , 2001 , 69, 341-50	11	127
280	Association between two unlinked loci at 8q24 and prostate cancer risk among European Americans. <i>Journal of the National Cancer Institute</i> , 2007 , 99, 1525-33	9.7	120
279	Effects of RNase L mutations associated with prostate cancer on apoptosis induced by 2R5R oligoadenylylates. <i>Cancer Research</i> , 2003 , 63, 6795-801	10.1	118
278	Macrophage inhibitory cytokine 1: a new prognostic marker in prostate cancer. <i>Clinical Cancer Research</i> , 2009 , 15, 6658-64	12.9	117
277	Carbohydrate restriction, prostate cancer growth, and the insulin-like growth factor axis. <i>Prostate</i> , 2008 , 68, 11-9	4.2	116
276	Sequencing of prostate cancers identifies new cancer genes, routes of progression and drug targets. <i>Nature Genetics</i> , 2018 , 50, 682-692	36.3	112
275	A novel role of myosin VI in human prostate cancer. <i>American Journal of Pathology</i> , 2006 , 169, 1843-54	5.8	112
274	Gene expression signature of benign prostatic hyperplasia revealed by cDNA microarray analysis. <i>Prostate</i> , 2002 , 51, 189-200	4.2	110
273	Allelic loss of the retinoblastoma gene in primary human prostatic adenocarcinomas. <i>Prostate</i> , 1995 , 26, 35-9	4.2	110
272	A meta-analysis of genome-wide association studies to identify prostate cancer susceptibility loci associated with aggressive and non-aggressive disease. <i>Human Molecular Genetics</i> , 2013 , 22, 408-15	5.6	109
271	Association of IL10 and other immune response- and obesity-related genes with prostate cancer in CLUE II. <i>Prostate</i> , 2009 , 69, 874-85	4.2	108
270	Structure and methylation-associated silencing of a gene within a homozygously deleted region of human chromosome band 8p22. <i>Genomics</i> , 1996 , 35, 55-65	4.3	108

269	Germline DNA-repair Gene Mutations and Outcomes in Men with Metastatic Castration-resistant Prostate Cancer Receiving First-line Abiraterone and Enzalutamide. <i>European Urology</i> , 2018 , 74, 218-225 ^{10.2}	107
268	Acute inflammatory proteins constitute the organic matrix of prostatic corpora amylacea and calculi in men with prostate cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009 , 106, 3443-8	11.5 107
267	Human polymorphisms at long non-coding RNAs (lncRNAs) and association with prostate cancer risk. <i>Carcinogenesis</i> , 2011 , 32, 1655-9	4.6 107
266	Role of Genetic Testing for Inherited Prostate Cancer Risk: Philadelphia Prostate Cancer Consensus Conference 2017. <i>Journal of Clinical Oncology</i> , 2018 , 36, 414-424	2.2 107
265	Associations between hOGG1 sequence variants and prostate cancer susceptibility. <i>Cancer Research</i> , 2002 , 62, 2253-7	10.1 105
264	Loss of PTEN is associated with aggressive behavior in ERG-positive prostate cancer. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2013 , 22, 2333-44	4 104
263	Explaining racial differences in prostate cancer in the United States: sociology or biology?. <i>Prostate</i> , 2005 , 62, 243-52	4.2 102
262	Physical mapping of chromosome 8p22 markers and their homozygous deletion in a metastatic prostate cancer. <i>Genomics</i> , 1996 , 35, 46-54	4.3 101
261	Sequence variants at 22q13 are associated with prostate cancer risk. <i>Cancer Research</i> , 2009 , 69, 10-5	10.1 100
260	H6D polymorphism in macrophage-inhibitory cytokine-1 gene associated with prostate cancer. <i>Journal of the National Cancer Institute</i> , 2004 , 96, 1248-54	9.7 99
259	Characterizing genetic risk at known prostate cancer susceptibility loci in African Americans. <i>PLoS Genetics</i> , 2011 , 7, e1001387	6 98
258	Decreased gene expression of steroid 5 alpha-reductase 2 in human prostate cancer: implications for finasteride therapy of prostate carcinoma. <i>Prostate</i> , 2003 , 57, 134-9	4.2 98
257	Polygenic risk score improves prostate cancer risk prediction: results from the Stockholm-1 cohort study. <i>European Urology</i> , 2011 , 60, 21-8	10.2 97
256	Alpha-methylacyl-CoA racemase: a variably sensitive immunohistochemical marker for the diagnosis of small prostate cancer foci on needle biopsy. <i>American Journal of Surgical Pathology</i> , 2003 , 27, 1128-33	6.7 97
255	Alpha-methylacyl-CoA racemase as an androgen-independent growth modifier in prostate cancer. <i>Cancer Research</i> , 2003 , 63, 7365-76	10.1 93
254	Fine mapping association study and functional analysis implicate a SNP in MSMB at 10q11 as a causal variant for prostate cancer risk. <i>Human Molecular Genetics</i> , 2009 , 18, 1368-75	5.6 91
253	Inherited genetic variant predisposes to aggressive but not indolent prostate cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010 , 107, 2136-40	11.5 90
252	Evaluation of linkage and association of HPC2/ELAC2 in patients with familial or sporadic prostate cancer. <i>American Journal of Human Genetics</i> , 2001 , 68, 901-11	11 87

251	The role of genetic markers in the management of prostate cancer. <i>European Urology</i> , 2012 , 62, 577-87	10.2	86
250	Individual and cumulative effect of prostate cancer risk-associated variants on clinicopathologic variables in 5,895 prostate cancer patients. <i>Prostate</i> , 2009 , 69, 1195-205	4.2	86
249	Homozygous deletions and recurrent amplifications implicate new genes involved in prostate cancer. <i>Neoplasia</i> , 2008 , 10, 897-907	6.4	86
248	COX-2 gene promoter haplotypes and prostate cancer risk. <i>Carcinogenesis</i> , 2004 , 25, 961-6	4.6	85
247	Modulation of CXCL14 (BRAK) expression in prostate cancer. <i>Prostate</i> , 2005 , 64, 67-74	4.2	85
246	Common sequence variants of the macrophage scavenger receptor 1 gene are associated with prostate cancer risk. <i>American Journal of Human Genetics</i> , 2003 , 72, 208-12	11	81
245	Implementation of Germline Testing for Prostate Cancer: Philadelphia Prostate Cancer Consensus Conference 2019. <i>Journal of Clinical Oncology</i> , 2020 , 38, 2798-2811	2.2	80
244	Validation of genome-wide prostate cancer associations in men of African descent. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2011 , 20, 23-32	4	79
243	Linkage of prostate cancer susceptibility loci to chromosome 1. <i>Human Genetics</i> , 2001 , 108, 335-45	6.3	79
242	MSH2 Loss in Primary Prostate Cancer. <i>Clinical Cancer Research</i> , 2017 , 23, 6863-6874	12.9	78
241	Focus on prostate cancer. <i>Cancer Cell</i> , 2002 , 2, 113-6	24.3	74
240	Genome-wide scan for prostate cancer susceptibility genes in the Johns Hopkins hereditary prostate cancer families. <i>Prostate</i> , 2003 , 57, 320-5	4.2	74
239	Potential impact of adding genetic markers to clinical parameters in predicting prostate biopsy outcomes in men following an initial negative biopsy: findings from the REDUCE trial. <i>European Urology</i> , 2012 , 62, 953-61	10.2	73
238	Assembly of inflammation-related genes for pathway-focused genetic analysis. <i>PLoS ONE</i> , 2007 , 2, e10357	5.7	73
237	Trefoil factor 3 overexpression in prostatic carcinoma: prognostic importance using tissue microarrays. <i>Prostate</i> , 2004 , 61, 215-27	4.2	73
236	Relation between aberrant alpha-catenin expression and loss of E-cadherin function in prostate cancer. <i>International Journal of Cancer</i> , 1997 , 74, 374-7	7.5	71
235	Germline Mutations in ATM and BRCA1/2 Are Associated with Grade Reclassification in Men on Active Surveillance for Prostate Cancer. <i>European Urology</i> , 2019 , 75, 743-749	10.2	71
234	A polymorphism in the CDKN1B gene is associated with increased risk of hereditary prostate cancer. <i>Cancer Research</i> , 2004 , 64, 1997-9	10.1	70

233	In Swedish families with hereditary prostate cancer, linkage to the HPC1 locus on chromosome 1q24-25 is restricted to families with early-onset prostate cancer. <i>American Journal of Human Genetics</i> , 1999 , 65, 134-40	11	70
232	Molecular and cellular changes associated with the acquisition of metastatic ability by prostatic cancer cells. <i>Prostate</i> , 1994 , 25, 249-65	4.2	70
231	DIAPH3 governs the cellular transition to the amoeboid tumour phenotype. <i>EMBO Molecular Medicine</i> , 2012 , 4, 743-60	12	69
230	Estimation of absolute risk for prostate cancer using genetic markers and family history. <i>Prostate</i> , 2009 , 69, 1565-72	4.2	69
229	Intraductal/ductal histology and lymphovascular invasion are associated with germline DNA-repair gene mutations in prostate cancer. <i>Prostate</i> , 2018 , 78, 401-407	4.2	68
228	Genetic markers associated with early cancer-specific mortality following prostatectomy. <i>Cancer</i> , 2013 , 119, 2405-12	6.4	68
227	XMRV: a new virus in prostate cancer?. <i>Cancer Research</i> , 2010 , 70, 10028-33	10.1	68
226	Association of a germ-line copy number variation at 2p24.3 and risk for aggressive prostate cancer. <i>Cancer Research</i> , 2009 , 69, 2176-9	10.1	68
225	Acne and risk of prostate cancer. <i>International Journal of Cancer</i> , 2007 , 121, 2688-92	7.5	66
224	Phenotypic characterization of telomerase-immortalized primary non-malignant and malignant tumor-derived human prostate epithelial cell lines. <i>Experimental Cell Research</i> , 2006 , 312, 831-43	4.2	66
223	Combined genome-wide scan for prostate cancer susceptibility genes. <i>Journal of the National Cancer Institute</i> , 2004 , 96, 1240-7	9.7	65
222	Evaluation of serum and seminal plasma markers in the diagnosis of canine prostatic disorders. <i>Journal of Veterinary Internal Medicine</i> , 1995 , 9, 149-53	3.1	65
221	Polymorphic GGC repeats in the androgen receptor gene are associated with hereditary and sporadic prostate cancer risk. <i>Human Genetics</i> , 2002 , 110, 122-9	6.3	64
220	Frequent loss of chromosome arms 8p and 13q in collecting duct carcinoma (CDC) of the kidney. <i>Genes Chromosomes and Cancer</i> , 1995 , 12, 76-80	5	64
219	The effects of basic fibroblast growth factor and suramin on cell motility and growth of rat prostate cancer cells. <i>Journal of Urology</i> , 1991 , 145, 199-202	2.5	63
218	Trans-ancestry genome-wide association meta-analysis of prostate cancer identifies new susceptibility loci and informs genetic risk prediction. <i>Nature Genetics</i> , 2021 , 53, 65-75	36.3	62
217	GOLPH2 and MYO6: putative prostate cancer markers localized to the Golgi apparatus. <i>Prostate</i> , 2008 , 68, 1387-95	4.2	61
216	Genome-wide association study identifies a new locus JMJD1C at 10q21 that may influence serum androgen levels in men. <i>Human Molecular Genetics</i> , 2012 , 21, 5222-8	5.6	60

215	Systematic replication study of reported genetic associations in prostate cancer: Strong support for genetic variation in the androgen pathway. <i>Prostate</i> , 2006 , 66, 1729-43	4.2	60
214	Titin, a huge, elastic sarcomeric protein with a probable role in morphogenesis. <i>BioEssays</i> , 1991 , 13, 157-61	4.1	60
213	BIOLOGICAL AGGRESSIVENESS OF HEREDITARY PROSTATE CANCER: LONG-TERM EVALUATION FOLLOWING RADICAL PROSTATECTOMY. <i>Journal of Urology</i> , 1998 , 160, 660-663	2.5	59
212	Genetic variants in the LEPR, CRY1, RNASEL, IL4, and ARVCF genes are prognostic markers of prostate cancer-specific mortality. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2011 , 20, 1928-36	4	58
211	Comprehensive assessment of DNA copy number alterations in human prostate cancers using Affymetrix 100K SNP mapping array. <i>Genes Chromosomes and Cancer</i> , 2006 , 45, 1018-32	5	58
210	Genomic organization of the human KAI1 metastasis-suppressor gene. <i>Genomics</i> , 1997 , 41, 25-32	4.3	57
209	Deletion of a small consensus region at 6q15, including the MAP3K7 gene, is significantly associated with high-grade prostate cancers. <i>Clinical Cancer Research</i> , 2007 , 13, 5028-33	12.9	57
208	Molecular advances in prostate cancer. <i>Current Opinion in Oncology</i> , 1997 , 9, 101-7	4.2	56
207	Identification of aryl hydrocarbon receptor as a putative Wnt/beta-catenin pathway target gene in prostate cancer cells. <i>Cancer Research</i> , 2004 , 64, 2523-33	10.1	56
206	Monocyte chemotactic protein-1 (MCP-1/CCL2) is associated with prostatic growth dysregulation and benign prostatic hyperplasia. <i>Prostate</i> , 2010 , 70, 473-81	4.2	55
205	Prostate cancer risk-associated variants reported from genome-wide association studies: meta-analysis and their contribution to genetic Variation. <i>Prostate</i> , 2010 , 70, 1729-38	4.2	55
204	Association of prostate cancer risk variants with clinicopathologic characteristics of the disease. <i>Clinical Cancer Research</i> , 2008 , 14, 5819-24	12.9	55
203	Stronger association between obesity and biochemical progression after radical prostatectomy among men treated in the last 10 years. <i>Clinical Cancer Research</i> , 2005 , 11, 2883-8	12.9	54
202	Prostate cancer risk associated loci in African Americans. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2009 , 18, 2145-9	4	53
201	Joint effect of HSD3B1 and HSD3B2 genes is associated with hereditary and sporadic prostate cancer susceptibility. <i>Cancer Research</i> , 2002 , 62, 1784-9	10.1	52
200	Generalizability of established prostate cancer risk variants in men of African ancestry. <i>International Journal of Cancer</i> , 2015 , 136, 1210-7	7.5	51
199	Sexually transmitted infections and prostatic inflammation/cell damage as measured by serum prostate specific antigen concentration. <i>Journal of Urology</i> , 2006 , 175, 1937-42	2.5	51
198	Identification of miR-30b-3p and miR-30d-5p as direct regulators of androgen receptor signaling in prostate cancer by complementary functional microRNA library screening. <i>Oncotarget</i> , 2016 , 7, 72593-72607	2.3	51

197	Increased gene copy number of ERG on chromosome 21 but not TMPRSS2-ERG fusion predicts outcome in prostatic adenocarcinomas. <i>Modern Pathology</i> , 2011 , 24, 1511-20	9.8	50
196	VITAMIN D RECEPTOR POLYMORPHISMS AND LETHAL PROSTATE CANCER. <i>Journal of Urology</i> , 1998 , 160, 1405-1409	2.5	49
195	Polymorphisms in the CYP1A1 gene are associated with prostate cancer risk. <i>International Journal of Cancer</i> , 2003 , 106, 375-8	7.5	49
194	LOSS OF HETEROZYGOSITY AT 12P12 13 IN PRIMARY AND METASTATIC PROSTATE ADENOCARCINOMA. <i>Journal of Urology</i> , 2000 , 164, 192-196	2.5	49
193	A novel prostate cancer susceptibility locus at 19q13. <i>Cancer Research</i> , 2009 , 69, 2720-3	10.1	48
192	Association of 17 prostate cancer susceptibility loci with prostate cancer risk in Chinese men. <i>Prostate</i> , 2010 , 70, 425-32	4.2	47
191	Genome-wide scan of 29,141 African Americans finds no evidence of directional selection since admixture. <i>American Journal of Human Genetics</i> , 2014 , 95, 437-44	11	46
190	Endoglin (CD105) as a urinary and serum marker of prostate cancer. <i>International Journal of Cancer</i> , 2009 , 124, 664-9	7.5	46
189	Evidence for a prostate cancer linkage to chromosome 20 in 159 hereditary prostate cancer families. <i>Human Genetics</i> , 2001 , 108, 430-5	6.3	46
188	Leveraging population admixture to characterize the heritability of complex traits. <i>Nature Genetics</i> , 2014 , 46, 1356-62	36.3	45
187	The G84E mutation of HOXB13 is associated with increased risk for prostate cancer: results from the REDUCE trial. <i>Carcinogenesis</i> , 2013 , 34, 1260-4	4.6	45
186	Cytokine profiling of prostatic fluid from cancerous prostate glands identifies cytokines associated with extent of tumor and inflammation. <i>Prostate</i> , 2008 , 68, 872-82	4.2	45
185	Interleukin-2 transfected prostate cancer cells generate a local antitumor effect in vivo. <i>Prostate</i> , 1994 , 24, 244-51	4.2	45
184	Evaluation of PPP2R2A as a prostate cancer susceptibility gene: a comprehensive germline and somatic study. <i>Cancer Genetics</i> , 2011 , 204, 375-81	2.3	44
183	Cumulative effect of five genetic variants on prostate cancer risk in multiple study populations. <i>Prostate</i> , 2008 , 68, 1257-62	4.2	44
182	Cyclin D1 Loss Distinguishes Prostatic Small-Cell Carcinoma from Most Prostatic Adenocarcinomas. <i>Clinical Cancer Research</i> , 2015 , 21, 5619-29	12.9	43
181	Design, synthesis, and in vitro testing of alpha-methylacyl-CoA racemase inhibitors. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 2700-7	8.3	43
180	A comprehensive association study for genes in inflammation pathway provides support for their roles in prostate cancer risk in the CAPS study. <i>Prostate</i> , 2006 , 66, 1556-64	4.2	43

179	Genetic variability in inflammation pathways and prostate cancer risk. <i>Urologic Oncology: Seminars and Original Investigations</i> , 2007 , 25, 250-9	2.8	42
178	Genome-wide screen for prostate cancer susceptibility genes in men with clinically significant disease. <i>Prostate</i> , 2005 , 64, 356-61	4.2	42
177	Linkage and association of CYP17 gene in hereditary and sporadic prostate cancer. <i>International Journal of Cancer</i> , 2001 , 95, 354-9	7.5	42
176	No evidence for a role of BRCA1 or BRCA2 mutations in Ashkenazi Jewish families with hereditary prostate cancer. <i>Prostate</i> , 1999 , 39, 280-4	4.2	42
175	Identification of new differentially methylated genes that have potential functional consequences in prostate cancer. <i>PLoS ONE</i> , 2012 , 7, e48455	3.7	42
174	Adding genetic risk score to family history identifies twice as many high-risk men for prostate cancer: Results from the prostate cancer prevention trial. <i>Prostate</i> , 2016 , 76, 1120-9	4.2	42
173	The HOXB13 G84E Mutation Is Associated with an Increased Risk for Prostate Cancer and Other Malignancies. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2015 , 24, 1366-72	4	41
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