William B Isaacs

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Association between pathogenic germline mutations in BRCA2 and ATM and tumor-infiltrating lymphocytes in primary prostate cancer. Cancer Immunology, Immunotherapy, 2022, 71, 943-951.	2.0	9
2	Association of germline rare pathogenic mutations in guidelineâ€recommended genes with prostate cancer progression: A metaâ€analysis. Prostate, 2022, 82, 107-119.	1.2	4
3	The HOXB13 variant X285K is associated with clinical significance and early age at diagnosis in African American prostate cancer patients. British Journal of Cancer, 2022, 126, 791-796.	2.9	13
4	Health inequity drives disease biology to create disparities in prostate cancer outcomes. Journal of Clinical Investigation, 2022, 132, .	3.9	17
5	KLK3 germline mutation I179T complements DNA repair genes for predicting prostate cancer progression. Prostate Cancer and Prostatic Diseases, 2022, , .	2.0	3
6	Germline <i>BRCA2</i> , <i>ATM</i> and <i>CHEK2</i> alterations shape somatic mutation landscapes in prostate cancer Journal of Clinical Oncology, 2022, 40, 148-148.	0.8	0
7	The role of genetic testing in prostate cancer screening, diagnosis, and treatment. Current Opinion in Oncology, 2022, Publish Ahead of Print, .	1.1	0
8	Inherited risk assessment and its clinical utility for predicting prostate cancer from diagnostic prostate biopsies. Prostate Cancer and Prostatic Diseases, 2022, 25, 422-430.	2.0	12
9	Identifying Phased Mutations and Complex Rearrangements in Human Prostate Cancer Cell Lines through Linked-Read Whole-Genome Sequencing. Molecular Cancer Research, 2022, 20, 1013-1020.	1.5	3
10	Two-stage Study of Familial Prostate Cancer by Whole-exome Sequencing and Custom Capture Identifies 10 Novel Genes Associated with the Risk of Prostate Cancer. European Urology, 2021, 79, 353-361.	0.9	28
11	Performance of Three Inherited Risk Measures for Predicting Prostate Cancer Incidence and Mortality: A Population-based Prospective Analysis. European Urology, 2021, 79, 419-426.	0.9	36
12	A novel method for detection of exfoliated prostate cancer cells in urine by RNA in situ hybridization. Prostate Cancer and Prostatic Diseases, 2021, 24, 220-232.	2.0	3
13	Trans-ancestry genome-wide association meta-analysis of prostate cancer identifies new susceptibility loci and informs genetic risk prediction. Nature Genetics, 2021, 53, 65-75.	9.4	264
14	Homologous recombination deficiency (HRD) score in germline BRCA2- versus ATM-altered prostate cancer. Modern Pathology, 2021, 34, 1185-1193.	2.9	61
15	Resistance to androgen receptor signaling inhibition does not necessitate development of neuroendocrine prostate cancer. JCI Insight, 2021, 6, .	2.3	22
16	Association of prostate cancer polygenic risk score with number and laterality of tumor cores in active surveillance patients. Prostate, 2021, 81, 703-709.	1.2	11
17	The somatic mutation landscape of germline <i>CHEK2-</i> altered prostate cancer Journal of Clinical Oncology, 2021, 39, 5084-5084.	0.8	2
18	Specific Detection of Prostate Cancer Cells in Urine by RNA In Situ Hybridization. Journal of Urology, 2021. 206. 37-43.	0.2	4

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19	Observed evidence for guidelineâ€recommended genes in predicting prostate cancer risk from a large populationâ€based cohort. Prostate, 2021, 81, 1002-1008.	1.2	10
20	Genetic Susceptibility for Low Testosterone in Men and Its Implications in Biology and Screening: Data from the UK Biobank. European Urology Open Science, 2021, 29, 36-46.	0.2	4
21	Prostate Cancer Predisposition. Urologic Clinics of North America, 2021, 48, 283-296.	0.8	12
22	Rare Germline Variants in ATM Predispose to Prostate Cancer: A PRACTICAL Consortium Study. European Urology Oncology, 2021, 4, 570-579.	2.6	38
23	Combined Longitudinal Clinical and Autopsy Phenomic Assessment in Lethal Metastatic Prostate Cancer: Recommendations for Advancing Precision Medicine. European Urology Open Science, 2021, 30, 47-62.	0.2	2
24	Incorporation of Polygenic Risk Score into Guidelines for Inherited Risk Assessment for Prostate Cancer. European Urology, 2021, 80, 139-141.	0.9	4
25	Inherited risk assessment of prostate cancer: it takes three to do it right. Prostate Cancer and Prostatic Diseases, 2020, 23, 59-61.	2.0	8
26	Germline <i>BLM</i> mutations and metastatic prostate cancer. Prostate, 2020, 80, 235-237.	1.2	15
27	Role of androgen receptor splice variant-7 (AR-V7) in prostate cancer resistance to 2nd-generation androgen receptor signaling inhibitors. Oncogene, 2020, 39, 6935-6949.	2.6	60
28	Genomic and Clinicopathologic Characterization of <i>ATM</i> -deficient Prostate Cancer. Clinical Cancer Research, 2020, 26, 4869-4881.	3.2	18
29	Validation of a prostate cancer polygenic risk score. Prostate, 2020, 80, 1314-1321.	1.2	23
30	Feasibility and performance of a novel probe panel to detect somatic DNA copy number alterations in clinical specimens for predicting prostate cancer progression. Prostate, 2020, 80, 1253-1262.	1.2	4
31	Germline HOXB13 G84E mutation carriers and risk to twenty common types of cancer: results from the UK Biobank. British Journal of Cancer, 2020, 123, 1356-1359.	2.9	11
32	A Germline Variant at 8q24 Contributes to Familial Clustering of Prostate Cancer in Men of African Ancestry. European Urology, 2020, 78, 316-320.	0.9	32
33	Implementation of Germline Testing for Prostate Cancer: Philadelphia Prostate Cancer Consensus Conference 2019. Journal of Clinical Oncology, 2020, 38, 2798-2811.	0.8	170
34	Use of Aspirin and Statins in Relation to Inflammation in Benign Prostate Tissue in the Placebo Arm of the Prostate Cancer Prevention Trial. Cancer Prevention Research, 2020, 13, 853-862.	0.7	8
35	Rare Germline Pathogenic Mutations of DNA Repair Genes Are Most Strongly Associated with Grade Group 5 Prostate Cancer. European Urology Oncology, 2020, 3, 224-230.	2.6	41
36	Distinct Genomic Alterations in Prostate Tumors Derived from African American Men. Molecular Cancer Research, 2020, 18, 1815-1824.	1.5	14

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37	<i>Trichomonas vaginalis</i> infection and prostateâ€specific antigen concentration: Insights into prostate involvement and prostate disease risk. Prostate, 2019, 79, 1622-1628.	1.2	11
38	Lactoferrin CpG Island Hypermethylation and Decoupling of mRNA and Protein Expression in the Early Stages of Prostate Carcinogenesis. American Journal of Pathology, 2019, 189, 2311-2322.	1.9	13
39	Concept and benchmarks for assessing narrowâ€sense validity of genetic risk score values. Prostate, 2019, 79, 1099-1105.	1.2	18
40	Current progress and questions in germline genetics of prostate cancer. Asian Journal of Urology, 2019, 6, 3-9.	0.5	11
41	Molecular Characterization and Clinical Outcomes of Primary Gleason Pattern 5 Prostate Cancer After Radical Prostatectomy. JCO Precision Oncology, 2019, 3, 1-13.	1.5	12
42	Single-Nucleotide Polymorphism–Based Genetic Risk Score and Patient Age at Prostate Cancer Diagnosis. JAMA Network Open, 2019, 2, e1918145.	2.8	20
43	Mannose Receptor–positive Macrophage Infiltration Correlates with Prostate Cancer Onset and Metastatic Castration-resistant Disease. European Urology Oncology, 2019, 2, 429-436.	2.6	46
44	HOXB13 interaction with MEIS1 modifies proliferation and gene expression in prostate cancer. Prostate, 2019, 79, 414-424.	1.2	39
45	Germline Mutations in ATM and BRCA1/2 Are Associated with Grade Reclassification in Men on Active Surveillance for Prostate Cancer. European Urology, 2019, 75, 743-749.	0.9	138
46	A systematic comparison of exercise training protocols on animal models of cardiovascular capacity. Life Sciences, 2019, 217, 128-140.	2.0	32
47	ATM loss in primary prostate cancer: Analysis of >1000 cases using a validated clinical-grade immunohistochemistry (IHC) assay Journal of Clinical Oncology, 2019, 37, 5069-5069.	0.8	3
48	Updated insights into genetic contribution to prostate cancer predisposition: focus on HOXB13. Canadian Journal of Urology, 2019, 26, 12-13.	0.0	5
49	A comprehensive evaluation of <i>CHEK2</i> germline mutations in men with prostate cancer. Prostate, 2018, 78, 607-615.	1.2	57
50	Sequencing of prostate cancers identifies new cancer genes, routes of progression and drug targets. Nature Genetics, 2018, 50, 682-692.	9.4	182
51	Germline DNA-repair Gene Mutations and Outcomes in Men with Metastatic Castration-resistant Prostate Cancer Receiving First-line Abiraterone and Enzalutamide. European Urology, 2018, 74, 218-225.	0.9	140
52	Intraductal/ductal histology and lymphovascular invasion are associated with germline DNAâ€repair gene mutations in prostate cancer. Prostate, 2018, 78, 401-407.	1.2	105
53	Genetic factors influencing prostate cancer risk in Norwegian men. Prostate, 2018, 78, 186-192.	1.2	11
54	Role of Genetic Testing for Inherited Prostate Cancer Risk: Philadelphia Prostate Cancer Consensus Conference 2017. Journal of Clinical Oncology, 2018, 36, 414-424.	0.8	155

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55	Differences in inherited risk among relatives of hereditary prostate cancer patients using genetic risk score. Prostate, 2018, 78, 1063-1068.	1.2	1
56	Germline mutations in <i>PPFIBP2</i> are associated with lethal prostate cancer. Prostate, 2018, 78, 1222-1228.	1.2	12
57	Constitutively active androgen receptor splice variants AR-V3, AR-V7 and AR-V9 are co-expressed in castration-resistant prostate cancer metastases. British Journal of Cancer, 2018, 119, 347-356.	2.9	63
58	Germline mutations in <scp>DNA</scp> repair genes are associated with bladder cancer risk and unfavourable prognosis. BJU International, 2018, 122, 808-813.	1.3	15
59	Sustained influence of infections on prostateâ€specific antigen concentration: An analysis of changes over 10 years of followâ€up. Prostate, 2018, 78, 1024-1034.	1.2	4
60	Effect of germline DNA repair gene mutations on outcomes in men with metastatic castration-resistant prostate cancer receiving first-line abiraterone and enzalutamide Journal of Clinical Oncology, 2018, 36, 221-221.	0.8	0
61	Donald S Coffey, a man who meant so much to so many. American Journal of Clinical and Experimental Urology, 2018, 6, 41-42.	0.4	Ο
62	gsSKAT: Rapid gene set analysis and multiple testing correction for rareâ€variant association studies using weighted linear kernels. Genetic Epidemiology, 2017, 41, 297-308.	0.6	9
63	Analytic, Preanalytic, and Clinical Validation of p53 IHC for Detection of <i>TP53</i> Missense Mutation in Prostate Cancer. Clinical Cancer Research, 2017, 23, 4693-4703.	3.2	62
64	Association between variants in genes involved in the immune response and prostate cancer risk in men randomized to the finasteride arm in the Prostate Cancer Prevention Trial. Prostate, 2017, 77, 908-919.	1.2	21
65	Germline Mutations in ATM and BRCA1/2 Distinguish Risk for Lethal and Indolent Prostate Cancer and are Associated with Early Age at Death. European Urology, 2017, 71, 740-747.	0.9	256
66	Insight into infectionâ€mediated prostate damage: Contrasting patterns of Câ€reactive protein and prostateâ€specific antigen levels during infection. Prostate, 2017, 77, 1325-1334.	1.2	8
67	MSH2 Loss in Primary Prostate Cancer. Clinical Cancer Research, 2017, 23, 6863-6874.	3.2	122
68	A genetic variant near <i>GATA3</i> implicated in inherited susceptibility and etiology of benign prostatic hyperplasia (BPH) and lower urinary tract symptoms (LUTS). Prostate, 2017, 77, 1213-1220.	1.2	19
69	What Do Myeloma, Breast Cancer, and Prostate Cancer Have in Common?. European Urology, 2017, 71, 166-167.	0.9	1
70	The expression of AURKA is androgen regulated in castration-resistant prostate cancer. Scientific Reports, 2017, 7, 17978.	1.6	38
71	Appraising the relevance of DNA copy number loss and gain in prostate cancer using whole genome DNA sequence data. PLoS Genetics, 2017, 13, e1007001.	1.5	34
72	Somatic molecular subtyping of prostate tumors from <i>HOXB13</i> G84E carriers. Oncotarget, 2017, 8, 22772-22782.	0.8	9

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73	Post hoc Analysis for Detecting Individual Rare Variant Risk Associations Using Probit Regression Bayesian Variable Selection Methods in Caseâ€Control Sequencing Studies. Genetic Epidemiology, 2016, 40, 461-469.	0.6	5
74	Adding genetic risk score to family history identifies twice as many high-risk men for prostate cancer: Results from the prostate cancer prevention trial. Prostate, 2016, 76, 1120-1129.	1.2	60
75	Genetic variants in cell cycle control pathway confer susceptibility to aggressive prostate carcinoma. Prostate, 2016, 76, 479-490.	1.2	12
76	Key genes involved in the immune response are generally not associated with intraprostatic inflammation in men without a prostate cancer diagnosis: Results from the prostate cancer prevention trial. Prostate, 2016, 76, 565-574.	1.2	5
77	Infectious mononucleosis, other infections and prostate-specific antigen concentration as a marker of prostate involvement during infection. International Journal of Cancer, 2016, 138, 2221-2230.	2.3	11
78	Peripheral Zone Inflammation Is Not Strongly Associated With Lower Urinary Tract Symptom Incidence and Progression in the Placebo Arm of the Prostate Cancer Prevention Trial. Prostate, 2016, 76, 1399-1408.	1.2	6
79	Integrated clinical, whole-genome, and transcriptome analysis of multisampled lethal metastatic prostate cancer. Journal of Physical Education and Sports Management, 2016, 2, a000752.	0.5	24
80	Inflammation, Microbiota, and Prostate Cancer. European Urology Focus, 2016, 2, 374-382.	1.6	40
81	REVEL: An Ensemble Method for Predicting the Pathogenicity of Rare Missense Variants. American Journal of Human Genetics, 2016, 99, 877-885.	2.6	1,555
82	Rare Variation in <i>TET2</i> Is Associated with Clinically Relevant Prostate Carcinoma in African Americans. Cancer Epidemiology Biomarkers and Prevention, 2016, 25, 1456-1463.	1.1	22
83	DNA-Repair Gene Mutations in Metastatic Prostate Cancer. New England Journal of Medicine, 2016, 375, 1802-1805.	13.9	26
84	Screening for familial and hereditary prostate cancer. International Journal of Cancer, 2016, 138, 2579-2591.	2.3	49
85	Cenome-wide association of familial prostate cancer cases identifies evidence for a rare segregating haplotype at 8q24.21. Human Genetics, 2016, 135, 923-938.	1.8	37
86	Germline Variants in Asporin Vary by Race, Modulate the Tumor Microenvironment, and Are Differentially Associated with Metastatic Prostate Cancer. Clinical Cancer Research, 2016, 22, 448-458.	3.2	29
87	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. Oncotarget, 2016, 7, 72593-72607.	0.8	71
88	Variation in genes involved in the immune response and prostate cancer risk in the placebo arm of the Prostate Cancer Prevention Trial. Prostate, 2015, 75, 1403-1418.	1.2	25
89	Polymorphisms Influencing Prostate-Specific Antigen Concentration May Bias Genome-Wide Association Studies on Prostate Cancer. Cancer Epidemiology Biomarkers and Prevention, 2015, 24, 88-93.	1.1	4
90	Generalizability of established prostate cancer risk variants in men of <scp>A</scp> frican ancestry. International Journal of Cancer, 2015, 136, 1210-1217.	2.3	62

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91	Associations of prostate cancer risk variants with disease aggressiveness: results of the NCI-SPORE Genetics Working Group analysis of 18,343 cases. Human Genetics, 2015, 134, 439-450.	1.8	45
92	The <i>HOXB13</i> G84E Mutation Is Associated with an Increased Risk for Prostate Cancer and Other Malignancies. Cancer Epidemiology Biomarkers and Prevention, 2015, 24, 1366-1372.	1.1	47
93	Integration of multiethnic fine-mapping and genomic annotation to prioritize candidate functional SNPs at prostate cancer susceptibility regions. Human Molecular Genetics, 2015, 24, 5603-5618.	1.4	50
94	The evolutionary history of lethal metastatic prostate cancer. Nature, 2015, 520, 353-357.	13.7	1,185
95	Cyclin D1 Loss Distinguishes Prostatic Small-Cell Carcinoma from Most Prostatic Adenocarcinomas. Clinical Cancer Research, 2015, 21, 5619-5629.	3.2	56
96	Large-scale association analysis in Asians identifies new susceptibility loci for prostate cancer. Nature Communications, 2015, 6, 8469.	5.8	51
97	Do Environmental Factors Modify the Genetic Risk of Prostate Cancer?. Cancer Epidemiology Biomarkers and Prevention, 2015, 24, 213-220.	1.1	12
98	Understanding the Mechanisms of Androgen Deprivation Resistance in Prostate Cancer at the Molecular Level. European Urology, 2015, 67, 470-479.	0.9	225
99	AR splice variant 7 (AR-V7) and response to taxanes in men with metastatic castration-resistant prostate cancer (mCRPC) Journal of Clinical Oncology, 2015, 33, 138-138.	0.8	14
100	Genome-Wide Association Scan for Variants Associated with Early-Onset Prostate Cancer. PLoS ONE, 2014, 9, e93436.	1.1	25
101	A Peripheral Circulating TH1 Cytokine Profile Is Inversely Associated with Prostate Cancer Risk in CLUE II. Cancer Epidemiology Biomarkers and Prevention, 2014, 23, 2561-2567.	1.1	18
102	Association analysis of 9,560 prostate cancer cases from the International Consortium of Prostate Cancer Genetics confirms the role of reported prostate cancer associated SNPs for familial disease. Human Genetics, 2014, 133, 347-356.	1.8	24
103	Prevalence of the <i><scp>HOXB13</scp></i> â€ <scp>C84E</scp> prostate cancer risk allele in men treated with radical prostatectomy. BJU International, 2014, 113, 830-835.	1.3	21
104	Leveraging population admixture to characterize the heritability of complex traits. Nature Genetics, 2014, 46, 1356-1362.	9.4	69
105	Rb Loss Is Characteristic of Prostatic Small Cell Neuroendocrine Carcinoma. Clinical Cancer Research, 2014, 20, 890-903.	3.2	275
106	Telomere length as a risk factor for hereditary prostate cancer. Prostate, 2014, 74, 359-364.	1.2	27
107	Extensive transduction of nonrepetitive DNA mediated by L1 retrotransposition in cancer genomes. Science, 2014, 345, 1251343.	6.0	348
108	A meta-analysis of 87,040 individuals identifies 23 new susceptibility loci for prostate cancer. Nature Genetics, 2014, 46, 1103-1109.	9.4	408

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109	AR-V7 and Resistance to Enzalutamide and Abiraterone in Prostate Cancer. New England Journal of Medicine, 2014, 371, 1028-1038.	13.9	2,233
110	Genome-wide Scan of 29,141 African Americans Finds No Evidence of Directional Selection since Admixture. American Journal of Human Genetics, 2014, 95, 437-444.	2.6	69
111	Identification of a novel germline <i>SPOP</i> mutation in a family with hereditary prostate cancer. Prostate, 2014, 74, 983-990.	1.2	18
112	Androgen receptor splice variant, AR-V7, and resistance to enzalutamide and abiraterone in men with metastatic castration-resistant prostate cancer (mCRPC) Journal of Clinical Oncology, 2014, 32, 5001-5001.	0.8	20
113	Association of the <i>HOXB13 G84E</i> mutation with increased risk for prostate cancer and other malignancies Journal of Clinical Oncology, 2014, 32, 1558-1558.	0.8	0
114	Genome-wide Association Study Identifies Loci at ATF7IP and KLK2 Associated with Percentage of Circulating Free PSA. Neoplasia, 2013, 15, 95-IN30.	2.3	11
115	Loss of PTEN Is Associated with Aggressive Behavior in ERG-Positive Prostate Cancer. Cancer Epidemiology Biomarkers and Prevention, 2013, 22, 2333-2344.	1.1	121
116	DNA Methylation Alterations Exhibit Intraindividual Stability and Interindividual Heterogeneity in Prostate Cancer Metastases. Science Translational Medicine, 2013, 5, 169ra10.	5.8	231
117	HOXB13 is a susceptibility gene for prostate cancer: results from the International Consortium for Prostate Cancer Genetics (ICPCG). Human Genetics, 2013, 132, 5-14.	1.8	166
118	Genome-wide Association Study Identifies Genetic Determinants of Urine PCA3 Levels in Men. Neoplasia, 2013, 15, 448-IN26.	2.3	7
119	Genetic markers associated with early cancerâ€specific mortality following prostatectomy. Cancer, 2013, 119, 2405-2412.	2.0	81
120	Nucleotide resolution analysis of <i><scp>TMPRSS2</scp></i> and <i><scp>ERG</scp></i> rearrangements in prostate cancer. Journal of Pathology, 2013, 230, 174-183.	2.1	41
121	A meta-analysis of genome-wide association studies to identify prostate cancer susceptibility loci associated with aggressive and non-aggressive disease. Human Molecular Genetics, 2013, 22, 408-415.	1.4	118
122	The G84E mutation of HOXB13 is associated with increased risk for prostate cancer: results from the REDUCE trial. Carcinogenesis, 2013, 34, 1260-1264.	1.3	50
123	Global Patterns of Prostate Cancer Incidence, Aggressiveness, and Mortality in Men of African Descent. Prostate Cancer, 2013, 2013, 1-12.	0.4	180
124	A Genome-Wide Assessment of Variability in Human Serum Metabolism. Human Mutation, 2013, 34, 515-524.	1.1	42
125	Tracking the clonal origin of lethal prostate cancer. Journal of Clinical Investigation, 2013, 123, 4918-4922.	3.9	440
126	Infections and inflammation in prostate cancer. American Journal of Clinical and Experimental Urology, 2013, 1, 3-11.	0.4	42

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127	Variation in <i>IL10</i> and Other Genes Involved in the Immune Response and in Oxidation and Prostate Cancer Recurrence. Cancer Epidemiology Biomarkers and Prevention, 2012, 21, 1774-1782.	1.1	49
128	A genome-wide search for loci interacting with known prostate cancer risk-associated genetic variants. Carcinogenesis, 2012, 33, 598-603.	1.3	38
129	Genome-wide association study identifies a new locus JMJD1C at 10q21 that may influence serum androgen levels in men. Human Molecular Genetics, 2012, 21, 5222-5228.	1.4	79
130	The Role of Genetic Markers in the Management of Prostate Cancer. European Urology, 2012, 62, 577-587.	0.9	92
131	Distinct Transcriptional Programs Mediated by the Ligand-Dependent Full-Length Androgen Receptor and Its Splice Variants in Castration-Resistant Prostate Cancer. Cancer Research, 2012, 72, 3457-3462.	0.4	518
132	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. European Urology, 2012, 62, 953-961.	0.9	85
133	Identification of a novel NBN truncating mutation in a family with hereditary prostate cancer. Familial Cancer, 2012, 11, 595-600.	0.9	15
134	Germline Mutations in <i>HOXB13</i> and Prostate-Cancer Risk. New England Journal of Medicine, 2012, 366, 141-149.	13.9	566
135	Association of prostate cancer risk with snps in regions containing androgen receptor binding sites captured by ChIPâ€Onâ€chip analyses. Prostate, 2012, 72, 376-385.	1.2	15
136	Chromosomes 4 and 8 implicated in a genome wide SNP linkage scan of 762 prostate cancer families collected by the ICPCG. Prostate, 2012, 72, 410-426.	1.2	14
137	DIAPH3 governs the cellular transition to the amoeboid tumour phenotype. EMBO Molecular Medicine, 2012, 4, 743-760.	3.3	92
138	Genome-wide two-locus epistasis scans in prostate cancer using two European populations. Human Genetics, 2012, 131, 1225-1234.	1.8	17
139	Validation of prostate cancer risk-related loci identified from genome-wide association studies using family-based association analysis: evidence from the International Consortium for Prostate Cancer Genetics (ICPCG). Human Genetics, 2012, 131, 1095-1103.	1.8	21
140	Inherited susceptibility for aggressive prostate cancer. Asian Journal of Andrology, 2012, 14, 415-418.	0.8	6
141	Identification of New Differentially Methylated Genes That Have Potential Functional Consequences in Prostate Cancer. PLoS ONE, 2012, 7, e48455.	1.1	65
142	Evaluation of PPP2R2A as a prostate cancer susceptibility gene: a comprehensive germline and somatic study. Cancer Genetics, 2011, 204, 375-381.	0.2	51
143	GENETIC BASIS FOR PROSTATE CANCER. , 2011, , 39-52.		0
144	Genome-wide association study of prostate cancer in men of African ancestry identifies a susceptibility locus at 17q21. Nature Genetics, 2011, 43, 570-573.	9.4	198

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145	Polygenic Risk Score Improves Prostate Cancer Risk Prediction: Results from the Stockholm-1 Cohort Study. European Urology, 2011, 60, 21-28.	0.9	117
146	Inherited genetic markers discovered to date are able to identify a significant number of men at considerably elevated risk for prostate cancer. Prostate, 2011, 71, 421-430.	1.2	38
147	Functional annotation of risk loci identified through genomeâ€wide association studies for prostate cancer. Prostate, 2011, 71, 955-963.	1.2	25
148	A snapshot of the expression signature of androgen receptor splicing variants and their distinctive transcriptional activities. Prostate, 2011, 71, 1656-1667.	1.2	177
149	Immunomodulatory ILâ€18 binding protein is produced by prostate cancer cells and its levels in urine and serum correlate with tumor status. International Journal of Cancer, 2011, 129, 424-432.	2.3	42
150	Genome-wide copy-number variation analysis identifies common genetic variants at 20p13 associated with aggressiveness of prostate cancer. Carcinogenesis, 2011, 32, 1057-1062.	1.3	33
151	Human polymorphisms at long non-coding RNAs (IncRNAs) and association with prostate cancer risk. Carcinogenesis, 2011, 32, 1655-1659.	1.3	132
152	High-Throughput Screen Identifies Novel Inhibitors of Cancer Biomarker α-Methylacyl Coenzyme A Racemase (AMACR/P504S). Molecular Cancer Therapeutics, 2011, 10, 825-838.	1.9	46
153	Genome-wide association study identifies new prostate cancer susceptibility loci. Human Molecular Genetics, 2011, 20, 3867-3875.	1.4	160
154	Fine mapping of a region of chromosome 11q13 reveals multiple independent loci associated with risk of prostate cancer. Human Molecular Genetics, 2011, 20, 2869-2878.	1.4	43
155	Prostate Cancer Predisposition Loci and Risk of Metastatic Disease and Prostate Cancer Recurrence. Clinical Cancer Research, 2011, 17, 1075-1081.	3.2	44
156	Large-scale fine mapping of the HNF1B locus and prostate cancer risk. Human Molecular Genetics, 2011, 20, 3322-3329.	1.4	28
157	The landscape of recombination in African Americans. Nature, 2011, 476, 170-175.	13.7	319
158	Increased gene copy number of ERG on chromosome 21 but not TMPRSS2–ERG fusion predicts outcome in prostatic adenocarcinomas. Modern Pathology, 2011, 24, 1511-1520.	2.9	57
159	A Genome-Wide Survey over the ChIP-On-Chip Identified Androgen Receptor-Binding Genomic Regions Identifies a Novel Prostate Cancer Susceptibility Locus at 12q13.13. Cancer Epidemiology Biomarkers and Prevention, 2011, 20, 2396-2403.	1.1	10
160	Validation of Genome-Wide Prostate Cancer Associations in Men of African Descent. Cancer Epidemiology Biomarkers and Prevention, 2011, 20, 23-32.	1.1	88
161	Germ-line sequence variants of PTEN do not have an important role in hereditary and non-hereditary prostate cancer susceptibility. Journal of Human Genetics, 2011, 56, 496-502.	1.1	10
162	PTEN Protein Loss by Immunostaining: Analytic Validation and Prognostic Indicator for a High Risk Surgical Cohort of Prostate Cancer Patients. Clinical Cancer Research, 2011, 17, 6563-6573.	3.2	309

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163	Genetic Variants in the <i>LEPR</i> , <i>CRY1</i> , <i>RNASEL</i> , <i>IL4</i> , and <i>ARVCF</i> Genes Are Prognostic Markers of Prostate Cancer-Specific Mortality. Cancer Epidemiology Biomarkers and Prevention, 2011, 20, 1928-1936.	1.1	68
164	Characterizing Genetic Risk at Known Prostate Cancer Susceptibility Loci in African Americans. PLoS Genetics, 2011, 7, e1001387.	1.5	117
165	A snapshot of the expression signature of androgen receptor splicing variants and their distinctive transcriptional activities. , 2011, 71, 1656.		1
166	Association of 17 prostate cancer susceptibility loci with prostate cancer risk in Chinese men. Prostate, 2010, 70, 425-432.	1.2	52
167	Monocyte chemotactic proteinâ€1 (MCPâ€1/CCL2) is associated with prostatic growth dysregulation and benign prostatic hyperplasia. Prostate, 2010, 70, 473-481.	1.2	62
168	Association of <i>CASP8 D302H</i> polymorphism with reduced risk of aggressive prostate carcinoma. Prostate, 2010, 70, 646-653.	1.2	18
169	Genomeâ€wide linkage analysis of 1,233 prostate cancer pedigrees from the International Consortium for prostate cancer Genetics using novel sumLINK and sumLOD analyses. Prostate, 2010, 70, 735-744.	1.2	22
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19

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