Jacques Simard

List of Publications by Year in descending order

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8172 7944 28,075 348 76 149 citations h-index g-index papers 362 362 362 24379 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Risks of Breast, Ovarian, and Contralateral Breast Cancer for <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers. JAMA - Journal of the American Medical Association, 2017, 317, 2402.	3.8	1,898
2	Association analysis identifies 65 new breast cancer risk loci. Nature, 2017, 551, 92-94.	13.7	1,099
3	Large-scale genotyping identifies 41 new loci associated with breast cancer risk. Nature Genetics, 2013, 45, 353-361.	9.4	960
4	Breast Cancer Risk Genes — Association Analysis in More than 113,000 Women. New England Journal of Medicine, 2021, 384, 428-439.	13.9	532
5	Pathology of Breast and Ovarian Cancers among <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers: Results from the Consortium of Investigators of Modifiers of <i>BRCA1</i> /i>/ <i>22/i> (CIMBA). Cancer Epidemiology Biomarkers and Prevention, 2012, 21, 134-147.</i>	1.1	513
6	Genome-wide association analysis of more than 120,000 individuals identifies 15 new susceptibility loci for breast cancer. Nature Genetics, 2015, 47, 373-380.	9.4	513
7	A candidate prostate cancer susceptibility gene at chromosome 17p. Nature Genetics, 2001, 27, 172-180.	9.4	504
8	Molecular Biology of the 3β-Hydroxysteroid Dehydrogenase/Δ5-Δ4 Isomerase Gene Family. Endocrine Reviews, 2005, 26, 525-582.	8.9	502
9	Endocrine and Intracrine Sources of Androgens in Women: Inhibition of Breast Cancer and Other Roles of Androgens and Their Precursor Dehydroepiandrosterone. Endocrine Reviews, 2003, 24, 152-182.	8.9	500
10	Multiple independent variants at the TERT locus are associated with telomere length and risks of breast and ovarian cancer. Nature Genetics, 2013, 45, 371-384.	9.4	493
11	The key role of 17β-hydroxysteroid dehydrogenases in sex steroid biology. Steroids, 1997, 62, 148-158.	0.8	448
12	Prediction of Breast Cancer Risk Based on Profiling With Common Genetic Variants. Journal of the National Cancer Institute, $2015,107,100$	3.0	428
13	BOADICEA: a comprehensive breast cancer risk prediction model incorporating genetic and nongenetic risk factors. Genetics in Medicine, 2019, 21, 1708-1718.	1.1	415
14	Association of Type and Location of <i>BRCA1 </i> BRCA2 Mutations With Risk of Breast and Ovarian Cancer. JAMA - Journal of the American Medical Association, 2015, 313, 1347.	3.8	390
15	Genome-wide association studies identify four ER negative–specific breast cancer risk loci. Nature Genetics, 2013, 45, 392-398.	9.4	374
16	Large-scale genomic analyses link reproductive aging to hypothalamic signaling, breast cancer susceptibility and BRCA1-mediated DNA repair. Nature Genetics, 2015, 47, 1294-1303.	9.4	357
17	Identification of 12 new susceptibility loci for different histotypes of epithelial ovarian cancer. Nature Genetics, 2017, 49, 680-691.	9.4	356
18	Structure and Expression of a New Complementary DNA Encoding the almost Exclusive 3β-Hydroxysteroid Dehydrogenase/Δ ⁵ -Δ ⁴ -lsomerase in Human Adrenals and Gonads. Molecular Endocrinology, 1991, 5, 1147-1157.	3.7	340

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19	DHEA and the Intracrine Formation of Androgens and Estrogens in Peripheral Target Tissues: Its Role during Aging. Steroids, 1998, 63, 322-328.	0.8	335
20	BRCA2 germline mutations in male breast cancer cases and breast cancer families. Nature Genetics, 1996, 13, 123-125.	9.4	315
21	A locus on 19p13 modifies risk of breast cancer in BRCA1 mutation carriers and is associated with hormone receptor–negative breast cancer in the general population. Nature Genetics, 2010, 42, 885-892.	9.4	309
22	DHEA and Its Transformation into Androgens and Estrogens in Peripheral Target Tissues: Intracrinology. Frontiers in Neuroendocrinology, 2001, 22, 185-212.	2.5	307
23	Identification of ten variants associated with risk of estrogen-receptor-negative breast cancer. Nature Genetics, 2017, 49, 1767-1778.	9.4	289
24	Characterization of cDNAs for Human Estradiol 17β-Dehydrogenase and Assignment of the Gene to Chromosome 17: Evidence of two mRNA Species with Distinct 5′-Termini in Human Placenta. Molecular Endocrinology, 1989, 3, 1301-1309.	3.7	282
25	The OncoArray Consortium: A Network for Understanding the Genetic Architecture of Common Cancers. Cancer Epidemiology Biomarkers and Prevention, 2017, 26, 126-135.	1.1	278
26	Genome-wide association study identifies 32 novel breast cancer susceptibility loci from overall and subtype-specific analyses. Nature Genetics, 2020, 52, 572-581.	9.4	265
27	Common Breast Cancer-Predisposition Alleles Are Associated with Breast Cancer Risk in BRCA1 and BRCA2 Mutation Carriers. American Journal of Human Genetics, 2008, 82, 937-948.	2.6	257
28	The emergence of an ethical duty to disclose genetic research results: international perspectives. European Journal of Human Genetics, 2006, 14, 1170-1178.	1.4	254
29	Genome-Wide Association Study in BRCA1 Mutation Carriers Identifies Novel Loci Associated with Breast and Ovarian Cancer Risk. PLoS Genetics, 2013, 9, e1003212.	1.5	244
30	DHEA and Peripheral Androgen and Estrogen Formation: Intracrinology. Annals of the New York Academy of Sciences, 1995, 774, 16-28.	1.8	243
31	Evaluation of Polygenic Risk Scores for Breast and Ovarian Cancer Risk Prediction in BRCA1 and BRCA2 Mutation Carriers. Journal of the National Cancer Institute, 2017, 109, .	3.0	242
32	Congenital adrenal hyperplasia due to point mutations in the type II 3β–hydroxysteroid dehydrogenase gene. Nature Genetics, 1992, 1, 239-245.	9.4	228
33	Mutational spectrum in a worldwide study of 29,700 families with <i>BRCA1</i> PRCA1SRCA2	1.1	224
34	Identification of six new susceptibility loci for invasive epithelial ovarian cancer. Nature Genetics, 2015, 47, 164-171.	9.4	221
35	RAD51 135Gâ†'C Modifies Breast Cancer Risk among BRCA2 Mutation Carriers: Results from a Combined Analysis of 19 Studies. American Journal of Human Genetics, 2007, 81, 1186-1200.	2.6	217
36	Functional Variants at the 11q13 Risk Locus for Breast Cancer Regulate Cyclin D1 Expression through Long-Range Enhancers. American Journal of Human Genetics, 2013, 92, 489-503.	2.6	201

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37	Effect of Chest X-Rays on the Risk of Breast Cancer Among BRCA1/2 Mutation Carriers in the International BRCA1/2 Carrier Cohort Study: A Report from the EMBRACE, GENEPSO, GEO-HEBON, and IBCCS Collaborators' Group. Journal of Clinical Oncology, 2006, 24, 3361-3366.	0.8	188
38	A transcriptome-wide association study of 229,000 women identifies new candidate susceptibility genes for breast cancer. Nature Genetics, 2018, 50, 968-978.	9.4	184
39	Structure of Two in Tandem Human $17\hat{l}^2$ -Hydroxysteroid Dehydrogenase Genes. Molecular Endocrinology, 1990, 4, 268-275.	3.7	183
40	Personalized early detection and prevention of breast cancer: ENVISION consensus statement. Nature Reviews Clinical Oncology, 2020, 17, 687-705.	12.5	178
41	<i>PALB2</i> , <i>CHEK2</i> and <i>ATM</i> rare variants and cancer risk: data from COGS. Journal of Medical Genetics, 2016, 53, 800-811.	1.5	174
42	Common Breast Cancer Susceptibility Alleles and the Risk of Breast Cancer for <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers: Implications for Risk Prediction. Cancer Research, 2010, 70, 9742-9754.	0.4	169
43	A <scp>RAD</scp> 51 assay feasible in routine tumor samples calls <scp>PARP</scp> inhibitor response beyond <scp>BRCA</scp> mutation. EMBO Molecular Medicine, 2018, 10, .	3.3	169
44	EM-652 (SCH 57068), a third generation SERM acting as pure antiestrogen in the mammary gland and endometrium. Journal of Steroid Biochemistry and Molecular Biology, 1999, 69, 51-84.	1.2	157
45	Genome-Wide Meta-Analyses of Breast, Ovarian, and Prostate Cancer Association Studies Identify Multiple New Susceptibility Loci Shared by at Least Two Cancer Types. Cancer Discovery, 2016, 6, 1052-1067.	7.7	157
46	Prediction of Breast and Prostate Cancer Risks in Male <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers Using Polygenic Risk Scores. Journal of Clinical Oncology, 2017, 35, 2240-2250.	0.8	152
47	Gonadotropin-Releasing Hormone Agonists in the Treatment of Prostate Cancer. Endocrine Reviews, 2005, 26, 361-379.	8.9	149
48	Tamoxifen and Risk of Contralateral Breast Cancer for <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers. Journal of Clinical Oncology, 2013, 31, 3091-3099.	0.8	148
49	(S)-(+)-4-[7-(2,2-Dimethyl-1-oxopro-poxy)-4-methyl-2-[4-[2-(1-piperidinyl)-ethoxy]phenyl]-2H-1-benzopyran-3-yl]- phenyl 2,2-Dimethylpropanoate (EM-800):Â A Highly Potent, Specific, and Orally Active Nonsteroidal Antiestrogen. Journal of Medicinal Chemistry, 1997, 40, 2117-2122.	2.9	143
50	A Combined Genomewide Linkage Scan of 1,233 Families for Prostate Cancer–Susceptibility Genes Conducted by the International Consortium for Prostate Cancer Genetics. American Journal of Human Genetics, 2005, 77, 219-229.	2.6	138
51	Reproductive and Hormonal Factors, and Ovarian Cancer Risk for <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers: Results from the International <i>BRCA1/2</i> Carrier Cohort Study. Cancer Epidemiology Biomarkers and Prevention, 2009, 18, 601-610.	1.1	130
52	Breast cancer risk variants at 6q25 display different phenotype associations and regulate ESR1, RMND1 and CCDC170. Nature Genetics, 2016, 48, 374-386.	9.4	125
53	Role of $17\hat{l}^2$ -Hydroxysteroid Dehydrogenases in Sex Steroid Formation in Peripheral Intracrine Tissues. Trends in Endocrinology and Metabolism, 2000, 11, 421-427.	3.1	124
54	Linkage Analysis of Chromosome 1q Markers in 136 Prostate Cancer Families. American Journal of Human Genetics, 1998, 62, 653-658.	2.6	123

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55	Fine-mapping of 150 breast cancer risk regions identifies 191 likely target genes. Nature Genetics, 2020, 52, 56-73.	9.4	120
56	Genetically Predicted Body Mass Index and Breast Cancer Risk: Mendelian Randomization Analyses of Data from 145,000 Women of European Descent. PLoS Medicine, 2016, 13, e1002105.	3.9	118
57	Regulation of Progesterone-Binding Breast Cyst Protein GCDFP-24 Secretion by Estrogens and Androgens in Human Breast Cancer Cells: A New Marker of Steroid Action in Breast Cancer*. Endocrinology, 1990, 126, 3223-3231.	1.4	110
58	Stimulation of Growth Hormone Release and Synthesis by Estrogens in Rat Anterior Pituitary Cells in Culture. Endocrinology, 1986, 119, 2004-2011.	1.4	105
59	Identification of a BRCA2-Specific Modifier Locus at 6p24 Related to Breast Cancer Risk. PLoS Genetics, 2013, 9, e1003173.	1.5	105
60	Evidence that breast cancer risk at the 2q35 locus is mediated through IGFBP5 regulation. Nature Communications, 2014, 5, 4999.	5.8	105
61	Down-Regulation of Estrogen Receptors by Androgens in the ZR-75-1 Human Breast Cancer Cell Line*. Endocrinology, 1989, 125, 392-399.	1.4	102
62	Regulation of Pro-Gonadotropin-Releasing Hormone Gene Expression by Sex Steroids in The Brain of Male and Female Rats. Molecular Endocrinology, 1989, 3, 1748-1756.	3.7	99
63	Common variants in LSP1, 2q35 and 8q24 and breast cancer risk for BRCA1 and BRCA2 mutation carriers. Human Molecular Genetics, 2009, 18, 4442-4456.	1.4	99
64	Height and Breast Cancer Risk: Evidence From Prospective Studies and Mendelian Randomization. Journal of the National Cancer Institute, 2015, 107, djv219.	3.0	99
65	Genetic discrimination and life insurance: a systematic review of the evidence. BMC Medicine, 2013, 11, 25.	2.3	98
66	Fine-Scale Mapping of the FGFR2 Breast Cancer Risk Locus: Putative Functional Variants Differentially Bind FOXA1 and E2F1. American Journal of Human Genetics, 2013, 93, 1046-1060.	2.6	98
67	Identification of four novel susceptibility loci for oestrogen receptor negative breast cancer. Nature Communications, 2016, 7, 11375.	5.8	93
68	STRUCTURE AND SEXUAL DIMORPHIC EXPRESSION OF A LIVER-SPECIFIC RAT 3β-HYDROXYSTEROID DEHYDROGENASE/ISOMERASE. Endocrinology, 1990, 127, 3237-3239.	1.4	92
69	Genome-wide association and transcriptome studies identify target genes and risk loci for breast cancer. Nature Communications, 2019, 10, 1741.	5.8	90
70	New Insight into the Molecular Basis of 3β-Hydroxysteroid Dehydrogenase Deficiency: Identification of Eight Mutations in the HSD3B2 Gene in Eleven Patients from Seven New Families and Comparison of the Functional Properties of Twenty-Five Mutant Enzymes1. Journal of Clinical Endocrinology and Metabolism, 1999, 84, 4410-4425.	1.8	88
71	Shared heritability and functional enrichment across six solid cancers. Nature Communications, 2019, 10, 431.	5.8	88
72	European polygenic risk score for prediction of breast cancer shows similar performance in Asian women. Nature Communications, 2020, 11, 3833.	5.8	88

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73	Combination therapy for prostate cancer. Endocrine and biologic basis of its choice as new standard first-line therapy. Cancer, 1993, 71, 1059-1067.	2.0	86
74	Stimulation of Androgen-Dependent Gene Expression by the Adrenal Precursors Dehydroepiandrosterone and Androstenedione in the Rat Ventral Prostate*. Endocrinology, 1989, 124, 2745-2754.	1.4	85
75	The Tumor Suppressor PALB2: Inside Out. Trends in Biochemical Sciences, 2019, 44, 226-240.	3.7	83
76	GATA Factors and the Nuclear Receptors, Steroidogenic Factor $1/\text{Liver}$ Receptor Homolog 1 , Are Key Mutual Partners in the Regulation of the Human 31^2 -Hydroxysteroid Dehydrogenase Type 2 Promoter. Molecular Endocrinology, 2005, 19 , 2358-2370.	3.7	82
77	Polygenic risk scores and breast and epithelial ovarian cancer risks for carriers of BRCA1 and BRCA2 pathogenic variants. Genetics in Medicine, 2020, 22, 1653-1666.	1.1	82
78	Molecular cloning, cDNA structure and predicted amino acid sequence of bovine 3β-hydroxy-5-ene steroid dehydrogenase/Δ5-Δ4isomerase. FEBS Letters, 1989, 259, 153-157.	1.3	81
79	Associations of obesity and circulating insulin and glucose with breast cancer risk: a Mendelian randomization analysis. International Journal of Epidemiology, 2019, 48, 795-806.	0.9	81
80	Association of Specific LDL Receptor Gene Mutations With Differential Plasma Lipoprotein Response to Simvastatin in Young French Canadians With Heterozygous Familial Hypercholesterolemia. Arteriosclerosis, Thrombosis, and Vascular Biology, 1998, 18, 1007-1012.	1.1	80
81	Incorporating truncating variants in PALB2, CHEK2, and ATM into the BOADICEA breast cancer risk model. Genetics in Medicine, 2016, 18, 1190-1198.	1.1	80
82	Common variants at 12p11, 12q24, 9p21, 9q31.2 and in ZNF365 are associated with breast cancer risk for BRCA1 and/or BRCA2mutation carriers. Breast Cancer Research, 2012, 14, R33.	2.2	78
83	Functional mechanisms underlying pleiotropic risk alleles at the 19p13.1 breast–ovarian cancer susceptibility locus. Nature Communications, 2016, 7, 12675.	5.8	78
84	BRCA2 Polymorphic Stop Codon K3326X and the Risk of Breast, Prostate, and Ovarian Cancers. Journal of the National Cancer Institute, 2016, 108, djv315.	3.0	77
85	Fine-Scale Mapping of the 5q11.2 Breast Cancer Locus Reveals at Least Three Independent Risk Variants Regulating MAP3K1. American Journal of Human Genetics, 2015, 96, 5-20.	2.6	76
86	BRCA1 and BRCA2 mutation predictions using the BOADICEA and BRCAPRO models and penetrance estimation in high-risk French-Canadian families. Breast Cancer Research, 2005, 8, R3.	2.2	75
87	Assessment of polygenic architecture and risk prediction based on common variants across fourteen cancers. Nature Communications, 2020, 11, 3353.	5.8	7 5
88	Evaluation of BRCA1 and BRCA2 mutation prevalence, risk prediction models and a multistep testing approach in French-Canadian families with high risk of breast and ovarian cancer. Journal of Medical Genetics, 2006, 44, 107-121.	1.5	72
89	Common breast cancer susceptibility alleles are associated with tumour subtypes in BRCA1 and BRCA2 mutation carriers: results from the Consortium of Investigators of Modifiers of BRCA1/2. Breast Cancer Research, 2011, 13, R110.	2.2	71
90	Molecular basis of human $3\hat{l}^2$ -hydroxysteroid dehydrogenase deficiency. Journal of Steroid Biochemistry and Molecular Biology, 1995, 53, 127-138.	1.2	70

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91	Structure-function relationships of 3β-hydroxysteroid dehydrogenase: Contribution made by the molecular genetics of 3β-hydroxysteroid dehydrogenase deficiency. Steroids, 1997, 62, 176-184.	0.8	68
92	Common alleles at $6q25.1$ and $1p11.2$ are associated with breast cancer risk for BRCA1 and BRCA2 mutation carriers. Human Molecular Genetics, 2011, 20, 3304-3321.	1.4	68
93	Induction of $3\hat{l}^2$ -Hydroxysteroid Dehydrogenase \hat{l} "5- \hat{l} "4 Isomerase Type 1 Gene Transcription in Human Breast Cancer Cell Lines and in Normal Mammary Epithelial Cells by Interleukin-4 and Interleukin-13. Molecular Endocrinology, 1999, 13, 66-81.	3.7	67
94	Multiple novel prostate cancer susceptibility signals identified by fine-mapping of known risk loci among Europeans. Human Molecular Genetics, 2015, 24, 5589-5602.	1.4	67
95	Localization of 3β-Hydroxysteroid DehydrogenaseΔ ₅ -Δ ₄ -Isomerase in Rat Gonads and Adrenal Glands by Immunocytochemistry and <i>in Situ</i> Hybridization. Endocrinology, 1990, 127, 1394-1403.	1.4	66
96	11 Structure, regulation and role of $3\hat{l}^2$ -hydroxysteroid dehydrogenase, $17\hat{l}^2$ -hydroxysteroid dehydrogenase and aromatase enzymes in the formation of sex steroids in classical and peripheral intracrine tissues. Bailliere's Clinical Endocrinology and Metabolism, 1994, 8, 451-474.	1.0	66
97	The Human Type II 17β-Hydroxysteroid Dehydrogenase Gene Encodes Two Alternatively Spliced mRNA Species. DNA and Cell Biology, 1995, 14, 849-861.	0.9	65
98	Characterization of macaque 3β-hydroxy-5-ene steroid dehydrogenase/Δ5-Δ4 isomerase: structure and expression in steroidogenic and peripheral tissues in primate. Molecular and Cellular Endocrinology, 1991, 75, 101-110.	1.6	63
99	Genetic mapping of the breast-ovarian cancer syndrome to a small interval on chromosome 17q12–21: exclusion of candidate genes EDH17B2 and RARA. Human Molecular Genetics, 1993, 2, 1193-1199.	1.4	63
100	Characterization of the effects of the novel non-steroidal antiestrogen EM-800 on basal and estrogen-induced proliferation of T-47D, ZR-75-1 and MCF-7 human breast cancer cellsin vitro. , 1997, 73, 104-112.		63
101	Inhibitory Effect of Estrogens on GCDFP-15 mRNA Levels and Secretion in ZR-75-1 Human Breast Cancer Cells. Molecular Endocrinology, 1989, 3, 694-702.	3.7	61
102	Congenital Adrenal Hyperplasia due to 3β-Hydroxysteroid Dehydrogenase/ ΰ5-ΰ4lsomerase Deficiency. Seminars in Reproductive Medicine, 2002, 20, 255-276.	0.5	59
103	Evidence that the 5p12 Variant rs10941679 Confers Susceptibility to Estrogen-Receptor-Positive Breast Cancer through FGF10 and MRPS30 Regulation. American Journal of Human Genetics, 2016, 99, 903-911.	2.6	59
104	Personalized Risk Assessment for Prevention and Early Detection of Breast Cancer: Integration and Implementation (PERSPECTIVE I& Detection of Personalized Medicine, 2021, 11, 511.	1.1	59
105	Regulation of 3β-Hydroxysteroid Dehydrogenase/Δ ⁵ -Δ ⁴ Isomerase Expression and Activity in the Hypophysectomized Rat Ovary: Interactions between the Stimulatory Effect of Human Chorionic Gonadotropin and the Luteolytic Effect of Prolactin*. Endocrinology, 1990, 127, 2726-2737.	1.4	58
106	Induction of 3Î ² -Hydroxysteroid Dehydrogenase/ Isomerase Type 1 Expression by Interleukin-4 in Human Normal Prostate Epithelial Cells, Immortalized Keratinocytes, Colon, and Cervix Cancer Cell Lines1. Endocrinology, 1999, 140, 4573-4584.	1.4	57
107	Pooled genome linkage scan of aggressive prostate cancer: results from the International Consortium for Prostate Cancer Genetics. Human Genetics, 2006, 120, 471-485.	1.8	57
108	Associations of common breast cancer susceptibility alleles with risk of breast cancer subtypes in BRCA1 and BRCA2 mutation carriers. Breast Cancer Research, 2014, 16, 3416.	2.2	57

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109	Novel Associations between Common Breast Cancer Susceptibility Variants and Risk-Predicting Mammographic Density Measures. Cancer Research, 2015, 75, 2457-2467.	0.4	55
110	A Transcriptome-Wide Association Study Among 97,898 Women to Identify Candidate Susceptibility Genes for Epithelial Ovarian Cancer Risk. Cancer Research, 2018, 78, 5419-5430.	0.4	54
111	Interleukin-4 and interleukin-13 inhibit estrogen-induced breast cancer cell proliferation and stimulate GCDFP-15 expression in human breast cancer cells. Molecular and Cellular Endocrinology, 1996, 121, 11-18.	1.6	53
112	Common non-synonymous SNPs associated with breast cancer susceptibility: findings from the Breast Cancer Association Consortium. Human Molecular Genetics, 2014, 23, 6096-6111.	1.4	53
113	Prediction of breast cancer risk based on common genetic variants in women of East Asian ancestry. Breast Cancer Research, 2016, 18, 124.	2.2	52
114	Genome-wide association study of germline variants and breast cancer-specific mortality. British Journal of Cancer, 2019, 120, 647-657.	2.9	52
115	Molecular and genealogical characterization of the R1443X BRCA1 mutation in high-risk French-Canadian breast/ovarian cancer families. Human Genetics, 2005, 117, 119-132.	1.8	51
116	Fineâ€scale mapping of 8q24 locus identifies multiple independent risk variants for breast cancer. International Journal of Cancer, 2016, 139, 1303-1317.	2.3	51
117	Perspective: Prostate Cancer Susceptibility Genes. Endocrinology, 2002, 143, 2029-2040.	1.4	49
118	MicroRNA Related Polymorphisms and Breast Cancer Risk. PLoS ONE, 2014, 9, e109973.	1.1	49
119	Factors Associated with an Individual's Decision to Withdraw from Genetic Testing for Breast And Ovarian Cancer Susceptibility: Implications for Counseling. Genetic Testing and Molecular Biomarkers, 2007, 11, 45-54.	1.7	48
120	Characterization of the Cancer Spectrum in Men With Germline <i>BRCA1</i> Aland <i>BRCA2</i> Frathogenic Variants. JAMA Oncology, 2020, 6, 1218.	3.4	48
121	Are ATM Mutations 7271T→G and IVS10-6T→G Really High-Risk Breast Cancer-Susceptibility Alleles?. Cancer Research, 2004, 64, 840-843.	0.4	47
122	Common Variants at the 19p13.1 and <i>ZNF365</i> Loci Are Associated with ER Subtypes of Breast Cancer and Ovarian Cancer Risk in <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers. Cancer Epidemiology Biomarkers and Prevention, 2012, 21, 645-657.	1.1	47
123	DNA Glycosylases Involved in Base Excision Repair May Be Associated with Cancer Risk in BRCA1 and BRCA2 Mutation Carriers. PLoS Genetics, 2014, 10, e1004256.	1.5	47
124	EM-652 (SCH57068), a pure SERM having complete antiestrogenic activity in the mammary gland and endometrium. Journal of Steroid Biochemistry and Molecular Biology, 2001, 79, 213-225.	1.2	46
125	No Evidence of False Reassurance among Women with an Inconclusive BRCA1/2 Genetic Test Result. Cancer Epidemiology Biomarkers and Prevention, 2005, 14, 2862-2867.	1.1	46
126	Identification of novel breast cancer susceptibility loci in meta-analyses conducted among Asian and European descendants. Nature Communications, 2020, 11, 1217.	5.8	46

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127	Mutation analysis and characterization of ATR sequence variants in breast cancer cases from high-risk French Canadian breast/ovarian cancer families. BMC Cancer, 2006, 6, 230.	1.1	45
128	No Evidence of BRCA1/2 Genomic Rearrangements in High-Risk French-Canadian Breast/Ovarian Cancer Families. Genetic Testing and Molecular Biomarkers, 2006, 10, 104-115.	1.7	45
129	Personalized medicine and access to health care: potential for inequitable access?. European Journal of Human Genetics, 2013, 21, 143-147.	1.4	45
130	Functional analysis of genetic variants in the high-risk breast cancer susceptibility gene PALB2. Nature Communications, 2019, 10, 5296.	5.8	45
131	Generation of a Transcription Map at the HSD17B Locus Centromeric to BRCA1 at 17q21. Genomics, 1995, 28, 530-542.	1.3	44
132	Multihormonal Control of Pre-Pro-Somatostatin mRNA Levels in the Periventricular Nucleus of the Male and Female Rat Hypothalamus. Neuroendocrinology, 1990, 52, 527-536.	1.2	43
133	Genetic predisposition to ductal carcinoma in situ of the breast. Breast Cancer Research, 2016, 18, 22.	2,2	43
134	Prospective evaluation of a breast-cancer risk model integrating classical risk factors and polygenic risk in 15 cohorts from six countries. International Journal of Epidemiology, 2022, 50, 1897-1911.	0.9	43
135	Localization of the Endogenous Benzodiazepine Ligand Octadecaneuropeptide in the Rat Testis. Endocrinology, 1990, 127, 1986-1994.	1.4	42
136	Inheritance of deleterious mutations at both BRCA1 and BRCA2 in an international sample of 32,295 women. Breast Cancer Research, 2016, 18, 112.	2,2	42
137	Crucial role of cytokines in sex steroid formation in normal and tumoral tissues. Molecular and Cellular Endocrinology, 2001, 171, 25-40.	1.6	41
138	Results of a genome-wide linkage analysis in prostate cancer families ascertained through the ACTANE consortium. Prostate, 2003, 57, 270-279.	1.2	41
139	Life insurance: genomic stratification and risk classification. European Journal of Human Genetics, 2014, 22, 575-579.	1.4	41
140	A Mendelian randomization analysis of circulating lipid traits and breast cancer risk. International Journal of Epidemiology, 2020, 49, 1117-1131.	0.9	41
141	Risk-reducing salpingo-oophorectomy, natural menopause, and breast cancer risk: an international prospective cohort of BRCA1 and BRCA2 mutation carriers. Breast Cancer Research, 2020, 22, 8.	2.2	41
142	Genetic Variation at 9p22.2 and Ovarian Cancer Risk for BRCA1 and BRCA2 Mutation Carriers. Journal of the National Cancer Institute, 2011, 103, 105-116.	3.0	40
143	Fine-mapping identifies two additional breast cancer susceptibility loci at 9q31.2. Human Molecular Genetics, 2015, 24, 2966-2984.	1.4	40
144	Functional characterization of 84 PALB2 variants of uncertain significance. Genetics in Medicine, 2020, 22, 622-632.	1.1	40

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145	Congenital adrenal hyperplasia caused by a novel homozygous frameshift mutation 273î"AA in type II 3β-hydroxysteroid dehydrogenase gene (HSD3B2) in three male patients of Afghan/Pakistani origin. Human Molecular Genetics, 1994, 3, 327-330.	1.4	39
146	Inverse relationships between cell proliferation and basal or androgen-stimulated apolipoprotein D secretion in LNCaP human prostate cancer cells. Journal of Steroid Biochemistry and Molecular Biology, 1994, 51, 167-174.	1.2	39
147	Interleukin-6 inhibits the potent stimulatory action of androgens, glucocorticoids and interleukin-1α on apolipoprotein D and GCDFP-15 expression in human breast cancer cells. International Journal of Cancer, 1995, 62, 732-737.	2.3	39
148	Comparison of in vitro effects of the pure antiandrogens OH-flutamide, casodex, and nilutamide on androgen-sensitive parameters. Urology, 1997, 49, 580-589.	0.5	39
149	A Novel A10E Homozygous Mutation in the <i>HSD3B2</i> Gene Causing Severe Salt-Wasting 3β-Hydroxysteroid Dehydrogenase Deficiency in 46,XX and 46,XY French-Canadians: Evaluation of Gonadal Function after Puberty ¹ . Journal of Clinical Endocrinology and Metabolism, 2000. 85, 1968-1974.	1.8	39
150	Genetic Predisposition to In Situ and Invasive Lobular Carcinoma of the Breast. PLoS Genetics, 2014, 10, e1004285.	1.5	39
151	Association of Genomic Domains in <i>BRCA1</i> and <i>BRCA2</i> with Prostate Cancer Risk and Aggressiveness. Cancer Research, 2020, 80, 624-638.	0.4	39
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