

Stig Bojesen

List of Publications by Citations

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292
papers

20,912
citations

73
h-index

139
g-index

326
ext. papers

25,866
ext. citations

10
avg, IF

6.29
L-index

#	Paper	IF	Citations
292	Genome-wide association study identifies novel breast cancer susceptibility loci. <i>Nature</i> , 2007 , 447, 1087-93	50.4	1957
291	Large-scale genotyping identifies 41 new loci associated with breast cancer risk. <i>Nature Genetics</i> , 2013 , 45, 353-61, 361e1-2	36.3	813
290	Statin use and reduced cancer-related mortality. <i>New England Journal of Medicine</i> , 2012 , 367, 1792-802	59.2	669
289	Association analysis identifies 65 new breast cancer risk loci. <i>Nature</i> , 2017 , 551, 92-94	50.4	643
288	A common coding variant in CASP8 is associated with breast cancer risk. <i>Nature Genetics</i> , 2007 , 39, 352-8	36.3	557
287	Associations of breast cancer risk factors with tumor subtypes: a pooled analysis from the Breast Cancer Association Consortium studies. <i>Journal of the National Cancer Institute</i> , 2011 , 103, 250-63	9.7	513
286	Multiple independent variants at the TERT locus are associated with telomere length and risks of breast and ovarian cancer. <i>Nature Genetics</i> , 2013 , 45, 371-84, 384e1-2	36.3	422
285	Identification of 23 new prostate cancer susceptibility loci using the iCOGS custom genotyping array. <i>Nature Genetics</i> , 2013 , 45, 385-91, 391e1-2	36.3	413
284	Genome-wide association analysis of more than 120,000 individuals identifies 15 new susceptibility loci for breast cancer. <i>Nature Genetics</i> , 2015 , 47, 373-80	36.3	406
283	Parent-of-origin-specific allelic associations among 106 genomic loci for age at menarche. <i>Nature</i> , 2014 , 514, 92-97	50.4	401
282	Newly discovered breast cancer susceptibility loci on 3p24 and 17q23.2. <i>Nature Genetics</i> , 2009 , 41, 585-90	36.3	393
281	C-reactive protein as a predictor of prognosis in chronic obstructive pulmonary disease. <i>American Journal of Respiratory and Critical Care Medicine</i> , 2007 , 175, 250-5	10.2	391
280	Polygenic Risk Scores for Prediction of Breast Cancer and Breast Cancer Subtypes. <i>American Journal of Human Genetics</i> , 2019 , 104, 21-34	11	363
279	Genome-wide association studies identify four ER negative-specific breast cancer risk loci. <i>Nature Genetics</i> , 2013 , 45, 392-8, 398e1-2	36.3	327
278	Prediction of breast cancer risk based on profiling with common genetic variants. <i>Journal of the National Cancer Institute</i> , 2015 , 107,	9.7	324
277	MicroRNA biomarkers in whole blood for detection of pancreatic cancer. <i>JAMA - Journal of the American Medical Association</i> , 2014 , 311, 392-404	27.4	315
276	Baseline C-reactive protein is associated with incident cancer and survival in patients with cancer. <i>Journal of Clinical Oncology</i> , 2009 , 27, 2217-24	2.2	305

275	Heterogeneity of breast cancer associations with five susceptibility loci by clinical and pathological characteristics. <i>PLoS Genetics</i> , 2008 , 4, e1000054	6	280
274	Genomic analyses identify hundreds of variants associated with age at menarche and support a role for puberty timing in cancer risk. <i>Nature Genetics</i> , 2017 , 49, 834-841	36.3	257
273	Large-scale association analysis identifies new lung cancer susceptibility loci and heterogeneity in genetic susceptibility across histological subtypes. <i>Nature Genetics</i> , 2017 , 49, 1126-1132	36.3	246
272	Seven prostate cancer susceptibility loci identified by a multi-stage genome-wide association study. <i>Nature Genetics</i> , 2011 , 43, 785-91	36.3	243
271	Genome-wide association analysis identifies three new breast cancer susceptibility loci. <i>Nature Genetics</i> , 2012 , 44, 312-8	36.3	237
270	Large-scale genomic analyses link reproductive aging to hypothalamic signaling, breast cancer susceptibility and BRCA1-mediated DNA repair. <i>Nature Genetics</i> , 2015 , 47, 1294-1303	36.3	226
269	CHEK2*1100delC genotyping for clinical assessment of breast cancer risk: meta-analyses of 26,000 patient cases and 27,000 controls. <i>Journal of Clinical Oncology</i> , 2008 , 26, 542-8	2.2	216
268	Peripheral blood leukocyte telomere length and mortality among 64,637 individuals from the general population. <i>Journal of the National Cancer Institute</i> , 2015 , 107, djv074	9.7	200
267	Low 25-hydroxyvitamin D and risk of type 2 diabetes: a prospective cohort study and metaanalysis. <i>Clinical Chemistry</i> , 2013 , 59, 381-91	5.5	189
266	Identification of ten variants associated with risk of estrogen-receptor-negative breast cancer. <i>Nature Genetics</i> , 2017 , 49, 1767-1778	36.3	186
265	The OncoArray Consortium: A Network for Understanding the Genetic Architecture of Common Cancers. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2017 , 26, 126-135	4	183
264	Genetically low vitamin D concentrations and increased mortality: Mendelian randomisation analysis in three large cohorts. <i>BMJ, The</i> , 2014 , 349, g6330	5.9	182
263	Short telomere length, cancer survival, and cancer risk in 47102 individuals. <i>Journal of the National Cancer Institute</i> , 2013 , 105, 459-68	9.7	168
262	Functional variants at the 11q13 risk locus for breast cancer regulate cyclin D1 expression through long-range enhancers. <i>American Journal of Human Genetics</i> , 2013 , 92, 489-503	11	167
261	Worldwide trends in hypertension prevalence and progress in treatment and control from 1990 to 2019: a pooled analysis of 1201 population-representative studies with 104 million participants. <i>Lancet, The</i> , 2021 , 398, 957-980	40	154
260	Short telomere length, myocardial infarction, ischemic heart disease, and early death. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2012 , 32, 822-9	9.4	152
259	Breast Cancer Risk Genes - Association Analysis in More than 113,000 Women. <i>New England Journal of Medicine</i> , 2021 , 384, 428-439	59.2	143
258	Low penetrance breast cancer susceptibility loci are associated with specific breast tumor subtypes: findings from the Breast Cancer Association Consortium. <i>Human Molecular Genetics</i> , 2011 , 20, 3289-303	5.6	140

257	CHEK2*1100delC heterozygosity in women with breast cancer associated with early death, breast cancer-specific death, and increased risk of a second breast cancer. <i>Journal of Clinical Oncology</i> , 2012 , 30, 4308-16	2.2	134
256	PALB2, CHEK2 and ATM rare variants and cancer risk: data from COGS. <i>Journal of Medical Genetics</i> , 2016 , 53, 800-811	5.8	121
255	Vitamin D concentration, obesity, and risk of diabetes: a mendelian randomisation study. <i>Lancet Diabetes and Endocrinology</i> , 2014 , 2, 298-306	18.1	120
254	Short telomere length, lung function and chronic obstructive pulmonary disease in 46,396 individuals. <i>Thorax</i> , 2013 , 68, 429-35	7.3	118
253	Telomere shortening unrelated to smoking, body weight, physical activity, and alcohol intake: 4,576 general population individuals with repeat measurements 10 years apart. <i>PLoS Genetics</i> , 2014 , 10, e1004191	6	112
252	Evidence of gene-environment interactions between common breast cancer susceptibility loci and established environmental risk factors. <i>PLoS Genetics</i> , 2013 , 9, e1003284	6	112
251	Reduced 25-hydroxyvitamin D and risk of Alzheimer's disease and vascular dementia. <i>Alzheimer's and Dementia</i> , 2014 , 10, 296-302	1.2	111
250	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
249	Age- and Tumor Subtype-Specific Breast Cancer Risk Estimates for CHEK2*1100delC Carriers. <i>Journal of Clinical Oncology</i> , 2016 , 34, 2750-60	2.2	107
248	A genome-wide association scan (GWAS) for mean telomere length within the COGS project: identified loci show little association with hormone-related cancer risk. <i>Human Molecular Genetics</i> , 2013 , 22, 5056-64	5.6	107
247	Genome-Wide Meta-Analyses of Breast, Ovarian, and Prostate Cancer Association Studies Identify Multiple New Susceptibility Loci Shared by at Least Two Cancer Types. <i>Cancer Discovery</i> , 2016 , 6, 1052-67	24.4	104
246	A transcriptome-wide association study of 229,000 women identifies new candidate susceptibility genes for breast cancer. <i>Nature Genetics</i> , 2018 , 50, 968-978	36.3	101
245	Tumor suppressor p53 Arg72Pro polymorphism and longevity, cancer survival, and risk of cancer in the general population. <i>Journal of Experimental Medicine</i> , 2007 , 204, 1295-301	16.6	100
244	Breast cancer risk variants at 6q25 display different phenotype associations and regulate ESR1, RMND1 and CCDC170. <i>Nature Genetics</i> , 2016 , 48, 374-86	36.3	93
243	19p13.1 is a triple-negative-specific breast cancer susceptibility locus. <i>Cancer Research</i> , 2012 , 72, 1795-803	33.1	93
242	Identification of a BRCA2-specific modifier locus at 6p24 related to breast cancer risk. <i>PLoS Genetics</i> , 2013 , 9, e1003173	6	90
241	Plasma YKL-40 levels in healthy subjects from the general population. <i>Clinica Chimica Acta</i> , 2011 , 412, 709-12	6.2	90
240	C-reactive protein and the risk of cancer: a mendelian randomization study. <i>Journal of the National Cancer Institute</i> , 2010 , 102, 202-6	9.7	90

239	Fine-mapping identifies multiple prostate cancer risk loci at 5p15, one of which associates with TERT expression. <i>Human Molecular Genetics</i> , 2013 , 22, 2520-8	5.6	88
238	Increased risk of breast cancer associated with CHEK2*1100delC. <i>Journal of Clinical Oncology</i> , 2007 , 25, 57-63	2.2	88
237	Evidence that breast cancer risk at the 2q35 locus is mediated through IGFBP5 regulation. <i>Nature Communications</i> , 2014 , 4, 4999	17.4	87
236	The JAK2 V617F somatic mutation, mortality and cancer risk in the general population. <i>Haematologica</i> , 2011 , 96, 450-3	6.6	85
235	Telomeres and human health. <i>Journal of Internal Medicine</i> , 2013 , 274, 399-413	10.8	84
234	Genome-wide association study identifies a common variant in RAD51B associated with male breast cancer risk. <i>Nature Genetics</i> , 2012 , 44, 1182-4	36.3	84
233	No evidence that protein truncating variants in BRIP1 are associated with breast cancer risk: implications for gene panel testing. <i>Journal of Medical Genetics</i> , 2016 , 53, 298-309	5.8	83
232	(cg05575921) hypomethylation marks smoking behaviour, morbidity and mortality. <i>Thorax</i> , 2017 , 72, 646-653	7.3	82
231	Refined histopathological predictors of BRCA1 and BRCA2 mutation status: a large-scale analysis of breast cancer characteristics from the BCAC, CIMBA, and ENIGMA consortia. <i>Breast Cancer Research</i> , 2014 , 16, 3419	8.3	82
230	Fine-scale mapping of the FGFR2 breast cancer risk locus: putative functional variants differentially bind FOXA1 and E2F1. <i>American Journal of Human Genetics</i> , 2013 , 93, 1046-60	11	80
229	Genetically Predicted Body Mass Index and Breast Cancer Risk: Mendelian Randomization Analyses of Data from 145,000 Women of European Descent. <i>PLoS Medicine</i> , 2016 , 13, e1002105	11.6	80
228	Risk of cancer among HIV-infected individuals compared to the background population: impact of smoking and HIV. <i>Aids</i> , 2014 , 28, 1499-508	3.5	77
227	Low plasma 25-hydroxyvitamin D and risk of tobacco-related cancer. <i>Clinical Chemistry</i> , 2013 , 59, 771-80	5.5	77
226	Genome-wide association study identifies 32 novel breast cancer susceptibility loci from overall and subtype-specific analyses. <i>Nature Genetics</i> , 2020 , 52, 572-581	36.3	76
225	Serum Biomarker Signature-Based Liquid Biopsy for Diagnosis of Early-Stage Pancreatic Cancer. <i>Journal of Clinical Oncology</i> , 2018 , 36, 2887-2894	2.2	76
224	Association of ESR1 gene tagging SNPs with breast cancer risk. <i>Human Molecular Genetics</i> , 2009 , 18, 1133-9	3.9	75
223	The role of genetic breast cancer susceptibility variants as prognostic factors. <i>Human Molecular Genetics</i> , 2012 , 21, 3926-39	5.6	75
222	Height and Breast Cancer Risk: Evidence From Prospective Studies and Mendelian Randomization. <i>Journal of the National Cancer Institute</i> , 2015 , 107,	9.7	74

221	Lymphopenia and risk of infection and infection-related death in 98,344 individuals from a prospective Danish population-based study. <i>PLoS Medicine</i> , 2018 , 15, e1002685	11.6	74
220	Height and body-mass index trajectories of school-aged children and adolescents from 1985 to 2019 in 200 countries and territories: a pooled analysis of 2181 population-based studies with 65 million participants. <i>Lancet, The</i> , 2020 , 396, 1511-1524	4.0	73
219	Short Telomere Length and Ischemic Heart Disease: Observational and Genetic Studies in 290 022 Individuals. <i>Clinical Chemistry</i> , 2016 , 62, 1140-9	5.5	70
218	Increased Risk for Other Cancers in Addition to Breast Cancer for CHEK2*1100delC Heterozygotes Estimated From the Copenhagen General Population Study. <i>Journal of Clinical Oncology</i> , 2016 , 34, 1208-16	2.2	70
217	Elevated plasma YKL-40 predicts increased risk of gastrointestinal cancer and decreased survival after any cancer diagnosis in the general population. <i>Journal of Clinical Oncology</i> , 2009 , 27, 572-8	2.2	70
216	Diagnostic value of JAK2 V617F somatic mutation for myeloproliferative cancer in 49 488 individuals from the general population. <i>British Journal of Haematology</i> , 2013 , 160, 70-9	4.5	68
215	The common germline Arg72Pro polymorphism of p53 and increased longevity in humans. <i>Cell Cycle</i> , 2008 , 7, 158-63	4.7	68
214	Prevalence of the HOXB13 G84E germline mutation in British men and correlation with prostate cancer risk, tumour characteristics and clinical outcomes. <i>Annals of Oncology</i> , 2015 , 26, 756-761	10.3	67
213	Associations of common variants at 1p11.2 and 14q24.1 (RAD51L1) with breast cancer risk and heterogeneity by tumor subtype: findings from the Breast Cancer Association Consortium. <i>Human Molecular Genetics</i> , 2011 , 20, 4693-706	5.6	66
212	BRCA2 Polymorphic Stop Codon K3326X and the Risk of Breast, Prostate, and Ovarian Cancers. <i>Journal of the National Cancer Institute</i> , 2016 , 108,	9.7	65
211	Increased body mass index, elevated C-reactive protein, and short telomere length. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014 , 99, E1671-5	5.6	65
210	Integrin beta3 Leu33Pro homozygosity and risk of cancer. <i>Journal of the National Cancer Institute</i> , 2003 , 95, 1150-7	9.7	65
209	Identification of four novel susceptibility loci for oestrogen receptor negative breast cancer. <i>Nature Communications</i> , 2016 , 7, 11375	17.4	64
208	Plasma 25-hydroxyvitamin D, lung function and risk of chronic obstructive pulmonary disease. <i>Thorax</i> , 2014 , 69, 24-31	7.3	63
207	Long telomeres and cancer risk among 95 568 individuals from the general population. <i>International Journal of Epidemiology</i> , 2016 , 45, 1634-1643	7.8	62
206	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
205	Association of clinical benign prostate hyperplasia with prostate cancer incidence and mortality revisited: a nationwide cohort study of 3,009,258 men. <i>European Urology</i> , 2011 , 60, 691-8	10.2	61
204	Fine-scale mapping of the 5q11.2 breast cancer locus reveals at least three independent risk variants regulating MAP3K1. <i>American Journal of Human Genetics</i> , 2015 , 96, 5-20	11	59

203	Statin use and reduced cancer-related mortality. <i>New England Journal of Medicine</i> , 2013 , 368, 576-7	59.2	58
202	Plasma YKL-40 and total and disease-specific mortality in the general population. <i>Clinical Chemistry</i> , 2010 , 56, 1580-91	5.5	56
201	Elevated plasma YKL-40 levels and ischemic stroke in the general population. <i>Annals of Neurology</i> , 2010 , 68, 672-80	9.4	56
200	Obesity, metabolic factors and risk of different histological types of lung cancer: A Mendelian randomization study. <i>PLoS ONE</i> , 2017 , 12, e0177875	3.7	56
199	Fine-mapping of 150 breast cancer risk regions identifies 191 likely target genes. <i>Nature Genetics</i> , 2020 , 52, 56-73	36.3	56
198	Combined genetic and splicing analysis of BRCA1 c.[594-2A>C; 641A>G] highlights the relevance of naturally occurring in-frame transcripts for developing disease gene variant classification algorithms. <i>Human Molecular Genetics</i> , 2016 , 25, 2256-2268	5.6	55
197	Platelet glycoprotein IIb/IIIa Pl(A2)/Pl(A2) homozygosity associated with risk of ischemic cardiovascular disease and myocardial infarction in young men: the Copenhagen City Heart Study. <i>Journal of the American College of Cardiology</i> , 2003 , 42, 661-7	15.1	55
196	Multiple novel prostate cancer susceptibility signals identified by fine-mapping of known risk loci among Europeans. <i>Human Molecular Genetics</i> , 2015 , 24, 5589-602	5.6	54
195	JAK2V617F somatic mutation in the general population: myeloproliferative neoplasm development and progression rate. <i>Haematologica</i> , 2014 , 99, 1448-55	6.6	54
194	Joint associations of a polygenic risk score and environmental risk factors for breast cancer in the Breast Cancer Association Consortium. <i>International Journal of Epidemiology</i> , 2018 , 47, 526-536	7.8	53
193	Functional mechanisms underlying pleiotropic risk alleles at the 19p13.1 breast-ovarian cancer susceptibility locus. <i>Nature Communications</i> , 2016 , 7, 12675	17.4	53
192	Associations of obesity and circulating insulin and glucose with breast cancer risk: a Mendelian randomization analysis. <i>International Journal of Epidemiology</i> , 2019 , 48, 795-806	7.8	52
191	Genetic modifiers of CHEK2*1100delC-associated breast cancer risk. <i>Genetics in Medicine</i> , 2017 , 19, 599-603	6.0	51
190	Inflammatory biomarkers and risk of cancer in 84,000 individuals from the general population. <i>International Journal of Cancer</i> , 2016 , 139, 1493-500	7.5	51
189	Hypomorphic Missense Variants Confer Moderate Risks of Breast Cancer. <i>Cancer Research</i> , 2017 , 77, 2789-2799	10.1	49
188	Common non-synonymous SNPs associated with breast cancer susceptibility: findings from the Breast Cancer Association Consortium. <i>Human Molecular Genetics</i> , 2014 , 23, 6096-111	5.6	48
187	Genome-wide association and transcriptome studies identify target genes and risk loci for breast cancer. <i>Nature Communications</i> , 2019 , 10, 1741	17.4	47
186	Hyperhomocysteinemia, methylenetetrahydrofolate reductase c.677C>T polymorphism and risk of cancer: cross-sectional and prospective studies and meta-analyses of 75,000 cases and 93,000 controls. <i>International Journal of Cancer</i> , 2011 , 128, 644-52	7.5	47

185	Smoking and Increased White and Red Blood Cells. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2019 , 39, 965-977	9.4	46
184	Shared heritability and functional enrichment across six solid cancers. <i>Nature Communications</i> , 2019 , 10, 431	17.4	45
183	Association between a germline OCA2 polymorphism at chromosome 15q13.1 and estrogen receptor-negative breast cancer survival. <i>Journal of the National Cancer Institute</i> , 2010 , 102, 650-62	9.7	45
182	Nicotinic acetylcholine receptor polymorphism, smoking behavior, and tobacco-related cancer and lung and cardiovascular diseases: a cohort study. <i>Journal of Clinical Oncology</i> , 2011 , 29, 2875-82	2.2	44
181	Shorter leukocyte telomere length is associated with higher risk of infections: a prospective study of 75,309 individuals from the general population. <i>Haematologica</i> , 2017 , 102, 1457-1465	6.6	43
180	Evidence that the 5p12 Variant rs10941679 Confers Susceptibility to Estrogen-Receptor-Positive Breast Cancer through FGF10 and MRPS30 Regulation. <i>American Journal of Human Genetics</i> , 2016 , 99, 903-911	11	43
179	CYP19A1 fine-mapping and Mendelian randomization: estradiol is causal for endometrial cancer. <i>Endocrine-Related Cancer</i> , 2016 , 23, 77-91	5.7	41
178	Plasma 25-hydroxyvitamin D and risk of non-melanoma and melanoma skin cancer: a prospective cohort study. <i>Journal of Investigative Dermatology</i> , 2013 , 133, 629-636	4.3	40
177	Identification of novel genetic markers of breast cancer survival. <i>Journal of the National Cancer Institute</i> , 2015 , 107,	9.7	38
176	Genetic predisposition to in situ and invasive lobular carcinoma of the breast. <i>PLoS Genetics</i> , 2014 , 10, e1004285	6	38
175	MicroRNA related polymorphisms and breast cancer risk. <i>PLoS ONE</i> , 2014 , 9, e109973	3.7	37
174	Fine-mapping identifies two additional breast cancer susceptibility loci at 9q31.2. <i>Human Molecular Genetics</i> , 2015 , 24, 2966-84	5.6	36
173	CHRNA3 genotype, nicotine dependence, lung function and disease in the general population. <i>European Respiratory Journal</i> , 2012 , 40, 1538-44	13.6	36
172	Identification and characterization of novel associations in the CASP8/ALS2CR12 region on chromosome 2 with breast cancer risk. <i>Human Molecular Genetics</i> , 2015 , 24, 285-98	5.6	35
171	DNA mismatch repair gene MSH6 implicated in determining age at natural menopause. <i>Human Molecular Genetics</i> , 2014 , 23, 2490-7	5.6	35
170	11q13 is a susceptibility locus for hormone receptor positive breast cancer. <i>Human Mutation</i> , 2012 , 33, 1123-32	4.7	33
169	Is high vitamin B12 status a cause of lung cancer?. <i>International Journal of Cancer</i> , 2019 , 145, 1499-1503	7.5	33
168	High tobacco consumption is causally associated with increased all-cause mortality in a general population sample of 55,568 individuals, but not with short telomeres: a Mendelian randomization study. <i>International Journal of Epidemiology</i> , 2014 , 43, 1473-83	7.8	32

167	Prostate-specific antigen and long-term prediction of prostate cancer incidence and mortality in the general population. <i>European Urology</i> , 2012 , 61, 865-74	10.2	32
166	Copy number variation in glutathione-S-transferase T1 and M1 predicts incidence and 5-year survival from prostate and bladder cancer, and incidence of corpus uteri cancer in the general population. <i>Pharmacogenomics Journal</i> , 2011 , 11, 292-9	3.5	32
165	Missense variants in ATM in 26,101 breast cancer cases and 29,842 controls. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2010 , 19, 2143-51	4	31
164	Genetic predisposition to ductal carcinoma in situ of the breast. <i>Breast Cancer Research</i> , 2016 , 18, 22	8.3	31
163	Elevated plasma YKL-40, lipids and lipoproteins, and ischemic vascular disease in the general population. <i>Stroke</i> , 2015 , 46, 329-35	6.7	29
162	Identification of susceptibility pathways for the role of chromosome 15q25.1 in modifying lung cancer risk. <i>Nature Communications</i> , 2018 , 9, 3221	17.4	29
161	Alcohol consumption and survival after a breast cancer diagnosis: a literature-based meta-analysis and collaborative analysis of data for 29,239 cases. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2014 , 23, 934-45	4	29
160	Genome-wide association study of germline variants and breast cancer-specific mortality. <i>British Journal of Cancer</i> , 2019 , 120, 647-657	8.7	28
159	A large-scale assessment of two-way SNP interactions in breast cancer susceptibility using 46,450 cases and 42,461 controls from the breast cancer association consortium. <i>Human Molecular Genetics</i> , 2014 , 23, 1934-46	5.6	28
158	Plasma testosterone in the general population, cancer prognosis and cancer risk: a prospective cohort study. <i>Annals of Oncology</i> , 2014 , 25, 712-718	10.3	28
157	Role of inflammatory marker YKL-40 in the diagnosis, prognosis and cause of cardiovascular and liver diseases. <i>Critical Reviews in Clinical Laboratory Sciences</i> , 2016 , 53, 396-408	9.4	28
156	Genetic insights into biological mechanisms governing human ovarian ageing. <i>Nature</i> , 2021 , 596, 393-397	30.4	28
155	Appraising the causal relevance of DNA methylation for risk of lung cancer. <i>International Journal of Epidemiology</i> , 2019 , 48, 1493-1504	7.8	27
154	Body mass index and breast cancer survival: a Mendelian randomization analysis. <i>International Journal of Epidemiology</i> , 2017 , 46, 1814-1822	7.8	27
153	Confirmation of 5p12 as a susceptibility locus for progesterone-receptor-positive, lower grade breast cancer. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2011 , 20, 2222-31	4	27
152	Missense polymorphisms in BRCA1 and BRCA2 and risk of breast and ovarian cancer. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2009 , 18, 2339-42	4	27
151	Cancer risk by combined levels of YKL-40 and C-reactive protein in the general population. <i>British Journal of Cancer</i> , 2012 , 106, 199-205	8.7	27
150	Polymorphisms in a Putative Enhancer at the 10q21.2 Breast Cancer Risk Locus Regulate NRBF2 Expression. <i>American Journal of Human Genetics</i> , 2015 , 97, 22-34	11	26

149	Multi-stage genome-wide association study identifies new susceptibility locus for testicular germ cell tumour on chromosome 3q25. <i>Human Molecular Genetics</i> , 2015 , 24, 1169-76	5.6	26
148	Reproductive profiles and risk of breast cancer subtypes: a multi-center case-only study. <i>Breast Cancer Research</i> , 2017 , 19, 119	8.3	26
147	Investigation of gene-environment interactions between 47 newly identified breast cancer susceptibility loci and environmental risk factors. <i>International Journal of Cancer</i> , 2015 , 136, E685-96	7.5	26
146	IgE and risk of cancer in 37 747 individuals from the general population. <i>Annals of Oncology</i> , 2015 , 26, 1784-90	10.3	26
145	Fine-scale mapping of 8q24 locus identifies multiple independent risk variants for breast cancer. <i>International Journal of Cancer</i> , 2016 , 139, 1303-1317	7.5	26
144	Patient survival and tumor characteristics associated with CHEK2:p.I157T - findings from the Breast Cancer Association Consortium. <i>Breast Cancer Research</i> , 2016 , 18, 98	8.3	26
143	Identification of independent association signals and putative functional variants for breast cancer risk through fine-scale mapping of the 12p11 locus. <i>Breast Cancer Research</i> , 2016 , 18, 64	8.3	25
142	An intergenic risk locus containing an enhancer deletion in 2q35 modulates breast cancer risk by deregulating IGFBP5 expression. <i>Human Molecular Genetics</i> , 2016 , 25, 3863-3876	5.6	24
141	Identification of new genetic susceptibility loci for breast cancer through consideration of gene-environment interactions. <i>Genetic Epidemiology</i> , 2014 , 38, 84-93	2.6	24
140	Common germline polymorphisms associated with breast cancer-specific survival. <i>Breast Cancer Research</i> , 2015 , 17, 58	8.3	24
139	Fine mapping of MHC region in lung cancer highlights independent susceptibility loci by ethnicity. <i>Nature Communications</i> , 2018 , 9, 3927	17.4	24
138	Arterial and venous thrombosis by high platelet count and high hematocrit: 108 521 individuals from the Copenhagen General Population Study. <i>Journal of Thrombosis and Haemostasis</i> , 2019 , 17, 1898-1911	15.4	23
137	Diagnostic and Prognostic Impact of Circulating YKL-40, IL-6, and CA 19.9 in Patients with Pancreatic Cancer. <i>PLoS ONE</i> , 2013 , 8, e67059	3.7	23
136	Secular trends in smoking in relation to prevalent and incident smoking-related disease: A prospective population-based study. <i>Tobacco Induced Diseases</i> , 2019 , 17, 72	3.2	23
135	The potential diagnostic value of serum microRNA signature in patients with pancreatic cancer. <i>International Journal of Cancer</i> , 2016 , 139, 2312-24	7.5	23
134	hypomethylation, lung function, lung function decline and respiratory symptoms. <i>European Respiratory Journal</i> , 2018 , 51,	13.6	22
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