# Stig Bojesen

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4340396/publications.pdf

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|---|---|---|---|
| 294 papers | 30,638 citations | 6124 83 h-index | 6686 161 g-index |
| 326 all docs | 326 docs citations | 326 times ranked | 36046 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Risk of ulcerative colitis and Crohnâ€™s disease in smokers lacks causal evidence. European Journal of Epidemiology, 2022, 37, 735-745. | 2.5 | 5 |
| 2 | Chronic lymphocytic leukaemia clones are detectable decades before diagnosis. British Journal of Haematology, 2022, 196, 784-787. | 1.2 | 3 |
| 3 | Rare germline copy number variants (CNVs) and breast cancer risk. Communications Biology, 2022, 5, 65. | 2.0 | 6 |
| 4 | AHRR (cg5575921) methylation safely improves specificity of lung cancer screening eligibility criteria: A cohort study. Cancer Epidemiology Biomarkers and Prevention, 2022, , cebp.1059.2021. | 1.1 | 10 |
| 5 | Epigenetic Regulation of F2RL3 Associates With Myocardial Infarction and Platelet Function. Circulation Research, 2022, 130, 384-400. | 2.0 | 10 |
| 6 | Genome-wide interaction analysis identified low-frequency variants with sex disparity in lung cancer risk. Human Molecular Genetics, 2022, 31, 2831-2843. | 1.4 | 4 |
| 7 | Geneâ€“gene interaction of AhRwith and within the Wntcascade affects susceptibility to lung cancer. European Journal of Medical Research, 2022, 27, 14. | 0.9 | 1 |
| 8 | Common variants in breast cancer risk loci predispose to distinct tumor subtypes. Breast Cancer Research, 2022, 24, 2. | 2.2 | 15 |
| 9 | Pathology of Tumors Associated With Pathogenic Germline Variants in 9 Breast Cancer Susceptibility Genes. JAMA Oncology, 2022, 8, e216744. | 3.4 | 51 |
| 10 | Development and validation of a model to predict incident chronic liver disease in the general population: The CLivD score. Journal of Hepatology, 2022, 77, 302-311. | 1.8 | 21 |
| 11 | Self-reported and genetically predicted coffee consumption and smoking in dementia: A Mendelian randomization study. Atherosclerosis, 2022, 348, 36-43. | 0.4 | 8 |
| 12 | Pre- and Perioperative Inflammatory Biomarkers in Older Patients Resected for Localized Colorectal Cancer: Associations with Complications and Prognosis. Cancers, 2022, 14, 161. | 1.7 | 9 |
| 13 | A Large-Scale Genome-Wide Gene-Gene Interaction Study of Lung Cancer Susceptibility in Europeans With a Trans-Ethnic Validation in Asians. Journal of Thoracic Oncology, 2022, 17, 974-990. | 0.5 | 18 |
| 14 | Breast cancer risks associated with missense variants in breast cancer susceptibility genes. Genome Medicine, 2022, 14, 51. | 3.6 | 19 |
| 15 | Allostatic load as predictor of mortality: a cohort study from Lolland-Falster, Denmark. BMJ Open, 2022, 12, e057136. | 0.8 | 5 |
| 16 | Associations of a breast cancer polygenic risk score with tumor characteristics and survival.. Journal of Clinical Oncology, 2022, 40, 563-563. | 0.8 | 1 |
| 17 | Combined Associations of a Polygenic Risk Score and Classical Risk Factors With Breast Cancer Risk. Journal of the National Cancer Institute, 2021, 113, 329-337. | 3.0 | 45 |
| 18 | Common Susceptibility Loci for Male Breast Cancer. Journal of the National Cancer Institute, 2021, 113, 453-461. | 3.0 | 12 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Evaluating the role of alcohol consumption in breast and ovarian cancer susceptibility using populationâ€based cohort studies and twoâ€sample Mendelian randomization analyses. International Journal of Cancer, 2021, 148, 1338-1350. | 2.3 | 9 |
| 20 | Integration of multiomic annotation data to prioritize and characterize inflammation and immuneâ€related risk variants in squamous cell lung cancer. Genetic Epidemiology, 2021, 45, 99-114. | 0.6 | 7 |
| 21 | Causal relationships between body mass index, smoking and lung cancer: Univariable and multivariable Mendelian randomization. International Journal of Cancer, 2021, 148, 1077-1086. | 2.3 | 73 |
| 22 | Myocardial Ischemia Induced by 5-Fluorouracil: A Prospective Electrocardiographic and Cardiac Biomarker Study. Oncologist, 2021, 26, e403-e413. | 1.9 | 18 |
| 23 | Comprehensive functional annotation of susceptibility variants identifies genetic heterogeneity between lung adenocarcinoma and squamous cell carcinoma. Frontiers of Medicine, 2021, 15, 275-291. | 1.5 | 21 |
| 24 | CYP3A7*1C allele: linking premenopausal oestrone and progesterone levels with risk of hormone receptor-positive breast cancers. British Journal of Cancer, 2021, 124, 842-854. | 2.9 | 5 |
| 25 | 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 |
| 26 | Assessing Lung Cancer Absolute Risk Trajectory Based on a Polygenic Risk Model. Cancer Research, 2021, 81, 1607-1615. | 0.4 | 50 |
| 27 | Reference intervals for 12 clinical laboratory tests in a Danish population: The Lolland-Falster Health Study. Scandinavian Journal of Clinical and Laboratory Investigation, 2021, 81, 104-111. | 0.6 | 2 |
| 28 | A case-only study to identify genetic modifiers of breast cancer risk for BRCA1/BRCA2 mutation carriers. Nature Communications, 2021, 12, 1078. | 5.8 | 19 |
| 29 | Breast Cancer Risk Genes â€" Association Analysis in More than 113,000 Women. New England Journal of Medicine, 2021, 384, 428-439. | 13.9 | 532 |
| 30 | Pre-treatment serum vitamin D deficiency is associated with increased inflammatory biomarkers and short overall survival in patients with pancreatic cancer. European Journal of Cancer, 2021, 144, 72-80. | 1.3 | 17 |
| 31 | Circulating Protein Biomarkers for Use in Pancreatic Ductal Adenocarcinoma Identification. Clinical Cancer Research, 2021, 27, 2592-2603. | 3.2 | 14 |
| 32 | Heterogeneous contributions of change in population distribution of body mass index to change in obesity and underweight. ELife, 2021, 10, . | 2.8 | 41 |
| 33 | Genome-wide association meta-analysis identifies pleiotropic risk loci for aerodigestive squamous cell cancers. PLoS Genetics, 2021, 17, e1009254. | 1.5 | 19 |
| 34 | Genetic predisposition to long telomeres is associated with increased mortality after melanoma: A study of 2101 melanoma patients from hospital clinics and the general population. Pigment Cell and Melanoma Research, 2021, 34, 946-954. | 1.5 | 4 |
| 35 | Gene-Environment Interactions Relevant to Estrogen and Risk of Breast Cancer: Can Gene-Environment Interactions Be Detected Only among Candidate SNPs from Genome-Wide Association Studies?. Cancers, 2021, 13, 2370. | 1.7 | 4 |
| 36 | Functional annotation of the 2q35 breast cancer risk locus implicates a structural variant in influencing activity of a long-range enhancer element. American Journal of Human Genetics, 2021, 108, 1190-1203. | 2.6 | 6 |

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|---|---|---|---|
| 37 | Association of germline genetic variants with breast cancer-specific survival in patient subgroups defined by clinic-pathological variables related to tumor biology and type of systemic treatment. Breast Cancer Research, 2021, 23, 86. | 2.2 | 7 |
| 38 | Mendelian randomisation study of smoking exposure in relation to breast cancer risk. British Journal of Cancer, 2021, 125, 1135-1145. | 2.9 | 9 |
| 39 | Genetic insights into biological mechanisms governing human ovarian ageing. Nature, 2021, 596, 393-397. | 13.7 | 183 |
| 40 | Detection and characterization of lung cancer using cell-free DNA fragmentomes. Nature Communications, 2021, 12, 5060. | 5.8 | 161 |
| 41 | Inflammatory Biomarker Score Identifies Patients with Six-Fold Increased Risk of One-Year Mortality after Pancreatic Cancer. Cancers, 2021, 13, 4599. | 1.7 | 5 |
| 42 | 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. Lancet, The, 2021, 398, 957-980. | 6.3 | 1,289 |
| 43 | Breast Cancer Risk Factors and Survival by Tumor Subtype: Pooled Analyses from the Breast Cancer Association Consortium. Cancer Epidemiology Biomarkers and Prevention, 2021, 30, 623-642. | 1.1 | 19 |
| 44 | Germline variants and breast cancer survival in patients with distant metastases at primary breast cancer diagnosis. Scientific Reports, 2021, 11, 19787. | 1.6 | 2 |
| 45 | Response to the letter entitled: Re: Pre-treatment serum vitamin D deficiency is associated with increased inflammatory biomarkers and short overall survival in patients with pancreatic cancer. European Journal of Cancer, 2021, 158, 248-250. | 1.3 | 0 |
| 46 | Observational and genetic studies of short telomeres and Alzheimerâ€™s disease in 67,000 and 152,000 individuals: a Mendelian randomization study. European Journal of Epidemiology, 2020, 35, 147-156. | 2.5 | 36 |
| 47 | Clinical value of serum hyaluronan and propeptide of type III collagen in patients with pancreatic cancer. International Journal of Cancer, 2020, 146, 2913-2922. | 2.3 | 41 |
| 48 | Transcriptomeâ€wide association study reveals candidate causal genes for lung cancer. International Journal of Cancer, 2020, 146, 1862-1878. | 2.3 | 33 |
| 49 | Prognostic Value of Combined Detection of Serum IL6, YKL-40, and C-reactive Protein in Patients with Unresectable Pancreatic Cancer. Cancer Epidemiology Biomarkers and Prevention, 2020, 29, 176-184. | 1.1 | 12 |
| 50 | Genomeâ€wide association study of INDELs identified four novel susceptibility loci associated with lung cancer risk. International Journal of Cancer, 2020, 146, 2855-2864. | 2.3 | 7 |
| 51 | Fine-mapping of 150 breast cancer risk regions identifies 191 likely target genes. Nature Genetics, 2020, 52, 56-73. | 9.4 | 120 |
| 52 | Immune-mediated genetic pathways resulting in pulmonary function impairment increase lung cancer susceptibility. Nature Communications, 2020, 11, 27. | 5.8 | 23 |
| 53 | Smoking, blood cells and myeloproliferative neoplasms: metaâ€analysis and Mendelian randomization of 2Â·3 million people. British Journal of Haematology, 2020, 189, 323-334. | 1.2 | 27 |
| 54 | Low high-density lipoprotein and increased risk of several cancers: 2 population-based cohort studies including 116,728 individuals. Journal of Hematology and Oncology, 2020, 13, 129. | 6.9 | 46 |

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|---|---|---|---|
| 55 | AHRR hypomethylation as an epigenetic marker of smoking history predicts risk of myocardial infarction in former smokers. Atherosclerosis, 2020, 312, 8-15. | 0.4 | 7 |
| 56 | Loss-of-function polymorphism in IL6R reduces risk of JAK2V617F somatic mutation and myeloproliferative neoplasm: A Mendelian randomization study. EClinicalMedicine, 2020, 21, 100280. | 3.2 | 19 |
| 57 | Breast cancer risk factors and their effects on survival: a Mendelian randomisation study. BMC Medicine, 2020, 18, 327. | 2.3 | 40 |
| 58 | Burden of prediabetes, undiagnosed, and poorly or potentially sub-controlled diabetes: Lolland-Falster health study. BMC Public Health, 2020, 20, 1711. | 1.2 | 8 |
| 59 | Breast Cancer Polygenic Risk Score and Contralateral Breast Cancer Risk. American Journal of Human Genetics, 2020, 107, 837-848. | 2.6 | 39 |
| 60 | 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. Lancet, The, 2020, 396, 1511-1524. | 6.3 | 219 |
| 61 | 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 |
| 62 | Protein-altering germline mutations implicate novel genes related to lung cancer development. Nature Communications, 2020, 11, 2220. | 5.8 | 31 |
| 63 | Germline HOXB13 mutations p.G84E and p.R217C do not confer an increased breast cancer risk. Scientific Reports, 2020, 10, 9688. | 1.6 | 2 |
| 64 | Reply to: Clinical impact of high platelet count and high hematocrit, by Marc Sorigue. Journal of Thrombosis and Haemostasis, 2020, 18, 522-523. | 1.9 | 0 |
| 65 | Two-fold risk of pneumonia and respiratory mortality in individuals with myeloproliferative neoplasm: A population-based cohort study. EClinicalMedicine, 2020, 21, 100295. | 3.2 | 5 |
| 66 | Tocilizumab and soluble interleukin-6 receptor in JAK2V617F somatic mutation and myeloproliferative neoplasm. EClinicalMedicine, 2020, 22, 100337. | 3.2 | 2 |
| 67 | Transcriptomeâ€wide association study of breast cancer risk by estrogenâ€receptor status. Genetic Epidemiology, 2020, 44, 442-468. | 0.6 | 32 |
| 68 | Incidental lymphopenia and mortality: a prospective cohort study. Cmaj, 2020, 192, E25-E33. | 0.9 | 34 |
| 69 | A network analysis to identify mediators of germline-driven differences in breast cancer prognosis. Nature Communications, 2020, 11, 312. | 5.8 | 30 |
| 70 | Elevated plasma YKL-40 and risk of infectious disease: a prospective study of 94665 individuals from the general population. Clinical Microbiology and Infection, 2020, 26, 1411.e1-1411.e9. | 2.8 | 8 |
| 71 | Prediction of contralateral breast cancer: external validation of risk calculators in 20 international cohorts. Breast Cancer Research and Treatment, 2020, 181, 423-434. | 1.1 | 14 |
| 72 | Association Analysis of Driver Geneâ€"Related Genetic Variants Identified Novel Lung Cancer Susceptibility Loci with 20,871 Lung Cancer Cases and 15,971 Controls. Cancer Epidemiology Biomarkers and Prevention, 2020, 29, 1423-1429. | 1.1 | 6 |

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|---|---|---|---|
| 73 | Arterial and venous thrombosis by high platelet count and high hematocrit: 108Â521 individuals from the Copenhagen General Population Study. Journal of Thrombosis and Haemostasis, 2019, 17, 1898-1911. | 1.9 | 46 |
| 74 | Lung Cancer Risk in Never-Smokers of European Descent is Associated With Genetic Variation in the 5p15.33 TERT-CLPTM1Ll Region. Journal of Thoracic Oncology, 2019, 14, 1360-1369. | 0.5 | 27 |
| 75 | Investigation of Leukocyte Telomere Length and Genetic Variants in Chromosome 5p15.33 as Prognostic Markers in Lung Cancer. Cancer Epidemiology Biomarkers and Prevention, 2019, 28, 1228-1237. | 1.1 | 11 |
| 76 | Measured and genetically predicted plasma YKL-40 levels and melanoma mortality. European Journal of Cancer, 2019, 121, 74-84. | 1.3 | 3 |
| 77 | The FANCM:p.Arg658* truncating variant is associated with risk of triple-negative breast cancer. Npj Breast Cancer, 2019, 5, 38. | 2.3 | 28 |
| 78 | Two truncating variants in FANCC and breast cancer risk. Scientific Reports, 2019, 9, 12524. | 1.6 | 5 |
| 79 | Appraising the causal relevance of DNA methylation for risk of lung cancer. International Journal of Epidemiology, 2019, 48, 1493-1504. | 0.9 | 53 |
| 80 | Shared heritability and functional enrichment across six solid cancers. Nature Communications, 2019, 10, 431. | 5.8 | 88 |
| 81 | Germline BRCA2 K3326X and CHEK2 I157T mutations increase risk for sporadic pancreatic ductal adenocarcinoma. International Journal of Cancer, 2019, 145, 686-693. | 2.3 | 20 |
| 82 | Elevated Platelet Count Appears to Be Causally Associated with Increased Risk of Lung Cancer: A Mendelian Randomization Analysis. Cancer Epidemiology Biomarkers and Prevention, 2019, 28, 935-942. | 1.1 | 21 |
| 83 | Physical activity and risk of instant and 28-day case-fatality in myocardial infarction. PLoS ONE, 2019, 14, e0217398. | 1.1 | 6 |
| 84 | Smoking does not accelerate leucocyte telomere attrition: a meta-analysis of 18 longitudinal cohorts. Royal Society Open Science, 2019, 6, 190420. | 1.1 | 33 |
| 85 | Genetic interaction analysis among oncogenesis-related genes revealed novel genes and networks in lung cancer development. Oncotarget, 2019, 10, 1760-1774. | 0.8 | 25 |
| 86 | Genome-wide association and transcriptome studies identify target genes and risk loci for breast cancer. Nature Communications, 2019, 10, 1741. | 5.8 | 90 |
| 87 | Smoking and Increased White and Red Blood Cells. Arteriosclerosis, Thrombosis, and Vascular Biology, 2019, 39, 965-977. | 1.1 | 98 |
| 88 | Genome-wide association study of germline variants and breast cancer-specific mortality. British Journal of Cancer, 2019, 120, 647-657. | 2.9 | 52 |
| 89 | AHRR (cg05575921) methylation extent of leukocyte DNA and lung cancer survival. PLoS ONE, 2019, 14, e0211745. | 1.1 | 9 |
| 90 | Prediction and clinical utility of a contralateral breast cancer risk model. Breast Cancer Research, 2019, 21, 144. | 2.2 | 24 |

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|---|---|---|---|
| 91 | JAK2-tree: a simple CBC-based decision rule to guide appropriate JAK2 V617F mutation testing. Journal of Clinical Pathology, 2019, 72, 172-176. | 1.0 | 8 |
| 92 | Polygenic Risk Scores for Prediction of Breast Cancer and Breast Cancer Subtypes. American Journal of Human Genetics, 2019, 104, 21-34. | 2.6 | 711 |
| 93 | Bone marrow mononuclear cell telomere length in acute myeloid leukaemia and highâ€risk myelodysplastic syndrome. European Journal of Haematology, 2019, 102, 218-226. | 1.1 | 6 |
| 94 | Systematic analyses of regulatory variants in DNase I hypersensitive sites identified two novel lung cancer susceptibility loci. Carcinogenesis, 2019, 40, 432-440. | 1.3 | 5 |
| 95 | Mononuclear Cell Telomere Attrition Is Associated with Overall Survival after Nonmyeloablative Allogeneic Hematopoietic Cell Transplantation for Hematologic Malignancies. Biology of Blood and Marrow Transplantation, 2019, 25, 496-504. | 2.0 | 4 |
| 96 | Mendelian Randomization and mediation analysis of leukocyte telomere length and risk of lung and head and neck cancers. International Journal of Epidemiology, 2019, 48, 751-766. | 0.9 | 32 |
| 97 | Is high vitamin B12 status a cause of lung cancer?. International Journal of Cancer, 2019, 145, 1499-1503. | 2.3 | 58 |
| 98 | 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 |
| 99 | Secular trends in smoking in relation to prevalent and incident smoking-related disease: A prospective population-based study. Tobacco Induced Diseases, 2019, 17, 72. | 0.3 | 39 |
| 100 | The BRCA2 c.68-7TÂ>ÂA variant is not pathogenic: A model for clinical calibration of spliceogenicity. Human Mutation, 2018, 39, 729-741. | 1.1 | 19 |
| 101 | Is smoking heaviness causally associated with alcohol use? A Mendelian randomization study in four European cohorts. International Journal of Epidemiology, 2018, 47, 1098-1105. | 0.9 | 17 |
| 102 | Genome-wide interaction study of smoking behavior and non-small cell lung cancer risk in Caucasian population. Carcinogenesis, 2018, 39, 336-346. | 1.3 | 29 |
| 103 | AHRR hypomethylation, lung function, lung function decline and respiratory symptoms. European Respiratory Journal, 2018, 51, 1701512. | 3.1 | 35 |
| 104 | Joint associations of a polygenic risk score and environmental risk factors for breast cancer in the Breast Cancer Association Consortium. International Journal of Epidemiology, 2018, 47, 526-536. | 0.9 | 88 |
| 105 | Serum Biomarker Signature-Based Liquid Biopsy for Diagnosis of Early-Stage Pancreatic Cancer. Journal of Clinical Oncology, 2018, 36, 2887-2894. | 0.8 | 108 |
| 106 | Fine mapping of MHC region in lung cancer highlights independent susceptibility loci by ethnicity. Nature Communications, 2018, 9, 3927. | 5.8 | 43 |
| 107 | Lymphopenia and risk of infection and infection-related death in 98,344 individuals from a prospective Danish population-based study. PLoS Medicine, 2018, 15, e1002685. | 3.9 | 119 |
| 108 | Genetic modifiers of radon-induced lung cancer risk: a genome-wide interaction study in former uranium miners. International Archives of Occupational and Environmental Health, 2018, 91, 937-950. | 1.1 | 27 |

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|---|---|---|---|
| 109 | Identification of susceptibility pathways for the role of chromosome 15q25.1 in modifying lung cancer risk. Nature Communications, 2018, 9, 3221. | 5.8 | 60 |
| 110 | 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 |
| 111 | BRCA2 Hypomorphic Missense Variants Confer Moderate Risks of Breast Cancer. Cancer Research, 2017, 77, 2789-2799. | 0.4 | 75 |
| 112 | Genomic analyses identify hundreds of variants associated with age at menarche and support a role for puberty timing in cancer risk. Nature Genetics, 2017, 49, 834-841. | 9.4 | 426 |
| 113 | TP53 Arg72Pro, mortality after cancer, and all-cause mortality in 105,200 individuals. Scientific Reports, 2017, 7, 336. | 1.6 | 8 |
| 114 | Shorter leukocyte telomere length is associated with higher risk of infections: a prospective study of 75,309 individuals from the general population. Haematologica, 2017, 102, 1457-1465. | 1.7 | 63 |
| 115 | Asthma, other atopic conditions and risk of infections in 105 519 general population never and ever smokers. Journal of Internal Medicine, 2017, 282, 254-267. | 2.7 | 25 |
| 116 | Large-scale association analysis identifies new lung cancer susceptibility loci and heterogeneity in genetic susceptibility across histological subtypes. Nature Genetics, 2017, 49, 1126-1132. | 9.4 | 472 |
| 117 | AHRR(cg05575921) hypomethylation marks smoking behaviour, morbidity and mortality. Thorax, 2017, 72, 646-653. | 2.7 | 147 |
| 118 | Telomere length and depression: Prospective cohort study and Mendelian randomisation study in 67 306 individuals. British Journal of Psychiatry, 2017, 210, 31-38. | 1.7 | 26 |
| 119 | Kringle IV Type 2, Not Low Lipoprotein(a), as a Cause of Diabetes: A Novel Genetic Approach Using SNPs Associated Selectively with Lipoprotein(a) Concentrations or with Kringle IV Type 2 Repeats. Clinical Chemistry, 2017, 63, 1866-1876. | 1.5 | 28 |
| 120 | Association analysis identifies 65 new breast cancer risk loci. Nature, 2017, 551, 92-94. | 13.7 | 1,099 |
| 121 | Identification of ten variants associated with risk of estrogen-receptor-negative breast cancer. Nature Genetics, 2017, 49, 1767-1778. | 9.4 | 289 |
| 122 | Geneâ€“environment interactions involving functional variants: Results from the Breast Cancer Association Consortium. International Journal of Cancer, 2017, 141, 1830-1840. | 2.3 | 20 |
| 123 | Common breast cancer risk alleles and risk assessment: a study on 35Â441 individuals from the Danish general population. Annals of Oncology, 2017, 28, 175-181. | 0.6 | 6 |
| 124 | Genetic modifiers of CHEK2*1100delC-associated breast cancer risk. Genetics in Medicine, 2017, 19, 599-603. | 1.1 | 67 |
| 125 | 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 |
| 126 | Body mass index and breast cancer survival: a Mendelian randomization analysis. International Journal of Epidemiology, 2017, 46, 1814-1822. | 0.9 | 45 |

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|---|---|---|---|
| 127 | Pleiotropy of genetic variants on obesity and smoking phenotypes: Results from the Oncoarray Project of The International Lung Cancer Consortium. PLoS ONE, 2017, 12, e0185660. | 1.1 | 11 |
| 128 | Reproductive profiles and risk of breast cancer subtypes: a multi-center case-only study. Breast Cancer Research, 2017, 19, 119. | 2.2 | 43 |
| 129 | The prognostic value of serum CA 19-9 in patients with metastatic colorectal cancer.. Journal of Clinical Oncology, 2017, 35, e15131-e15131. | 0.8 | 1 |
| 130 | Obesity, metabolic factors and risk of different histological types of lung cancer: A Mendelian randomization study. PLoS ONE, 2017, 12, e0177875. | 1.1 | 79 |
| 131 | PHIP- a novel candidate breast cancer susceptibility locus on 6q14.1. Oncotarget, 2017, 8, 102769-102782. | 0.8 | 9 |
| 132 | The prognostic value of serum IL-6 and YKL-40 in patients with metastatic colorectal cancer.. Journal of Clinical Oncology, 2017, 35, e15060-e15060. | 0.8 | 0 |
| 133 | Association of breast cancer risk with genetic variants showing differential allelic expression: Identification of a novel breast cancer susceptibility locus at 4q21. Oncotarget, 2016, 7, 80140-80163. | 0.8 | 31 |
| 134 | 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 |
| 135 | Fine-Mapping of the 1p11.2 Breast Cancer Susceptibility Locus. PLoS ONE, 2016, 11, e0160316. | 1.1 | 12 |
| 136 | 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 |
| 137 | Inflammatory biomarkers and risk of cancer in 84,000 individuals from the general population. International Journal of Cancer, 2016, 139, 1493-1500. | 2.3 | 73 |
| 138 | PALB2,CHEK2andATMrare variants and cancer risk: data from COGS. Journal of Medical Genetics, 2016, 53, 800-811. | 1.5 | 174 |
| 139 | Patient survival and tumor characteristics associated with CHEK2:p.I157T â€“ findings from the Breast Cancer Association Consortium. Breast Cancer Research, 2016, 18, 98. | 2.2 | 39 |
| 140 | Identification of independent association signals and putative functional variants for breast cancer risk through fine-scale mapping of the 12p11 locus. Breast Cancer Research, 2016, 18, 64. | 2.2 | 31 |
| 141 | Role of inflammatory marker YKL-40 in the diagnosis, prognosis and cause of cardiovascular and liver diseases. Critical Reviews in Clinical Laboratory Sciences, 2016, 53, 396-408. | 2.7 | 50 |
| 142 | Genetic predisposition to ductal carcinoma in situ of the breast. Breast Cancer Research, 2016, 18, 22. | 2.2 | 43 |
| 143 | Association of genetic susceptibility variants for type 2 diabetes with breast cancer risk in women of European ancestry. Cancer Causes and Control, 2016, 27, 679-693. | 0.8 | 21 |
| 144 | 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 |

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|---|---|---|---|
| 145 | The potential diagnostic value of serum microRNA signature in patients with pancreatic cancer. International Journal of Cancer, 2016, 139, 2312-2324. | 2.3 | 33 |
| 146 | An intergenic risk locus containing an enhancer deletion in 2q35 modulates breast cancer risk by deregulating IGFBP5 expression. Human Molecular Genetics, 2016, 25, 3863-3876. | 1.4 | 33 |
| 147 | Long telomeres and cancer risk among 95â€‰568 individuals from the general population. International Journal of Epidemiology, 2016, 45, 1634-1643. | 0.9 | 90 |
| 148 | rs2735383, located at a microRNA binding site in the 3â€™UTR of NBS1, is not associated with breast cancer risk. Scientific Reports, 2016, 6, 36874. | 1.6 | 2 |
| 149 | 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 |
| 150 | Identification of four novel susceptibility loci for oestrogen receptor negative breast cancer. Nature Communications, 2016, 7, 11375. | 5.8 | 93 |
| 151 | Functional mechanisms underlying pleiotropic risk alleles at the 19p13.1 breastâ€“ovarian cancer susceptibility locus. Nature Communications, 2016, 7, 12675. | 5.8 | 78 |
| 152 | Fine scale mapping of the 17q22 breast cancer locus using dense SNPs, genotyped within the Collaborative Oncological Gene-Environment Study (COGs). Scientific Reports, 2016, 6, 32512. | 1.6 | 19 |
| 153 | Short Telomere Length and Ischemic Heart Disease: Observational and Genetic Studies in 290 022 Individuals. Clinical Chemistry, 2016, 62, 1140-1149. | 1.5 | 93 |
| 154 | Age- and Tumor Subtypeâ€“Specific Breast Cancer Risk Estimates for CHEK2*1100delC Carriers. Journal of Clinical Oncology, 2016, 34, 2750-2760. | 0.8 | 152 |
| 155 | CYP19A1 fine-mapping and Mendelian randomization: estradiol is causal for endometrial cancer. Endocrine-Related Cancer, 2016, 23, 77-91. | 1.6 | 62 |
| 156 | Observationally and Genetically High YKL-40 and Risk of Venous Thromboembolism in the General Population. Arteriosclerosis, Thrombosis, and Vascular Biology, 2016, 36, 1030-1036. | 1.1 | 18 |
| 157 | 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. Human Molecular Genetics, 2016, 25, 2256-2268. | 1.4 | 106 |
| 158 | Increased Risk for Other Cancers in Addition to Breast Cancer for CHEK2*1100delC Heterozygotes Estimated From the Copenhagen General Population Study. Journal of Clinical Oncology, 2016, 34, 1208-1216. | 0.8 | 97 |
| 159 | No evidence that protein truncating variants inBRIP1are associated with breast cancer risk: implications for gene panel testing. Journal of Medical Genetics, 2016, 53, 298-309. | 1.5 | 94 |
| 160 | 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 |
| 161 | Genetic variation in the immunosuppression pathway genes and breast cancer susceptibility: a pooled analysis of 42,510 cases and 40,577 controls from the Breast Cancer Association Consortium. Human Genetics, 2016, 135, 137-154. | 1.8 | 8 |
| 162 | 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 |

| # | Article | IF | Citations |
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