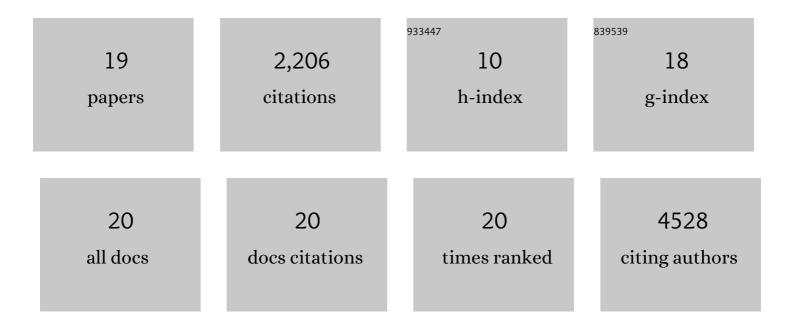
Carolina Ellberg

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Association analysis identifies 65 new breast cancer risk loci. Nature, 2017, 551, 92-94.	27.8	1,099
2	Polygenic Risk Scores for Prediction of Breast Cancer and Breast Cancer Subtypes. American Journal of Human Genetics, 2019, 104, 21-34.	6.2	711
3	Genome-wide association and transcriptome studies identify target genes and risk loci for breast cancer. Nature Communications, 2019, 10, 1741.	12.8	90
4	Shared heritability and functional enrichment across six solid cancers. Nature Communications, 2019, 10, 431.	12.8	88
5	Genome-wide association study of germline variants and breast cancer-specific mortality. British Journal of Cancer, 2019, 120, 647-657.	6.4	52
6	Oral contraceptive use and ovarian cancer risk for BRCA1/2 mutation carriers: an international cohort study. American Journal of Obstetrics and Gynecology, 2021, 225, 51.e1-51.e17.	1.3	34
7	Transcriptomeâ€wide association study of breast cancer risk by estrogenâ€receptor status. Genetic Epidemiology, 2020, 44, 442-468.	1.3	32
8	A network analysis to identify mediators of germline-driven differences in breast cancer prognosis. Nature Communications, 2020, 11, 312.	12.8	30
9	Accuracy of self-reported family history of cancer, mutation status and tumor characteristics in patients with early onset breast cancer. Acta Oncológica, 2018, 57, 595-603.	1.8	19
10	Association of a history of autoimmunity or hypersensitivity with overall survival in breast cancer Journal of Clinical Oncology, 2012, 30, 1594-1594.	1.6	11
11	Current smoking is associated with a larger waist circumference and a more androgenic profile in young healthy women from high-risk breast cancer families. Cancer Causes and Control, 2018, 29, 243-251.	1.8	7
12	Breast cancer patients with lobular cancer more commonly have a father than a mother diagnosed with cancer. BMC Cancer, 2011, 11, 497.	2.6	6
13	Impact of a paternal origin of germline <i>BRCA1</i> /2 mutations on the age at breast and ovarian cancer diagnosis in a Southern Swedish cohort. Genes Chromosomes and Cancer, 2015, 54, 39-50.	2.8	6
14	Can a phenotype for recessive inheritance in breast cancer be defined?. Familial Cancer, 2010, 9, 525-530.	1.9	4
15	Autoimmune diseases and hypersensitivities improve the prognosis in ER-negative breast cancer. SpringerPlus, 2013, 2, 357.	1.2	4
16	Breast cancer and spider telangiectasias at diagnosis and its relation to histopathology and prognosis: a population-based study. Breast Cancer Research and Treatment, 2012, 131, 177-186.	2.5	1
17	Variations in the Referral Pattern for Genetic Counseling of Patients with Early-Onset Breast Cancer: A Population-Based Study in Southern Sweden. Public Health Genomics, 2020, 23, 100-109.	1.0	1
18	Genetic testing in women with early-onset breast cancer: a Traceback pilot study. Breast Cancer Research and Treatment, 2021, 190, 307-315.	2.5	1

#	Article	IF	CITATIONS
19	Abstract A127: Parental influence on breast cancer penetrance inBRCA1/2mutation carriers: Impact of oral contraceptive use before age 20 years. , 2013, , .		0