Kari Hemminki

List of Publications by Citations

587 papers	23,689 citations	67 h-index	132 g-index
607 ext. papers	26,602 ext. citations	5.8 avg, IF	6.86 L-index

#	Paper	IF	Citations
587	Environmental and heritable factors in the causation of cancer--analyses of cohorts of twins from Sweden, Denmark, and Finland. *New England Journal of Medicine*, **2000**, 343, 78-85	59.2	3019
586	TERT promoter mutations in familial and sporadic melanoma. *Science*, **2013**, 339, 959-61	33.3	1261
585	Genome-wide association study identifies five susceptibility loci for glioma. *Nature Genetics*, **2009**, 41, 899-904	36.3	640
584	IPCS guidelines for the monitoring of genotoxic effects of carcinogens in humans. International Programme on Chemical Safety. *Mutation Research - Reviews in Mutation Research*, **2000**, 463, 111-72	7	536
583	Sequence variants at the TERT-CLPTM1L locus associate with many cancer types. *Nature Genetics*, **2009**, 41, 221-7	36.3	509
582	A genome-wide association study identifies colorectal cancer susceptibility loci on chromosomes 10p14 and 8q23.3. *Nature Genetics*, **2008**, 40, 623-30	36.3	463
581	Patterns of metastasis in colon and rectal cancer. *Scientific Reports*, **2016**, 6, 29765	4.9	409
580	Environmental and heritable causes of cancer among 9.6 million individuals in the Swedish Family-Cancer Database. *International Journal of Cancer*, **2002**, 99, 260-6	7.5	364
579	Polymorphisms in DNA repair and metabolic genes in bladder cancer. *Carcinogenesis*, **2004**, 25, 729-34	4.6	257
578	Variation in CDKN2A at 9p21.3 influences childhood acute lymphoblastic leukemia risk. *Nature Genetics*, **2010**, 42, 492-4	36.3	214
577	Vascular endothelial growth factor polymorphisms in relation to breast cancer development and prognosis. *Clinical Cancer Research*, **2005**, 11, 3647-53	12.9	198
576	Modification of cancer risks in offspring by sibling and parental cancers from 2,112,616 nuclear families. *International Journal of Cancer*, **2001**, 92, 144-150	7.5	182
575	The XPD variant alleles are associated with increased aromatic DNA adduct level and lung cancer risk. *Carcinogenesis*, **2002**, 23, 599-603	4.6	175
574	Familial risk of lymphoproliferative tumors in families of patients with chronic lymphocytic leukemia: results from the Swedish Family-Cancer Database. *Blood*, **2004**, 104, 1850-4	2.2	173
573	Risk of second cancer among women with breast cancer. *International Journal of Cancer*, **2006**, 118, 2285-92	7.9	170
572	TERT promoter mutations in cancer development. *Current Opinion in Genetics and Development*, **2014**, 24, 30-7	4.9	167
571	A genome-wide association study of Hodgkin's lymphoma identifies new susceptibility loci at 2p16.1 (REL), 8q24.21 and 10p14 (GATA3). *Nature Genetics*, **2010**, 42, 1126-1130	36.3	158

570	Metastatic spread in patients with gastric cancer. *Oncotarget*, **2016**, 7, 52307-52316	3.3	158
569	Familial associations of rheumatoid arthritis with autoimmune diseases and related conditions. *Arthritis and Rheumatism*, **2009**, 60, 661-8		153
568	Organic solvents and cancer. *Cancer Causes and Control*, **1997**, 8, 406-19	2.8	153
567	Association of DNA repair polymorphisms with DNA repair functional outcomes in healthy human subjects. *Carcinogenesis*, **2007**, 28, 657-64	4.6	147
566	Autoimmunity and susceptibility to Hodgkin lymphoma: a population-based case-control study in Scandinavia. *Journal of the National Cancer Institute*, **2006**, 98, 1321-30	9.7	146
565	Cancer risks in first-generation immigrants to Sweden. *International Journal of Cancer*, **2002**, 99, 218-28	7.5	141
564	Familial aggregation of Hodgkin lymphoma and related tumors. *Cancer*, **2004**, 100, 1902-8	6.4	140
563	The balance between heritable and environmental aetiology of human disease. *Nature Reviews Genetics*, **2006**, 7, 958-65	30.1	137
562	Risk of cancer following hospitalization for type 2 diabetes. *Oncologist*, **2010**, 15, 548-55	5.7	136
561	The epidemiology of metastases in neuroendocrine tumors. *International Journal of Cancer*, **2016**, 139, 2679-2686	7.5	132
560	Telomerase reverse transcriptase promoter mutations in primary cutaneous melanoma. *Nature Communications*, **2014**, 5, 3401	17.4	132
559	Second primary neoplasms in 633,964 cancer patients in Sweden, 1958-1996. *International Journal of Cancer*, **2001**, 93, 155-61	7.5	132
558	Familial risks for nonmedullary thyroid cancer. *Journal of Clinical Endocrinology and Metabolism*, **2005**, 90, 5747-53	5.6	131
557	Chromosome 7p11.2 (EGFR) variation influences glioma risk. *Human Molecular Genetics*, **2011**, 20, 2897-904	5.8	129
556	TERT promoter mutations: a novel independent prognostic factor in primary glioblastomas. *Neuro-Oncology*, **2015**, 17, 45-52	1	123
555	Common variation at 3p22.1 and 7p15.3 influences multiple myeloma risk. *Nature Genetics*, **2011**, 44, 58-61	36.3	122
554	Low expression of hexokinase-2 is associated with false-negative FDG-positron emission tomography in multiple myeloma. *Blood*, **2017**, 130, 30-34	2.2	120
553	Common variation at 3q26.2, 6p21.33, 17p11.2 and 22q13.1 influences multiple myeloma risk. *Nature Genetics*, **2013**, 45, 1221-1225	36.3	119

552	Single nucleotide polymorphisms in breast cancer. *Oncology Reports*, **2004**, 11, 917-22	3.5	113
551	Familial characteristics of autoimmune and hematologic disorders in 8,406 multiple myeloma patients: a population-based case-control study. *International Journal of Cancer*, **2006**, 118, 3095-8	7.5	109
550	Cancer risks in second-generation immigrants to Sweden. *International Journal of Cancer*, **2002**, 99, 229-37	7.5	108
549	Verification of the susceptibility loci on 7p12.2, 10q21.2, and 14q11.2 in precursor B-cell acute lymphoblastic leukemia of childhood. *Blood*, **2010**, 115, 1765-7	2.2	107
548	Familial and second primary pancreatic cancers: a nationwide epidemiologic study from Sweden. *International Journal of Cancer*, **2003**, 103, 525-30	7.5	107
547	Familial carcinoid tumors and subsequent cancers: a nation-wide epidemiologic study from Sweden. *International Journal of Cancer*, **2001**, 94, 444-8	7.5	103
546	Genome-wide association study identifies multiple susceptibility loci for multiple myeloma. *Nature Communications*, **2016**, 7, 12050	17.4	101
545	Familial risks for type 2 diabetes in Sweden. *Diabetes Care*, **2010**, 33, 293-7	14.6	99
544	The epidemiology of Graves' disease: evidence of a genetic and an environmental contribution. *Journal of Autoimmunity*, **2010**, 34, J307-13	15.5	99
543	Parental Age As a Risk Factor of Childhood Leukemia and Brain Cancer in Offspring. *Epidemiology*, **1999**, 10, 271-275	3.1	98
542	Risk factors and age-incidence relationships for contralateral breast cancer. *International Journal of Cancer*, **2000**, 88, 998-1002	7.5	97
541	Familial risk for non-Hodgkin lymphoma and other lymphoproliferative malignancies by histopathologic subtype: the Swedish Family-Cancer Database. *Blood*, **2005**, 106, 668-72	2.2	89
540	Familial risk of cancer: data for clinical counseling and cancer genetics. *International Journal of Cancer*, **2004**, 108, 109-14	7.5	88
539	The Swedish Family-Cancer Database 2009: prospects for histology-specific and immigrant studies. *International Journal of Cancer*, **2010**, 126, 2259-67	7.5	85
538	Comparability of cancer identification among Death Registry, Cancer Registry and Hospital Discharge Registry. *International Journal of Cancer*, **2012**, 131, 2085-93	7.5	80
537	The CCND1 c.870G>A polymorphism is a risk factor for t(11;14)(q13;q32) multiple myeloma. *Nature Genetics*, **2013**, 45, 522-525	36.3	79
536	Somatic alterations in the melanoma genome: a high-resolution array-based comparative genomic hybridization study. *Genes Chromosomes and Cancer*, **2010**, 49, 733-45	5	79
535	TERT promoter mutations in melanoma survival. *International Journal of Cancer*, **2016**, 139, 75-84	7.5	79

534	Socioeconomic factors in cancer in Sweden. *International Journal of Cancer*, **2003**, 105, 692-700	7.5	78
533	Copy number variant in the candidate tumor suppressor gene MTUS1 and familial breast cancer risk. *Carcinogenesis*, **2007**, 28, 1442-5	4.6	77
532	Familial risk of cancer by site and histopathology. *International Journal of Cancer*, **2003**, 103, 105-9	7.5	77
531	Age-specific familial risks in common cancers of the offspring. *International Journal of Cancer*, **1998**, 78, 172-5	7.5	74
530	Risk of second malignant neoplasms after childhood leukemia and lymphoma: an international study. *Journal of the National Cancer Institute*, **2007**, 99, 790-800	9.7	73
529	Familial risks in testicular cancer as aetiological clues. *Journal of Developmental and Physical Disabilities*, **2006**, 29, 205-10		72
528	Risk of inflammatory bowel disease in first- and second-generation immigrants in Sweden: a nationwide follow-up study. *Inflammatory Bowel Diseases*, **2011**, 17, 1784-91	4.5	71
527	Association of HLA-DRB1, interleukin-6 and cyclin D1 polymorphisms with cervical cancer in the Swedish population--a candidate gene approach. *International Journal of Cancer*, **2009**, 125, 1851-8	7.5	71
526	Markers of individual susceptibility and DNA repair rate in workers exposed to xenobiotics in a tire plant. *Environmental and Molecular Mutagenesis*, **2004**, 44, 283-92	3.2	70
525	Single nucleotide polymorphisms in DNA repair genes and basal cell carcinoma of skin. *Carcinogenesis*, **2006**, 27, 1676-81	4.6	70
524	Promoter polymorphisms in matrix metalloproteinases and their inhibitors: few associations with breast cancer susceptibility and progression. *Breast Cancer Research and Treatment*, **2007**, 103, 61-9	4.4	68
523	Single nucleotide polymorphisms in the XPG gene: determination of role in DNA repair and breast cancer risk. *International Journal of Cancer*, **2003**, 103, 671-5	7.5	68
522	Familial risks in nervous-system tumours: a histology-specific analysis from Sweden and Norway. *Lancet Oncology, The*, **2009**, 10, 481-8	21.7	67
521	Association between genetic polymorphisms and biomarkers in styrene-exposed workers. *Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis*, **2001**, 482, 89-103	3.3	67
520	Familial risks in cervical cancer: is there a hereditary component?. *International Journal of Cancer*, **1999**, 82, 775-81	7.5	67
519	Consanguinity and genetic diseases in North Africa and immigrants to Europe. *European Journal of Public Health*, **2014**, 24 Suppl 1, 57-63	2.1	66
518	Influence of education level on breast cancer risk and survival in Sweden between 1990 and 2004. *International Journal of Cancer*, **2008**, 122, 165-9	7.5	66
517	Familial risks and temporal incidence trends of multiple myeloma. *European Journal of Cancer*, **2006** , 42, 1661-70	7.5	66

516	The impact of type 2 diabetes mellitus on cancer-specific survival: a follow-up study in Sweden. *Cancer*, **2012**, 118, 1353-61	6.4	65
515	Mutations in TERT promoter and FGFR3 and telomere length in bladder cancer. *International Journal of Cancer*, **2015**, 137, 1621-9	7.5	65
514	Tumor location and patient characteristics of colon and rectal adenocarcinomas in relation to survival and TNM classes. *BMC Cancer*, **2010**, 10, 688	4.8	65
513	Genome-wide association study on differentiated thyroid cancer. *Journal of Clinical Endocrinology and Metabolism*, **2013**, 98, E1674-81	5.6	64
512	Incidence trends of squamous cell and rare skin cancers in the Swedish national cancer registry point to calendar year and age-dependent increases. *Journal of Investigative Dermatology*, **2010**, 130, 1323-8	4.3	64
511	Cancer risk in patients hospitalized with polymyalgia rheumatica and giant cell arteritis: a follow-up study in Sweden. *Rheumatology*, **2010**, 49, 1158-63	3.9	64
510	Cancer risks in ulcerative colitis patients. *International Journal of Cancer*, **2008**, 123, 1417-21	7.5	64
509	Effect of autoimmune diseases on risk and survival in female cancers. *Gynecologic Oncology*, **2012**, 127, 180-5	4.9	63
508	Incidence and survival in non-hereditary amyloidosis in Sweden. *BMC Public Health*, **2012**, 12, 974	4.1	62
507	Familial risk and familial survival in prostate cancer. *World Journal of Urology*, **2012**, 30, 143-8	4	62
506	Familial relationships in thyroid cancer by histo-pathological type. *International Journal of Cancer*, **2000**, 85, 201-205	7.5	60
505	TERT promoter mutations and telomere length in adult malignant gliomas and recurrences. *Oncotarget*, **2015**, 6, 10617-33	3.3	60
504	Update on genetic predisposition to colorectal cancer and polyposis. *Molecular Aspects of Medicine*, **2019**, 69, 10-26	16.7	59
503	How common is familial cancer?. *Annals of Oncology*, **2008**, 19, 163-7	10.3	59
502	Familial risks of cancer as a guide to gene identification and mode of inheritance. *International Journal of Cancer*, **2004**, 110, 291-4	7.5	59
501	Inherited predisposition to early onset lung cancer according to histological type. *International Journal of Cancer*, **2004**, 112, 451-7	7.5	58
500	Identification of multiple risk loci and regulatory mechanisms influencing susceptibility to multiple myeloma. *Nature Communications*, **2018**, 9, 3707	17.4	57
499	Melanocortin receptor 1 variants and melanoma risk: a study of 2 European populations. *International Journal of Cancer*, **2009**, 125, 1868-75	7.5	56

498	The rare ERBB2 variant Ile654Val is associated with an increased familial breast cancer risk. *Carcinogenesis*, **2005**, 26, 643-7	4.6	56
497	Familial and attributable risks in cutaneous melanoma: effects of proband and age. *Journal of Investigative Dermatology*, **2003**, 120, 217-23	4.3	56
496	Risk of subsequent solid tumors after non-Hodgkin's lymphoma: effect of diagnostic age and time since diagnosis. *Journal of Clinical Oncology*, **2008**, 26, 1850-7	2.2	55
495	Patterns of autoimmunity and subsequent chronic lymphocytic leukemia in Nordic countries. *Blood*, **2006**, 108, 292-6	2.2	55
494	Defining the genetic susceptibility to cervical neoplasia-A genome-wide association study. *PLoS Genetics*, **2017**, 13, e1006866	6	55
493	Basal cell carcinoma and variants in genes coding for immune response, DNA repair, folate and iron metabolism. *Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis*, **2005**, 574, 105-11	3.3	54
492	Subsequent cancers after in situ and invasive squamous cell carcinoma of the skin. *Archives of Dermatology*, **2000**, 136, 647-51		54
491	Subsequent autoimmune or related disease in asthma patients: clustering of diseases or medical care?. *Annals of Epidemiology*, **2010**, 20, 217-22	6.4	53
490	What do prostate cancer patients die of?. *Oncologist*, **2011**, 16, 175-81	5.7	53
489	Selective deletion of exon 1 beta of the p19ARF gene in metastatic melanoma cell lines. *Genes Chromosomes and Cancer*, **1998**, 23, 273-7	5	53
488	Subsequent primary malignancies after endometrial carcinoma and ovarian carcinoma. *Cancer*, **2003**, 97, 2432-9	6.4	53
487	Second primary cancer after in situ and invasive cervical cancer. *Epidemiology*, **2000**, 11, 457-61	3.1	53
486	Cancer risks to spouses and offspring in the Family-Cancer Database. *Genetic Epidemiology*, **2001**, 20, 247-57	2.6	52
485	Risk for multiple sclerosis in relatives and spouses of patients diagnosed with autoimmune and related conditions. *Neurogenetics*, **2009**, 10, 5-11	3	51
484	Familial risks in cancer of unknown primary: tracking the primary sites. *Journal of Clinical Oncology*, **2011**, 29, 435-40	2.2	51
483	Estimation of genetic and environmental components in colorectal and lung cancer and melanoma. *Genetic Epidemiology*, **2001**, 20, 107-116	2.6	51
482	Concordance of survival in family members with prostate cancer. *Journal of Clinical Oncology*, **2008**, 26, 1705-9	2.2	50
481	Serum oncoproteins and growth factors in asbestosis and silicosis patients. *International Journal of Cancer*, **1992**, 50, 881-5	7.5	50

480	MC1R variants associated susceptibility to basal cell carcinoma of skin: interaction with host factors and XRCC3 polymorphism. *International Journal of Cancer*, **2008**, 122, 1787-93	7.5	49
479	Age specific and attributable risks of familial prostate carcinoma from the family-cancer database. *Cancer*, **2002**, 95, 1346-53	6.4	49
478	Variation at 3p24.1 and 6q23.3 influences the risk of Hodgkin's lymphoma. *Nature Communications*, **2013**, 4, 2549	17.4	48
477	High familial risks for cerebral palsy implicate partial heritable aetiology. *Paediatric and Perinatal Epidemiology*, **2007**, 21, 235-41	2.7	48
476	The CASP8 -652 6N del promoter polymorphism and breast cancer risk: a multicenter study. *Breast Cancer Research and Treatment*, **2008**, 111, 139-44	4.4	48
475	Age-specific risk of incident prostate cancer and risk of death from prostate cancer defined by the number of affected family members. *European Urology*, **2010**, 58, 275-80	10.2	47
474	Polymorphisms in the KDR and POSTN genes: association with breast cancer susceptibility and prognosis. *Breast Cancer Research and Treatment*, **2007**, 101, 83-93	4.4	47
473	Associations of genetic variants in the estrogen receptor coactivators PPARGC1A, PPARGC1B and EP300 with familial breast cancer. *Carcinogenesis*, **2006**, 27, 2201-8	4.6	47
472	A genetic study of Hodgkin's lymphoma: an estimate of heritability and anticipation based on the familial cancer database in Sweden. *Human Genetics*, **2000**, 106, 553-6	6.3	47
471	Risk of Second Cancer in Hodgkin Lymphoma Survivors and Influence of Family History. *Journal of Clinical Oncology*, **2017**, 35, 1584-1590	2.2	46
470	Cancer risk in patients with type 2 diabetes mellitus and their relatives. *International Journal of Cancer*, **2015**, 137, 903-10	7.5	46
469	Polymorphisms in the IGF-1 and IGFBP 3 promoter and the risk of breast cancer. *Breast Cancer Research and Treatment*, **2005**, 92, 133-40	4.4	46
468	Deciphering the 8q24.21 association for glioma. *Human Molecular Genetics*, **2013**, 22, 2293-302	5.6	45
467	Familial association of inflammatory bowel diseases with other autoimmune and related diseases. *American Journal of Gastroenterology*, **2010**, 105, 139-47	0.7	45
466	Familial lung cancer and aggregation of smoking habits: a simulation of the effect of shared environmental factors on the familial risk of cancer. *Cancer Epidemiology Biomarkers and Prevention* , **2005**, 14, 1738-40	4	45
465	The population impact of familial cancer, a major cause of cancer. *International Journal of Cancer*, **2014**, 134, 1899-906	7.5	44
464	Effect of autoimmune diseases on risk and survival in histology-specific lung cancer. *European Respiratory Journal*, **2012**, 40, 1489-95	13.6	44
463	Number of siblings and the risk of lymphoma, leukemia, and myeloma by histopathology. *Cancer Epidemiology Biomarkers and Prevention*, **2006**, 15, 1281-6	4	44

462	Familial risk of cancer shortly after diagnosis of the first familial tumor. *Journal of the National Cancer Institute*, **2005**, 97, 1575-9	9.7	44
461	Effect of type 2 diabetes predisposing genetic variants on colorectal cancer risk. *Journal of Clinical Endocrinology and Metabolism*, **2012**, 97, E845-51	5.6	43
460	Association of prolactin and its receptor gene regions with familial breast cancer. *Journal of Clinical Endocrinology and Metabolism*, **2006**, 91, 1513-9	5.6	43
459	Association of the CASP10 V410I variant with reduced familial breast cancer risk and interaction with the CASP8 D302H variant. *Carcinogenesis*, **2006**, 27, 606-9	4.6	43
458	Kidney cancer in the Swedish Family Cancer Database: familial risks and second primary malignancies. *Kidney International*, **2002**, 61, 1806-13	9.9	43
457	Multiple primary cancers of the colon, breast and skin (melanoma) as models for polygenic cancers. *International Journal of Cancer*, **2001**, 92, 883-7	7.5	42
456	The 'common disease-common variant' hypothesis and familial risks. *PLoS ONE*, **2008**, 3, e2504	3.7	42
455	Frequent DPH3 promoter mutations in skin cancers. *Oncotarget*, **2015**, 6, 35922-30	3.3	42
454	Mutations in the CDKN2A (p16INK4a) gene in microdissected sporadic primary melanomas. *International Journal of Cancer*, **1998**, 75, 193-8	7.5	41
453	Familial risk of ischemic and hemorrhagic stroke: a large-scale study of the Swedish population. *Stroke*, **2006**, 37, 1668-73	6.7	41
452	Genetic epidemiology of cancer: from families to heritable genes. *International Journal of Cancer*, **2004**, 111, 944-50	7.5	41
451	Concordant and discordant familial cancer: Familial risks, proportions and population impact. *International Journal of Cancer*, **2017**, 140, 1510-1516	7.5	40
450	Risk of breast cancer in families of multiple affected women and men. *Breast Cancer Research and Treatment*, **2012**, 132, 723-8	4.4	40
449	Mutations, tissue accumulations, and serum levels of p53 in patients with occupational cancers from asbestos and silica exposure. *Environmental and Molecular Mutagenesis*, **1997**, 30, 224-230	3.2	40
448	Interaction of Werner and Bloom syndrome genes with p53 in familial breast cancer. *Carcinogenesis* , **2006**, 27, 1655-60	4.6	40
447	Attributable risks for familial breast cancer by proband status and morphology: a nationwide epidemiologic study from Sweden. *International Journal of Cancer*, **2002**, 100, 214-9	7.5	40
446	Familial breast cancer: scope for more susceptibility genes?. *Breast Cancer Research and Treatment*, **2003**, 82, 17-22	4.4	40
445	Chromosomal damage among medical staff occupationally exposed to volatile anesthetics, antineoplastic drugs, and formaldehyde. *Scandinavian Journal of Work, Environment and Health*, **2013**, 39, 618-30	4.3	40

444	Genome-wide association study identifies susceptibility loci for B-cell childhood acute lymphoblastic leukemia. *Nature Communications*, **2018**, 9, 1340	17.4	39
443	Increased risk of hepatobiliary cancers after hospitalization for autoimmune disease. *Clinical Gastroenterology and Hepatology*, **2014**, 12, 1038-45.e7	6.9	39
442	Chromosomal damage in peripheral blood lymphocytes of newly diagnosed cancer patients and healthy controls. *Carcinogenesis*, **2010**, 31, 1238-41	4.6	39
441	Familial risks for cervical tumors in full and half siblings: etiologic apportioning. *Cancer Epidemiology Biomarkers and Prevention*, **2006**, 15, 1413-4	4	39
440	Quantifying the heritability of testicular germ cell tumour using both population-based and genomic approaches. *Scientific Reports*, **2015**, 5, 13889	4.9	38
439	Shared familial aggregation of susceptibility to autoimmune diseases. *Arthritis and Rheumatism*, **2009**, 60, 2845-7		38
438	Incidence and familial risks in pituitary adenoma and associated tumors. *Endocrine-Related Cancer*, **2007**, 14, 103-9	5.7	38
437	Cancer characteristics in Swedish families fulfilling criteria for hereditary nonpolyposis colorectal cancer. *Gastroenterology*, **2005**, 129, 1889-99	13.3	38
436	Risk factors for cancers of unknown primary site: Results from the prospective EPIC cohort. *International Journal of Cancer*, **2014**, 135, 2475-81	7.5	36
435	Site-specific survival rates for cancer of unknown primary according to location of metastases. *International Journal of Cancer*, **2013**, 133, 182-9	7.5	36
434	Single nucleotide polymorphisms in chromosomal instability genes and risk and clinical outcome of breast cancer: a Swedish prospective case-control study. *European Journal of Cancer*, **2009**, 45, 435-42	7.5	36
433	Familial risks for cancer as the basis for evidence-based clinical referral and counseling. *Oncologist*, **2008**, 13, 239-47	5.7	36
432	Survival in ovarian cancer patients by histology and family history. *Acta Oncologica*, **2008**, 47, 1133-9	3.2	36
431	Risk of second malignant neoplasms among lymphoma patients with a family history of cancer. *International Journal of Cancer*, **2007**, 120, 1099-102	7.5	36
430	Constraints for genetic association studies imposed by attributable fraction and familial risk. *Carcinogenesis*, **2007**, 28, 648-56	4.6	36
429	Risks for familial and contralateral breast cancer interact multiplicatively and cause a high risk. *Cancer Research*, **2007**, 67, 868-70	10.1	36
428	Risk of thyroid cancer in first-degree relatives of patients with non-medullary thyroid cancer by histology type and age at diagnosis: a joint study from five Nordic countries. *Journal of Medical Genetics*, **2013**, 50, 373-82	5.8	35
427	FAMILIAL BLADDER CANCER IN THE NATIONAL SWEDISH FAMILY CANCER DATABASE. *Journal of Urology*, **2001**, 166, 2129-2133	2.5	35

426	Cancer risks in twins: results from the Swedish family-cancer database. *International Journal of Cancer*, **2002**, 99, 873-8	7.5	34
425	Re: Association of a common variant of the CASP8 gene with reduced risk of breast cancer. *Journal of the National Cancer Institute*, **2005**, 97, 1012; author reply 1012-3	9.7	34
424	Familial colorectal adenocarcinoma from the Swedish Family-Cancer Database. *International Journal of Cancer*, **2001**, 94, 743-8	7.5	34
423	Does the breast cancer age at diagnosis differ by ethnicity? A study on immigrants to Sweden. *Oncologist*, **2011**, 16, 146-54	5.7	33
422	Familial and second lung cancers: a nation-wide epidemiologic study from Sweden. *Lung Cancer*, **2003**, 39, 255-63	5.9	33
421	Prostate cancer screening, changing age-specific incidence trends and implications on familial risk. *International Journal of Cancer*, **2005**, 113, 312-5	7.5	33
420	Second primary cancers after anogenital, skin, oral, esophageal and rectal cancers: etiological links?. *International Journal of Cancer*, **2001**, 93, 294-8	7.5	33
419	DNA adducts of 1,3-butadiene in humans: relationships to exposure, GST genotypes, single-strand breaks, and cytogenetic end points. *Environmental and Molecular Mutagenesis*, **2001**, 37, 226-30	3.2	33
418	Genetic epidemiology of multistage carcinogenesis. *Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis*, **2001**, 473, 11-21	3.3	33
417	Detection of increased amounts of the extracellular domain of the c-erbB-2 oncoprotein in serum during pulmonary carcinogenesis in humans. *International Journal of Cancer*, **1994**, 56, 383-6	7.5	33
416	Do discordant cancers share familial susceptibility?. *European Journal of Cancer*, **2012**, 48, 1200-7	7.5	32
415	Effect of autoimmune diseases on incidence and survival in subsequent multiple myeloma. *Journal of Hematology and Oncology*, **2012**, 5, 59	22.4	32
414	Autoimmune disease and subsequent urological cancer. *Journal of Urology*, **2013**, 189, 2262-8	2.5	32
413	The updated Swedish family-cancer database used to assess familial risks of prostate cancer during rapidly increasing incidence. *Hereditary Cancer in Clinical Practice*, **2006**, 4, 186-92	2.3	32
412	Use of pyrosequencing to detect clinically relevant polymorphisms of genes in basal cell carcinoma. *Clinica Chimica Acta*, **2004**, 342, 137-43	6.2	32
411	Genetic Epidemiology--science and ethics on familial cancers. *Acta Oncologica*, **2001**, 40, 439-44	3.2	32
410	Cancer Risk in Relatives of Testicular Cancer Patients by Histology Type and Age at Diagnosis: A Joint Study from Five Nordic Countries. *European Urology*, **2015**, 68, 283-9	10.2	31
409	Nasopharyngeal and hypopharyngeal carcinoma risk among immigrants in Sweden. *International Journal of Cancer*, **2010**, 127, 2888-92	7.5	31

408	Association between number of siblings and nervous system tumors suggests an infectious etiology. *Neurology*, **2006**, 67, 1979-83	6.5	31
407	Time trends and occupational risk factors for pleural mesothelioma in Sweden. *Journal of Occupational and Environmental Medicine*, **2003**, 45, 456-61	2	31
406	Attributable risks of familial cancer from the Family-Cancer Database. *Cancer Epidemiology Biomarkers and Prevention*, **2002**, 11, 1638-44	4	31
405	Genome-wide analysis associates familial colorectal cancer with increases in copy number variations and a rare structural variation at 12p12.3. *Carcinogenesis*, **2014**, 35, 315-23	4.6	30
404	Familial risks for common diseases: etiologic clues and guidance to gene identification. *Mutation Research - Reviews in Mutation Research*, **2008**, 658, 247-58	7	30
403	Association of genetic variants in the Rho guanine nucleotide exchange factor AKAP13 with familial breast cancer. *Carcinogenesis*, **2006**, 27, 593-8	4.6	30
402	Cancer in husbands of cervical cancer patients. *Epidemiology*, **2000**, 11, 347-9	3.1	30
401	Structural chromosomal aberrations as potential risk markers in incident cancer patients. *Mutagenesis*, **2015**, 30, 557-63	2.8	29
400	Analysis of 153 115 patients with hematological malignancies refines the spectrum of familial risk. *Blood*, **2019**, 134, 960-969	2.2	29
399	Survival in breast cancer is familial. *Breast Cancer Research and Treatment*, **2008**, 110, 177-82	4.4	29
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