Hanne Meijers-Heijboer

List of Publications by Year
in descending order

30 papers	3,973 citations	430874 18 h-index	526287 27 g-index
30 all docs	30 docs citations	30 times ranked	8374 citing authors

#	Article	IF	Citations
1	Association analysis identifies 65 new breast cancer risk loci. Nature, 2017, 551, 92-94.	27.8	1,099
2	Large-scale genotyping identifies 41 new loci associated with breast cancer risk. Nature Genetics, 2013, 45, 353-361.	21.4	960
3	Genome-wide association analysis of more than 120,000 individuals identifies 15 new susceptibility loci for breast cancer. Nature Genetics, 2015, 47, 373-380.	21.4	513
4	Swarm Intelligence-Enhanced Detection of Non-Small-Cell Lung Cancer Using Tumor-Educated Platelets. Cancer Cell, 2017, 32, 238-252.e9.	16.8	235
5	A transcriptome-wide association study of 229,000 women identifies new candidate susceptibility genes for breast cancer. Nature Genetics, 2018, 50, 968-978.	21.4	184
6	A novel Fanconi anaemia subtype associated with a dominant-negative mutation in RAD51. Nature Communications, 2015, 6, 8829.	12.8	130
7	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.	8.4	118
8	Characterization of pathogenic SORL1 genetic variants for association with Alzheimerâ€™s disease: a clinical interpretation strategy. European Journal of Human Genetics, 2017, 25, 973-981.	2.8	102
9	Associations of obesity and circulating insulin and glucose with breast cancer risk: a Mendelian randomization analysis. International Journal of Epidemiology, 2019, 48, 795-806.	1.9	81
10	Somatic genomic alterations in retinoblastoma beyond RB1 are rare and limited to copy number changes. Scientific Reports, 2016, 6, 25264.	3.3	75
11	RB1mutation spectrum in a comprehensive nationwide cohort of retinoblastoma patients. Journal of Medical Genetics, 2014, 51, 366-374.	3.2	69
12	A Meta-Analysis of Retinoblastoma Copy Numbers Refines the List of Possible Driver Genes Involved in Tumor Progression. PLoS ONE, 2016, 11, e0153323.	2.5	55
13	Loss of photoreceptorness and gain of genomic alterations in retinoblastoma reveal tumor progression. EBioMedicine, 2015, 2, 660-670.	6.1	54
14	MicroRNA Related Polymorphisms and Breast Cancer Risk. PLoS ONE, 2014, 9, e109973.	2.5	49
15	Genomic profiling of CHEK2*1100delC-mutated breast carcinomas. BMC Cancer, 2015, 15, 877.	2.6	44
16	Identification of a Dutch founder mutation in MUSK causing fetal akinesia deformation sequence. European Journal of Human Genetics, 2015, 23, 1151-1157.	2.8	42
17	Addition of a 161-SNP polygenic risk score to family history-based risk prediction: impact on clinical management in non-BRCA1/2breast cancer families. Journal of Medical Genetics, 2019, 56, 581-589.	3.2	35
18	Targeted carrier screening for four recessive disorders: High detection rate within a founder population. European Journal of Medical Genetics, 2015, 58, 123-128.	1.3	31

#	Article	IF	Citations
19	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.	1.8	21
20	Proper genomic profiling of (BRCA1â€mutated) basalâ€like breast carcinomas requires prior removal of tumor infiltrating lymphocytes. Molecular Oncology, 2015, 9, 877-888.	4.6	16
21	Uptake of prenatal diagnostic testing for retinoblastoma compared to other hereditary cancer syndromes in the Netherlands. Familial Cancer, 2017, 16, 271-277.	1.9	16
22	First steps in exploring prospective exome sequencing of consanguineous couples. European Journal of Medical Genetics, 2014, 57, 613-616.	1.3	11
23	The Effect of Predictive Testing in Adultâ€Onset Neurodegenerative Diseases on Social and Personal Life. Journal of Genetic Counseling, 2018, 27, 947-954.	1.6	10
24	A novel CCM2 variant in a family with nonâ€progressive cognitive complaints and cerebral microbleeds. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2017, 174, 220-226.	1.7	6
25	Coregulation of FANCA and BRCA1 in human cells. SpringerPlus, 2014, 3, 381.	1.2	5
26	Genomic landscape of retinoblastoma in Rb^{â^'/â^'}p130^{â^'/â^'} mice resembles human retinoblastoma. Genes Chromosomes and Cancer, 2017, 56, 231-242.	2.8	5
27	Direct-to-consumer carrier screening for cystic fibrosis via a hospital website: a 6-year evaluation. Journal of Community Genetics, 2019, 10, 249-257.	1.2	5
28	S4-01-01: Pathology of cognitively healthy centenarians. , 2015, 11, P257-P257.		2
29	P2â€134: THE ADDED VALUE OF EXTREME PHENOTYPES IN ALZHEIMER'S DISEASE CASEâ€CONTROL STUDIES. Alzheimer's and Dementia, 2018, 14, P719.	0.8	0
30	P4â€038: IS SORL1 AN AUTOSOMAL DOMINANT ALZHEIMER GENE?. Alzheimer's and Dementia, 2018, 14, P1447.	0.8	0