Cora M Aalfs

List of Publications by Year
in descending order

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40 papers	1,955 citations	430874 18 h-index	315739 38 g-index
41 all docs	41 docs citations	41 times ranked	4666 citing authors

#	Article	IF	Citations
1	Reproductive decision-making in the context of hereditary cancer: the effects of an online decision aid on informed decision-making. Journal of Community Genetics, 2021, 12, 101-110.	1.2	6
2	Effect of a health literacy training program for surgical oncologists and specialized nurses on disparities in referral to breast cancer genetic testing. Breast, 2021, 58, 80-87.	2.2	5
3	Lack of genotype-phenotype correlation in basal cell nevus syndrome: A Dutch multicenter retrospective cohort study. Journal of the American Academy of Dermatology, 2020, 83, 604-607.	1.2	7
4	â€˜We donâ€™t know for sureâ€™: discussion of uncertainty concerning multigene panel testing during initial cancer genetic consultations. Familial Cancer, 2020, 19, 65-76.	1.9	10
5	Prophylactic total pancreatectomy in individuals at high risk of pancreatic ductal adenocarcinoma (PROPAN): systematic review and shared decisionâ€making programme using decision tables. United European Gastroenterology Journal, 2020, 8, 865-877.	3.8	11
6	Addition of an online, validated family history questionnaire to the Dutch FIT-based screening programme did not improve its diagnostic yield. British Journal of Cancer, 2020, 122, 1865-1871.	6.4	4
7	Uncertainty related to multigene panel testing for cancer: a qualitative study on counsellorsâ€™ and counseleesâ€™ views. Journal of Community Genetics, 2019, 10, 303-312.	1.2	18
8	Online decision support for persons having a genetic predisposition to cancer and their partners during reproductive decisionâ€making. Journal of Genetic Counseling, 2019, 28, 533-542.	1.6	14
9	Height and Body Mass Index as Modifiers of Breast Cancer Risk in BRCA1/2 Mutation Carriers: A Mendelian Randomization Study. Journal of the National Cancer Institute, 2019, 111, 350-364.	6.3	30
10	The development of an online decision aid to support persons having a genetic predisposition to cancer and their partners during reproductive decision-making: a usability and pilot study. Familial Cancer, 2019, 18, 137-146.	1.9	17
11	Mutational spectrum in a worldwide study of 29,700 families withBRCA1orBRCA2mutations. Human Mutation, 2018, 39, 593-620.	2.5	224
12	Role of germline aberrations affecting CTNNA1, MAP3K6 and MYD88 in gastric cancer susceptibility. Journal of Medical Genetics, 2018, 55, 669-674.	3.2	37
13	Duodenal Adenomas in Patients With Multiple Colorectal Adenomas Without Germline APC or MUTYH Mutations. Diseases of the Colon and Rectum, 2018, 61, 58-66.	1.3	14
14	How to support cancer genetics counselees in informing at-risk relatives? Lessons from a randomized controlled trial. Patient Education and Counseling, 2018, 101, 1611-1619.	2.2	22
15	Uncertainty in consultations about genetic testing for cancer: an explorative observational study. Patient Education and Counseling, 2018, 101, 2083-2089.	2.2	10
16	Childhood tumours with a high probability of being part of a tumour predisposition syndrome; reason for referral for genetic consultation. European Journal of Cancer, 2017, 80, 48-54.	2.8	19
17	Identification of 12 new susceptibility loci for different histotypes of epithelial ovarian cancer. Nature Genetics, 2017, 49, 680-691.	21.4	356
18	Identification of ten variants associated with risk of estrogen-receptor-negative breast cancer. Nature Genetics, 2017, 49, 1767-1778.	21.4	289

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19	Unraveling genetic predisposition to familial or early onset gastric cancer using germline whole-exome sequencing. European Journal of Human Genetics, 2017, 25, 1246-1252.	2.8	34
20	Colorectal Cancer Risk in Patients With Lynch Syndrome andÂInflammatory Bowel Disease. Clinical Gastroenterology and Hepatology, 2017, 15, 454-458.e1.	4.4	20
21	Factors associated with cancer worries in individuals participating in annual pancreatic cancer surveillance. Familial Cancer, 2017, 16, 143-151.	1.9	25
22	An E-Learning Module to Improve Nongenetic Health Professionalsâ€™ Assessment of Colorectal Cancer Genetic Risk: Feasibility Study. JMIR Medical Education, 2017, 3, e24.	2.6	8
23	Design and Feasibility of an Intervention to Support Cancer Genetic Counselees in Informing their Atâ€Risk Relatives. Journal of Genetic Counseling, 2016, 25, 1179-1187.	1.6	21
24	Repeated participation in pancreatic cancer surveillance by highâ€risk individuals imposes low psychological burden. Psycho-Oncology, 2016, 25, 971-978.	2.3	33
25	Bias Explains Most of the Parent-of-Origin Effect on Breast Cancer Risk in BRCA1/2 Mutation Carriers. Cancer Epidemiology Biomarkers and Prevention, 2016, 25, 1251-1258.	2.5	9
26	Gatekeeper role of gastroenterologists and surgeons in recognising and discussing familial colorectal cancer. Familial Cancer, 2016, 15, 231-240.	1.9	6
27	An original phylogenetic approach identified mitochondrial haplogroup T1a1 as inversely associated with breast cancer risk in BRCA2 mutation carriers. Breast Cancer Research, 2015, 17, 61.	5.0	26
28	Assessing Associations between the AURKA-HMMR-TPX2-TUBG1 Functional Module and Breast Cancer Risk in BRCA1/2 Mutation Carriers. PLoS ONE, 2015, 10, e0120020.	2.5	34
29	Accuracy of Hereditary Diffuse Gastric Cancer Testing Criteria and Outcomes in Patients With a Germline Mutation in CDH1. Gastroenterology, 2015, 149, 897-906.e19.	1.3	70
30	Association of Type and Location ofBRCA1andBRCA2Mutations With Risk of Breast and Ovarian Cancer. JAMA - Journal of the American Medical Association, 2015, 313, 1347.	7.4	390
31	Pancreatic cancer-associated gene polymorphisms in a nation-wide cohort of p16-Leiden germline mutation carriers; a caseâ€“control study. BMC Research Notes, 2015, 8, 264.	1.4	10
32	Brain Tumors and Syndromes in Children. Neuropediatrics, 2014, 45, 137-161.	0.6	15
33	Counselorâ€“counselee interaction in reproductive genetic counseling: Does a pregnancy in the counselee make a difference?. Patient Education and Counseling, 2006, 60, 80-90.	2.2	11
34	Referral for genetic counselling during pregnancy: limited alertness and awareness about genetic risk factors among GPs. Family Practice, 2003, 20, 135-141.	1.9	35
35	A case of methemoglobinemia type II due to NADH-cytochrome b5 reductase deficiency: Determination of the molecular basis. Human Mutation, 2000, 16, 18-22.	2.5	36
36	A case of methemoglobinemia type II due to NADH-cytochrome b5 reductase deficiency: Determination of the molecular basis. Human Mutation, 2000, 16, 18.	2.5	5

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37	Molecular analysis of a translocation (6;11)(p21;q25) in a girl with Jacobsen syndrome. , 1999, 86, 398-400.		6
38	Asn540Thr substitution in the fibroblast growth factor receptor 3 tyrosine kinase domain causing hypochondroplasia. Human Mutation, 1998, 11, S62-S65.	2.5	34
39	Further delineation of the acro-renal-ocular syndrome. , 1996, 62, 276-281.		23
40	Two supernumerary marker chromosomes, derived from chromosome 6 and 9, in a boy with mild developmental delay. Clinical Genetics, 1996, 49, 42-45.	2.0	11