

Nicoline Hoogerbrugge

List of Publications by Year in descending order

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Version: 2024-02-01

308
papers

16,840
citations

16437

64
h-index

20343

116
g-index

314
all docs

314
docs citations

314
times ranked

18595
citing authors

#	ARTICLE	IF	CITATIONS
1	Healthcare professionalsâ€™ perspectives on implementation of universal tumor DNA testing in ovarian cancer patients: multidisciplinary focus groups. <i>Familial Cancer</i> , 2023, 22, 1-11.	0.9	3
2	Probability of detecting germline BRCA1/2 pathogenic variants in histological subtypes of ovarian carcinoma. A meta-analysis. <i>Gynecologic Oncology</i> , 2022, 164, 221-230.	0.6	11
3	Genetic Cancer Susceptibility in Adolescents and Adults 25 Years or Younger With Colorectal Cancer. <i>Gastroenterology</i> , 2022, 162, 969-974.e6.	0.6	2
4	Gastric cancer genetic predisposition and clinical presentations: Established heritable causes and potential candidate genes. <i>European Journal of Medical Genetics</i> , 2022, 65, 104401.	0.7	10
5	Cancer worry among BRCA1/2 pathogenic variant carriers choosing surgery to prevent tubal/ovarian cancer: course over time and associated factors. <i>Supportive Care in Cancer</i> , 2022, 30, 3409-3418.	1.0	1
6	Risk of Peritoneal Carcinomatosis After Risk-Reducing Salpingo-Oophorectomy: A Systematic Review and Individual Patient Data Meta-Analysis. <i>Journal of Clinical Oncology</i> , 2022, 40, 1879-1891.	0.8	25
7	Solving the genetic aetiology of hereditary gastrointestinal tumour syndromesâ€” a collaborative multicentre endeavour within the project Solve-RD. <i>European Journal of Medical Genetics</i> , 2022, 65, 104475.	0.7	2
8	Germline MBD4 deficiency causes a multi-tumor predisposition syndrome. <i>American Journal of Human Genetics</i> , 2022, 109, 953-960.	2.6	23
9	Catch them if you are aware: PTEN postzygotic mosaicism in clinically suspicious patients with PTEN Hamartoma Tumour Syndrome and literature review. <i>European Journal of Medical Genetics</i> , 2022, , 104533.	0.7	4
10	The yield and effectiveness of breast cancer surveillance in women with <sc>PTEN</sc> Hamartoma Tumor Syndrome. <i>Cancer</i> , 2022, 128, 2883-2891.	2.0	4
11	A review on ageâ€related cancer risks in <sc>PTEN</sc> hamartoma tumor syndrome. <i>Clinical Genetics</i> , 2021, 99, 219-225.	1.0	42
12	Somatic Nonepigenetic Mismatch Repair Gene Aberrations Underly Most Mismatch Repairâ€Deficient Lynch-Like Tumors. <i>Gastroenterology</i> , 2021, 160, 1414-1416.e3.	0.6	13
13	No signs of subclinical atherosclerosis after risk-reducing salpingo-oophorectomy in BRCA1/2 mutation carriers. <i>Journal of Cardiology</i> , 2021, 77, 570-575.	0.8	3
14	<i>RNF43</i> mutation analysis in serrated polyposis, sporadic serrated polyps and Lynch syndrome polyps. <i>Histopathology</i> , 2021, 78, 749-758.	1.6	10
15	Continue rare cancers collaboration with European Reference Networks after Brexit. <i>Lancet</i> , The, 2021, 397, 793.	6.3	2
16	European Reference Networks: challenges and opportunities. <i>Journal of Community Genetics</i> , 2021, 12, 217-229.	0.5	21
17	Breast cancer surveillance in women with PTEN Hamartoma Tumour Syndrome (PHTS). <i>Breast</i> , 2021, 56, S38.	0.9	0
18	Challenges of Neoantigen Targeting in Lynch Syndrome and Constitutional Mismatch Repair Deficiency Syndrome. <i>Cancers</i> , 2021, 13, 2345.	1.7	3

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19	A mosaic PIK3CA variant in a young adult with diffuse gastric cancer: case report. <i>European Journal of Human Genetics</i> , 2021, 29, 1354-1358.	1.4	9
20	Solve-RD: systematic pan-European data sharing and collaborative analysis to solve rare diseases. <i>European Journal of Human Genetics</i> , 2021, 29, 1325-1331.	1.4	49
21	Universal genetic assessment for women with ovarian cancer not yet achieved: the promises of universal tumor DNA testing. <i>Gynecologic Oncology Reports</i> , 2021, 38, 100825.	0.3	0
22	Variation in the risk of colorectal cancer in families with Lynch syndrome: a retrospective cohort study. <i>Lancet Oncology</i> , The, 2021, 22, 1014-1022.	5.1	58
23	Association of Salpingectomy With Delayed Oophorectomy Versus Salpingo-oophorectomy With Quality of Life in BRCA1/2 Pathogenic Variant Carriers. <i>JAMA Oncology</i> , 2021, 7, 1203.	3.4	27
24	Evaluation of a patient decision aid for BRCA1/2 pathogenic variant carriers choosing an ovarian cancer prevention strategy. <i>Gynecologic Oncology</i> , 2021, 163, 371-377.	0.6	2
25	Red flags for early recognition of adult patients with PTEN Hamartoma Tumour Syndrome. <i>European Journal of Medical Genetics</i> , 2021, 64, 104364.	0.7	7
26	Overview of hereditary breast and ovarian cancer (HBOC) guidelines across Europe. <i>European Journal of Medical Genetics</i> , 2021, 64, 104350.	0.7	22
27	Universal Tumor DNA BRCA1/2 Testing of Ovarian Cancer: Prescreening PARPi Treatment and Genetic Predisposition. <i>Journal of the National Cancer Institute</i> , 2020, 112, 161-169.	3.0	47
28	Response to Tomao, Panici, and Tomao. <i>Journal of the National Cancer Institute</i> , 2020, 112, 425-425.	3.0	0
29	Microsatellite instability screening in colorectal adenomas to detect Lynch syndrome patients? A systematic review and meta-analysis. <i>European Journal of Human Genetics</i> , 2020, 28, 277-286.	1.4	22
30	“Patient Journeys” improving care by patient involvement. <i>European Journal of Human Genetics</i> , 2020, 28, 141-143.	1.4	18
31	Candidate Gene Discovery in Hereditary Colorectal Cancer and Polyposis Syndromes—Considerations for Future Studies. <i>International Journal of Molecular Sciences</i> , 2020, 21, 8757.	1.8	7
32	Hereditary diffuse gastric cancer: updated clinical practice guidelines. <i>Lancet Oncology</i> , The, 2020, 21, e386-e397.	5.1	237
33	Monoallelic NTHL1 Loss-of-Function Variants and Risk of Polyposis and Colorectal Cancer. <i>Gastroenterology</i> , 2020, 159, 2241-2243.e6.	0.6	20
34	Recommendations on Surveillance for Differentiated Thyroid Carcinoma in Children with PTEN Hamartoma Tumor Syndrome. <i>European Thyroid Journal</i> , 2020, 9, 234-242.	1.2	14
35	Cancer Surveillance Guideline for individuals with PTEN hamartoma tumour syndrome. <i>European Journal of Human Genetics</i> , 2020, 28, 1387-1393.	1.4	63
36	Evaluation of yield and experiences of age-related molecular investigation for heritable and nonheritable causes of mismatch repair deficient colorectal cancer to identify Lynch syndrome. <i>International Journal of Cancer</i> , 2020, 147, 2150-2158.	2.3	15

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37	Alcohol Consumption, Cigarette Smoking, and Risk of Breast Cancer for <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers: Results from The BRCA1 and BRCA2 Cohort Consortium. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2020, 29, 368-378.	1.1	24
38	<i>TRIM28</i> haploinsufficiency predisposes to Wilms tumor. <i>International Journal of Cancer</i> , 2019, 145, 941-951.	2.3	45
39	Effect of PTEN inactivating germline mutations on innate immune cell function and thyroid cancer-induced macrophages in patients with PTEN hamartoma tumor syndrome. <i>Oncogene</i> , 2019, 38, 3743-3755.	2.6	20
40	Addition of a 161-SNP polygenic risk score to family history-based risk prediction: impact on clinical management in non- <i>BRCA1/2</i> breast cancer families. <i>Journal of Medical Genetics</i> , 2019, 56, 581-589.	1.5	35
41	De Novo and Inherited Pathogenic Variants in <i>KDM3B</i> Cause Intellectual Disability, Short Stature, and Facial Dysmorphism. <i>American Journal of Human Genetics</i> , 2019, 104, 758-766.	2.6	34
42	Romidepsin suppresses monosodium urate crystal-induced cytokine production through upregulation of suppressor of cytokine signaling 1 expression. <i>Arthritis Research and Therapy</i> , 2019, 21, 50.	1.6	25
43	Mutational Signature Analysis Reveals <i>NTHL1</i> Deficiency to Cause a Multi-tumor Phenotype. <i>Cancer Cell</i> , 2019, 35, 256-266.e5.	7.7	123
44	Cancer-related distress in unselected women with newly diagnosed breast or ovarian cancer undergoing <i>BRCA1/2</i> testing without pretest genetic counseling. <i>Acta Oncologica</i> , 2019, 58, 175-181.	0.8	6
45	Boosting care and knowledge about hereditary cancer: European Reference Network on Genetic Tumour Risk Syndromes. <i>Familial Cancer</i> , 2019, 18, 281-284.	0.9	17
46	PTEN Hamartoma Tumor Syndrome and Immune Dysregulation. <i>Translational Oncology</i> , 2019, 12, 361-367.	1.7	33
47	High Yield of Pathogenic Germline Mutations Causative or Likely Causative of the Cancer Phenotype in Selected Children with Cancer. <i>Clinical Cancer Research</i> , 2018, 24, 1594-1603.	3.2	52
48	A patient decision aid for risk-reducing surgery in premenopausal <i>BRCA1/2</i> mutation carriers: Development process and pilot testing. <i>Health Expectations</i> , 2018, 21, 659-667.	1.1	17
49	Role of germline aberrations affecting <i>CTNNA1</i> , <i>MAP3K6</i> and <i>MYD88</i> in gastric cancer susceptibility. <i>Journal of Medical Genetics</i> , 2018, 55, 669-674.	1.5	37
50	Peritoneal carcinomatosis after risk-reducing surgery in <i>BRCA1/2</i> mutation carriers. <i>Cancer</i> , 2018, 124, 952-959.	2.0	27
51	Self-compassion, physical fitness and climacteric symptoms in oophorectomized <i>BRCA1/2</i> mutation carriers. <i>Maturitas</i> , 2018, 108, 13-17.	1.0	3
52	Outcomes of screening gastroscopy in first-degree relatives of patients fulfilling hereditary diffuse gastric cancer criteria. <i>Gastrointestinal Endoscopy</i> , 2018, 87, 397-404.e2.	0.5	28
53	High demoralization in a minority of oophorectomized <i>BRCA1/2</i> mutation carriers influences quality of life. <i>Journal of Psychosomatic Obstetrics and Gynaecology</i> , 2018, 39, 96-104.	1.1	7
54	Increasing awareness and knowledge of lifestyle recommendations for cancer prevention in Lynch syndrome carriers: Randomized controlled trial. <i>Clinical Genetics</i> , 2018, 93, 67-77.	1.0	9

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55	Influence of Risk Category and Screening Round on the Performance of an MR Imaging and Mammography Screening Program in Carriers of the <i>BRCA</i> Mutation and Other Women at Increased Risk. <i>Radiology</i> , 2018, 286, 443-451.	3.6	48
56	<i>NTHL1</i> and <i>MUTYH</i> polyposis syndromes: two sides of the same coin?. <i>Journal of Pathology</i> , 2018, 244, 135-142.	2.1	63
57	Cancer Risks for <i>PMS2</i> -Associated Lynch Syndrome. <i>Journal of Clinical Oncology</i> , 2018, 36, 2961-2968.	0.8	147
58	The added value of mammography in different age-groups of women with and without BRCA mutation screened with breast MRI. <i>Breast Cancer Research</i> , 2018, 20, 84.	2.2	40
59	An online self-test added to colorectal cancer screening can increase the effectiveness of familial cancer risk assessment without increasing distress. <i>Colorectal Disease</i> , 2018, 20, 897-904.	0.7	2
60	Cancer prevention by aspirin in children with Constitutional Mismatch Repair Deficiency (CMMRD). <i>European Journal of Human Genetics</i> , 2018, 26, 1417-1423.	1.4	20
61	Online self-test identifies women at high familial breast cancer risk in population-based breast cancer screening without inducing anxiety or distress. <i>European Journal of Cancer</i> , 2017, 78, 45-52.	1.3	14
62	Surveillance of Women with the <i>BRCA1</i> or <i>BRCA2</i> Mutation by Using Biannual Automated Breast US, MR Imaging, and Mammography. <i>Radiology</i> , 2017, 285, 376-388.	3.6	61
63	Childhood neuroendocrine tumours: a descriptive study revealing clues for genetic predisposition. <i>British Journal of Cancer</i> , 2017, 116, 163-168.	2.9	16
64	Lifestyle Risk Factors for Breast Cancer in <i>BRCA1/2</i> Mutation Carriers Around Childbearing Age. <i>Journal of Genetic Counseling</i> , 2017, 26, 785-791.	0.9	6
65	Unraveling genetic predisposition to familial or early onset gastric cancer using germline whole-exome sequencing. <i>European Journal of Human Genetics</i> , 2017, 25, 1246-1252.	1.4	34
66	Self-compassion and climacteric symptoms in postmenopausal <i>BRCA1/2</i> mutation carriers. <i>Maturitas</i> , 2017, 100, 130.	1.0	0
67	Immunotherapy holds the key to cancer treatment and prevention in constitutional mismatch repair deficiency (CMMRD) syndrome. <i>Cancer Letters</i> , 2017, 403, 159-164.	3.2	37
68	Novel <i>BRCA1</i> and <i>BRCA2</i> Tumor Test as Basis for Treatment Decisions and Referral for Genetic Counselling of Patients with Ovarian Carcinomas. <i>Human Mutation</i> , 2017, 38, 226-235.	1.1	55
69	Colorectal Cancer Risk in Patients With Lynch Syndrome and Inflammatory Bowel Disease. <i>Clinical Gastroenterology and Hepatology</i> , 2017, 15, 454-458.e1.	2.4	20
70	Germline activating <i>TYK2</i> mutations in pediatric patients with two primary acute lymphoblastic leukemia occurrences. <i>Leukemia</i> , 2017, 31, 821-828.	3.3	35
71	A molecular inversion probe-based next-generation sequencing panel to detect germline mutations in Chinese early-onset colorectal cancer patients. <i>Oncotarget</i> , 2017, 8, 24533-24547.	0.8	12
72	Determinants of adherence to recommendations for cancer prevention among Lynch Syndrome mutation carriers: A qualitative exploration. <i>PLoS ONE</i> , 2017, 12, e0178205.	1.1	10

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73	Preventive dendritic cell vaccination in healthy Lynch syndrome mutation carriers. <i>Annals of Oncology</i> , 2016, 27, vi362.	0.6	4
74	Prevalence of germline mutations in the spindle assembly checkpoint gene BUB1B in individuals with early-onset colorectal cancer. <i>Genes Chromosomes and Cancer</i> , 2016, 55, 855-863.	1.5	30
75	Opportunities for immunotherapy in microsatellite instable colorectal cancer. <i>Cancer Immunology, Immunotherapy</i> , 2016, 65, 1249-1259.	2.0	67
76	Prospective Dutch colorectal cancer cohort: an infrastructure for long-term observational, prognostic, predictive and (randomized) intervention research. <i>Acta Oncologica</i> , 2016, 55, 1273-1280.	0.8	62
77	Finding all BRCA pathogenic mutation carriers: best practice models. <i>European Journal of Human Genetics</i> , 2016, 24, S19-S26.	1.4	15
78	Very high uptake of risk-reducing salpingo-oophorectomy in BRCA1/2 mutation carriers: A single-center experience. <i>Gynecologic Oncology</i> , 2016, 143, 113-119.	0.6	23
79	The genetic heterogeneity of colorectal cancer predisposition - guidelines for gene discovery. <i>Cellular Oncology (Dordrecht)</i> , 2016, 39, 491-510.	2.1	34
80	Salpingectomy With Delayed Oophorectomy in BRCA1/2 Mutation Carriers. <i>Obstetrics and Gynecology</i> , 2016, 127, 1054-1063.	1.2	21
81	Cardiovascular risk of BRCA1/2 mutation carriers: A review. <i>Maturitas</i> , 2016, 91, 135-139.	1.0	28
82	Health risks for ataxia-telangiectasia mutated heterozygotes: a systematic review, meta-analysis and evidence-based guideline. <i>Clinical Genetics</i> , 2016, 90, 105-117.	1.0	143
83	Recurrent candidiasis and early-onset gastric cancer in a patient with a genetically defined partial MYD88 defect. <i>Familial Cancer</i> , 2016, 15, 289-296.	0.9	13
84	Colorectal cancer risk variants at 8q23.3 and 11q23.1 are associated with disease phenotype in APC mutation carriers. <i>Familial Cancer</i> , 2016, 15, 563-570.	0.9	8
85	Recognition of genetic predisposition in pediatric cancer patients: An easy-to-use selection tool. <i>European Journal of Medical Genetics</i> , 2016, 59, 116-125.	0.7	125
86	High Satisfaction and Low Distress in Breast Cancer Patients One Year after BRCA Mutation Testing without Prior Face-to-Face Genetic Counseling. <i>Journal of Genetic Counseling</i> , 2016, 25, 504-514.	0.9	30
87	BRCA1/2 testing in newly diagnosed breast and ovarian cancer patients without prior genetic counselling: the DNA-BONus study. <i>European Journal of Human Genetics</i> , 2016, 24, 881-888.	1.4	58
88	The effect of genotypes and parent of origin on cancer risk and age of cancer development in PMS2 mutation carriers. <i>Genetics in Medicine</i> , 2016, 18, 405-409.	1.1	15
89	Peer support and additional information in group medical consultations (GMCs) for BRCA1/2 mutation carriers: A randomized controlled trial. <i>Acta Oncologica</i> , 2016, 55, 178-187.	0.8	5
90	Identification of Novel Candidate Genes for Early-Onset Colorectal Cancer Susceptibility. <i>PLoS Genetics</i> , 2016, 12, e1005880.	1.5	52

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91	Abstract IA44: Cancer prevention: Dendritic cell enhanced immune responses towards neoantigens in patients with Lynch syndrome. , 2016, , .		0
92	Deleterious Germline BLM Mutations and the Risk for Early-onset Colorectal Cancer. Scientific Reports, 2015, 5, 14060.	1.6	67
93	Breast Self-examination Education for BRCA Mutation Carriers by Clinical Nurse Specialists. Clinical Nurse Specialist, 2015, 29, E1-E7.	0.3	11
94	Poor prognosis of constitutive γ -H2AX expressing triple-negative breast cancers is associated with telomere length. Biomarkers in Medicine, 2015, 9, 383-390.	0.6	17
95	Risk-reducing salpingectomy with delayed oophorectomy in BRCA1/2 mutation carriers: Patients' and professionals' perspectives. Gynecologic Oncology, 2015, 136, 305-310.	0.6	31
96	No effects of atorvastatin (10mg/d or 80mg/d) on nitric oxide, prostacyclin, thromboxane and oxidative stress in type 2 diabetes mellitus patients of the DALI study. Pharmacological Research, 2015, 94, 1-8.	3.1	11
97	Gastric cancer in three relatives of a patient with a biallelic IL12RB1 mutation. Familial Cancer, 2015, 14, 89-94.	0.9	14
98	Identification of six new susceptibility loci for invasive epithelial ovarian cancer. Nature Genetics, 2015, 47, 164-171.	9.4	221
99	Germline deletions in the tumour suppressor gene <i>FOCAD</i> are associated with polyposis and colorectal cancer development. Journal of Pathology, 2015, 236, 155-164.	2.1	28
100	Hereditary diffuse gastric cancer: updated clinical guidelines with an emphasis on germline <i>CDH1</i> mutation carriers. Journal of Medical Genetics, 2015, 52, 361-374.	1.5	479
101	Highlights from the seventh European Multidisciplinary Colorectal Cancer Congress (EMCCC) 2014. Ecanermedicalscience, 2015, 9, 497.	0.6	0
102	How medical choices influence quality of life of women carrying a BRCA mutation. Critical Reviews in Oncology/Hematology, 2015, 96, 555-568.	2.0	22
103	Accuracy of Hereditary Diffuse Gastric Cancer Testing Criteria and Outcomes in Patients With a Germline Mutation in CDH1. Gastroenterology, 2015, 149, 897-906.e19.	0.6	70
104	Candidate colorectal cancer predisposing gene variants in Chinese early-onset and familial cases. World Journal of Gastroenterology, 2015, 21, 4136.	1.4	10
105	A germline homozygous mutation in the base-excision repair gene NTHL1 causes adenomatous polyposis and colorectal cancer. Nature Genetics, 2015, 47, 668-671.	9.4	311
106	Improving recognition and referral of patients with an increased familial risk of colorectal cancer: results from a randomized controlled trial. Colorectal Disease, 2015, 17, 499-510.	0.7	10
107	Low prevalence of serrated polyposis syndrome in screening populations: a systematic review. Endoscopy, 2015, 47, 1043-1049.	1.0	28
108	Early salpingectomy (TUBectomy) with delayed oophorectomy to improve quality of life as alternative for risk-reducing salpingo-oophorectomy in BRCA1/2 mutation carriers (TUBA study): a prospective non-randomised multicentre study. BMC Cancer, 2015, 15, 593.	1.1	88

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109	Lynch Syndrome Caused by Germline <i>PMS2</i> Mutations: Delineating the Cancer Risk. <i>Journal of Clinical Oncology</i> , 2015, 33, 319-325.	0.8	177
110	Patient experiences with gene panels based on exome sequencing in clinical diagnostics: high acceptance and low distress. <i>Clinical Genetics</i> , 2015, 87, 319-326.	1.0	23
111	Candidate Genetic Modifiers for Breast and Ovarian Cancer Risk in <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2015, 24, 308-316.	1.1	22
112	<i>NTHL1</i> defines novel cancer syndrome. <i>Oncotarget</i> , 2015, 6, 34069-34070.	0.8	21
113	More breast cancer patients prefer BRCA-mutation testing without prior face-to-face genetic counseling. <i>Familial Cancer</i> , 2014, 13, 143-51.	0.9	33
114	DNA Glycosylases Involved in Base Excision Repair May Be Associated with Cancer Risk in <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers. <i>PLoS Genetics</i> , 2014, 10, e1004256.	1.5	47
115	Germline <i>MUTYH</i> gene mutations are not frequently found in unselected patients with papillary breast carcinoma. <i>Hereditary Cancer in Clinical Practice</i> , 2014, 12, 21.	0.6	4
116	A multiplex method for the detection of serum antibodies against in silico-predicted tumor antigens. <i>Cancer Immunology, Immunotherapy</i> , 2014, 63, 1251-1259.	2.0	6
117	Breast and ovarian cancer risks in a large series of clinically ascertained families with a high proportion of <i>BRCA1</i> and <i>BRCA2</i> Dutch founder mutations. <i>Journal of Medical Genetics</i> , 2014, 51, 98-107.	1.5	74
118	Easy-to-Use Decision Aids for Improved Cancer Family History Collection and Use Among Oncology Practices. <i>Journal of Clinical Oncology</i> , 2014, 32, 3343-3343.	0.8	5
119	Easy-to-use online referral test detects most patients with a high familial risk of colorectal cancer. <i>Colorectal Disease</i> , 2014, 16, O26-34.	0.7	12
120	Relevance and efficacy of breast cancer screening in <i>BRCA1</i> and <i>BRCA2</i> mutation carriers above 60 years: A national cohort study. <i>International Journal of Cancer</i> , 2014, 135, 2940-2949.	2.3	13
121	Colorectal cancer risk variants on 11q23 and 15q13 are associated with unexplained adenomatous polyposis. <i>Journal of Medical Genetics</i> , 2014, 51, 55-60.	1.5	21
122	Somatic Mutations in <i>MLH1</i> and <i>MSH2</i> Are a Frequent Cause of Mismatch-Repair Deficiency in Lynch Syndrome-Like Tumors. <i>Gastroenterology</i> , 2014, 146, 643-646.e8.	0.6	294
123	Fourfold increased detection of Lynch syndrome by raising age limit for tumour genetic testing from 50 to 70 years is cost-effective. <i>Annals of Oncology</i> , 2014, 25, 2001-2007.	0.6	26
124	<i>HNF4A</i> immunohistochemistry facilitates distinction between primary and metastatic breast and gastric carcinoma. <i>Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin</i> , 2014, 464, 673-679.	1.4	26
125	<i>BRCA1/2</i> mutation carriers are potentially at higher cardiovascular risk. <i>Critical Reviews in Oncology/Hematology</i> , 2014, 91, 159-171.	2.0	31
126	<i>EPCAM</i> deletion carriers constitute a unique subgroup of Lynch syndrome patients. <i>Familial Cancer</i> , 2013, 12, 169-174.	0.9	100

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127	Germline Mutations in the Spindle Assembly Checkpoint Genes BUB1 and BUB3 Are Risk Factors for Colorectal Cancer. <i>Gastroenterology</i> , 2013, 145, 544-547.	0.6	86
128	Cost-Effectiveness of Screening Women With Familial Risk for Breast Cancer With Magnetic Resonance Imaging. <i>Journal of the National Cancer Institute</i> , 2013, 105, 1314-1321.	3.0	57
129	Clinical utility gene card for: Hereditary diffuse gastric cancer (HDGC). <i>European Journal of Human Genetics</i> , 2013, 21, 891-891.	1.4	22
130	Breast cancer size estimation with MRI in BRCA mutation carriers and other high risk patients. <i>European Journal of Radiology</i> , 2013, 82, 1416-1422.	1.2	18
131	Can we test for hereditary cancer at 18 years when we start surveillance at 25? Patient reported outcomes. <i>Familial Cancer</i> , 2013, 12, 675-682.	0.9	15
132	Familial colorectal cancer risk assessment needs improvement for more effective cancer prevention in relatives. <i>Colorectal Disease</i> , 2013, 15, e175-85; discussion p.e185.	0.7	19
133	Identification of germline mutations in the cancer predisposing gene CDH1 in patients with orofacial clefts. <i>Human Molecular Genetics</i> , 2013, 22, 919-926.	1.4	55
134	Focusing on Patient Needs and Preferences May Improve Genetic Counseling for Colorectal Cancer. <i>Journal of Genetic Counseling</i> , 2013, 22, 118-124.	0.9	10
135	Comparability versus statistical correctness. <i>European Journal of Radiology</i> , 2013, 82, e908.	1.2	1
136	Added Value of Family History in Counseling About Risk of BRCA1/2 Mutation in Early-Onset Epithelial Ovarian Cancer. <i>International Journal of Gynecological Cancer</i> , 2013, 23, 1406-1410.	1.2	4
137	Exome Sequencing of Germline DNA from Non-BRCA1/2 Familial Breast Cancer Cases Selected on the Basis of aCGH Tumor Profiling. <i>PLoS ONE</i> , 2013, 8, e55734.	1.1	29
138	A Nonsynonymous Polymorphism in <i>IRS1</i> Modifies Risk of Developing Breast and Ovarian Cancers in <i>BRCA1</i> and Ovarian Cancer in <i>BRCA2</i> Mutation Carriers. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2012, 21, 1362-1370.	1.1	23
139	Risks of Less Common Cancers in Proven Mutation Carriers With Lynch Syndrome. <i>Journal of Clinical Oncology</i> , 2012, 30, 4409-4415.	0.8	262
140	Differences in Natural History between Breast Cancers in <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers and Effects of MRI Screening-MRISC, MARIBS, and Canadian Studies Combined. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2012, 21, 1458-1468.	1.1	79
141	Pathology of Breast and Ovarian Cancers among <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers: Results from the Consortium of Investigators of Modifiers of <i>BRCA1/2</i> (CIMBA). <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2012, 21, 134-147.	1.1	513
142	Higher cytoplasmic and nuclear poly(ADP-ribose) polymerase expression in familial than in sporadic breast cancer. <i>Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin</i> , 2012, 461, 425-431.	1.4	4
143	Rare variants in XRCC2 as breast cancer susceptibility alleles: Table 1. <i>Journal of Medical Genetics</i> , 2012, 49, 618-620.	1.5	49
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300	Monocytes from patients with combined hypercholesterolemia-hypertriglyceridemia and isolated hypercholesterolemia show an increased adhesion to endothelial cells in vitro: II. Influence of intrinsic and extrinsic factors on monocyte binding. <i>Metabolism: Clinical and Experimental</i> , 1995, 44, 374-378.	1.5	22
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302	Hyperglycemia in the acute phase of stroke is not caused by stress.. <i>Stroke</i> , 1993, 24, 1129-1132.	1.0	126
303	The effect of growth hormone administration in growth hormone deficient adults on bone, protein, carbohydrate and lipid homeostasis, as well as on body composition. <i>Clinical Endocrinology</i> , 1992, 37, 79-87.	1.2	328
304	Patients with combined hypercholesterolemia-hypertriglyceridemia show an increased monocyte-endothelial cell adhesion in vitro: Triglyceride level as a major determinant. <i>Metabolism: Clinical and Experimental</i> , 1991, 40, 1119-1121.	1.5	34
305	The efficacy and safety of pravastatin, compared to and in combination with bile acid binding resins, in familial hypercholesterolaemia. <i>Journal of Internal Medicine</i> , 1990, 228, 261-266.	2.7	37
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