A M Lucassen

List of Publications by Year in Descending Order

182 papers	7,976 citations	43 h-index	86 g-index
195 ext. papers	9,061 ext. citations	6.7 avg, IF	5.61 L-index

#	Paper	IF	Citations
182	Re-imagining 'the patient': Linked lives and lessons from genomic medicine.. *Social Science and Medicine*, **2022**, 297, 114806	5.1	1
181	Public Trust and Trustworthiness in Biobanking: The Need for More Reflexivity.. *Biopreservation and Biobanking*, **2022**,	2.1	1
180	Sustainable biobanks: a case study for a green global bioethics.. *Global Bioethics*, **2022**, 33, 50-64	2.8	3
179	Ethical Considerations in Research with Genomic Data.. *New Bioethics*, **2022**, 1-15	1	
178	GA4GH: International policies and standards for data sharing across genomic research and healthcare.. *Cell Genomics*, **2021**, 1, 100029-100029		20
177	Care of men with cancer-predisposing variants. *BMJ, The*, **2021**, 375, n2376	5.9	0
176	Using a biomarker acutely to identify babies at risk of serious adverse effects from antibiotics: where is the 'Terrible Moral and Medical Dilemma'?. *Journal of Medical Ethics*, **2021**, 47, 117-118	2.5	
175	Mitochondrial DNA variants in genomic data: diagnostic uplifts and predictive implications. *Nature Reviews Genetics*, **2021**, 22, 547-548	30.1	
174	Couple-based expanded carrier screening provided by general practitioners to couples in the Dutch general population: psychological outcomes and reproductive intentions. *Genetics in Medicine*, **2021**, 23, 1761-1768	8.1	1
173	What is the meaning of a 'genomic result' in the context of pregnancy?. *European Journal of Human Genetics*, **2021**, 29, 225-230	5.3	1
172	Variation in the risk of colorectal cancer in families with Lynch syndrome: a retrospective cohort study. *Lancet Oncology, The*, **2021**, 22, 1014-1022	21.7	5
171	In the family: access to, and communication of, familial information in clinical practice. *Human Genetics*, **2021**, 1	6.3	1
170	From Beyond the Grave: Use of Medical Information from the Deceased to Guide Care of Living Relatives. *Current Genetic Medicine Reports*, **2020**, 8, 1-7	2.2	0
169	Cognitive and affective outcomes of genetic counselling in the Netherlands at group and individual level: a personalized approach seems necessary. *European Journal of Human Genetics*, **2020**, 28, 1187-1195	5.3	4
168	Old consent and new developments: health professionals should ask and not presume. *Journal of Medical Ethics*, **2020**, 46, 412-413	2.5	1
167	Exploring broad consent in the context of the 100,000 Genomes Project: a mixed methods study. *European Journal of Human Genetics*, **2020**, 28, 732-741	5.3	8
166	The Need for Machine-Processable Agreements in Health Data Management. *Algorithms*, **2020**, 13, 87	1.8	1

165	International perspectives on the implementation of reproductive carrier screening. *Prenatal Diagnosis*, **2020**, 40, 301-310	3.2	29
164	Using biomarkers in acute medicine to prevent hearing loss: should this require specific consent?. *Journal of Medical Ethics*, **2020**, 46, 536-537	2.5	4
163	Genome sequencing in healthcare: understanding the UK general public's views and implications for clinical practice. *European Journal of Human Genetics*, **2020**, 28, 155-164	5.3	3
162	GP-provided couple-based expanded preconception carrier screening in the Dutch general population: who accepts the test-offer and why?. *European Journal of Human Genetics*, **2020**, 28, 182-192	5.3	6
161	A validated PROM in genetic counselling: the psychometric properties of the Dutch version of the Genetic Counselling Outcome Scale. *European Journal of Human Genetics*, **2019**, 27, 681-690	5.3	6
160	Unpacking the Concept of a Genomic Result. *American Journal of Bioethics*, **2019**, 19, 70-71	1.1	4
159	Familial genetic risks: how can we better navigate patient confidentiality and appropriate risk disclosure to relatives?. *Journal of Medical Ethics*, **2019**, 45, 504-507	2.5	18
158	Recent developments in genetic/genomic medicine. *Clinical Science*, **2019**, 133, 697-708	6.5	42
157	The moral argument for heritable genome editing requires an inappropriately deterministic view of genetics. *Journal of Medical Ethics*, **2019**, 45, 526-527	2.5	3
156	Feasibility of couple-based expanded carrier screening offered by general practitioners. *European Journal of Human Genetics*, **2019**, 27, 691-700	5.3	20
155	When genomic medicine reveals misattributed genetic relationships-the debate about disclosure revisited. *Genetics in Medicine*, **2019**, 21, 97-101	8.1	4
154	Consent and Autonomy in the Genomics Era. *Current Genetic Medicine Reports*, **2019**, 7, 85-91	2.2	16
153	European recommendations integrating genetic testing into multidisciplinary management of sudden cardiac death. *European Journal of Human Genetics*, **2019**, 27, 1763-1773	5.3	31
152	Genomic testing in healthcare: a hybrid space where clinical practice and research need to co-exist. *Expert Review of Molecular Diagnostics*, **2019**, 19, 963-967	3.8	3
151	Genomic variant sharing: a position statement. *Wellcome Open Research*, **2019**, 4, 22	4.8	16
150	Potential for diagnosis of infectious disease from the 100,000 Genomes Project Metagenomic Dataset: Recommendations for reporting results. *Wellcome Open Research*, **2019**, 4, 155	4.8	4
149	Direct-to-consumer genetic testing with third party interpretation: beware of spurious results. *Emerging Topics in Life Sciences*, **2019**, 3, 669-674	3.5	4
148	Expanded carrier screening for autosomal recessive conditions in health care: Arguments for a couple-based approach and examination of couples' views. *Prenatal Diagnosis*, **2019**, 39, 369-378	3.2	17

147	Is it acceptable to contact an anonymous egg donor to facilitate diagnostic genetic testing for the donor-conceived child?. *Journal of Medical Ethics*, **2019**, 45, 357-360	2.5	6
146	Reinterpretation, reclassification, and its downstream effects: challenges for clinical laboratory geneticists. *BMC Medical Genomics*, **2019**, 12, 170	3.7	15
145	Recontacting patients in clinical genetics services: recommendations of the European Society of Human Genetics. *European Journal of Human Genetics*, **2019**, 27, 169-182	5.3	34
144	Fostering trust in healthcare: Participants' experiences, views, and concerns about the 100,000 genomes project. *European Journal of Medical Genetics*, **2019**, 62, 335-341	2.6	5
143	Recontacting or not recontacting? A survey of current practices in clinical genetics centres in Europe. *European Journal of Human Genetics*, **2018**, 26, 946-954	5.3	19
142	Towards a national genomics medicine service: the challenges facing clinical-research hybrid practices and the case of the 100 000 genomes project. *Journal of Medical Ethics*, **2018**, 44, 397-403	2.5	18
141	The Road to Clinical Fantasy: A UK Perspective. *American Journal of Bioethics*, **2018**, 18, 26-27	1.1	2
140	How do clinical genetics consent forms address the familial approach to confidentiality and incidental findings? A mixed-methods study. *Familial Cancer*, **2018**, 17, 155-166	3	9
139	Using a genetic test result in the care of family members: how does the duty of confidentiality apply?. *European Journal of Human Genetics*, **2018**, 26, 955-959	5.3	17
138	Limitations and Pitfalls of Using Family Letters to Communicate Genetic Risk: a Qualitative Study with Patients and Healthcare Professionals. *Journal of Genetic Counseling*, **2018**, 27, 689-701	2.5	24
137	Disclosure of genetic information to relatives: balancing confidentiality and relatives' interests. *Journal of Medical Genetics*, **2018**, 55, 285-286	5.8	8
136	Dimensions of responsibility in medical genetics: exploring the complexity of the duty to recontact *New Genetics and Society*, **2018**, 37, 187-206	1.9	3
135	In Defense of Best Interests: When Parents and Clinicians Disagree. *American Journal of Bioethics*, **2018**, 18, 67-69	1.1	2
134	Alerting relatives about heritable risks: the limits of confidentiality. *BMJ, The*, **2018**, 361, k1409	5.9	14
133	Predictive Genetic Testing of Children for Adult-Onset Conditions: Negotiating Requests with Parents. *Journal of Genetic Counseling*, **2017**, 26, 244-250	2.5	8
132	Recontacting in clinical genetics and genomic medicine? We need to talk about it. *European Journal of Human Genetics*, **2017**, 25, 520-521	5.3	13
131	Expanded carrier screening: what determines intended participation and can this be influenced by message framing and narrative information?. *European Journal of Human Genetics*, **2017**, 25, 793-800	5.3	11
130	A 'joint venture' model of recontacting in clinical genomics: challenges for responsible implementation. *European Journal of Medical Genetics*, **2017**, 60, 403-409	2.6	23

129	Recontacting in clinical practice: an investigation of the views of healthcare professionals and clinical scientists in the United Kingdom. *European Journal of Human Genetics*, **2017**, 25, 275-279	5.3	24
128	Approaching confidentiality at a familial level in genomic medicine: a focus group study with healthcare professionals. *BMJ Open*, **2017**, 7, e012443	3	27
127	Healthcare professionals' and patients' perspectives on consent to clinical genetic testing: moving towards a more relational approach. *BMC Medical Ethics*, **2017**, 18, 47	2.9	34
126	Ethical issues in genetic medicine. *InnovAiT*, **2017**, 10, 481-488	0	1
125	Recontacting in clinical practice: the views and expectations of patients in the United Kingdom. *European Journal of Human Genetics*, **2017**, 25, 1106-1112	5.3	17
124	Health-care professionals' responsibility to patients' relatives in genetic medicine: a systematic review and synthesis of empirical research. *Genetics in Medicine*, **2016**, 18, 290-301	8.1	56
123	Population-based preconception carrier screening: how potential users from the general population view a test for 50 serious diseases. *European Journal of Human Genetics*, **2016**, 24, 1417-23	5.3	50
122	Responsible implementation of expanded carrier screening. *European Journal of Human Genetics*, **2016**, 24, e1-e12	5.3	167
121	Relative Risk and Relatives' Risks in Genomic Medicine. *American Journal of Bioethics*, **2016**, 16, 25-7	1.1	8
120	Telemedicine uptake among Genetics Professionals in Europe: room for expansion. *European Journal of Human Genetics*, **2016**, 24, 157-63	5.3	26
119	Genomic Analysis in Clinical Practice: What Are the Challenges? **2016**, 191-199		
118	Management of Incidental Findings in Clinical Genomic Sequencing Studies **2016**, 1-7		5
117	What results to disclose, when, and who decides? Healthcare professionals' views on prenatal chromosomal microarray analysis. *Prenatal Diagnosis*, **2016**, 36, 252-9	3.2	24
116	'Is this knowledge mine and nobody else's? I don't feel that.' Patient views about consent, confidentiality and information-sharing in genetic medicine. *Journal of Medical Ethics*, **2016**, 42, 174-9	2.5	43
115	Recontact in clinical practice: a survey of clinical genetics services in the United Kingdom. *Genetics in Medicine*, **2016**, 18, 876-81	8.1	29
114	Is there a duty to recontact in light of new genetic technologies? A systematic review of the literature. *Genetics in Medicine*, **2015**, 17, 668-78	8.1	55
113	Rescue obligations and collective approaches: complexities in genomics. *American Journal of Bioethics*, **2015**, 15, 23-5	1.1	2
112	Non-invasive prenatal testing for aneuploidy and beyond: challenges of responsible innovation in prenatal screening. Summary and recommendations. *European Journal of Human Genetics*, **2015**,	5.3	9

111	Non-invasive prenatal testing for aneuploidy and beyond: challenges of responsible innovation in prenatal screening. *European Journal of Human Genetics*, **2015**, 23, 1438-50	5.3	189
110	Genetic testing of children for adult-onset conditions: opinions of the British adult population and implications for clinical practice. *European Journal of Human Genetics*, **2015**, 23, 1281-5	5.3	16
109	I Had Genetic Testing for Alzheimer's Disease Without My Consent. *Narrative Inquiry in Bioethics*, **2015**, 5, 214-6	0.2	2
108	Prenatal diagnosis of chromosomal imbalances. *Archives of Disease in Childhood: Fetal and Neonatal Edition*, **2014**, 99, F338-41	4.7	4
107	Rare variants in NR2F2 cause congenital heart defects in humans. *American Journal of Human Genetics*, **2014**, 94, 574-85	11	115
106	The challenges of genome analysis in the health care setting. *Genes*, **2014**, 5, 576-85	4.2	20
105	Genetic testing of children: the need for a family perspective. *American Journal of Bioethics*, **2014**, 14, 26-8	1.1	7
104	Defining and managing incidental findings in genetic and genomic practice. *Journal of Medical Genetics*, **2014**, 51, 715-23	5.8	46
103	A more fitting term in the incidental findings debate: one term does not fit all situations. *European Journal of Human Genetics*, **2014**, 22, 957	5.3	8
102	Genetic medicine and incidental findings: it is more complicated than deciding whether to disclose or not. *Genetics in Medicine*, **2013**, 15, 896-9	8.1	26
101	Next generation diagnostics in inherited arrhythmia syndromes : a comparison of two approaches. *Journal of Cardiovascular Translational Research*, **2013**, 6, 94-103	3.3	28
100	Germline mutations affecting the proofreading domains of POLE and POLD1 predispose to colorectal adenomas and carcinomas. *Nature Genetics*, **2013**, 45, 136-44	36.3	686
99	Developing a policy for paediatric biobanks: principles for good practice. *European Journal of Human Genetics*, **2013**, 21, 2-7	5.3	52
98	A primary care specialist genetics service: a cluster-randomised factorial trial. *British Journal of General Practice*, **2012**, 62, e191-7	1.6	8
97	Genomic medicine: challenges and opportunities for physicians. *Clinical Medicine*, **2012**, 12, 416-9	1.9	14
96	Testing children for adult onset conditions: the importance of contextual clinical judgement. *Journal of Medical Ethics*, **2012**, 38, 531-2; discussion 533-4	2.5	8
95	The impact of cancer pathology confirmation on clinical management of a family history of cancer. *Familial Cancer*, **2011**, 10, 373-80	3	6
94	No evidence of RET germline mutations in familial pituitary adenoma. *Journal of Molecular Endocrinology*, **2011**, 46, 1-8	4.5	32

93	The shifting sands of patient autonomy and public interest considerations in health care. *Clinical Ethics*, **2011**, 6, 203-206	1	2
92	Mainstreaming genetics: the potential for miscommunication. *Clinical Ethics*, **2011**, 6, 159-161	1	
91	Clinical Ethics Committee Case 14: How should we transfer a euthanasia request between general practice and a hospital setting?. *Clinical Ethics*, **2011**, 6, 58-63	1	1
90	Meta-analysis of three genome-wide association studies identifies susceptibility loci for colorectal cancer at 1q41, 3q26.2, 12q13.13 and 20q13.33. *Nature Genetics*, **2010**, 42, 973-7	36.3	301
89	Legal implications of tissue. *Annals of the Royal College of Surgeons of England*, **2010**, 92, 189-92	1.4	1
88	Family history and adoption in the UK: conflicts of interest in medical disclosure. *Archives of Disease in Childhood*, **2010**, 95, 7-11	2.2	6
87	An investigation of patients' motivations for their participation in genetics-related research. *Journal of Medical Ethics*, **2010**, 36, 37-45	2.5	84
86	'Ethnicity testing' before adoption: a help or hindrance?. *Archives of Disease in Childhood*, **2010**, 95, 404-5	2.2	2
85	Lay and Professional Understandings of Research and Clinical Activities in Cancer Genetics and Their Implications for Informed Consent. *American Journal of Bioethics Primary Research*, **2010**, 1, 25-34		3
84	Confidentiality and sharing genetic information with relatives. *Lancet, The*, **2010**, 375, 1507-9	40	22
83	Guidelines for colorectal cancer screening and surveillance in moderate and high risk groups (update from 2002). *Gut*, **2010**, 59, 666-89	19.2	843
82	What facilitates or impedes family communication following genetic testing for cancer risk? A systematic review and meta-synthesis of primary qualitative research. *Journal of Genetic Counseling*, **2010**, 19, 330-42	2.5	109
81	Predictive genetic testing in a young child: a case report. *Familial Cancer*, **2010**, 9, 61-4	3	2
80	Predictive genetic testing in children: where are we now? An overview and a UK perspective. *Familial Cancer*, **2010**, 9, 3-7	3	7
79	Genetic testing from different angles. Introduction. *Familial Cancer*, **2010**, 9, 1	3	1
78	Ethicolegal aspects of genetics in surgical practice. *Annals of the Royal College of Surgeons of England*, **2009**, 91, 451-5	1.4	
77	'Over-the-counter' genetic testing: what does it really mean for primary care?. *British Journal of General Practice*, **2009**, 59, 283-7	1.6	31
76	Healthcare professionals' and researchers' understanding of cancer genetics activities: a qualitative interview study. *Journal of Medical Ethics*, **2009**, 35, 113-9	2.5	19

75	Distinguishing research from clinical care in cancer genetics: theoretical justifications and practical strategies. *Social Science and Medicine*, **2009**, 68, 2010-7	5.1	39
74	Missed threads. The impact of pre-mRNA splicing defects on clinical practice. *EMBO Reports*, **2009**, 10, 810-6	6.5	89
73	Risk reducing mastectomy: outcomes in 10 European centres. *Journal of Medical Genetics*, **2009**, 46, 254-8	5.8	73
72	Two children with subtelomeric 11q deletions: a description and interpretation of their clinical presentations and molecular genetic findings. *Clinical Dysmorphology*, **2009**, 18, 98-102	0.9	2
71	Deciphering the genetics of hereditary non-syndromic colorectal cancer. *European Journal of Human Genetics*, **2008**, 16, 1477-86	5.3	29
70	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
69	Refinement of the basis and impact of common 11q23.1 variation to the risk of developing colorectal cancer. *Human Molecular Genetics*, **2008**, 17, 3720-7	5.6	57
68	Unregulated smooth-muscle myosin in human intestinal neoplasia. *Proceedings of the National Academy of Sciences of the United States of America*, **2008**, 105, 5513-8	11.5	64
67	Ethical Issues in Genetic Medicine. *InnovAiT*, **2008**, 1, 589-595	0	
66	Recall of participation in research projects in cancer genetics: some implications for research ethics. *Clinical Ethics*, **2008**, 3, 180-184	1	9
65	Ethics and research governance: the views of researchers, health-care professionals and other stakeholders. *Clinical Ethics*, **2008**, 3, 85-90	1	10
64	Disclosure of genetic information within families: a case report. *Clinical Ethics*, **2008**, 3, 7-10	1	
63	Large genomic deletions in AIP in pituitary adenoma predisposition. *Journal of Clinical Endocrinology and Metabolism*, **2008**, 93, 4146-51	5.6	66
62	Should families own genetic information? Yes. *BMJ, The*, **2007**, 335, 22	5.9	11
61	Surveillance for familial breast cancer: Differences in outcome according to BRCA mutation status. *International Journal of Cancer*, **2007**, 121, 1017-20	7.5	76
60	Germline CDKN1B/p27Kip1 mutation in multiple endocrine neoplasia. *Journal of Clinical Endocrinology and Metabolism*, **2007**, 92, 3321-5	5.6	230
59	Cowden syndrome and Bannayan Riley Ruvalcaba syndrome represent one condition with variable expression and age-related penetrance: results of a clinical study of PTEN mutation carriers. *Journal of Medical Genetics*, **2007**, 44, 579-85	5.8	128
58	Implications of data protection legislation for family history. *BMJ, The*, **2006**, 332, 299-301	5.9	11

57	Genetic testing without consent: the implications of the new Human Tissue Act 2004. *Journal of Medical Ethics*, **2006**, 32, 690-2	2.5	6
56	Evidence for a colorectal cancer susceptibility locus on chromosome 3q21-q24 from a high-density SNP genome-wide linkage scan. *Human Molecular Genetics*, **2006**, 15, 2903-10	5.6	50
55	The UK Genethics Club: clinical ethics support for genetic services. *Clinical Ethics*, **2006**, 1, 219-223	1	2
54	Applying a cognitive behavioral model of health anxiety in a cancer genetics service. *Health Psychology*, **2006**, 25, 171-80	5	22
53	Guilt, blame and responsibility: men's understanding of their role in the transmission of BRCA1/2 mutations within their family. *Sociology of Health and Illness*, **2006**, 28, 969-88	3	65
52	Feasibility and acceptability of providing nurse counsellor genetics clinics in primary care. *Journal of Advanced Nursing*, **2006**, 53, 591-604	3.1	19
51	Exonic STK11 deletions are not a rare cause of Peutz-Jeghers syndrome. *Journal of Medical Genetics* , **2006**, 43, e15	5.8	46
50	Family history of breast cancer. *BMJ, The*, **2005**, 330, 26	5.9	3
49	Deletions involving long-range conserved nongenic sequences upstream and downstream of FOXL2 as a novel disease-causing mechanism in blepharophimosis syndrome. *American Journal of Human Genetics*, **2005**, 77, 205-18	11	112
48	Communication about genetic testing in families of male BRCA1/2 carriers and non-carriers: patterns, priorities and problems. *Clinical Genetics*, **2005**, 67, 492-502	4	91
47	Dilemma still not resolved. *European Journal of Human Genetics*, **2005**, 13, 399-400	5.3	1
46	Genetic professionals' reports of nondisclosure of genetic risk information within families. *European Journal of Human Genetics*, **2005**, 13, 556-62	5.3	96
45	Sequence changes in predicted promoter elements of STK11/LKB1 are unlikely to contribute to Peutz-Jeghers syndrome. *BMC Genomics*, **2005**, 6, 38	4.5	11
44	Men's decision-making about predictive BRCA1/2 testing: the role of family. *Journal of Genetic Counseling*, **2005**, 14, 207-17	2.5	63
43	Families and genetic testing: the case of Jane and Phyllis **2005**, 7-26		2
42	Interpretation and dialogue in hermeneutic ethics **2005**, 57-76		10
41	Reading the genes **2005**, 95-114		
40	A virtue-ethics approach **2005**, 45-56		5

39	Response to ethical dissections of the case **2005**, 213-224		
38	Screening for familial ovarian cancer: failure of current protocols to detect ovarian cancer at an early stage according to the international Federation of gynecology and obstetrics system. *Journal of Clinical Oncology*, **2005**, 23, 5588-96	2.2	134
37	Role of next of kin in accessing health records of deceased relatives. *BMJ, The*, **2004**, 328, 952-3	5.9	5
36	Genetic information: a joint account?. *BMJ, The*, **2004**, 329, 165-7	5.9	91
35	Confidentiality and serious harm in genetics - preserving the confidentiality of one patient and preventing harm to relatives. *European Journal of Human Genetics*, **2004**, 12, 93-7	5.3	31
34	Mapping of a translocation breakpoint in a Peutz-Jeghers hamartoma to the putative PJS locus at 19q13.4 and mutation analysis of candidate genes in polyp and STK11-negative PJS cases. *Genes Chromosomes and Cancer*, **2004**, 41, 163-9	5	26
33	RNA analysis reveals splicing mutations and loss of expression defects in MLH1 and BRCA1. *Human Mutation*, **2004**, 24, 272	4.7	44
32	Improving the ascertainment of families at high risk of colorectal cancer: a prospective GP register study. *British Journal of General Practice*, **2004**, 54, 267-71	1.6	10
31	Further observations on LKB1/STK11 status and cancer risk in Peutz-Jeghers syndrome. *British Journal of Cancer*, **2003**, 89, 308-13	8.7	126
30	Concern for families and individuals in clinical genetics. *Journal of Medical Ethics*, **2003**, 29, 70-3	2.5	38
29	What motivates interest in attending a familial cancer genetics clinic?. *Familial Cancer*, **2003**, 2, 159-68	3	26
28	Education improves general practitioner (GP) management of familial breast/ovarian cancer: findings from a cluster randomised controlled trial. *Journal of Medical Genetics*, **2002**, 39, 779-81	5.8	17
27	Working towards ethical management of genetic testing. *Lancet, The*, **2002**, 360, 1685-8	40	19
26	Evidence based case report: Advice about mammography for a young woman with a family history of breast cancer. *BMJ: British Medical Journal*, **2001**, 322, 1040-2		13
25	Cystic fibrosis: A further case of an asymptomatic compound heterozygote. *American Journal of Medical Genetics Part A*, **2001**, 103, 342-343		4
24	A study of GP referrals to a family cancer clinic for breast/ovarian cancer. *Family Practice*, **2001**, 18, 131-4	1.9	44
23	Guidelines for referral to a regional genetics service: GPs respond by referring more appropriate cases. *Family Practice*, **2001**, 18, 135-40	1.9	54
22	Referral of patients with a family history of breast/ovarian cancer--GPs' knowledge and expectations. *Family Practice*, **2001**, 18, 487-90	1.9	43

21	Revealing false paternity: some ethical considerations. *Lancet, The*, **2001**, 357, 1033-5	40	76
20	Common hereditary cancers and implications for primary care. *Lancet, The*, **2001**, 358, 56-63	40	62
19	The Opinions, Expectations and Experiences of Women with a Family History of Breast Cancer Who Consult Their GP and Are Referred to Secondary Care. *Public Health Genomics*, **2001**, 4, 239-243	1.9	5
18	Evaluation of the impact of two educational interventions on GP management of familial breast/ovarian cancer cases: a cluster randomised controlled trial. *British Journal of General Practice* , **2001**, 51, 817-21	1.6	49
17	A comparison of methods currently used in clinical practice to estimate familial breast cancer risks. *Annals of Oncology*, **2000**, 11, 451-4	10.3	13
16	Clinical geneticists' attitudes and practice towards testing for breast cancer susceptibility genes. *Journal of Medical Genetics*, **2000**, 37, 157-60	5.8	7
15	Hereditary cancer. *Lancet Oncology, The*, **2000**, 1, 12-13	21.7	1
14	Hereditary cancer the evidence for current recommended management. *Lancet Oncology, The*, **2000**, 1, 9-16	21.7	14
13	Development and Evaluation of Educational Materials for Primary Care on Familial Breast and/or Ovarian Cancer. *Disease Markers*, **1999**, 15, 156-156	3.2	78
12	Ethical issues in genetics of mental disorders. *Lancet, The*, **1998**, 352, 1004-5	40	13
11	Will gene testing cut risk of familial colorectal cancer?. *Practitioner*, **1998**, 242, 306-10, 314		
10	The insulin gene region and susceptibility to insulin-dependent diabetes mellitus in four races; new insights from Afro-Caribbean race-specific haplotypes. *Autoimmunity*, **1997**, 26, 11-22	3	6
9	Genetic Screening for Breast Cancer?. *The Journal of the British Menopause Society*, **1997**, 3, 20-24		
8	Breast cancer: who is at risk?. *Practitioner*, **1997**, 241, 757-9, 762		
7	Susceptibility to human type 1 diabetes at IDDM2 is determined by tandem repeat variation at the insulin gene minisatellite locus. *Nature Genetics*, **1995**, 9, 284-92	36.3	640
6	Reply to Insulin expression: is VNTR allele 698 really anomalous? *Nature Genetics*, **1995**, 10, 379-380	36.3	5
5	Regulation of insulin gene expression by the IDDM associated, insulin locus haplotype. *Human Molecular Genetics*, **1995**, 4, 501-6	5.6	85
4	Multiple DNA variant association analysis: application to the insulin gene region in type I diabetes. *American Journal of Human Genetics*, **1994**, 55, 1247-54	11	31

3	Susceptibility to insulin dependent diabetes mellitus maps to a 4.1 kb segment of DNA spanning the insulin gene and associated VNTR. *Nature Genetics*, **1993**, 4, 305-10	36.3	226
2	Missense glucokinase mutation in maturity-onset diabetes of the young and mutation screening in late-onset diabetes. *Nature Genetics*, **1992**, 2, 153-6	36.3	131
1	Genomic variant sharing: a position statement. *Wellcome Open Research*,4, 22	4.8	5