

# Hans F A Vasen

## List of Publications by Year in descending order

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126  
papers

13,823  
citations

44444

50  
h-index

23841

115  
g-index

132  
all docs

132  
docs citations

132  
times ranked

11228  
citing authors

#	ARTICLE	IF	CITATIONS
1	Revised guidelines for the clinical management of Lynch syndrome (HNPCC): recommendations by a group of European experts. <i>Gut</i> , 2013, 62, 812-823.	6.1	630
2	Cancer risk in hereditary nonpolyposis colorectal cancer due to MSH6 mutations: impact on counseling and surveillance. <i>Gastroenterology</i> , 2004, 127, 17-25.	0.6	536
3	The risk of extra-colonic, extra-endometrial cancer in the Lynch syndrome. <i>International Journal of Cancer</i> , 2008, 123, 444-449.	2.3	481
4	Mutations Associated with HNPCC Predisposition – Update of ICG-HNPCC/INSiGHT Mutation Database. <i>Disease Markers</i> , 2004, 20, 269-276.	0.6	416
5	Cancer incidence and survival in Lynch syndrome patients receiving colonoscopic and gynaecological surveillance: first report from the prospective Lynch syndrome database. <i>Gut</i> , 2017, 66, 464-472.	6.1	411
6	Cancer risk and survival in <i>path_MMR</i> carriers by gene and gender up to 75 years of age: a report from the Prospective Lynch Syndrome Database. <i>Gut</i> , 2018, 67, 1306-1316.	6.1	410
7	Familial endometrial cancer in female carriers of MSH6 germline mutations. <i>Nature Genetics</i> , 1999, 23, 142-144.	9.4	378
8	Cancer risks by gene, age, and gender in 6350 carriers of pathogenic mismatch repair variants: findings from the Prospective Lynch Syndrome Database. <i>Genetics in Medicine</i> , 2020, 22, 15-25.	1.1	365
9	Management of patients with increased risk for familial pancreatic cancer: updated recommendations from the International Cancer of the Pancreas Screening (CAPS) Consortium. <i>Gut</i> , 2020, 69, 7-17.	6.1	357
10	Diagnostic criteria for constitutional mismatch repair deficiency syndrome: suggestions of the European consortium – Care for CMMRD™ (C4CMMRD). <i>Journal of Medical Genetics</i> , 2014, 51, 355-365.	1.5	351
11	Benefit of Surveillance for Pancreatic Cancer in High-Risk Individuals: Outcome of Long-Term Prospective Follow-Up Studies From Three European Expert Centers. <i>Journal of Clinical Oncology</i> , 2016, 34, 2010-2019.	0.8	280
12	Effect of Aspirin or Resistant Starch on Colorectal Neoplasia in the Lynch Syndrome. <i>New England Journal of Medicine</i> , 2008, 359, 2567-2578.	13.9	273
13	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
14	Decrease in Mortality in Lynch Syndrome Families Because of Surveillance. <i>Gastroenterology</i> , 2006, 130, 665-671.	0.6	246
15	One to 2-Year Surveillance Intervals Reduce Risk of Colorectal Cancer in Families With Lynch Syndrome. <i>Gastroenterology</i> , 2010, 138, 2300-2306.	0.6	219
16	The outcome of endometrial carcinoma surveillance by ultrasound scan in women at risk of hereditary nonpolyposis colorectal carcinoma and familial colorectal carcinoma. <i>Cancer</i> , 2002, 94, 1708-1712.	2.0	217
17	MSH2 genomic deletions are a frequent cause of HNPCC. <i>Nature Genetics</i> , 1998, 20, 326-328.	9.4	216
18	Risk and Epidemiological Time Trends of Gastric Cancer in Lynch Syndrome Carriers in The Netherlands. <i>Gastroenterology</i> , 2010, 138, 487-492.	0.6	201

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19	Microsatellite Instability, Immunohistochemistry, and Additional PMS2 Staining in Suspected Hereditary Nonpolyposis Colorectal Cancer. <i>Clinical Cancer Research</i> , 2004, 10, 972-980.	3.2	200
20	The role of mismatch repair gene defects in the development of adenomas in patients with HNPCC. <i>Gastroenterology</i> , 2004, 126, 42-48.	0.6	199
21	The tumour spectrum in hereditary non-polyposis colorectal cancer: A study of 24 kindreds in the Netherlands. <i>International Journal of Cancer</i> , 1990, 46, 31-34.	2.3	182
22	Prediction of a mismatch repair gene defect by microsatellite instability and immunohistochemical analysis in endometrial tumours from HNPCC patients. <i>Journal of Pathology</i> , 2000, 192, 328-335.	2.1	168
23	MUTYH-associated polyposis (MAP). <i>Critical Reviews in Oncology/Hematology</i> , 2011, 79, 1-16.	2.0	163
24	Conventional and Tissue Microarray Immunohistochemical Expression Analysis of Mismatch Repair in Hereditary Colorectal Tumors. <i>American Journal of Pathology</i> , 2003, 162, 469-477.	1.9	159
25	A nationwide study comparing sporadic and familial adenomatous polyposis-related desmoid-type fibromatoses. <i>International Journal of Cancer</i> , 2011, 129, 256-261.	2.3	154
26	Magnetic Resonance Imaging Surveillance Detects Early-Stage Pancreatic Cancer in Carriers of a p16-Leiden Mutation. <i>Gastroenterology</i> , 2011, 140, 850-856.	0.6	148
27	Cancer Risks for PMS2-Associated Lynch Syndrome. <i>Journal of Clinical Oncology</i> , 2018, 36, 2961-2968.	0.8	147
28	Heterozygous Mutations in PMS2 Cause Hereditary Nonpolyposis Colorectal Carcinoma (Lynch) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 3	0.6	132
29	Incidence of and survival after subsequent cancers in carriers of pathogenic MMR variants with previous cancer: a report from the prospective Lynch syndrome database. <i>Gut</i> , 2017, 66, 1657-1664.	6.1	127
30	Characteristics of small bowel carcinoma in hereditary nonpolyposis colorectal carcinoma. , 1998, 83, 240-244.		125
31	Survival analysis of endometrial carcinoma associated with hereditary nonpolyposis colorectal cancer. <i>International Journal of Cancer</i> , 2002, 102, 198-200.	2.3	119
32	Cancer risk and genotype-phenotype correlations in PTEN hamartoma tumor syndrome. <i>Familial Cancer</i> , 2014, 13, 57-63.	0.9	119
33	Family History, Surgery, and APC Mutation Are Risk Factors for Desmoid Tumors in Familial Adenomatous Polyposis: An International Cohort Study. <i>Diseases of the Colon and Rectum</i> , 2011, 54, 1229-1234.	0.7	115
34	Survival after adjuvant 5-FU treatment for stage III colon cancer in hereditary nonpolyposis colorectal cancer. <i>International Journal of Cancer</i> , 2004, 109, 468-471.	2.3	105
35	Germline variants in POLE are associated with early onset mismatch repair deficient colorectal cancer. <i>European Journal of Human Genetics</i> , 2015, 23, 1080-1084.	1.4	101
36	Clinical management of hereditary colorectal cancer syndromes. <i>Nature Reviews Gastroenterology and Hepatology</i> , 2015, 12, 88-97.	8.2	99

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37	Functional Outcome After Colectomy and Ileorectal Anastomosis Compared With Proctocolectomy and Ileal Pouch–Anal Anastomosis in Familial Adenomatous Polyposis. <i>Annals of Surgery</i> , 1999, 230, 648.	2.1	99
38	Long-term effect of resistant starch on cancer risk in carriers of hereditary colorectal cancer: an analysis from the CAPP2 randomised controlled trial. <i>Lancet Oncology</i> , The, 2012, 13, 1242-1249.	5.1	95
39	Desmoid Tumors in a Dutch Cohort of Patients With Familial Adenomatous Polyposis. <i>Clinical Gastroenterology and Hepatology</i> , 2008, 6, 215-219.	2.4	83
40	Chromosome 8q23.3 and 11q23.1 Variants Modify Colorectal Cancer Risk in Lynch Syndrome. <i>Gastroenterology</i> , 2009, 136, 131-137.	0.6	80
41	Quality of Life After Surgery for Colon Cancer in Patients With Lynch Syndrome. <i>Diseases of the Colon and Rectum</i> , 2012, 55, 653-659.	0.7	80
42	Long Term Follow-up of HNPCC Gene Mutation Carriers: Compliance with Screening and Satisfaction with Counseling and Screening Procedures. <i>Familial Cancer</i> , 2005, 4, 295-300.	0.9	79
43	The Natural Course of Multiple Endocrine Neoplasia Type iib. <i>Archives of Internal Medicine</i> , 1992, 152, 1250.	4.3	78
44	Diagnosis of Constitutional Mismatch Repair-Deficiency Syndrome Based on Microsatellite Instability and Lymphocyte Tolerance to Methylating Agents. <i>Gastroenterology</i> , 2015, 149, 1017-1029.e3.	0.6	76
45	Prospective Results of Surveillance Colonoscopy in Dominant Familial Colorectal Cancer With and Without Lynch Syndrome. <i>Gastroenterology</i> , 2006, 130, 1995-2000.	0.6	72
46	Small-bowel cancer in Lynch syndrome: is it time for surveillance?. <i>Lancet Oncology</i> , The, 2008, 9, 901-905.	5.1	65
47	Body Mass Index Increases Risk of Colorectal Adenomas in Men With Lynch Syndrome: The GEOLynch Cohort Study. <i>Journal of Clinical Oncology</i> , 2010, 28, 4346-4353.	0.8	62
48	Survival in women with MMR mutations and ovarian cancer: a multicentre study in Lynch syndrome kindreds. <i>Journal of Medical Genetics</i> , 2010, 47, 99-102.	1.5	61
49	Surveillance for hereditary cancer: Does the benefit outweigh the psychological burden?â€”A systematic review. <i>Critical Reviews in Oncology/Hematology</i> , 2012, 83, 329-340.	2.0	60
50	Variation in Precursor Lesions of Pancreatic Cancer among High-Risk Groups. <i>Clinical Cancer Research</i> , 2013, 19, 442-449.	3.2	58
51	The “unnatural” history of colorectal cancer in Lynch syndrome: Lessons from colonoscopy surveillance. <i>International Journal of Cancer</i> , 2021, 148, 800-811.	2.3	55
52	Colorectal cancer incidence in path_MLH1 carriers subjected to different follow-up protocols: a Prospective Lynch Syndrome Database report. <i>Hereditary Cancer in Clinical Practice</i> , 2017, 15, 18.	0.6	49
53	Prevalence of small-bowel neoplasia in Lynch syndrome assessed by video capsule endoscopy. <i>Gut</i> , 2015, 64, 1578-1583.	6.1	47
54	Smoking Increases the Risk for Colorectal Adenomas in Patients With Lynch Syndrome. <i>Gastroenterology</i> , 2012, 142, 241-247.	0.6	44

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55	Disease expression in juvenile polyposis syndrome: a retrospective survey on a cohort of 221 European patients and comparison with a literature-derived cohort of 473 SMAD4/BMPR1A pathogenic variant carriers. <i>Genetics in Medicine</i> , 2020, 22, 1524-1532.	1.1	44
56	Germ Line Mutations of Mismatch Repair Genes in Hereditary Nonpolyposis Colorectal Cancer Patients with Small Bowel Cancer: International Society for Gastrointestinal Hereditary Tumours Collaborative Study: Table 1.. <i>Clinical Cancer Research</i> , 2006, 12, 3389-3393.	3.2	42
57	Lack of association between screening interval and cancer stage in Lynch syndrome may be accounted for by over-diagnosis; a prospective Lynch syndrome database report. <i>Hereditary Cancer in Clinical Practice</i> , 2019, 17, 8.	0.6	42
58	Attitudes toward genetic testing in childhood and reproductive decision-making for familial adenomatous polyposis. <i>European Journal of Human Genetics</i> , 2010, 18, 186-193.	1.4	38
59	Loss-of-Function Mutations in the Cell-Cycle Control Gene <i>CDKN2A</i> Impact on Glucose Homeostasis in Humans. <i>Diabetes</i> , 2016, 65, 527-533.	0.3	38
60	Dietary patterns and colorectal adenomas in Lynch syndrome. <i>Cancer</i> , 2013, 119, 512-521.	2.0	37
61	Colorectal surveillance in Lynch syndrome families. <i>Familial Cancer</i> , 2013, 12, 261-265.	0.9	36
62	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
63	Combined analysis of three lynch syndrome cohorts confirms the modifying effects of 8q23.3 and 11q23.1 in MLH1 mutation carriers. <i>International Journal of Cancer</i> , 2013, 132, 1556-1564.	2.3	33
64	Prospective risk of cancer and the influence of tobacco use in carriers of the p16-Leiden germline variant. <i>European Journal of Human Genetics</i> , 2015, 23, 711-714.	1.4	29
65	Risk-reducing hysterectomy and bilateral salpingo-oophorectomy in female heterozygotes of pathogenic mismatch repair variants: a Prospective Lynch Syndrome Database report. <i>Genetics in Medicine</i> , 2021, 23, 705-712.	1.1	28
66	Hereditary cancer registries improve the care of patients with a genetic predisposition to cancer: contributions from the Dutch Lynch syndrome registry. <i>Familial Cancer</i> , 2016, 15, 429-435.	0.9	25
67	Recurrent Coding Sequence Variation Explains Only A Small Fraction of the Genetic Architecture of Colorectal Cancer. <i>Scientific Reports</i> , 2015, 5, 16286.	1.6	24
68	Targeted next-generation sequencing of FNA-derived DNA in pancreatic cancer. <i>Journal of Clinical Pathology</i> , 2017, 70, 174-178.	1.0	24
69	A pooled analysis of the outcome of prospective colonoscopic surveillance for familial colorectal cancer. <i>International Journal of Cancer</i> , 2014, 134, 939-947.	2.3	22
70	Pancreatic carcinoma in carriers of a specific 19 base pair deletion of <i>CDKN2A/p16</i> (p16-leiden). <i>Clinical Cancer Research</i> , 2003, 9, 3598-605.	3.2	22
71	The natural history of a combined defect in <i>MSH6</i> and <i>MUTYH</i> in a HNPCC family. <i>Familial Cancer</i> , 2007, 6, 43-51.	0.9	21
72	Randomized Comparison of Surveillance Intervals in Familial Colorectal Cancer. <i>Journal of Clinical Oncology</i> , 2015, 33, 4188-4193.	0.8	21

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73	Duodenal Adenomas and Cancer in MUTYH-associated Polyposis: An International Cohort Study. <i>Gastroenterology</i> , 2021, 160, 952-954.e4.	0.6	20
74	Equivalent <i>Helicobacter pylori</i> infection rates in Lynch syndrome mutation carriers with and without a first-degree relative with gastric cancer. <i>International Journal of Colorectal Disease</i> , 2016, 31, 693-697.	1.0	18
75	Cost-utility analysis of genetic screening in families of patients with germline MUTYH mutations. <i>BMC Medical Genetics</i> , 2007, 8, 42.	2.1	17
76	Transanal minimally invasive surgery (TAMIS) versus endoscopic submucosal dissection (ESD) for resection of non-pedunculated rectal lesions (TRIASSIC study): study protocol of a European multicenter randomised controlled trial. <i>BMC Gastroenterology</i> , 2020, 20, 225.	0.8	17
77	Incidence of small bowel neoplasia in Lynch syndrome assessed by video capsule endoscopy. <i>Endoscopy International Open</i> , 2017, 05, E622-E626.	0.9	16
78	Lynch syndrome—how should colorectal cancer be managed?. <i>Nature Reviews Gastroenterology and Hepatology</i> , 2011, 8, 184-186.	8.2	15
79	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
80	Psychological distress and quality of life following positive fecal occult blood testing in colorectal cancer screening. <i>Psycho-Oncology</i> , 2020, 29, 1084-1091.	1.0	15
81	Skin self-examination of persons from families with familial atypical multiple mole melanoma (FAMMM). <i>Patient Education and Counseling</i> , 2009, 75, 251-255.	1.0	14
82	Clinical Perspective on Proteomic and Glycomic Biomarkers for Diagnosis, Prognosis, and Prediction of Pancreatic Cancer. <i>International Journal of Molecular Sciences</i> , 2021, 22, 2655.	1.8	14
83	Dietary B vitamin and methionine intake and MTHFR C677T genotype on risk of colorectal tumors in Lynch syndrome: the GEOLynch cohort study. <i>Cancer Causes and Control</i> , 2014, 25, 1119-1129.	0.8	13
84	CM-Score: a validated scoring system to predict <i>CDKN2A</i> germline mutations in melanoma families from Northern Europe. <i>Journal of Medical Genetics</i> , 2018, 55, 661-668.	1.5	13
85	Genotype-phenotype correlations for pancreatic cancer risk in Dutch melanoma families with pathogenic <i>CDKN2A</i> variants. <i>Journal of Medical Genetics</i> , 2021, 58, 264-269.	1.5	13
86	Declining detection rates for APC and biallelic MUTYH variants in polyposis patients, implications for DNA testing policy. <i>European Journal of Human Genetics</i> , 2020, 28, 222-230.	1.4	12
87	Bias in detection of instability of the (C)8 mononucleotide repeat of MSH6 in tumours from HNPCC patients. <i>Oncogene</i> , 2001, 20, 6241-6244.	2.6	11
88	Uptake of hysterectomy and bilateral salpingo-oophorectomy in carriers of pathogenic mismatch repair variants: a Prospective Lynch Syndrome Database report. <i>European Journal of Cancer</i> , 2021, 148, 124-133.	1.3	11
89	No Difference in Penetrance between Truncating and Missense/Aberrant Splicing Pathogenic Variants in MLH1 and MSH2: A Prospective Lynch Syndrome Database Study. <i>Journal of Clinical Medicine</i> , 2021, 10, 2856.	1.0	11
90	Clinical utility gene card for: Familial adenomatous polyposis (FAP) and attenuated FAP (AFAP). <i>European Journal of Human Genetics</i> , 2011, 19, 832-832.	1.4	10

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91	Quality of colonoscopy in Lynch syndrome. <i>Endoscopy International Open</i> , 2014, 02, E252-E255.	0.9	10
92	Pancreatic cancer-associated gene polymorphisms in a nation-wide cohort of p16-Leiden germline mutation carriers; a case-control study. <i>BMC Research Notes</i> , 2015, 8, 264.	0.6	10
93	The importance of a well-structured pancreatic screening program for familial and hereditary pancreatic cancer. <i>Familial Cancer</i> , 2018, 17, 1-3.	0.9	10
94	Clinical Utility Gene Card for: Familial adenomatous polyposis (FAP) and attenuated FAP (AFAP) - update 2014. <i>European Journal of Human Genetics</i> , 2015, 23, 889-889.	1.4	9
95	Application of a Serum Protein Signature for Pancreatic Cancer to Separate Cases from Controls in a Pancreatic Surveillance Cohort. <i>Translational Oncology</i> , 2016, 9, 242-247.	1.7	9
96	Diagnostic value of targeted next-generation sequencing in patients with suspected pancreatic or periampullary cancer. <i>Journal of Clinical Pathology</i> , 2018, 71, 246-252.	1.0	9
97	Risk of multiple pancreatic cancers in CDKN2A-p16-Leiden mutation carriers. <i>European Journal of Human Genetics</i> , 2018, 26, 1227-1229.	1.4	9
98	Serum peptide signatures for pancreatic cancer based on mass spectrometry: a comparison to CA19-9 levels and routine imaging techniques. <i>Journal of Cancer Research and Clinical Oncology</i> , 2015, 141, 531-541.	1.2	8
99	A new hereditary colorectal cancer network in the Middle East and eastern mediterranean countries to improve care for high-risk families. <i>Familial Cancer</i> , 2018, 17, 209-212.	0.9	8
100	Low frequency of POLD1 and POLE exonuclease domain variants in patients with multiple colorectal polyps. <i>Molecular Genetics &amp; Genomic Medicine</i> , 2019, 7, e00603.	0.6	8
101	Detection of pancreatic cancer using serum protein profiling. <i>Hpb</i> , 2013, 15, 602-610.	0.1	7
102	Dietary Supplement Use and Colorectal Adenoma Risk in Individuals with Lynch Syndrome: The GEOLynch Cohort Study. <i>PLoS ONE</i> , 2013, 8, e66819.	1.1	7
103	Colonoscopy in Lynch syndrome: the need for a new quality score. <i>Familial Cancer</i> , 2017, 16, 239-241.	0.9	7
104	Optimizing the timing of colorectal surgery in patients with familial adenomatous polyposis in clinical practice. <i>Scandinavian Journal of Gastroenterology</i> , 2019, 54, 733-739.	0.6	7
105	Dilatation of the main pancreatic duct as first manifestation of small pancreatic ductal adenocarcinomas detected in a hereditary pancreatic cancer surveillance program. <i>Hpb</i> , 2019, 21, 1371-1375.	0.1	7
106	Value-based healthcare in Lynch syndrome. <i>Familial Cancer</i> , 2013, 12, 347-354.	0.9	6
107	A genetic variant in telomerase reverse transcriptase (TERT) modifies cancer risk in Lynch syndrome patients harbouring pathogenic MSH2 variants. <i>Scientific Reports</i> , 2021, 11, 11401.	1.6	6
108	8q23.3 and 11q23.1 as modifying loci influencing the risk for CRC in Lynch syndrome. <i>European Journal of Human Genetics</i> , 2012, 20, 487-488.	1.4	5

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109	Identification of familial colorectal cancer and hereditary colorectal cancer syndromes through the Dutch population-screening program: results of a pilot study. <i>Scandinavian Journal of Gastroenterology</i> , 2016, 51, 1227-1232.	0.6	5
110	High Growth Rate of Pancreatic Ductal Adenocarcinoma in <i>CDKN2A-p16-Leiden</i> Mutation Carriers. <i>Cancer Prevention Research</i> , 2018, 11, 551-556.	0.7	5
111	A hundred years of Lynch syndrome research (1913–2013). <i>Familial Cancer</i> , 2013, 12, 141-142.	0.9	4
112	Progress Report: New insights into the prevention of CRC by colonoscopic surveillance in Lynch syndrome. <i>Familial Cancer</i> , 2022, 21, 49-56.	0.9	4
113	An Overview of the Lynch Syndrome (Hereditary Non-polyposis Colorectal Cancer)., 2010, , 271-299.		4
114	Constitutional or biallelic? Settling on a name for a recessively inherited cancer susceptibility syndrome. <i>Journal of Medical Genetics</i> , 2016, 53, 226-226.	1.5	3
115	Dilemmas in the management of screen-detected lesions in patients at high risk for pancreatic cancer. <i>Familial Cancer</i> , 2017, 16, 111-115.	0.9	3
116	Identification and management of Lynch syndrome in the Middle East and North African countries: outcome of a survey in 12 countries. <i>Familial Cancer</i> , 2020, 20, 215-221.	0.9	3
117	Is a colorectal neoplasm diagnosis a trigger to change dietary and other lifestyle habits for persons with Lynch syndrome? A prospective cohort study. <i>Familial Cancer</i> , 2021, 20, 125-135.	0.9	3
118	Endoscopic full thickness resection for early colon cancer in Lynch syndrome. <i>Familial Cancer</i> , 2019, 18, 349-352.	0.9	2
119	Diet quality and colorectal tumor risk in persons with Lynch syndrome. <i>Cancer Epidemiology</i> , 2020, 69, 101809.	0.8	2
120	Kluwer Academic Publishers. <i>Annals of Oncology</i> , 2001, 12, 735-735.	0.6	1
121	Clinical evidence for an association between familial adenomatous polyposis and type II diabetes. <i>International Journal of Cancer</i> , 2012, 131, 1488-1489.	2.3	1
122	Preface. <i>Bailliere's Best Practice and Research in Clinical Gastroenterology</i> , 2009, 23, 125-126.	1.0	0
123	Reply to Win and Jenkins. <i>International Journal of Cancer</i> , 2013, 133, 1764-1764.	2.3	0
124	Reply to V. Bonadona et al. <i>Journal of Clinical Oncology</i> , 2013, 31, 2230-2230.	0.8	0
125	Familial Pancreatic Cancer: To Screen or not to Screen?. <i>EBioMedicine</i> , 2015, 2, 1858-1859.	2.7	0
126	Screening of Individuals at High Risk for Pancreatic Cancer. <i>Clinical Gastroenterology and Hepatology</i> , 2019, 17, 1916-1917.	2.4	0