Malcolm Dunlop

List of Publications by Year in descending order

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269 papers 28,862 citations

76 h-index 162 g-index

287 all docs

287 docs citations

times ranked

287

31283 citing authors

#	Article	IF	CITATIONS
1	Bidirectional Mendelian randomisation analysis of the relationship between circulating vitamin D concentration and colorectal cancer risk. International Journal of Cancer, 2022, 150, 303-307.	5.1	13
2	Disease consequences of higher adiposity uncoupled from its adverse metabolic effects using Mendelian randomisation. ELife, 2022, 11, .	6.0	10
3	Alcohol consumption, <scp>DNA</scp> methylation and colorectal cancer risk: Results from pooled cohort studies and Mendelian randomization analysis. International Journal of Cancer, 2022, 151, 83-94.	5.1	22
4	A systematic review of microbial markers for risk prediction of colorectal neoplasia. British Journal of Cancer, 2022, 126, 1318-1328.	6.4	26
5	Phenome-wide association study (PheWAS) of colorectal cancer risk SNP effects on health outcomes in UK Biobank. British Journal of Cancer, 2022, 126, 822-830.	6.4	4
6	Vitamin D treatment induces in vitro and ex vivo transcriptomic changes indicating antiâ€ŧumor effects. FASEB Journal, 2022, 36, e22082.	0.5	6
7	RNA splicing is a key mediator of tumour cell plasticity and a therapeutic vulnerability in colorectal cancer. Nature Communications, 2022, 13, 2791.	12.8	11
8	Replication of 15 loci involved in human plasma protein N-glycosylation in 4802 samples from four cohorts. Glycobiology, 2021, 31, 82-88.	2.5	15
9	Aspirin Rescues Wnt-Driven Stem-like Phenotype in Human Intestinal Organoids and Increases the Wnt Antagonist Dickkopf-1. Cellular and Molecular Gastroenterology and Hepatology, 2021, 11, 465-489.	4.5	15
10	Colorectal cancer risk variants rs10161980 and rs7495132 are associated with cancer survival outcome by a recessive mode of inheritance. International Journal of Cancer, 2021, 148, 2774-2778.	5.1	7
11	Genetically predicted physical activity levels are associated with lower colorectal cancer risk: a Mendelian randomisation study. British Journal of Cancer, 2021, 124, 1330-1338.	6.4	17
12	Implementation of a risk mitigating COVID-adapted colorectal cancer pathway. BMJ Open Quality, 2021, 10, e001135.	1.1	2
13	Nurse-led telephone outreach for a COVID-adapted suspected colorectal cancer pathway. Gastrointestinal Nursing, 2021, 19, 22-26.	0.1	3
14	Shortâ€ŧerm outcomes of a COVIDâ€adapted triage pathway for colorectal cancer detection. Colorectal Disease, 2021, 23, 1639-1648.	1.4	10
15	RAC1B modulates intestinal tumourigenesis via modulation of WNT and EGFR signalling pathways. Nature Communications, 2021, 12, 2335.	12.8	20
16	Differential genetic influences over colorectal cancer risk and gene expression in large bowel mucosa. International Journal of Cancer, 2021, 149, 1100-1108.	5.1	7
17	Risk of missing colorectal cancer with a COVID-adapted diagnostic pathway using quantitative faecal immunochemical testing. BJS Open, 2021, 5, .	1.7	4
18	Development of a customised data management system for a COVID-19-adapted colorectal cancer pathway. BMJ Health and Care Informatics, 2021, 28, e100307.	3.0	0

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19	Oral vitamin D supplementation induces transcriptomic changes in rectal mucosa that are linked to anti-tumour effects. BMC Medicine, 2021, 19, 174.	5.5	7
20	An observational and Mendelian randomisation study on vitamin D and COVID-19 risk in UK Biobank. Scientific Reports, 2021, $11,18262$.	3.3	13
21	Characteristics of Early-Onset vs Late-Onset Colorectal Cancer. JAMA Surgery, 2021, 156, 865.	4.3	110
22	A genome-wide search for determinants of survival in 1926 patients with advanced colorectal cancer with follow-up in over 22,000 patients. European Journal of Cancer, 2021, 159, 247-258.	2.8	6
23	Gene Co-Expression Network Analysis Identifies Vitamin D-Associated Gene Modules in Adult Normal Rectal Epithelium Following Supplementation. Frontiers in Genetics, 2021, 12, 783970.	2.3	3
24	Low plasma vitamin D is associated with adverse colorectal cancer survival after surgical resection, independent of systemic inflammatory response. Gut, 2020, 69, 103-111.	12.1	44
25	Modifiable pathways for colorectal cancer: a mendelian randomisation analysis. The Lancet Gastroenterology and Hepatology, 2020, 5, 55-62.	8.1	79
26	Risk mitigation for suspected colorectal cancer diagnostic pathway during COVID-19 pandemic. British Journal of Surgery, 2020, 107, e361-e362.	0.3	14
27	The effect of vitamin D supplementation on survival in patients with colorectal cancer: systematic review and meta-analysis of randomised controlled trials. British Journal of Cancer, 2020, 123, 1705-1712.	6.4	67
28	Nonâ€genetic biomarkers and colorectal cancer risk: Umbrella review and evidence triangulation. Cancer Medicine, 2020, 9, 4823-4835.	2.8	12
29	Prediction of colorectal cancer risk based on profiling with common genetic variants. International Journal of Cancer, 2020, 147, 3431-3437.	5.1	17
30	Systematic Evaluation of Normalization Methods for Glycomics Data Based on Performance of Network Inference. Metabolites, 2020, 10, 271.	2.9	13
31	Risk factors and risk prediction models for colorectal cancer metastasis and recurrence: an umbrella review of systematic reviews and meta-analyses of observational studies. BMC Medicine, 2020, 18, 172.	5.5	66
32	Glycosylation of immunoglobulin G is regulated by a large network of genes pleiotropic with inflammatory diseases. Science Advances, 2020, 6, eaax0301.	10.3	90
33	Systematic meta-analyses, field synopsis and global assessment of the evidence of genetic association studies in colorectal cancer. Gut, 2020, 69, 1460-1471.	12.1	27
34	Guidelines for the management of hereditary colorectal cancer from the British Society of Gastroenterology (BSG)/Association of Coloproctology of Great Britain and Ireland (ACPGBI)/United Kingdom Cancer Genetics Group (UKCGG). Gut, 2020, 69, 411-444.	12.1	263
35	A Systematic Analysis of Interactions between Environmental Risk Factors and Genetic Variation in Susceptibility to Colorectal Cancer. Cancer Epidemiology Biomarkers and Prevention, 2020, 29, 1145-1153.	2.5	16
36	Physical activity and COVID-19: an observational and Mendelian randomisation study. Journal of Global Health, 2020, 10, 020514.	2.7	24

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37	Effects of common genetic variants associated with colorectal cancer risk on survival outcomes after diagnosis: A large populationâ€based cohort study. International Journal of Cancer, 2019, 145, 2427-2432.	5.1	11
38	Phenome-wide Mendelian-randomization study of genetically determined vitamin D on multiple health outcomes using the UK Biobank study. International Journal of Epidemiology, 2019, 48, 1425-1434.	1.9	61
39	British Society of Gastroenterology consensus guidelines on the management of inflammatory bowel disease in adults. Gut, 2019, 68, s1-s106.	12.1	1,353
40	Colorectal cancer: management. Medicine, 2019, 47, 405-409.	0.4	0
41	Flexible and scalable diagnostic filtering of genomic variants using G2P with Ensembl VEP. Nature Communications, 2019, 10, 2373.	12.8	86
42	Performance of prediction models on survival outcomes of colorectal cancer with surgical resection: A systematic review and meta-analysis. Surgical Oncology, 2019, 29, 196-202.	1.6	20
43	Association analyses identify 31 new risk loci for colorectal cancer susceptibility. Nature Communications, 2019, 10, 2154.	12.8	172
44	Whether vitamin D supplementation protects against colorectal cancer risk remains an open question. European Journal of Cancer, 2019, 115, 1-3.	2.8	7
45	Defining the genetic control of human blood plasma N-glycome using genome-wide association study. Human Molecular Genetics, 2019, 28, 2062-2077.	2.9	40
46	Higher Post-Operative Serum Vitamin D Level is Associated with Better Survival Outcome in Colorectal Cancer Patients. Nutrition and Cancer, 2019, 71, 1078-1085.	2.0	18
47	Head-to-Head Comparison of Family History of Colorectal Cancer and a Genetic Risk Score for Colorectal Cancer Risk Stratification. Clinical and Translational Gastroenterology, 2019, 10, e00106.	2.5	4
48	A Comprehensive Study of the Effect on Colorectal Cancer Survival of Common Germline Genetic Variation Previously Linked with Cancer Prognosis. Cancer Epidemiology Biomarkers and Prevention, 2019, 28, 1944-1946.	2.5	4
49	Genetic predisposition to mosaic Y chromosome loss in blood. Nature, 2019, 575, 652-657.	27.8	198
50	Gene–environment interactions and colorectal cancer risk: An umbrella review of systematic reviews and metaâ€analyses of observational studies. International Journal of Cancer, 2019, 145, 2315-2329.	5.1	47
51	Genome-wide association study in 79,366 European-ancestry individuals informs the genetic architecture of 25-hydroxyvitamin D levels. Nature Communications, 2018, 9, 260.	12.8	295
52	Glycosylation of Immunoglobulin G Associates With Clinical Features of Inflammatory Bowel Diseases. Gastroenterology, 2018, 154, 1320-1333.e10.	1.3	116
53	Genomeâ€wide association study and metaâ€analysis in Northern European populations replicate multiple colorectal cancer risk loci. International Journal of Cancer, 2018, 142, 540-546.	5.1	26
54	The Association of Coloproctology of Great Britain and Ireland consensus guidelines in surgery for inflammatory bowel disease. Colorectal Disease, 2018, 20, 3-117.	1.4	52

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55	Coding variants in NOD-like receptors: An association study on risk and survival of colorectal cancer. PLoS ONE, 2018, 13, e0199350.	2.5	6
56	Genome-wide scan of the effect of common nsSNPs on colorectal cancer survival outcome. British Journal of Cancer, 2018, 119, 988-993.	6.4	10
57	Exploring causality in the association between circulating 25-hydroxyvitamin D and colorectal cancer risk: a large Mendelian randomisation study. BMC Medicine, 2018, 16, 142.	5.5	62
58	Plasma N-glycans in colorectal cancer risk. Scientific Reports, 2018, 8, 8655.	3.3	57
59	Recurrent, low-frequency coding variants contributing to colorectal cancer in the Swedish population. PLoS ONE, 2018, 13, e0193547.	2.5	10
60	Mendelian randomisation implicates hyperlipidaemia as a risk factor for colorectal cancer. International Journal of Cancer, 2017, 140, 2701-2708.	5.1	76
61	The impact of vitamin D pathway genetic variation and circulating 25-hydroxyvitamin D on cancer outcome: systematic review and meta-analysis. British Journal of Cancer, 2017, 116, 1092-1110.	6.4	115
62	Establishing Key Performance Indicators [KPIs] and Their Importance for the Surgical Management of Inflammatory Bowel Disease–Results From a Pan-European, Delphi Consensus Study. Journal of Crohn's and Colitis, 2017, 11, 1362-1368.	1.3	28
63	Pro-inflammatory fatty acid profile and colorectal cancer risk: A Mendelian randomisation analysis. European Journal of Cancer, 2017, 84, 228-238.	2.8	81
64	Evidence for genetic association between chromosome 1q loci and predisposition to colorectal neoplasia. British Journal of Cancer, 2017, 117, 1215-1223.	6.4	10
65	Reply to †Comment on †The impact of vitamin D pathway genetic variation and circulating 25-hydroxyvitamin D on cancer outcome: systematic review and meta-analysis†M†M. British Journal of Cancer, 2017, 117, e4-e4.	6.4	0
66	Validation of Recently Proposed Colorectal Cancer Susceptibility Gene Variants in an Analysis of Families and Patients—a Systematic Review. Gastroenterology, 2017, 152, 75-77.e4.	1.3	80
67	Farming, Foreign Holidays, and Vitamin D in Orkney. PLoS ONE, 2016, 11, e0155633.	2.5	5
68	Mendelian randomisation analysis strongly implicates adiposity with risk of developing colorectal cancer. British Journal of Cancer, 2016, 115, 266-272.	6.4	57
69	Correspondence: SEMA4A variation and risk of colorectal cancer. Nature Communications, 2016, 7, 10611.	12.8	7
70	IgG Glycome in Colorectal Cancer. Clinical Cancer Research, 2016, 22, 3078-3086.	7.0	111
71	Five endometrial cancer risk loci identified through genome-wide association analysis. Nature Genetics, 2016, 48, 667-674.	21.4	77
72	Mercaptopurine versus placebo to prevent recurrence of Crohn's disease after surgical resection (TOPPIC): a multicentre, double-blind, randomised controlled trial. The Lancet Gastroenterology and Hepatology, 2016, 1, 273-282.	8.1	91

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73	Glycosylation of plasma IgG in colorectal cancer prognosis. Scientific Reports, 2016, 6, 28098.	3.3	84
74	Rare disruptive mutations and their contribution to the heritable risk of colorectal cancer. Nature Communications, 2016, 7, 11883.	12.8	122
75	The contributions of adjusted ambient ultraviolet B radiation at place of residence and other determinants to serum 25-hydroxyvitamin D concentrations. British Journal of Dermatology, 2016, 174, 1068-1078.	1.5	23
76	Variation at 2q35 (<i>PNKD</i> and <i>TMBIM1</i>) influences colorectal cancer risk and identifies a pleiotropic effect with inflammatory bowel disease. Human Molecular Genetics, 2016, 25, 2349-2359.	2.9	37
77	Investigation of gene–environment interactions between vitamin D and colorectal cancer susceptibility genetic variants in large bowel epithelium. Lancet, The, 2016, 387, S102.	13.7	1
78	Systematic meta-analyses and field synopsis of genetic association studies in colorectal adenomas. International Journal of Epidemiology, 2016, 45, 186-205.	1.9	21
79	Recurrent Coding Sequence Variation Explains Only A Small Fraction of the Genetic Architecture of Colorectal Cancer. Scientific Reports, 2015, 5, 16286.	3.3	24
80	Meta-analysis of genome-wide association studies identifies common susceptibility polymorphisms for colorectal and endometrial cancer near SH2B3 and TSHZ1. Scientific Reports, 2015, 5, 17369.	3.3	35
81	PWE-322ÂAre we over-treating polyp cancers?. Gut, 2015, 64, A352.1-A352.	12.1	0
82	Inflammatory Bowel Disease Associates with Proinflammatory Potential of the Immunoglobulin G Glycome. Inflammatory Bowel Diseases, 2015, 21, 1.	1.9	161
83	A simple method to overcome the inhibitory effect of heparin on DNA amplification. Cellular Oncology (Dordrecht), 2015, 38, 493-495.	4.4	9
84	A new GWAS and meta-analysis with 1000Genomes imputation identifies novel risk variants for colorectal cancer. Scientific Reports, 2015, 5, 10442.	3.3	109
85	Reply to F.J.S. Conway et al. Journal of Clinical Oncology, 2015, 33, 224-225.	1.6	1
86	Exome Sequencing to Detect Rare Variants Associated With General Cognitive Ability: A Pilot Study. Twin Research and Human Genetics, 2015, 18, 117-125.	0.6	7
87	Colorectal cancer: management. Medicine, 2015, 43, 303-307.	0.4	1
88	Cardiometabolic effects of genetic upregulation of the interleukin 1 receptor antagonist: a Mendelian randomisation analysis. Lancet Diabetes and Endocrinology,the, 2015, 3, 243-253.	11.4	115
89	Diverticular disease in Scotland: 2000–2010. Colorectal Disease, 2015, 17, 329-334.	1.4	30
90	Obesity, Aspirin, and Risk of Colorectal Cancer in Carriers of Hereditary Colorectal Cancer: A Prospective Investigation in the CAPP2 Study. Journal of Clinical Oncology, 2015, 33, 3591-3597.	1.6	91

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91	Modulation of Genetic Associations with Serum Urate Levels by Body-Mass-Index in Humans. PLoS ONE, 2015, 10, e0119752.	2.5	64
92	Associations between dietary and lifestyle risk factors and colorectal cancer in the Scottish population. European Journal of Cancer Prevention, 2014, 23, 8-17.	1.3	39
93	Identification of susceptibility loci for colorectal cancer in a genome-wide meta-analysis. Human Molecular Genetics, 2014, 23, 4729-4737.	2.9	128
94	Plasma Vitamin D Concentration Influences Survival Outcome After a Diagnosis of Colorectal Cancer. Journal of Clinical Oncology, 2014, 32, 2430-2439.	1.6	128
95	Re: Role of the Oxidative DNA Damage Repair Gene OGG1 in Colorectal Tumorigenesis. Journal of the National Cancer Institute, 2014, 106 , .	6.3	9
96	The MLH1 c.1852_1853delinsGC (p.K618A) Variant in Colorectal Cancer: Genetic Association Study in 18,723 Individuals. PLoS ONE, 2014, 9, e95022.	2.5	7
97	Cumulative impact of common genetic variants and other risk factors on colorectal cancer risk in $42\hat{a}\in 103$ individuals. Gut, 2013 , 62 , $871-881$.	12.1	117
98	Deciphering the genetic architecture of low-penetrance susceptibility to colorectal cancer. Human Molecular Genetics, 2013, 22, 5075-5082.	2.9	19
99	The <i><scp>MSH2</scp></i> c.388_389del mutation shows a founder effect in Portuguese Lynch syndrome families. Clinical Genetics, 2013, 84, 244-250.	2.0	13
100	Genome-wide association analyses identify 18 new loci associated with serum urate concentrations. Nature Genetics, 2013, 45, 145-154.	21.4	675
101	Long term effect of surgery and radiotherapy for colorectal cancer on defecatory function and quality of life. European Journal of Oncology Nursing, 2013, 17, 570-577.	2.1	54
102	Causal Relationship between Obesity and Vitamin D Status: Bi-Directional Mendelian Randomization Analysis of Multiple Cohorts. PLoS Medicine, 2013, 10, e1001383.	8.4	753
103	Inference of identity by descent in population isolates and optimal sequencing studies. European Journal of Human Genetics, 2013, 21, 1140-1145.	2.8	14
104	Genome-wide association study identifies genetic risk underlying primary rhegmatogenous retinal detachment. Human Molecular Genetics, 2013, 22, 3174-3185.	2.9	34
105	Germline Variants and Advanced Colorectal Adenomas: Adenoma Prevention with Celecoxib Trial Genome-wide Association Study. Clinical Cancer Research, 2013, 19, 6430-6437.	7.0	9
106	Model Selection Approach Suggests Causal Association between 25-Hydroxyvitamin D and Colorectal Cancer. PLoS ONE, 2013, 8, e63475.	2.5	10
107	Meta-Analysis of Mismatch Repair Polymorphisms within the Cogent Consortium for Colorectal Cancer Susceptibility. PLoS ONE, 2013, 8, e72091.	2.5	19
108	SMAD7 Variant rs4939827 Is Associated with Colorectal Cancer Risk in Croatian Population. PLoS ONE, 2013, 8, e74042.	2.5	12

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109	Evidence of Inbreeding Depression on Human Height. PLoS Genetics, 2012, 8, e1002655.	3.5	79
110	Investigation of the effects of DNA repair gene polymorphisms on the risk of colorectal cancer. Mutagenesis, 2012, 27, 219-223.	2.6	29
111	<i>TERC</i> polymorphisms are associated both with susceptibility to colorectal cancer and with longer telomeres. Gut, 2012, 61, 248-254.	12.1	94
112	The TERT variant rs2736100 is associated with colorectal cancer risk. British Journal of Cancer, 2012, 107, 1001-1008.	6.4	50
113	Genome-wide association study of age-related macular degeneration identifies associated variants in the TNXB–FKBPL–NOTCH4 region of chromosome 6p21.3. Human Molecular Genetics, 2012, 21, 4138-4150.	2.9	80
114	Refinement of the associations between risk of colorectal cancer and polymorphisms on chromosomes 1q41 and 12q13.13. Human Molecular Genetics, 2012, 21, 934-946.	2.9	19
115	In vitro stability of APC gene sequences and the influence of DNA repair status. Mutagenesis, 2012, 27, 233-238.	2.6	3
116	The interleukin-6 receptor as a target for prevention of coronary heart disease: a mendelian randomisation analysis. Lancet, The, 2012, 379, 1214-1224.	13.7	886
117	Common variation near CDKN1A, POLD3 and SHROOM2 influences colorectal cancer risk. Nature Genetics, 2012, 44, 770-776.	21.4	210
118	Systematic Meta-Analyses and Field Synopsis of Genetic Association Studies in Colorectal Cancer. Journal of the National Cancer Institute, 2012, 104, 1433-1457.	6.3	91
119	Statin use and association with colorectal cancer survival and risk: case control study with prescription data linkage. BMC Cancer, 2012, 12, 487.	2.6	39
120	Aspirin Inhibits mTOR Signaling, Activates AMP-Activated Protein Kinase, and Induces Autophagy in Colorectal Cancer Cells. Gastroenterology, 2012, 142, 1504-1515.e3.	1.3	356
121	Long-term effect of resistant starch on cancer risk in carriers of hereditary colorectal cancer: an analysis from the CAPP2 randomised controlled trial. Lancet Oncology, The, 2012, 13, 1242-1249.	10.7	95
122	Instrumental Variable Estimation of the Causal Effect of Plasma 25-Hydroxy-Vitamin D on Colorectal Cancer Risk: A Mendelian Randomization Analysis. PLoS ONE, 2012, 7, e37662.	2.5	51
123	The Association of Dietary Intake of Purine-Rich Vegetables, Sugar-Sweetened Beverages and Dairy with Plasma Urate, in a Cross-Sectional Study. PLoS ONE, 2012, 7, e38123.	2.5	106
124	A Novel Test for Gene-Ancestry Interactions in Genome-Wide Association Data. PLoS ONE, 2012, 7, e48687.	2.5	3
125	Diet, Environmental Factors, and Lifestyle Underlie the High Prevalence of Vitamin D Deficiency in Healthy Adults in Scotland, and Supplementation Reduces the Proportion That Are Severely Deficient. Journal of Nutrition, 2011, 141, 1535-1542.	2.9	75
126	Long-term effect of aspirin on cancer risk in carriers of hereditary colorectal cancer: an analysis from the CAPP2 randomised controlled trial. Lancet, The, 2011, 378, 2081-2087.	13.7	849

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127	A doubling of admissions due to diverticular disease in Scottish hospitals, in the last 14 years. Gut, 2011, 60, A7-A7.	12.1	0
128	SNP mistyping in genotyping arrays-an important cause of spurious association in case-control studies. Genetic Epidemiology, 2011, 35, 423-426.	1.3	5
129	The value of FDG positron emission tomography/computerised tomography (PET/CT) in pre-operative staging of colorectal cancer: a systematic review and economic evaluation Health Technology Assessment, 2011, 15, 1-192, iii-iv.	2.8	100
130	A Randomized Placebo-Controlled Prevention Trial of Aspirin and/or Resistant Starch in Young People with Familial Adenomatous Polyposis. Cancer Prevention Research, 2011, 4, 655-665.	1.5	193
131	c-Src dependency of NSAID-induced effects on NF-ÂB-mediated apoptosis in colorectal cancer cells. Carcinogenesis, 2011, 32, 1069-1077.	2.8	19
132	Fine-mapping of colorectal cancer susceptibility loci at 8q23.3, 16q22.1 and 19q13.11: refinement of association signals and use of in silico analysis to suggest functional variation and unexpected candidate target genes. Human Molecular Genetics, 2011, 20, 2879-2888.	2.9	56
133	Multiple Common Susceptibility Variants near BMP Pathway Loci GREM1, BMP4, and BMP2 Explain Part of the Missing Heritability of Colorectal Cancer. PLoS Genetics, 2011, 7, e1002105.	3.5	188
134	Characterisation of Genome-Wide Association Epistasis Signals for Serum Uric Acid in Human Population Isolates. PLoS ONE, 2011, 6, e23836.	2.5	15
135	Cigarette Smoke Extract (CSE) Delays NOD2 Expression and Affects NOD2/RIPK2 Interactions in Intestinal Epithelial Cells. PLoS ONE, 2011, 6, e24715.	2.5	22
136	Report of the Combined Meeting of the International Society for Gastrointestinal Hereditary Tumours, the Human Variome Project and the National Cancer Institute Colon Cancer Family Registry, Duesseldorf, Germany, 24 June 2009. Familial Cancer, 2010, 9, 705-711.	1.9	5
137	<i>MLH1</i> Differential Allelic Expression in Mutation Carriers and Controls. Annals of Human Genetics, 2010, 74, 479-488.	0.8	12
138	COGENT (COlorectal cancer GENeTics): an international consortium to study the role of polymorphic variation on the risk of colorectal cancer. British Journal of Cancer, 2010, 102, 447-454.	6.4	43
139	Association studies on 11 published colorectal cancer risk loci. British Journal of Cancer, 2010, 103, 575-580.	6.4	61
140	A large-scale meta-analysis to refine colorectal cancer risk estimates associated with MUTYH variants. British Journal of Cancer, 2010, 103, 1875-1884.	6.4	107
141	Genome-wide association study identifies variants at CSF1, OPTN and TNFRSF11A as genetic risk factors for Paget's disease of bone. Nature Genetics, 2010, 42, 520-524.	21.4	258
142	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, 973-977.	21.4	335
143	Comprehensive assessment of variation at the transforming growth factor \hat{l}^2 type 1 receptor locus and colorectal cancer predisposition. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 7858-7862.	7.1	26
144	Ten Common Genetic Variants Associated with Colorectal Cancer Risk Are Not Associated with Survival after Diagnosis. Clinical Cancer Research, 2010, 16, 3754-3759.	7.0	36

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145	Nucleolar Targeting of RelA(p65) Is Regulated by COMMD1-Dependent Ubiquitination. Cancer Research, 2010, 70, 139-149.	0.9	61
146	Risks of Lynch Syndrome Cancers for MSH6 Mutation Carriers. Journal of the National Cancer Institute, 2010, 102, 193-201.	6.3	328
147	Association between common mtDNA variants and all-cause or colorectal cancer mortality. Carcinogenesis, 2010, 31, 296-301.	2.8	20
148	Colorectal Cancer Susceptibility Loci in a Population-Based Study. American Journal of Pathology, 2010, 177, 2688-2693.	3.8	16
149	Guidelines for colorectal cancer screening and surveillance in moderate and high risk groups (update from 2002). Gut, 2010, 59, 666-689.	12.1	1,000
150	Effect of aspirin and NSAIDs on risk and survival from colorectal cancer. Gut, 2010, 59, 1670-1679.	12.1	254
151	Polyposis Syndromes and Colorectal Cancer Predisposition., 2010,, 545-559.		0
152	The Search for Gene-Gene Interactions in Colorectal Cancer: Using HPC to Overcome Computational Barriers. , 2009, , .		1
153	Common variants in the JAZF1 gene associated with height identified by linkage and genome-wide association analysis. Human Molecular Genetics, 2009, 18, 373-380.	2.9	88
154	Singleâ€cell expression profiling of dopaminergic neurons combined with association analysis identifies pyridoxal kinase as Parkinson's disease gene. Annals of Neurology, 2009, 66, 792-798.	5.3	49
155	Common genetic variants at the MC4R locus are associated with obesity, but not with dietary energy intake or colorectal cancer in the Scottish population. International Journal of Obesity, 2009, 33, 284-288.	3.4	27
156	New insights into the aetiology of colorectal cancer from genome-wide association studies. Nature Reviews Genetics, 2009, 10, 353-358.	16.3	355
157	Fecal Calprotectin Predicts the Clinical Course of Acute Severe Ulcerative Colitis. American Journal of Gastroenterology, 2009, 104, 673-678.	0.4	143
158	MUTYH-Associated Polyposis and Colorectal Cancer. Surgical Oncology Clinics of North America, 2009, 18, 599-610.	1.5	10
159	Cascade genetic testing for mismatch repair gene mutations. Familial Cancer, 2008, 7, 293-301.	1.9	2
160	Classification of ambiguous mutations in DNA mismatch repair genes identified in a population-based study of colorectal cancer. Human Mutation, 2008, 29, 367-374.	2.5	68
161	Modification of the inverse association between dietary vitamin D intake and colorectal cancer risk by a <i>Fok</i> I variant supports a chemoprotective action of Vitamin D intake mediated through VDR binding. International Journal of Cancer, 2008, 123, 2170-2179.	5.1	54
162	Runs of Homozygosity in European Populations. American Journal of Human Genetics, 2008, 83, 359-372.	6.2	958

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163	The proapoptotic effects of sulindac, sulindac sulfone and indomethacin are mediated by nucleolar translocation of the RelA(p65) subunit of NF-ÎB. Oncogene, 2008, 27, 2648-2655.	5.9	54
164	A genome-wide association study identifies colorectal cancer susceptibility loci on chromosomes 10p14 and 8q23.3. Nature Genetics, 2008, 40, 623-630.	21.4	514
165	Genome-wide association scan identifies a colorectal cancer susceptibility locus on $11q23$ and replicates risk loci at $8q24$ and $18q21$. Nature Genetics, 2008, 40, 631-637.	21.4	542
166	Common genetic variants at the CRAC1 (HMPS) locus on chromosome 15q13.3 influence colorectal cancer risk. Nature Genetics, 2008, 40, 26-28.	21.4	277
167	Meta-analysis of genome-wide association data identifies four new susceptibility loci for colorectal cancer. Nature Genetics, 2008, 40, 1426-1435.	21.4	498
168	Modification of the associations between lifestyle, dietary factors and colorectal cancer risk by APC variants. Carcinogenesis, 2008, 29, 1774-1780.	2.8	25
169	Dietary Vitamin B6 Intake and the Risk of Colorectal Cancer. Cancer Epidemiology Biomarkers and Prevention, 2008, 17, 171-182.	2.5	78
170	Effect of Aspirin or Resistant Starch on Colorectal Neoplasia in the Lynch Syndrome. New England Journal of Medicine, 2008, 359, 2567-2578.	27.0	273
171	Analysis of Germline GLI1 Variation Implicates Hedgehog Signalling in the Regulation of Intestinal Inflammatory Pathways. PLoS Medicine, 2008, 5, e239.	8.4	63
172	Dietary Fatty Acids and Colorectal Cancer: A Case-Control Study. American Journal of Epidemiology, 2007, 166, 181-195.	3.4	120
173	CDK4 Inhibitors and Apoptosis: A Novel Mechanism Requiring Nucleolar Targeting of RelA. Cell Cycle, 2007, 6, 1293-1297.	2.6	42
174	Dietary Flavonoids and the Risk of Colorectal Cancer. Cancer Epidemiology Biomarkers and Prevention, 2007, 16, 684-693.	2.5	207
175	p38-Mediated Inactivation of Cyclin D1/Cyclin-Dependent Kinase 4 Stimulates Nucleolar Translocation of RelA and Apoptosis in Colorectal Cancer Cells. Cancer Research, 2007, 67, 1660-1669.	0.9	105
176	Peutz–Jeghers syndrome: A case report and discussion of surveillance recommendations. European Journal of Radiology Extra, 2007, 62, 81-84.	0.1	3
177	Contribution of the NOD1/CARD4 insertion/deletion polymorphism +32656 to inflammatory bowel disease in Northern Europe#. Inflammatory Bowel Diseases, 2007, 13, 882-889.	1.9	29
178	Genome-wide association scan identifies a colorectal cancer susceptibility locus on chromosome 8q24. Nature Genetics, 2007, 39, 989-994.	21.4	676
179	Germline mutation prevalence in the base excision repair gene, <i>MYH</i> , in patients with endometrial cancer. Clinical Genetics, 2007, 72, 551-555.	2.0	42
180	Outcomes of the rectal remnant following colectomy for ulcerative colitis. Colorectal Disease, 2007, 10, 070621084454038-???.	1.4	28

#	Article	lF	Citations
181	Chromatin structure and evolution in the human genome. BMC Evolutionary Biology, 2007, 7, 72.	3.2	80
182	Reply to Webb et al American Journal of Human Genetics, 2006, 79, 771.	6.2	12
183	Association of MUTYH and colorectal cancer. British Journal of Cancer, 2006, 95, 239-242.	6.4	87
184	Association of DLG5 and inflammatory bowel disease across populations. European Journal of Human Genetics, 2006, 14, 259-260.	2.8	15
185	Reply to Daly and Rioux response. European Journal of Human Genetics, 2006, 14, 261-261.	2.8	1
186	Validity of tagging SNPs across populations for association studies. European Journal of Human Genetics, 2006, 14, 357-363.	2.8	13
187	Identification and Survival of Carriers of Mutations in DNA Mismatch-Repair Genes in Colon Cancer. New England Journal of Medicine, 2006, 354, 2751-2763.	27.0	424
188	Aspirin activates the NF-ÂB signalling pathway and induces apoptosis in intestinal neoplasia in two in vivo models of human colorectal cancer. Carcinogenesis, 2006, 28, 968-976.	2.8	121
189	Disease severity and genetic pathways in attenuated familial adenomatous polyposis vary greatly but depend on the site of the germline mutation. Gut, 2006, 55, 1440-1448.	12.1	87
190	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 Clinical Cancer Research, 2006, 12, 3389-3393.	7.0	42
191	Aspirin-induced nuclear translocation of NFîºB and apoptosis in colorectal cancer is independent of p53 status and DNA mismatch repair proficiency. British Journal of Cancer, 2005, 92, 1137-1143.	6.4	39
192	Inheritance of colorectal cancer susceptibility. British Journal of Surgery, 2005, 77, 245-245.	0.3	3
193	Vacuum drainage of groin wounds after vascular surgery: A controlled trial. British Journal of Surgery, 2005, 77, 562-563.	0.3	29
194	Screening for large bowel neoplasms in individuals with a family history of colorectal cancer. British Journal of Surgery, 2005, 79, 488-494.	0.3	40
195	Molecular genetic basis of colorectal cancer susceptibility. British Journal of Surgery, 2005, 83, 321-329.	0.3	34
196	Molecular genetic basis of colorectal cancer susceptibility. British Journal of Surgery, 2005, 83, 1161-1162.	0.3	0
197	Prevalence of family history of colorectal cancer in the general population. British Journal of Surgery, 2005, 92, 1161-1164.	0.3	36
198	Screening for exonic copy number mutations at MSH2 and MLH1 by MAPH. Familial Cancer, 2005, 4, 145-149.	1.9	10

#	Article	IF	Citations
199	Impact of UK Colorectal Cancer Screening Pilot on hospital diagnostic services. Journal of Public Health, 2005, 27, 246-253.	1.8	18
200	Molecular genetics in gastroenterology: from research success to clinical application?. Nature Reviews Gastroenterology & Hepatology, 2005, 2, 118-119.	1.7	1
201	Nucleolar Sequestration of RelA (p65) Regulates NF-κB-Driven Transcription and Apoptosis. Molecular and Cellular Biology, 2005, 25, 5985-6004.	2.3	117
202	Aurora- A/STK15 T $+$ 91A is a general low penetrance cancer susceptibility gene: a meta-analysis of multiple cancer types. Carcinogenesis, 2005, 26, 1368-1373.	2.8	132
203	Re: Association Between Biallelic and Monoallelic Germline MYH Gene Mutations and Colorectal Cancer Risk. Journal of the National Cancer Institute, 2005, 97, 320-321.	6.3	16
204	Germline Susceptibility to Colorectal Cancer Due to Base-Excision Repair Gene Defects. American Journal of Human Genetics, 2005, 77, 112-119.	6.2	268
205	Accuracy of reporting of family history of colorectal cancer. Gut, 2004, 53, 291-295.	12.1	160
206	Assessment of outcomes after colorectal cancer resection in the elderly as a rationale for screening and early detection. British Journal of Surgery, 2004, 91, 1345-1351.	0.3	53
207	Evidence for colorectal cancer cell specificity of aspirin effects on NFκB signalling and apoptosis. British Journal of Cancer, 2004, 91, 381-388.	6.4	95
208	The contribution of DNA mismatch repair gene defects to the burden of gynecological cancer. International Journal of Gynecological Cancer, 2003, 13, 262-277.	2.5	23
209	Is it acceptable to approach colorectal cancer patients at diagnosis to discuss genetic testing? A pilot study. British Journal of Cancer, 2003, 89, 1400-1402.	6.4	17
210	Colonoscopy surveillance of individuals at risk of familial colorectal cancer. Gut, 2003, 52, 1748-1751.	12.1	29
211	Whole-gene APC deletions cause classical familial adenomatous polyposis, but not attenuated polyposis or "multiple" colorectal adenomas. Proceedings of the National Academy of Sciences of the United States of America, 2002, 99, 2954-2958.	7.1	127
212	Guidance on gastrointestinal surveillance for hereditary non-polyposis colorectal cancer, familial adenomatous polypolis, juvenile polyposis, and Peutz-Jeghers syndrome. Gut, 2002, 51, v21-v27.	12.1	194
213	Guidance on large bowel surveillance for people with two first degree relatives with colorectal cancer or one first degree relative diagnosed with colorectal cancer under 45 years. Gut, 2002, 51, v17-v20.	12.1	45
214	Mismatch Repair Genes hMLH1 and hMSH2 and Colorectal Cancer: A HuGE Review. American Journal of Epidemiology, 2002, 156, 885-902.	3.4	128
215	Evidence for an age-related influence of microsatellite instability on colorectal cancer survival. International Journal of Cancer, 2002, 98, 844-850.	5.1	45
216	Prognosis in DNA Mismatch Repair Deficient Colorectal Cancer: are all MSI Tumours Equivalent?. Familial Cancer, 2002, 3, 85-91.	1.9	37

#	Article	IF	CITATIONS
217	Germline Mutations in BMPR1A/ALK3 Cause a Subset of Cases of Juvenile Polyposis Syndrome and of Cowden and Bannayan-Riley-Ruvalcaba Syndromes*. American Journal of Human Genetics, 2001, 69, 704-711.	6.2	236
218	Early-onset colorectal cancer with stable microsatellite DNA and near-diploid chromosomes. Oncogene, 2001, 20, 4871-4876.	5.9	65
219	Mutation frequency in coding and non-coding repeat sequences in mismatch repair deficient cells derived from normal human tissue. Oncogene, 2001, 20, 7464-7471.	5.9	11
220	Hypermutability at a poly(A/T) tract in the human germline. Nucleic Acids Research, 2001, 29, 4405-4413.	14.5	25
221	Aspirin-induced activation of the NF-kappaB signaling pathway: a novel mechanism for aspirin-mediated apoptosis in colon cancer cells. FASEB Journal, 2001, 15, 1273-5.	0.5	63
222	NF-?B kinetics predetermine TNF-? sensitivity of colorectal cancer cells. Journal of Gene Medicine, 2000, 2, 334-343.	2.8	27
223	Population carrier frequency of hMSH2 and hMLH1 mutations. British Journal of Cancer, 2000, 83, 1643-1645.	6.4	66
224	Sequence interruptions confer differential stability at microsatellite alleles in mismatch repair-deficient cells. Human Molecular Genetics, 2000, 9, 2707-2713.	2.9	28
225	Analysis of genetic and phenotypic heterogeneity in juvenile polyposis. Gut, 2000, 46, 656-660.	12.1	117
226	NF–κB kinetics predetermine TNF-α sensitivity of colorectal cancer cells. , 2000, 2, 334.		1
227	Molecular Markers of Prognosis in Colorectal Cancer. Journal of the National Cancer Institute, 1999, 91, 1267-1269.	6.3	26
228	Mosaicism and Sporadic Familial Adenomatous Polyposis. American Journal of Human Genetics, 1999, 64, 653-658.	6.2	21
229	Systematic Analysis of hMSH2 and hMLH1 in Young Colon Cancer Patients and Controls. American Journal of Human Genetics, 1998, 63, 749-759.	6.2	159
230	GENE THERAPY FOR COLON CANCER. Hematology/Oncology Clinics of North America, 1998, 12, 595-615.	2.2	28
231	Mutations in DPC4 (SMAD4) cause juvenile polyposis syndrome, but only account for a minority of cases. Human Molecular Genetics, 1998, 7, 1907-1912.	2.9	142
232	The case for surveillance of 'high-risk' families. European Journal of Gastroenterology and Hepatology, 1998, 10, 229-234.	1.6	3
233	Cancer risk associated with germline DNA mismatch repair gene mutations. Human Molecular Genetics, 1997, 6, 105-110.	2.9	593
234	Gene Therapy. Edited by N. R. Lemoine and D. N. Cooper. Bios Scientific Publishers, 1996. 343+xiv pages. Price £60, hard cover. ISBN 185996205 X. Gene Therapy Protocols. Edited by Paul D. Robbins. Humana Press, 1997. 432+xiv pages. Price \$74.50 ISBN 0896033074 Genetical Research, 1997, 69, 159-162.	0.9	1

#	Article	IF	CITATIONS
235	APC EXPRESSION IN NORMAL HUMAN TISSUES. , 1997, 181, 426-433.		86
236	Screening for people with a family history of colorectal cancer. BMJ: British Medical Journal, 1997, 314, 1779-1779.	2.3	25
237	Science, medicine, and the future: Colorectal Cancer. BMJ: British Medical Journal, 1997, 314, 1882-1882.	2.3	27
238	Exclusion of PTEN and 10q22-24 as the susceptibility locus for juvenile polyposis syndrome. Cancer Research, 1997, 57, 5017-21.	0.9	58
239	Mutator genes and mosaicism in colorectal cancer. Current Opinion in Genetics and Development, 1996, 6, 76-81.	3.3	17
240	APC mutations in colorectal tumors with mismatch repair deficiency Proceedings of the National Academy of Sciences of the United States of America, 1996, 93, 9049-9054.	7.1	294
241	Extracolonic features of familial adenomatous polyposis in patients with sporadic colorectal cancer. British Journal of Cancer, 1996, 74, 1789-1795.	6.4	10
242	Cranial desmoid tumor associated with homozygous inactivation of the adenomatous polyposis coli gene in a 2-year-old girl with familial adenomatous polyposis. Cancer, 1996, 77, 972-976.	4.1	19
243	Analysis of mismatch repair genes in hereditary non–polyposis colorectal cancer patients. Nature Medicine, 1996, 2, 169-174.	30.7	892
244	The genetics of familial colon cancer. , 1996, , 306-319.		2
244	The genetics of familial colon cancer., 1996,, 306-319. Detailed physical and deletion mapping of 8p with isolation of YAC clones from tumour suppressor loci involved in colorectal cancer. Oncogene, 1996, 12, 1803-8.	5.9	2 34
	Detailed physical and deletion mapping of 8p with isolation of YAC clones from tumour suppressor	5.9 5.9	
245	Detailed physical and deletion mapping of 8p with isolation of YAC clones from tumour suppressor loci involved in colorectal cancer. Oncogene, 1996, 12, 1803-8. Microsatellite instability and the role of hMSH2 in sporadic colorectalcancer. Oncogene, 1996, 12,		34
245	Detailed physical and deletion mapping of 8p with isolation of YAC clones from tumour suppressor loci involved in colorectal cancer. Oncogene, 1996, 12, 1803-8. Microsatellite instability and the role of hMSH2 in sporadic colorectalcancer. Oncogene, 1996, 12, 2641-9. Genetic instability occurs in the majority of young patients with colorectal cancer. Nature Medicine,	5.9	34 179
245 246 247	Detailed physical and deletion mapping of 8p with isolation of YAC clones from tumour suppressor loci involved in colorectal cancer. Oncogene, 1996, 12, 1803-8. Microsatellite instability and the role of hMSH2 in sporadic colorectalcancer. Oncogene, 1996, 12, 2641-9. Genetic instability occurs in the majority of young patients with colorectal cancer. Nature Medicine, 1995, 1, 348-352. Suggested Screening Guidelines for Familial Colorectal Cancer. Journal of Medical Screening, 1995, 2,	5.9 30.7	34 179 355
245 246 247 248	Detailed physical and deletion mapping of 8p with isolation of YAC clones from tumour suppressor loci involved in colorectal cancer. Oncogene, 1996, 12, 1803-8. Microsatellite instability and the role of hMSH2 in sporadic colorectalcancer. Oncogene, 1996, 12, 2641-9. Genetic instability occurs in the majority of young patients with colorectal cancer. Nature Medicine, 1995, 1, 348-352. Suggested Screening Guidelines for Familial Colorectal Cancer. Journal of Medical Screening, 1995, 2, 45-51.	5.9 30.7 2.3	34 179 355 23
245 246 247 248	Detailed physical and deletion mapping of 8p with isolation of YAC clones from tumour suppressor loci involved in colorectal cancer. Oncogene, 1996, 12, 1803-8. Microsatellite instability and the role of hMSH2 in sporadic colorectalcancer. Oncogene, 1996, 12, 2641-9. Genetic instability occurs in the majority of young patients with colorectal cancer. Nature Medicine, 1995, 1, 348-352. Suggested Screening Guidelines for Familial Colorectal Cancer. Journal of Medical Screening, 1995, 2, 45-51. Genetics of colorectal cancer. British Medical Bulletin, 1994, 50, 640-655.	5.9 30.7 2.3 6.9	34 179 355 23 22

#	Article	IF	Citations
253	Exclusion of constitutional p53 mutations as a cause of genetic susceptibility to colorectal cancer. British Journal of Cancer, 1993, 68, 712-714.	6.4	4
254	Deletion mapping in colorectal cancer of a putative tumour suppressor gene in 8p22-p21.3. Oncogene, 1993, 8, 1391-6.	5.9	52
255	Presymptomatic diagnosis of polyposis coli by DNA. Gastroenterology, 1992, 102, 374.	1.3	0
256	Colorectal cancer genetics. Seminars in Cancer Biology, 1992, 3, 131-40.	9.6	9
257	Allele losses and onco-suppressor genes. Journal of Pathology, 1991, 163, 1-5.	4.5	7
258	Linked DNA markers for presymptomatic diagnosis of familial adenomatous polyposis. Lancet, The, 1991, 337, 313-316.	13.7	60
259	Telomere loss and cancer. Nature, 1991, 350, 197-197.	27.8	4
260	Telomere reduction in human colorectal carcinoma and with ageing. Nature, 1990, 346, 866-868.	27.8	1,612
261	Status of the APC Gene in Familial and Sporadic Colorectal Tumours as Determined by Closely Flanking Markers. , 1990, , 453-456.		3
262	Genetic linkage map of six polymorphic DNA markers around the gene for familial adenomatous polyposis on chromosome 5. American Journal of Human Genetics, 1990, 47, 982-7.	6.2	25
263	Linkage analysis in familial adenomatous polyposis: Order of C11P11 (D5S71) and ?227 (D5S37) loci at the apc gene. Genomics, 1989, 5, 350-353.	2.9	23
264	High frequency of APC loss in sporadic colorectal carcinoma due to breaks clustered in 5q21-22. Oncogene, 1989, 4, 1169-74.	5.9	95
265	Guidelines for selective radiological assessment of inversion ankle injuries: Authors' reply. BMJ: British Medical Journal, 1986, 293, 957-957.	2.3	0
266	Distal arterial emboli following seat belt injury of the aorta. Injury, 1986, 17, 370-371.	1.7	8
267	Guidelines for selective radiological assessment of inversion ankle injuries BMJ: British Medical Journal, 1986, 293, 603-605.	2.3	62
268	Genetics of colorectal cancer., 0,, 245-267.		0
269	FDG PET-CT imaging for pre operative staging in patients with colorectal cancer. The Cochrane Library, 0, , .	2.8	4