

Daniel D Buchanan

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

237
papers

11,343
citations

58
h-index

100
g-index

273
ext. papers

13,803
ext. citations

6.6
avg, IF

5.4
L-index

#	Paper	IF	Citations
237	CpG island methylator phenotype underlies sporadic microsatellite instability and is tightly associated with BRAF mutation in colorectal cancer. <i>Nature Genetics</i> , 2006 , 38, 787-93	36.3	1514
236	Identification of Lynch syndrome among patients with colorectal cancer. <i>JAMA - Journal of the American Medical Association</i> , 2012 , 308, 1555-65	27.4	323
235	Risks of Lynch syndrome cancers for MSH6 mutation carriers. <i>Journal of the National Cancer Institute</i> , 2010 , 102, 193-201	9.7	279
234	Association between molecular subtypes of colorectal cancer and patient survival. <i>Gastroenterology</i> , 2015 , 148, 77-87.e2	13.3	273
233	Colorectal and other cancer risks for carriers and noncarriers from families with a DNA mismatch repair gene mutation: a prospective cohort study. <i>Journal of Clinical Oncology</i> , 2012 , 30, 958-64	2.2	245
232	Colorectal cancer cell lines are representative models of the main molecular subtypes of primary cancer. <i>Cancer Research</i> , 2014 , 74, 3238-47	10.1	240
231	Advanced colorectal polyps with the molecular and morphological features of serrated polyps and adenomas: concept of a fusion pathway to colorectal cancer. <i>Histopathology</i> , 2006 , 49, 121-31	7.3	214
230	Correlation of tumour BRAF mutations and MLH1 methylation with germline mismatch repair (MMR) gene mutation status: a literature review assessing utility of tumour features for MMR variant classification. <i>Journal of Medical Genetics</i> , 2012 , 49, 151-7	5.8	200
229	Prevalence and Penetrance of Major Genes and Polygenes for Colorectal Cancer. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2017 , 26, 404-412	4	185
228	Discovery of common and rare genetic risk variants for colorectal cancer. <i>Nature Genetics</i> , 2019 , 51, 76-83	36.3	177
227	GSK3B polymorphisms alter transcription and splicing in Parkinson disease. <i>Annals of Neurology</i> , 2005 , 58, 829-39	9.4	172
226	Cancer risks for MLH1 and MSH2 mutation carriers. <i>Human Mutation</i> , 2013 , 34, 490-7	4.7	171
225	Hyperplastic polyposis syndrome: phenotypic presentations and the role of MBD4 and MYH. <i>Gastroenterology</i> , 2006 , 131, 30-9	13.3	166
224	KRAS-mutation status in relation to colorectal cancer survival: the joint impact of correlated tumour markers. <i>British Journal of Cancer</i> , 2013 , 108, 1757-64	8.7	163
223	Risks of primary extracolonic cancers following colorectal cancer in lynch syndrome. <i>Journal of the National Cancer Institute</i> , 2012 , 104, 1363-72	9.7	158
222	Tumor mismatch repair immunohistochemistry and DNA MLH1 methylation testing of patients with endometrial cancer diagnosed at age younger than 60 years optimizes triage for population-level germline mismatch repair gene mutation testing. <i>Journal of Clinical Oncology</i> , 2014 , 32, 90-100	2.2	149
221	Tau haplotypes regulate transcription and are associated with Parkinson disease. <i>Annals of Neurology</i> , 2004 , 55, 329-34	9.4	147

220	Determining Risk of Colorectal Cancer and Starting Age of Screening Based on Lifestyle, Environmental, and Genetic Factors. <i>Gastroenterology</i> , 2018 , 154, 2152-2164.e19	13.3	131
219	Risk of colorectal cancer for carriers of mutations in MUTYH, with and without a family history of cancer. <i>Gastroenterology</i> , 2014 , 146, 1208-11.e1-5	13.3	128
218	Extensive DNA methylation in normal colorectal mucosa in hyperplastic polyposis. <i>Gut</i> , 2006 , 55, 1467-74.9.2		120
217	Prediction of overall survival in stage II and III colon cancer beyond TNM system: a retrospective, pooled biomarker study. <i>Annals of Oncology</i> , 2017 , 28, 1023-1031	10.3	116
216	Evidence for BRAF mutation and variable levels of microsatellite instability in a syndrome of familial colorectal cancer. <i>Clinical Gastroenterology and Hepatology</i> , 2005 , 3, 254-63	6.9	113
215	BRAFV600E immunohistochemistry facilitates universal screening of colorectal cancers for Lynch syndrome. <i>American Journal of Surgical Pathology</i> , 2013 , 37, 1592-602	6.7	112
214	Identification of susceptibility loci for colorectal cancer in a genome-wide meta-analysis. <i>Human Molecular Genetics</i> , 2014 , 23, 4729-37	5.6	107
213	Colorectal carcinomas with KRAS mutation are associated with distinctive morphological and molecular features. <i>Modern Pathology</i> , 2013 , 26, 825-34	9.8	106
212	BRAF mutation status and survival after colorectal cancer diagnosis according to patient and tumor characteristics. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2012 , 21, 1792-8	4	103
211	PIK3CA activating mutation in colorectal carcinoma: associations with molecular features and survival. <i>PLoS ONE</i> , 2013 , 8, e65479	3.7	102
210	Cancer Risks for PMS2-Associated Lynch Syndrome. <i>Journal of Clinical Oncology</i> , 2018 , 36, 2961-2968	2.2	102
209	Expression of MUC2, MUC5AC, MUC5B, and MUC6 mucins in colorectal cancers and their association with the CpG island methylator phenotype. <i>Modern Pathology</i> , 2013 , 26, 1642-56	9.8	99
208	Quality assessment and correlation of microsatellite instability and immunohistochemical markers among population- and clinic-based colorectal tumors results from the Colon Cancer Family Registry. <i>Journal of Molecular Diagnostics</i> , 2011 , 13, 271-81	5.1	95
207	A new GWAS and meta-analysis with 1000Genomes imputation identifies novel risk variants for colorectal cancer. <i>Scientific Reports</i> , 2015 , 5, 10442	4.9	94
206	Ethnicity and risk for colorectal cancers showing somatic BRAF V600E mutation or CpG island methylator phenotype. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2008 , 17, 1774-80	4	89
205	Risk of metachronous colon cancer following surgery for rectal cancer in mismatch repair gene mutation carriers. <i>Annals of Surgical Oncology</i> , 2013 , 20, 1829-36	3.1	87
204	DNA methylation-based biological aging and cancer risk and survival: Pooled analysis of seven prospective studies. <i>International Journal of Cancer</i> , 2018 , 142, 1611-1619	7.5	83
203	Association analyses identify 31 new risk loci for colorectal cancer susceptibility. <i>Nature Communications</i> , 2019 , 10, 2154	17.4	81

202	Telomere length varies by DNA extraction method: implications for epidemiologic research. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2013 , 22, 2047-54	4	81
201	Molecular, pathologic, and clinical features of early-onset endometrial cancer: identifying presumptive Lynch syndrome patients. <i>Clinical Cancer Research</i> , 2008 , 14, 1692-700	12.9	81
200	Lynch syndrome-associated breast cancers: clinicopathologic characteristics of a case series from the colon cancer family registry. <i>Clinical Cancer Research</i> , 2010 , 16, 2214-24	12.9	80
199	Risks of colorectal and other cancers after endometrial cancer for women with Lynch syndrome. <i>Journal of the National Cancer Institute</i> , 2013 , 105, 274-9	9.7	78
198	The prognostic impact of consensus molecular subtypes (CMS) and its predictive effects for bevacizumab benefit in metastatic colorectal cancer: molecular analysis of the AGITG MAX clinical trial. <i>Annals of Oncology</i> , 2018 , 29, 2240-2246	10.3	77
197	Phenotype and polyp landscape in serrated polyposis syndrome: a series of 100 patients from genetics clinics. <i>American Journal of Surgical Pathology</i> , 2012 , 36, 876-82	6.7	73
196	Variations in the monoamine oxidase B (MAOB) gene are associated with Parkinson disease. <i>Movement Disorders</i> , 1999 , 14, 219-24	7	73
195	Mutational Signature Analysis Reveals NTHL1 Deficiency to Cause a Multi-tumor Phenotype. <i>Cancer Cell</i> , 2019 , 35, 256-266.e5	24.3	72
194	Identification of nine new susceptibility loci for endometrial cancer. <i>Nature Communications</i> , 2018 , 9, 3166	17.4	70
193	A multifactorial likelihood model for MMR gene variant classification incorporating probabilities based on sequence bioinformatics and tumor characteristics: a report from the Colon Cancer Family Registry. <i>Human Mutation</i> , 2013 , 34, 200-9	4.7	70
192	Cancer risks for relatives of patients with serrated polyposis. <i>American Journal of Gastroenterology</i> , 2012 , 107, 770-8	0.7	69
191	Germline TP53 Mutations in Patients With Early-Onset Colorectal Cancer in the Colon Cancer Family Registry. <i>JAMA Oncology</i> , 2015 , 1, 214-21	13.4	68
190	Risk of extracolonic cancers for people with biallelic and monoallelic mutations in MUTYH. <i>International Journal of Cancer</i> , 2016 , 139, 1557-63	7.5	67
189	Novel Common Genetic Susceptibility Loci for Colorectal Cancer. <i>Journal of the National Cancer Institute</i> , 2019 , 111, 146-157	9.7	67
188	Aspirin, Ibuprofen, and the Risk of Colorectal Cancer in Lynch Syndrome. <i>Journal of the National Cancer Institute</i> , 2015 , 107,	9.7	66
187	Cancer risks for monoallelic MUTYH mutation carriers with a family history of colorectal cancer. <i>International Journal of Cancer</i> , 2011 , 129, 2256-62	7.5	66
186	Promoter methylation of Wnt antagonists DKK1 and SFRP1 is associated with opposing tumor subtypes in two large populations of colorectal cancer patients. <i>Carcinogenesis</i> , 2011 , 32, 741-7	4.6	66
185	Cross-Cancer Genome-Wide Analysis of Lung, Ovary, Breast, Prostate, and Colorectal Cancer Reveals Novel Pleiotropic Associations. <i>Cancer Research</i> , 2016 , 76, 5103-14	10.1	66

184	Association of DNA Methylation-Based Biological Age With Health Risk Factors and Overall and Cause-Specific Mortality. <i>American Journal of Epidemiology</i> , 2018 , 187, 529-538	3.8	61
183	Immunohistochemical testing of conventional adenomas for loss of expression of mismatch repair proteins in Lynch syndrome mutation carriers: a case series from the Australasian site of the colon cancer family registry. <i>Modern Pathology</i> , 2012 , 25, 722-30	9.8	61
182	Risk factors for colorectal cancer in patients with multiple serrated polyps: a cross-sectional case series from genetics clinics. <i>PLoS ONE</i> , 2010 , 5, e11636	3.7	60
181	Association of the colorectal CpG island methylator phenotype with molecular features, risk factors, and family history. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2015 , 24, 512-519	4	59
180	Body size and risk for colorectal cancers showing BRAF mutations or microsatellite instability: a pooled analysis. <i>International Journal of Epidemiology</i> , 2012 , 41, 1060-72	7.8	59
179	Pooled analysis of iron-related genes in Parkinson® disease: association with transferrin. <i>Neurobiology of Disease</i> , 2014 , 62, 172-8	7.5	57
178	Pro-inflammatory fatty acid profile and colorectal cancer risk: A Mendelian randomisation analysis. <i>European Journal of Cancer</i> , 2017 , 84, 228-238	7.5	56
177	Classifying MLH1 and MSH2 variants using bioinformatic prediction, splicing assays, segregation, and tumor characteristics. <i>Human Mutation</i> , 2009 , 30, 757-70	4.7	55
176	Female Hormonal Factors and the Risk of Endometrial Cancer in Lynch Syndrome. <i>JAMA - Journal of the American Medical Association</i> , 2015 , 314, 61-71	27.4	53
175	Associations of alcohol intake, smoking, physical activity and obesity with survival following colorectal cancer diagnosis by stage, anatomic site and tumor molecular subtype. <i>International Journal of Cancer</i> , 2018 , 142, 238-250	7.5	53
174	Clinical problems of colorectal cancer and endometrial cancer cases with unknown cause of tumor mismatch repair deficiency (suspected Lynch syndrome). <i>The Application of Clinical Genetics</i> , 2014 , 7, 183-93	3.1	53
173	Identification of BRCA1 missense substitutions that confer partial functional activity: potential moderate risk variants?. <i>Breast Cancer Research</i> , 2007 , 9, R82	8.3	51
172	Mendelian randomisation implicates hyperlipidaemia as a risk factor for colorectal cancer. <i>International Journal of Cancer</i> , 2017 , 140, 2701-2708	7.5	50
171	Tumour MLH1 promoter region methylation testing is an effective prescreen for Lynch Syndrome (HNPCC). <i>Journal of Medical Genetics</i> , 2014 , 51, 789-96	5.8	50
170	The Cys282Tyr polymorphism in the HFE gene in Australian Parkinson® disease patients. <i>Neuroscience Letters</i> , 2002 , 327, 91-4	3.3	48
169	Screening PARK genes for mutations in early-onset Parkinson® disease patients from Queensland, Australia. <i>Parkinsonism and Related Disorders</i> , 2009 , 15, 105-9	3.6	47
168	Cumulative Burden of Colorectal Cancer-Associated Genetic Variants Is More Strongly Associated With Early-Onset vs Late-Onset Cancer. <i>Gastroenterology</i> , 2020 , 158, 1274-1286.e12	13.3	47
167	Shared heritability and functional enrichment across six solid cancers. <i>Nature Communications</i> , 2019 , 10, 431	17.4	45

166	Circulating Levels of Insulin-like Growth Factor 1 and Insulin-like Growth Factor Binding Protein 3 Associate With Risk of Colorectal Cancer Based on Serologic and Mendelian Randomization Analyses. <i>Gastroenterology</i> , 2020 , 158, 1300-1312.e20	13.3	45
165	KRAS mutations in ovarian low-grade endometrioid adenocarcinoma: association with concurrent endometriosis. <i>Human Pathology</i> , 2012 , 43, 1177-83	3.7	43
164	Update on Hereditary Colorectal Cancer: Improving the Clinical Utility of Multigene Panel Testing. <i>Clinical Colorectal Cancer</i> , 2018 , 17, e293-e305	3.8	42
163	Phenotypic diversity in patients with multiple serrated polyps: a genetics clinic study. <i>International Journal of Colorectal Disease</i> , 2010 , 25, 703-12	3	42
162	High prevalence of mismatch repair deficiency in prostate cancers diagnosed in mismatch repair gene mutation carriers from the colon cancer family registry. <i>Familial Cancer</i> , 2014 , 13, 573-82	3	40
161	Body mass index in early adulthood and colorectal cancer risk for carriers and non-carriers of germline mutations in DNA mismatch repair genes. <i>British Journal of Cancer</i> , 2011 , 105, 162-9	8.7	40
160	Confirmation of linkage to and localization of familial colon cancer risk haplotype on chromosome 9q22. <i>Cancer Research</i> , 2010 , 70, 5409-18	10.1	40
159	A frame-shift mutation of PMS2 is a widespread cause of Lynch syndrome. <i>Journal of Medical Genetics</i> , 2008 , 45, 340-5	5.8	40
158	Should the grading of colorectal adenocarcinoma include microsatellite instability status?. <i>Human Pathology</i> , 2014 , 45, 2077-84	3.7	39
157	Mendelian randomisation analysis strongly implicates adiposity with risk of developing colorectal cancer. <i>British Journal of Cancer</i> , 2016 , 115, 266-72	8.7	39
156	Association between body mass index and mortality for colorectal cancer survivors: overall and by tumor molecular phenotype. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2015 , 24, 1229-38	4	38
155	The intestinal epithelial cell differentiation marker intestinal alkaline phosphatase (ALPi) is selectively induced by histone deacetylase inhibitors (HDACi) in colon cancer cells in a Kruppel-like factor 5 (KLF5)-dependent manner. <i>Journal of Biological Chemistry</i> , 2014 , 289, 25306-16	5.4	38
154	Identification of novel variants in colorectal cancer families by high-throughput exome sequencing. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2013 , 22, 1239-51	4	36
153	Association between hypermethylation of DNA repetitive elements in white blood cell DNA and early-onset colorectal cancer. <i>Epigenetics</i> , 2013 , 8, 748-55	5.7	36
152	Physical activity and risks of breast and colorectal cancer: a Mendelian randomisation analysis. <i>Nature Communications</i> , 2020 , 11, 597	17.4	36
151	Lack of evidence for germline mutations in patients with serrated polyposis syndrome from a large multinational study. <i>Gut</i> , 2017 , 66, 1170-1172	19.2	35
150	Multiplicity and molecular heterogeneity of colorectal carcinomas in individuals with serrated polyposis. <i>American Journal of Surgical Pathology</i> , 2013 , 37, 434-42	6.7	35
149	Mutation deep within an intron of MSH2 causes Lynch syndrome. <i>Familial Cancer</i> , 2011 , 10, 297-301	3	34

148	Risk of colorectal cancer for carriers of a germ-line mutation in POLE or POLD1. <i>Genetics in Medicine</i> , 2018 , 20, 890-895	8.1	34
147	Characterisation of familial colorectal cancer Type X, Lynch syndrome, and non-familial colorectal cancer. <i>British Journal of Cancer</i> , 2014 , 111, 598-602	8.7	33
146	Determining the frequency of de novo germline mutations in DNA mismatch repair genes. <i>Journal of Medical Genetics</i> , 2011 , 48, 530-4	5.8	32
145	Role of tumour molecular and pathology features to estimate colorectal cancer risk for first-degree relatives. <i>Gut</i> , 2015 , 64, 101-10	19.2	31
144	Genome-wide Modeling of Polygenic Risk Score in Colorectal Cancer Risk. <i>American Journal of Human Genetics</i> , 2020 , 107, 432-444	11	31
143	Quantifying the utility of single nucleotide polymorphisms to guide colorectal cancer screening. <i>Future Oncology</i> , 2016 , 12, 503-13	3.6	30
142	Promoter methylation of Wnt5a is associated with microsatellite instability and BRAF V600E mutation in two large populations of colorectal cancer patients. <i>British Journal of Cancer</i> , 2011 , 104, 1906-12	8.7	30
141	Genome-wide search for gene-gene interactions in colorectal cancer. <i>PLoS ONE</i> , 2012 , 7, e52535	3.7	29
140	Alcohol Consumption and the Risk of Colorectal Cancer for Mismatch Repair Gene Mutation Carriers. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2017 , 26, 366-375	4	28
139	Sequence variation in the proximity of IDE may impact age at onset of both Parkinson disease and Alzheimer disease. <i>Neurogenetics</i> , 2004 , 5, 115-9	3	28
138	Tumor testing to identify lynch syndrome in two Australian colorectal cancer cohorts. <i>Journal of Gastroenterology and Hepatology (Australia)</i> , 2017 , 32, 427-438	4	27
137	Variation at 2q35 (PNKD and TMBIM1) influences colorectal cancer risk and identifies a pleiotropic effect with inflammatory bowel disease. <i>Human Molecular Genetics</i> , 2016 , 25, 2349-2359	5.6	27
136	Meta-analysis of genome-wide association studies identifies common susceptibility polymorphisms for colorectal and endometrial cancer near SH2B3 and TSHZ1. <i>Scientific Reports</i> , 2015 , 5, 17369	4.9	27
135	Dysfunctional epigenetic aging of the normal colon and colorectal cancer risk. <i>Clinical Epigenetics</i> , 2020 , 12, 5	7.7	27
134	Are the common genetic variants associated with colorectal cancer risk for DNA mismatch repair gene mutation carriers?. <i>European Journal of Cancer</i> , 2013 , 49, 1578-87	7.5	26
133	Novel associations between blood DNA methylation and body mass index in middle-aged and older adults. <i>International Journal of Obesity</i> , 2018 , 42, 887-896	5.5	25
132	Improving identification of lynch syndrome patients: a comparison of research data with clinical records. <i>International Journal of Cancer</i> , 2013 , 132, 2876-83	7.5	25
131	Prediagnostic Physical Activity and Colorectal Cancer Survival: Overall and Stratified by Tumor Characteristics. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2015 , 24, 1130-7	4	25

130	A novel colorectal cancer risk locus at 4q32.2 identified from an international genome-wide association study. <i>Carcinogenesis</i> , 2014 , 35, 2512-9	4.6	25
129	Germline mutations in PMS2 and MLH1 in individuals with solitary loss of PMS2 expression in colorectal carcinomas from the Colon Cancer Family Registry Cohort. <i>BMJ Open</i> , 2016 , 6, e010293	3	24
128	Comparison of Prediction Models for Lynch Syndrome Among Individuals With Colorectal Cancer. <i>Journal of the National Cancer Institute</i> , 2016 , 108,	9.7	24
127	Dietary intake of one-carbon metabolism nutrients and DNA methylation in peripheral blood. <i>American Journal of Clinical Nutrition</i> , 2018 , 108, 611-621	7	24
126	Cohort Profile: The Colon Cancer Family Registry Cohort (CCFRC). <i>International Journal of Epidemiology</i> , 2018 , 47, 387-388i	7.8	23
125	Colorectal cancer linkage on chromosomes 4q21, 8q13, 12q24, and 15q22. <i>PLoS ONE</i> , 2012 , 7, e38175	3.7	23
124	Association of APOE with Parkinson disease age-at-onset in women. <i>Neuroscience Letters</i> , 2007 , 411, 185-8	3.3	23
123	Potential impact of family history-based screening guidelines on the detection of early-onset colorectal cancer. <i>Cancer</i> , 2020 , 126, 3013-3020	6.4	23
122	Family history of colorectal cancer in BRAF p.V600E-mutated colorectal cancer cases. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2013 , 22, 917-26	4	22
121	Genome-wide association study and meta-analysis in Northern European populations replicate multiple colorectal cancer risk loci. <i>International Journal of Cancer</i> , 2018 , 142, 540-546	7.5	21
120	Family history of cancer predicts endometrial cancer risk independently of Lynch Syndrome: Implications for genetic counselling. <i>Gynecologic Oncology</i> , 2017 , 147, 381-387	4.9	21
119	The association of telomere length with colorectal cancer differs by the age of cancer onset. <i>Clinical and Translational Gastroenterology</i> , 2014 , 5, e52	4.2	21
118	Germline HOXB13 p.Gly84Glu mutation and risk of colorectal cancer. <i>Cancer Epidemiology</i> , 2013 , 37, 424-428		21
117	Multivitamin, calcium and folic acid supplements and the risk of colorectal cancer in Lynch syndrome. <i>International Journal of Epidemiology</i> , 2016 , 45, 940-53	7.8	21
116	Lifetime alcohol intake is associated with an increased risk of KRAS+ and BRAF-/KRAS- but not BRAF+ colorectal cancer. <i>International Journal of Cancer</i> , 2017 , 140, 1485-1493	7.5	20
115	Targeted sequencing of 36 known or putative colorectal cancer susceptibility genes. <i>Molecular Genetics & Genomic Medicine</i> , 2017 , 5, 553-569	2.3	20
114	Germline Mutations in the Polyposis-Associated Genes BMPR1A, SMAD4, PTEN, MUTYH and GREM1 Are Not Common in Individuals with Serrated Polyposis Syndrome. <i>PLoS ONE</i> , 2013 , 8, e66705	3.7	20
113	Hyperplastic polyp of the duodenum: a report of 9 cases with immunohistochemical and molecular findings. <i>Human Pathology</i> , 2011 , 42, 1953-9	3.7	20

112	Endometrial cancer risk and survival by tumor MMR status. <i>Journal of Gynecologic Oncology</i> , 2018 , 29, e39	4	19
111	Analysis of families with Lynch syndrome complicated by advanced serrated neoplasia: the importance of pathology review and pedigree analysis. <i>Familial Cancer</i> , 2009 , 8, 313-23	3	19
110	The ACE deletion polymorphism is not associated with Parkinson® disease. <i>European Neurology</i> , 1999 , 41, 103-6	2.1	19
109	Risk factors for metachronous colorectal cancer following a primary colorectal cancer: A prospective cohort study. <i>International Journal of Cancer</i> , 2016 , 139, 1081-90	7.5	19
108	Targeted sequencing of established and candidate colorectal cancer genes in the Colon Cancer Family Registry Cohort. <i>Oncotarget</i> , 2017 , 8, 93450-93463	3.3	18
107	SNP rs16906252C>T Is an Expression and Methylation Quantitative Trait Locus Associated with an Increased Risk of Developing MGMT-Methylated Colorectal Cancer. <i>Clinical Cancer Research</i> , 2016 , 22, 6266-6277	12.9	18
106	Do serrated neoplasms of the small intestine represent a distinct entity? Pathological findings and molecular alterations in a series of 13 cases. <i>Histopathology</i> , 2015 , 66, 333-42	7.3	18
105	Mendelian randomization analysis of C-reactive protein on colorectal cancer risk. <i>International Journal of Epidemiology</i> , 2019 , 48, 767-780	7.8	18
104	Ability of known susceptibility SNPs to predict colorectal cancer risk for persons with and without a family history. <i>Familial Cancer</i> , 2019 , 18, 389-397	3	17
103	Association Between Molecular Subtypes of Colorectal Tumors and Patient Survival, Based on Pooled Analysis of 7 International Studies. <i>Gastroenterology</i> , 2020 , 158, 2158-2168.e4	13.3	17
102	Linkage to chromosome 2q32.2-q33.3 in familial serrated neoplasia (Jass syndrome). <i>Familial Cancer</i> , 2011 , 10, 245-54	3	17
101	Adiposity, metabolites, and colorectal cancer risk: Mendelian randomization study. <i>BMC Medicine</i> , 2020 , 18, 396	11.4	17
100	Genetic variant predictors of gene expression provide new insight into risk of colorectal cancer. <i>Human Genetics</i> , 2019 , 138, 307-326	6.3	17
99	Clinicopathologic Risk Factor Distributions for MLH1 Promoter Region Methylation in CIMP-Positive Tumors. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2016 , 25, 68-75	4	16
98	Landscape of somatic single nucleotide variants and indels in colorectal cancer and impact on survival. <i>Nature Communications</i> , 2020 , 11, 3644	17.4	16
97	Immunophenotypic analysis of ovarian endometrioid adenocarcinoma: correlation with KRAS mutation and the presence of endometriosis. <i>Pathology</i> , 2013 , 45, 559-66	1.6	15
96	The parkin gene S/N167 polymorphism in Australian Parkinson® disease patients and controls. <i>Parkinsonism and Related Disorders</i> , 2001 , 7, 89-91	3.6	15
95	Characteristics of Early-Onset vs Late-Onset Colorectal Cancer: A Review. <i>JAMA Surgery</i> , 2021 , 156, 865-874	8.4	15

94	Circulating 25-Hydroxyvitamin D Concentration and Risk of Breast, Prostate, and Colorectal Cancers: The Melbourne Collaborative Cohort Study. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2019 , 28, 900-908	4	14
93	DNA mismatch repair protein deficient non-neoplastic colonic crypts: a novel indicator of Lynch syndrome. <i>Modern Pathology</i> , 2018 , 31, 1608-1618	9.8	14
92	Association between hypermethylation of DNA repetitive elements in white blood cell DNA and pancreatic cancer. <i>Cancer Epidemiology</i> , 2014 , 38, 576-82	2.8	14
91	A perspective on bi-allelic MUTYH mutations in patients with hyperplastic polyposis syndrome. <i>Gastroenterology</i> , 2009 , 136, 2407-8	13.3	14
90	Lack of association between CYP1A1 polymorphism and Parkinson disease in a Chinese population. <i>Journal of Neural Transmission</i> , 2002 , 109, 35-9	4.3	14
89	Consequences of germline variation disrupting the constitutional translational initiation codon start sites of MLH1 and BRCA2: Use of potential alternative start sites and implications for predicting variant pathogenicity. <i>Molecular Carcinogenesis</i> , 2015 , 54, 513-22	5	13
88	Detection of large scale 3Q deletions in the PMS2 gene amongst Colon-CFR participants: have we been missing anything?. <i>Familial Cancer</i> , 2013 , 12, 563-6	3	13
87	Mendelian randomization analyses suggest a role for cholesterol in the development of endometrial cancer. <i>International Journal of Cancer</i> , 2021 , 148, 307-319	7.5	13
86	Phenotypic confirmation of oligodontia, colorectal polyposis and cancer in a family carrying an exon 7 nonsense variant in the AXIN2 gene. <i>Familial Cancer</i> , 2019 , 18, 311-315	3	11
85	Systematic meta-analyses, field synopsis and global assessment of the evidence of genetic association studies in colorectal cancer. <i>Gut</i> , 2020 , 69, 1460-1471	19.2	11
84	Somatic mutations of the coding microsatellites within the beta-2-microglobulin gene in mismatch repair-deficient colorectal cancers and adenomas. <i>Familial Cancer</i> , 2018 , 17, 91-100	3	11
83	Lynch syndrome and cervical cancer. <i>International Journal of Cancer</i> , 2015 , 137, 2757-61	7.5	11
82	Methylation of Breast Cancer Predisposition Genes in Early-Onset Breast Cancer: Australian Breast Cancer Family Registry. <i>PLoS ONE</i> , 2016 , 11, e0165436	3.7	11
81	Biological Aging Measures Based on Blood DNA Methylation and Risk of Cancer: A Prospective Study. <i>JNCI Cancer Spectrum</i> , 2021 , 5, pkaa109	4.6	11
80	Medically actionable pathogenic variants in a population of 13,131 healthy elderly individuals. <i>Genetics in Medicine</i> , 2020 , 22, 1883-1886	8.1	10
79	Monoallelic NTHL1 Loss-of-Function Variants and Risk of Polyposis and Colorectal Cancer. <i>Gastroenterology</i> , 2020 , 159, 2241-2243.e6	13.3	10
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34	Associations between Genetically Predicted Circulating Protein Concentrations and Endometrial Cancer Risk. <i>Cancers</i> , 2021 , 13,	6.6	3
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25	Overall lack of replication of associations between dietary intake of folate and vitamin B-12 and DNA methylation in peripheral blood. <i>American Journal of Clinical Nutrition</i> , 2020 , 111, 228-230	7	2
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16	Evaluating the utility of tumour mutational signatures for identifying hereditary colorectal cancer and polyposis syndrome carriers		1
15	Biological aging measures based on blood DNA methylation and risk of cancer: a prospective study		1
14	A combined proteomics and Mendelian randomization approach to investigate the effects of aspirin-targeted proteins on colorectal cancer		1
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