Daniel D Buchanan

List of Publications by Year in Descending Order

Source: https://exaly.com/author-pdf/8246655/daniel-d-buchanan-publications-by-year.pdf

Version: 2024-04-19

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

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

58 11,343 100 237 h-index g-index citations papers 6.6 13,803 273 5.4 L-index avg, IF ext. citations ext. papers

#	Paper	IF	Citations
237	Genome-wide association study identifies tumor anatomical site-specific risk variants for colorectal cancer survival <i>Scientific Reports</i> , 2022 , 12, 127	4.9	2
236	Impact of microsatellite status in early-onset colonic cancer British Journal of Surgery, 2022,	5.3	1
235	Beyond GWAS of Colorectal Cancer: Evidence of Interaction with Alcohol Consumption and Putative Causal Variant for the 10q24.2 Region <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2022 , OF1-OF13	4	O
234	Cross-Cancer Genome-Wide Association Study of Endometrial Cancer and Epithelial Ovarian Cancer Identifies Genetic Risk Regions Associated with Risk of Both Cancers. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2021 , 30, 217-228	4	7
233	A Combined Proteomics and Mendelian Randomization Approach to Investigate the Effects of Aspirin-Targeted Proteins on Colorectal Cancer. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2021 , 30, 564-575	4	2
232	Molecular and Pathology Features of Colorectal Tumors and Patient Outcomes Are Associated with and Its Subspecies. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2021 ,	4	1
231	Salicylic Acid and Risk of Colorectal Cancer: A Two-Sample Mendelian Randomization Study. <i>Nutrients</i> , 2021 , 13,	6.7	1
230	Rare germline variants in the AXIN2 gene in families with colonic polyposis and colorectal cancer. <i>Familial Cancer</i> , 2021 , 1	3	0
229	A genome-wide search for determinants of survival in 1926 patients with advanced colorectal cancer with follow-up in over 22,000 patients. <i>European Journal of Cancer</i> , 2021 , 159, 247-258	7.5	1
228	"Left in limbo": Exploring how patients with colorectal cancer interpret and respond to a suspected Lynch syndrome diagnosis. <i>Hereditary Cancer in Clinical Practice</i> , 2021 , 19, 43	2.3	О
227	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
226	Germline and Tumor Sequencing as a Diagnostic Tool To Resolve Suspected Lynch Syndrome. Journal of Molecular Diagnostics, 2021 , 23, 358-371	5.1	1
225	Response to Li and Hopper. American Journal of Human Genetics, 2021, 108, 527-529	11	1
224	Assessment of a Polygenic Risk Score for Colorectal Cancer to Predict Risk of Lynch Syndrome Colorectal Cancer. <i>JNCI Cancer Spectrum</i> , 2021 , 5, pkab022	4.6	2
223	Epigenetic Drift Association with Cancer Risk and Survival, and Modification by Sex. <i>Cancers</i> , 2021 , 13,	6.6	4
222	Associations between Genetically Predicted Circulating Protein Concentrations and Endometrial Cancer Risk. <i>Cancers</i> , 2021 , 13,	6.6	3
221	DNA Methylation Signatures and the Contribution of Age-Associated Methylomic Drift to Carcinogenesis in Early-Onset Colorectal Cancer. <i>Cancers</i> , 2021 , 13,	6.6	3

(2020-2021)

220	Genetically Predicted Circulating C-Reactive Protein Concentration and Colorectal Cancer Survival: A Mendelian Randomization Consortium Study. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2021 , 30, 1349-1358	4	1
219	Association Between Smoking and Molecular Subtypes of Colorectal Cancer. <i>JNCI Cancer Spectrum</i> , 2021 , 5, pkab056	4.6	2
218	No Difference in Penetrance between Truncating and Missense/Aberrant Splicing Pathogenic Variants in and : A Prospective Lynch Syndrome Database Study. <i>Journal of Clinical Medicine</i> , 2021 , 10,	5.1	1
217	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
216	An integrated mass spectrometry imaging and digital pathology workflow for objective detection of colorectal tumours by unique atomic signatures. <i>Chemical Science</i> , 2021 , 12, 10321-10333	9.4	5
215	Evaluating the utility of tumour mutational signatures for identifying hereditary colorectal cancer and polyposis syndrome carriers. <i>Gut</i> , 2021 , 70, 2138-2149	19.2	6
214	Genetic architectures of proximal and distal colorectal cancer are partly distinct. <i>Gut</i> , 2021 , 70, 1325-13	34 9.2	7
213	Rare Variants in the DNA Repair Pathway and the Risk of Colorectal Cancer. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2021 , 30, 895-903	4	1
212	Genetic analyses of gynecological disease identify genetic relationships between uterine fibroids and endometrial cancer, and a novel endometrial cancer genetic risk region at the WNT4 1p36.12 locus. <i>Human Genetics</i> , 2021 , 140, 1353-1365	6.3	5
211	Variation in the risk of colorectal cancer in families with Lynch syndrome: a retrospective cohort study. <i>Lancet Oncology, The</i> , 2021 , 22, 1014-1022	21.7	5
210	Informed choice and attitudes regarding a genomic test to predict risk of colorectal cancer in general practice. <i>Patient Education and Counseling</i> , 2021 ,	3.1	1
209	Characteristics of Early-Onset vs Late-Onset Colorectal Cancer: A Review. <i>JAMA Surgery</i> , 2021 , 156, 865	-8.744	15
208	DNA methylation-based signature of CD8+ tumor-infiltrating lymphocytes enables evaluation of immune response and prognosis in colorectal cancer 2021 , 9,		7
207	Functional informed genome-wide interaction analysis of body mass index, diabetes and colorectal cancer risk. <i>Cancer Medicine</i> , 2020 , 9, 3563-3573	4.8	4
206	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
205	The MLH1 polymorphism rs1800734 and risk of endometrial cancer with microsatellite instability. <i>Clinical Epigenetics</i> , 2020 , 12, 102	7.7	2
204	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
203	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

202	Genetic Variants in the Regulatory T cell-Related Pathway and Colorectal Cancer Prognosis. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2020 , 29, 2719-2728	4	
201	Physical activity and risks of breast and colorectal cancer: a Mendelian randomisation analysis. Nature Communications, 2020, 11, 597	17.4	36
200	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
199	Dysfunctional epigenetic aging of the normal colon and colorectal cancer risk. <i>Clinical Epigenetics</i> , 2020 , 12, 5	7.7	27
198	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
197	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
196	Postmenopausal Hormone Therapy and Colorectal Cancer Risk by Molecularly Defined Subtypes and Tumor Location. <i>JNCI Cancer Spectrum</i> , 2020 , 4, pkaa042	4.6	2
195	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
194	Genome-wide Modeling of Polygenic Risk Score in Colorectal Cancer Risk. <i>American Journal of Human Genetics</i> , 2020 , 107, 432-444	11	31
193	Stochastic Epigenetic Mutations Are Associated with Risk of Breast Cancer, Lung Cancer, and Mature B-cell Neoplasms. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2020 , 29, 2026-2037	4	6
192	A Genomic Test for Colorectal Cancer Risk: Is This Acceptable and Feasible in Primary Care?. <i>Public Health Genomics</i> , 2020 , 23, 110-121	1.9	3
191	Monoallelic NTHL1 Loss-of-Function Variants and Risk of Polyposis and Colorectal Cancer. <i>Gastroenterology</i> , 2020 , 159, 2241-2243.e6	13.3	10
190	Intake of Dietary Fruit, Vegetables, and Fiber and Risk of Colorectal Cancer According to Molecular Subtypes: A Pooled Analysis of 9 Studies. <i>Cancer Research</i> , 2020 , 80, 4578-4590	10.1	8
189	Adiposity, metabolites, and colorectal cancer risk: Mendelian randomization study. <i>BMC Medicine</i> , 2020 , 18, 396	11.4	17
188	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
187	Hi-Plex2: a simple and robust approach to targeted sequencing-based genetic screening. <i>BioTechniques</i> , 2019 , 67, 118-122	2.5	6
186	Type 2 diabetes mellitus, blood cholesterol, triglyceride and colorectal cancer risk in Lynch syndrome. <i>British Journal of Cancer</i> , 2019 , 121, 869-876	8.7	4
185	Shared heritability and functional enrichment across six solid cancers. <i>Nature Communications</i> , 2019 , 10, 431	17.4	45

(2018-2019)

184	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
183	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
182	Tumor mutational signatures in sebaceous skin lesions from individuals with Lynch syndrome. <i>Molecular Genetics & Cenomic Medicine</i> , 2019 , 7, e00781	2.3	6
181	Association analyses identify 31 new risk loci for colorectal cancer susceptibility. <i>Nature Communications</i> , 2019 , 10, 2154	17.4	81
180	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
179	Mutational Signature Analysis Reveals NTHL1 Deficiency to Cause a Multi-tumor Phenotype. <i>Cancer Cell</i> , 2019 , 35, 256-266.e5	24.3	72
178	Novel Common Genetic Susceptibility Loci for Colorectal Cancer. <i>Journal of the National Cancer Institute</i> , 2019 , 111, 146-157	9.7	67
177	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
176	Mendelian randomization analysis of C-reactive protein on colorectal cancer risk. <i>International Journal of Epidemiology</i> , 2019 , 48, 767-780	7.8	18
175	Risks of Colorectal Cancer and Cancer-Related Mortality in Familial Colorectal Cancer Type X and Lynch Syndrome Families. <i>Journal of the National Cancer Institute</i> , 2019 , 111, 675-683	9.7	5
174	Clinico-pathological predictors of mismatch repair deficiency in sebaceous neoplasia: A large case series from a single Australian private pathology service. <i>Australasian Journal of Dermatology</i> , 2019 , 60, 126-133	1.3	6
173	Discovery of common and rare genetic risk variants for colorectal cancer. <i>Nature Genetics</i> , 2019 , 51, 76-	-83 6.3	177
172	Assessing the ProMCol classifier as a prognostic marker for non-metastatic colorectal cancer within the Melbourne Collaborative Cohort Study. <i>Gut</i> , 2019 , 68, 761-762	19.2	
171	Cohort Profile: The Colon Cancer Family Registry Cohort (CCFRC). <i>International Journal of Epidemiology</i> , 2018 , 47, 387-388i	7.8	23
170	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
169	Costs and outcomes of Lynch syndrome screening in the Australian colorectal cancer population. Journal of Gastroenterology and Hepatology (Australia), 2018, 33, 1737-1744	4	8
168	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
167	DUSP5 is methylated in CIMP-high colorectal cancer but is not a major regulator of intestinal cell proliferation and tumorigenesis. <i>Scientific Reports</i> , 2018 , 8, 1767	4.9	7

166	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
165	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
164	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
163	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
162	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
161	Endometrial cancer risk and survival by tumor MMR status. <i>Journal of Gynecologic Oncology</i> , 2018 , 29, e39	4	19
160	Utility of immunohistochemistry for mismatch repair proteins on colorectal polyps in the familial cancer clinic. <i>Internal Medicine Journal</i> , 2018 , 48, 1325-1330	1.6	2
159	Family history-based colorectal cancer screening in Australia: A modelling study of the costs, benefits, and harms of different participation scenarios. <i>PLoS Medicine</i> , 2018 , 15, e1002630	11.6	2
158	Identification of nine new susceptibility loci for endometrial cancer. <i>Nature Communications</i> , 2018 , 9, 3166	17.4	70
157	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
156	Physical activity and the risk of colorectal cancer in Lynch syndrome. <i>International Journal of Cancer</i> , 2018 , 143, 2250-2260	7.5	9
155	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
154	RNF43 is mutated less frequently in Lynch Syndrome compared with sporadic microsatellite unstable colorectal cancers. <i>Familial Cancer</i> , 2018 , 17, 63-69	3	9
153	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
152	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
151	Cancer Risks for PMS2-Associated Lynch Syndrome. <i>Journal of Clinical Oncology</i> , 2018 , 36, 2961-2968	2.2	102
150	Risk and prognostic factors for endometrial carcinoma after diagnosis of breast or Lynch-associated cancers-A population-based analysis. <i>Cancer Medicine</i> , 2018 , 7, 6411-6422	4.8	8
149	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

(2016-2018)

148	Current mismatch repair deficiency tumor testing practices and capabilities: A survey of Australian pathology providers. <i>Asia-Pacific Journal of Clinical Oncology</i> , 2018 , 14, 417-425	1.9	6
147	Mendelian randomisation study of age at menarche and age at menopause and the risk of colorectal cancer. <i>British Journal of Cancer</i> , 2018 , 118, 1639-1647	8.7	7
146	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
145	Modeling of successive cancer risks in Lynch syndrome families in the presence of competing risks using copulas. <i>Biometrics</i> , 2017 , 73, 271-282	1.8	4
144	Mendelian randomisation implicates hyperlipidaemia as a risk factor for colorectal cancer. <i>International Journal of Cancer</i> , 2017 , 140, 2701-2708	7.5	50
143	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
142	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
141	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
140	Risk factors for metachronous colorectal cancer or polyp: A systematic review and meta-analysis. Journal of Gastroenterology and Hepatology (Australia), 2017, 32, 301-326	4	8
139	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
138	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
137	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
136	Targeted sequencing of 36 known or putative colorectal cancer susceptibility genes. <i>Molecular Genetics & Molecular Genetics & Molecula</i>	2.3	20
135	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
134	Prevalence and Penetrance of Major Genes and Polygenes for Colorectal Cancer. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2017 , 26, 404-412	4	185
133	Germline miRNA DNA variants and the risk of colorectal cancer by subtype. <i>Genes Chromosomes and Cancer</i> , 2017 , 56, 177-184	5	6
132	Reducing the polyp burden in serrated polyposis by serial colonoscopy: the impact of nationally coordinated community surveillance. <i>New Zealand Medical Journal</i> , 2017 , 130, 57-67	0.8	4
131	Promoter methylation of ITF2, but not APC, is associated with microsatellite instability in two populations of colorectal cancer patients. <i>BMC Cancer</i> , 2016 , 16, 113	4.8	6

130	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
129	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
128	Determining the familial risk distribution of colorectal cancer: a data mining approach. <i>Familial Cancer</i> , 2016 , 15, 241-51	3	5
127	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
126	Quantifying the utility of single nucleotide polymorphisms to guide colorectal cancer screening. <i>Future Oncology</i> , 2016 , 12, 503-13	3.6	30
125	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
124	GWASeq: targeted re-sequencing follow up to GWAS. BMC Genomics, 2016, 17, 176	4.5	7
123	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
122	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
121	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
120	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
119	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
118	Cholecystectomy and the risk of colorectal cancer by tumor mismatch repair deficiency status. <i>International Journal of Colorectal Disease</i> , 2016 , 31, 1451-7	3	5
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	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
115	Risk of colorectal cancer for people with a mutation in both a MUTYH and a DNA mismatch repair gene. <i>Familial Cancer</i> , 2015 , 14, 575-83	3	8
114	Aspirin, Ibuprofen, and the Risk of Colorectal Cancer in Lynch Syndrome. <i>Journal of the National Cancer Institute</i> , 2015 , 107,	9.7	66
113	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

(2014-2015)

112	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
111	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
110	Childhood cancers in families with and without Lynch syndrome. Familial Cancer, 2015, 14, 545-51	3	4
109	Association between molecular subtypes of colorectal cancer and patient survival. <i>Gastroenterology</i> , 2015 , 148, 77-87.e2	13.3	273
108	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
107	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
106	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
105	Lynch syndrome and cervical cancer. <i>International Journal of Cancer</i> , 2015 , 137, 2757-61	7.5	11
104	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
103	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
102	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
101	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
100	Genetic variants within the hTERT gene and the risk of colorectal cancer in Lynch syndrome. <i>Genes and Cancer</i> , 2015 , 6, 445-51	2.9	5
99	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
98	Pooled analysis of iron-related genes in Parkinson® disease: association with transferrin. <i>Neurobiology of Disease</i> , 2014 , 62, 172-8	7.5	57
97	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
96	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
95	Should the grading of colorectal adenocarcinoma include microsatellite instability status?. <i>Human Pathology</i> , 2014 , 45, 2077-84	3.7	39

94	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
93	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
92	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
91	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
90	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
89	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
88	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
87	Reply to J. Moline et al. <i>Journal of Clinical Oncology</i> , 2014 , 32, 2278-9	2.2	3
86	Re: Microsatellite instability and BRAF mutation testing in colorectal cancer prognostication. <i>Journal of the National Cancer Institute</i> , 2014 , 106,	9.7	6
85	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
84	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
83	Molecular subtypes of colorectal cancer in relation to disease survival <i>Journal of Clinical Oncology</i> , 2014 , 32, 451-451	2.2	
82	Lynch syndrome-associated breast cancers do not overexpress chromosome 11-encoded mucins. <i>Modern Pathology</i> , 2013 , 26, 944-54	9.8	1
81	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
80	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
79	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
78	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
77	Risks of colorectal and other cancers after endometrial cancer for women with Lynch syndrome. Journal of the National Cancer Institute, 2013, 105, 274-9	9.7	78

(2012-2013)

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
Cancer risks for MLH1 and MSH2 mutation carriers. <i>Human Mutation</i> , 2013 , 34, 490-7	4.7	171
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
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
Colorectal carcinomas with KRAS mutation are associated with distinctive morphological and molecular features. <i>Modern Pathology</i> , 2013 , 26, 825-34	9.8	106
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
Germline HOXB13 p.Gly84Glu mutation and risk of colorectal cancer. Cancer Epidemiology, 2013 , 37, 42	?.4 ₂7 .8	21
Telomere length varies by DNA extraction method: implications for epidemiologic research. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2013 , 22, 2047-54	4	81
Detection of large scale 3Odeletions in the PMS2 gene amongst Colon-CFR participants: have we been missing anything?. <i>Familial Cancer</i> , 2013 , 12, 563-6	3	13
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
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
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
Absence of PMS2 mutations in colon-CFR participants whose colorectal cancers demonstrate unexplained loss of MLH1 expression. <i>Clinical Genetics</i> , 2013 , 83, 591-3	4	8
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
PIK3CA activating mutation in colorectal carcinoma: associations with molecular features and survival. <i>PLoS ONE</i> , 2013 , 8, e65479	3.7	102
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
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
KRAS mutations in ovarian low-grade endometrioid adenocarcinoma: association with concurrent endometriosis. <i>Human Pathology</i> , 2012 , 43, 1177-83	3.7	43
	based on sequence bioinformatics and tumor characteristics: a report from the Colon Cancer Family Registry. Human Mutation, 2013, 34, 200-9 Cancer risks for MLH1 and MSH2 mutation carriers. Human Mutation, 2013, 34, 490-7 KRAS-mutation status in relation to colorectal cancer survival: the joint impact of correlated tumour markers. British Journal of Cancer, 2013, 108, 1757-64 Risk of metachronous colon cancer following surgery for rectal cancer in mismatch repair gene mutation carriers. Annals of Surgical Oncology, 2013, 20, 1829-36 Colorectal carcinomas with KRAS mutation are associated with distinctive morphological and molecular features. Modern Pathology, 2013, 26, 825-34 Association between hypermethylation of DNA repetitive elements in white blood cell DNA and early-onset colorectal cancer. Epigenetics, 2013, 8, 748-55 Germline HOXB13 p.Gly84Glu mutation and risk of colorectal cancer. Cancer Epidemiology, 2013, 37, 42 Telomere length varies by DNA extraction method: implications for epidemiologic research. Cancer Epidemiology Biomarkers and Prevention, 2013, 22, 2047-54 Detection of large scale 3Gleletions in the PMS2 gene amongst Colon-CFR participants: have we been missing anything?. Familial Cancer, 2013, 12, 563-6 Family history of colorectal cancer in BRAF p.V600E-mutated colorectal cancer cases. Cancer Epidemiology Biomarkers and Prevention, 2013, 22, 917-26 Immunophenotypic analysis of ovarian endometrioid adenocarcinoma: correlation with KRAS mutation and the presence of endometriosis. Pathology, 2013, 37, 1592-602 Absence of PMS2 mutations in colon-CFR participants whose colorectal cancers demonstrate unexplained loss of MLH1 expression. Clinical Genetics, 2013, 83, 591-3 Multiplicity and molecular heterogeneity of colorectal carcinomas in individuals with serrated polyposis. American Journal of Surgical Pathology, 2013, 37, 434-42 PIK3CA activating mutations in the Polyposis-Associated Cenes BMPR1A, SMADA, PTEN, MUTYH and GREM1 Are Not Common in Individuals with Serrated Polyposis S	Registry. Human Mutation, 2013, 34, 200-9 Cancer risks for MLH1 and MSH2 mutation carriers. Human Mutation, 2013, 34, 490-7 KRAS-mutation status in relation to colorectal cancer survival: the joint impact of correlated tumour markers. British Journal of Cancer, 2013, 108, 1757-64 Risk of metachronous colon cancer following surgery for rectal cancer in mismatch repair gene mutation carriers. Annals of Surgical Oncology, 2013, 20, 1829-36 Colorectal carcinomas with KRAS mutation are associated with distinctive morphological and molecular features. Modern Pathology, 2013, 26, 825-34 Association between hypermethylation of DNA repetitive elements in white blood cell DNA and early-onset colorectal cancer. Epigenetics, 2013, 8, 748-55 Germline HOXB13 p. Gly84Glu mutation and risk of colorectal cancer. Cancer Epidemiology, 2013, 37, 424-278 Telomere length varies by DNA extraction method: implications for epidemiologic research. Cancer Epidemiology Biomarkers and Prevention, 2013, 22, 2047-54 Detection of large scale 3Caleletions in the PMS2 gene amongst Colon-CFR participants: have we been missing anything. Familial Cancer, 2013, 12, 563-6 Family history of colorectal cancer in BRAF p./600E-mutated colorectal cancer cases. Cancer Epidemiology Biomarkers and Prevention, 2013, 22, 917-26 Immunophenotypic analysis of ovarian endometrioid adenocarcinoma: correlation with KRAS mutation and the presence of endometrioiss. Pathology, 2013, 45, 559-66 BRAFV600E immunohistochemistry facilitates universal screening of colorectal cancers for Lynch syndrome. American Journal of Surgical Pathology, 2013, 37, 1392-602 Absence of PMS2 mutations in colon-CFR participants whose colorectal cancers demonstrate unexplained loss of MLH1 expression. Clinical Genetics, 2013, 83, 591-3 Multiplicity and molecular heterogeneity of colorectal carcinomas in individuals with serrated polyposis. American Journal of Surgical Pathology, 2013, 37, 434-412 PIKS3CA activating mutation in colorectal carcinoma: associations with molecular f

58	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
57	Cancer risks for relatives of patients with serrated polyposis. <i>American Journal of Gastroenterology</i> , 2012 , 107, 770-8	0.7	69
56	Colorectal cancer linkage on chromosomes 4q21, 8q13, 12q24, and 15q22. <i>PLoS ONE</i> , 2012 , 7, e38175	3.7	23
55	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
54	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
53	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
52	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
51	Cancer risks for the relatives of colorectal cancer cases with a methylated MLH1 promoter region: data from the Colorectal Cancer Family Registry. <i>Cancer Prevention Research</i> , 2012 , 5, 328-35	3.2	9
50	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
49	Using tumour pathology to identify people at high genetic risk of breast and colorectal cancers. <i>Pathology</i> , 2012 , 44, 89-98	1.6	6
48	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
47	Genome-wide search for gene-gene interactions in colorectal cancer. <i>PLoS ONE</i> , 2012 , 7, e52535	3.7	29
46	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
45	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
44	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
43	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
42	Linkage to chromosome 2q32.2-q33.3 in familial serrated neoplasia (Jass syndrome). <i>Familial Cancer</i> , 2011 , 10, 245-54	3	17
41	Mutation deep within an intron of MSH2 causes Lynch syndrome. Familial Cancer, 2011 , 10, 297-301	3	34

(2007-2011)

40	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
39	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
38	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
37	Cigarette pack labelling in 12 countries at different levels of economic development. <i>Journal of Public Health Policy</i> , 2011 , 32, 146-64	2.9	6
36	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
35	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
34	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
33	Risks of Lynch syndrome cancers for MSH6 mutation carriers. <i>Journal of the National Cancer Institute</i> , 2010 , 102, 193-201	9.7	279
32	Lessons from Lynch syndrome: a tumor biology-based approach to familial colorectal cancer. <i>Future Oncology</i> , 2010 , 6, 539-49	3.6	5
31	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
30	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
29	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
28	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
27	A perspective on bi-allelic MUTYH mutations in patients with hyperplastic polyposis syndrome. <i>Gastroenterology</i> , 2009 , 136, 2407-8	13.3	14
26	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
25	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
24	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
23	Stability of BAT26 in Lynch syndrome colorectal tumours. <i>European Journal of Human Genetics</i> , 2007 , 15, 139-41; author reply 141-2	5.3	5

22	Association of APOE with Parkinson disease age-at-onset in women. <i>Neuroscience Letters</i> , 2007 , 411, 185-8	3.3	23
21	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
20	Extensive DNA methylation in normal colorectal mucosa in hyperplastic polyposis. <i>Gut</i> , 2006 , 55, 1467-	74 9.2	120
19	Hyperplastic polyposis syndrome: phenotypic presentations and the role of MBD4 and MYH. <i>Gastroenterology</i> , 2006 , 131, 30-9	13.3	166
18	Advanced colorectal polyps with the molecular and morphological features of serrated polyps and adenomas: concept of a © usionOpathway to colorectal cancer. <i>Histopathology</i> , 2006 , 49, 121-31	7.3	214
17	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
16	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
15	GSK3B polymorphisms alter transcription and splicing in Parkinson@ disease. <i>Annals of Neurology</i> , 2005 , 58, 829-39	9.4	172
14	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
13	Tau haplotypes regulate transcription and are associated with Parkinson@ disease. <i>Annals of Neurology</i> , 2004 , 55, 329-34	9.4	147
12	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
11	The Cys282Tyr polymorphism in the HFE gene in Australian Parkinson@ disease patients. <i>Neuroscience Letters</i> , 2002 , 327, 91-4	3.3	48
10	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
9	The ACE deletion polymorphism is not associated with Parkinson@ disease. <i>European Neurology</i> , 1999 , 41, 103-6	2.1	19
8	Variations in the monoamine oxidase B (MAOB) gene are associated with Parkinson@ disease. <i>Movement Disorders</i> , 1999 , 14, 219-24	7	73
7	Variations in the monoamine oxidase B lpar;MAOB) gene are associated with Parkinson@ disease 1999 , 14, 219		3
6	Mutational Signature Analysis Reveals NTHL1 Deficiency to Cause a Multi-Tumor Phenotype Including a Predisposition to Colon and Breast Cancer. <i>SSRN Electronic Journal</i> ,	1	1
5	Evaluating the utility of tumour mutational signatures for identifying hereditary colorectal cancer and polyposis syndrome carriers		1

LIST OF PUBLICATIONS

4	Adiposity, metabolites, and colorectal cancer risk: Mendelian randomization study	2
3	Biological aging measures based on blood DNA methylation and risk of cancer: a prospective study	1
2	A combined proteomics and Mendelian randomization approach to investigate the effects of aspirin-targeted proteins on colorectal cancer	1
1	Germline loss-of-function variants in the base-excision repair gene MBD4 cause a Mendelian recessive syndrome of adenomatous colorectal polyposis and acute myeloid leukaemia	3