Aung K Win

List of Publications by Year
in descending order

186	9,176	35864	50	50037	85
papers	citations		h-index		g-index

312	312	312	12753
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Discovery of common and rare genetic risk variants for colorectal cancer. Nature Genetics, 2019, 51, 76-87.	20.2	409
2	Cancer risks by gene, age, and gender in 6350 carriers of pathogenic mismatch repair variants: findings from the Prospective Lynch Syndrome Database. Genetics in Medicine, 2020, 22, 15-25.	2.4	400
3	Prevalence and Penetrance of Major Genes and Polygenes for Colorectal Cancer. Cancer Epidemiology Biomarkers and Prevention, 2017, 26, 404-412.	1.9	368
4	Colorectal and Other Cancer Risks for Carriers and Noncarriers From Families With a DNA Mismatch Repair Gene Mutation: A Prospective Cohort Study. Journal of Clinical Oncology, 2012, 30, 958-964.	5.4	292
5	Metachronous colorectal cancer risk for mismatch repair gene mutation carriers: the advantage of more extensive colon surgery. Gut, 2011, 60, 950-957.	13.5	236
6	Cancer Risks forMLH1andMSH2Mutation Carriers. Human Mutation, 2013, 34, 490-497.	2.7	206
7	Risks of Primary Extracolonic Cancers Following Colorectal Cancer in Lynch Syndrome. Journal of the National Cancer Institute, 2012, 104, 1363-1372.	6.3	198
8	Risk of Colorectal Cancer for Carriers of Mutations in MUTYH, WithÂand Without a Family History of Cancer. Gastroenterology, 2014, 146, 1208-1211.e5.	1.4	184
9	Association analyses identify 31 new risk loci for colorectal cancer susceptibility. Nature Communications, 2019, 10, 2154.	13.0	183
10	Incidence and prevalence of nonâ€melanoma skin cancer in <scp>A</scp>ustralia: A systematic review. Australasian Journal of Dermatology, 2015, 56, 258-267.	0.8	171
11	Cancer Risks for PMS2-Associated Lynch Syndrome. Journal of Clinical Oncology, 2018, 36, 2961-2968.	5.4	156
12	Risk Prediction Models for Colorectal Cancer: A Systematic Review. Cancer Prevention Research, 2016, 9, 13-26.	1.6	150
13	Novel Common Genetic Susceptibility Loci for Colorectal Cancer. Journal of the National Cancer Institute, 2019, 111, 146-157.	6.3	136
14	Genome-wide Modeling of Polygenic Risk Score in Colorectal Cancer Risk. American Journal of Human Genetics, 2020, 107, 432-444.	6.1	135
15	Expression of MUC2, MUC5AC, MUC5B, and MUC6 mucins in colorectal cancers and their association with the CpG island methylator phenotype. Modern Pathology, 2013, 26, 1642-1656.	5.6	131
16	Colorectal carcinomas with KRAS mutation are associated with distinctive morphological and molecular features. Modern Pathology, 2013, 26, 825-834.	5.6	130
17	Identification of susceptibility loci for colorectal cancer in a genome-wide meta-analysis. Human Molecular Genetics, 2014, 23, 4729-4737.	3.0	130
18	BRAFV600E Immunohistochemistry Facilitates Universal Screening of Colorectal Cancers for Lynch Syndrome. American Journal of Surgical Pathology, 2013, 37, 1592-1602.	3.9	127

#	Article	IF	Citations
19	Rising incidence of earlyâ€onset colorectal cancer in <scp>A</scp>ustralia over two decades: Report and review. Journal of Gastroenterology and Hepatology (Australia), 2015, 30, 6-13.	2.8	126
20	PIK3CA Activating Mutation in Colorectal Carcinoma: Associations with Molecular Features and Survival. PLoS ONE, 2013, 8, e65479.	2.5	120
21	Cumulative Burden of Colorectal Cancerâ€"Associated Genetic Variants Is More Strongly Associated With Early-Onset vs Late-Onset Cancer. Gastroenterology, 2020, 158, 1274-1286.e12.	1.4	119
22	BRAF Mutation Status and Survival after Colorectal Cancer Diagnosis According to Patient and Tumor Characteristics. Cancer Epidemiology Biomarkers and Prevention, 2012, 21, 1792-1798.	1.9	117
23	Risk of extracolonic cancers for people with biallelic and monoallelic mutations in MUTYH. International Journal of Cancer, 2016, 139, 1557-1563.	5.4	114
24	Risk of breast cancer in Lynch syndrome: a systematic review. Breast Cancer Research, 2013, 15, R27.	5.1	113
25	A new GWAS and meta-analysis with 1000Genomes imputation identifies novel risk variants for colorectal cancer. Scientific Reports, 2015, 5, 10442.	3.4	112
26	Risk of Prostate Cancer in Lynch Syndrome: A Systematic Review and Meta-analysis. Cancer Epidemiology Biomarkers and Prevention, 2014, 23, 437-449.	1.9	109
27	Risk of Metachronous Colon Cancer Following Surgery for Rectal Cancer in Mismatch Repair Gene Mutation Carriers. Annals of Surgical Oncology, 2013, 20, 1829-1836.	1.9	107
28	Cancer risks for monoallelic MUTYH mutation carriers with a family history of colorectal cancer. International Journal of Cancer, 2011, 129, 2256-2262.	5.4	95
29	Risks of Colorectal and Other Cancers After Endometrial Cancer for Women With Lynch Syndrome. Journal of the National Cancer Institute, 2013, 105, 274-279.	6.3	95
30	Risk Prediction Models for Colorectal Cancer: A Review. Cancer Epidemiology Biomarkers and Prevention, 2012, 21, 398-410.	1.9	87
31	Pro-inflammatory fatty acid profile and colorectal cancer risk: A Mendelian randomisation analysis. European Journal of Cancer, 2017, 84, 228-238.	2.9	86
32	Phenotype and Polyp Landscape in Serrated Polyposis Syndrome. American Journal of Surgical Pathology, 2012, 36, 876-882.	3.9	85
33	Colon and Rectal Cancer Survival by Tumor Location and Microsatellite Instability. Diseases of the Colon and Rectum, 2013, 56, 937-944.	1.5	85
34	Increased Cancer Risks in Myotonic Dystrophy. Mayo Clinic Proceedings, 2012, 87, 130-135.	2.8	84
35	Immunohistochemical pitfalls and the importance of glypican 3 and arginase in the diagnosis of scirrhous hepatocellular carcinoma. Modern Pathology, 2013, 26, 782-791.	5.6	83
36	Cancer Risks for Relatives of Patients With Serrated Polyposis. American Journal of Gastroenterology, 2012, 107, 770-778.	0.4	81

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37	Aspirin, Ibuprofen, and the Risk of Colorectal Cancer in Lynch Syndrome. Journal of the National Cancer Institute, 2015, 107, djv170.	6.3	80
38	Mendelian randomisation implicates hyperlipidaemia as a risk factor for colorectal cancer. International Journal of Cancer, 2017, 140, 2701-2708.	5.4	80
39	Family History and Risk of Endometrial Cancer. Obstetrics and Gynecology, 2015, 125, 89-98.	2.3	75
40	Female Hormonal Factors and the Risk of Endometrial Cancer in Lynch Syndrome. JAMA - Journal of the American Medical Association, 2015, 314, 61.	7.0	75
41	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. Modern Pathology, 2012, 25, 722-730.	5.6	73
42	Mendelian Randomization Study of Body Mass Index and Colorectal Cancer Risk. Cancer Epidemiology Biomarkers and Prevention, 2015, 24, 1024-1031.	1.9	71
43	Clinical problems of colorectal cancer and endometrial cancer cases with unknown cause of tumor mismatch repair deficiency (suspected Lynch syndrome). The Application of Clinical Genetics, 2014, 7, 183.	2.8	70
44	Deciphering colorectal cancer genetics through multi-omic analysis of 100,204 cases and 154,587 controls of European and east Asian ancestries. Nature Genetics, 2023, 55, 89-99.	20.2	70
45	Risk Factors for Colorectal Cancer in Patients with Multiple Serrated Polyps: A Cross-Sectional Case Series from Genetics Clinics. PLoS ONE, 2010, 5, e11636.	2.5	69
46	Variation in the risk of colorectal cancer in families with Lynch syndrome: a retrospective cohort study. Lancet Oncology, The, 2021, 22, 1014-1022.	10.7	67
47	The Exponentiated Generalized Class of Distributions. Journal of Data Science, 2013, 11, 1-27.	0.9	64
48	Enhancing antibody-dependent cell-mediated cytotoxicity: a strategy for improving antibody-based immunotherapy. Antibody Therapeutics, 2018, 1, 7-12.	1.9	62
49	Association between monoallelic MUTYH mutation and colorectal cancer risk: a meta-regression analysis. Familial Cancer, 2011, 10, 1-9.	2.0	61
50	Mendelian randomisation analysis strongly implicates adiposity with risk of developing colorectal cancer. British Journal of Cancer, 2016, 115, 266-272.	6.5	60
51	Mendelian randomization study of height and risk of colorectal cancer. International Journal of Epidemiology, 2015, 44, 662-672.	2.0	56
52	Targeting ACSS2 with a Transition-State Mimetic Inhibits Triple-Negative Breast Cancer Growth. Cancer Research, 2021, 81, 1252-1264.	0.9	55
53	Potential impact of family historyâ€“based screening guidelines on the detection of earlyâ€onset colorectal cancer. Cancer, 2020, 126, 3013-3020.	4.1	53
54	PMS2 monoallelic mutation carriers: the known unknown. Genetics in Medicine, 2016, 18, 13-19.	2.4	52

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55	Mutation Spectrum and Risk of Colorectal Cancer in African American Families with Lynch Syndrome. Gastroenterology, 2015, 149, 1446-1453.	1.4	50
56	Genetics and genetic testing for age-related macular degeneration. Eye, 2018, 32, 849-857.	2.3	50
57	Risk of colorectal cancer for carriers of a germ-line mutation in POLE or POLD1. Genetics in Medicine, 2018, 20, 890-895.	2.4	50
58	Phenotypic diversity in patients with multiple serrated polyps: a genetics clinic study. International Journal of Colorectal Disease, 2010, 25, 703-712.	2.3	48
59	Tumor testing to identify lynch syndrome in two Australian colorectal cancer cohorts. Journal of Gastroenterology and Hepatology (Australia), 2017, 32, 427-438.	2.8	48
60	Association between Body Mass Index and Mortality for Colorectal Cancer Survivors: Overall and by Tumor Molecular Phenotype. Cancer Epidemiology Biomarkers and Prevention, 2015, 24, 1229-1238.	1.9	47
61	High prevalence of mismatch repair deficiency in prostate cancers diagnosed in mismatch repair gene mutation carriers from the colon cancer family registry. Familial Cancer, 2014, 13, 573-582.	2.0	46
62	Subtraction of Epsteinâ€“Barr virus and microsatellite instability genotypes from the Lauren histotypes: Combined molecular and histologic subtyping with clinicopathological and prognostic significance validated in a cohort of 1,248 cases. International Journal of Cancer, 2019, 145, 3218-3230.	5.4	45
63	Should the grading of colorectal adenocarcinoma include microsatellite instability status?. Human Pathology, 2014, 45, 2077-2084.	2.3	44
64	DNA methylation-based signature of CD8+ tumor-infiltrating lymphocytes enables evaluation of immune response and prognosis in colorectal cancer. , 2021, 9, e002671.		44
65	Nongenetic Determinants of Risk forÂEarly-Onset Colorectal Cancer. JNCI Cancer Spectrum, 2021, 5, pkab029.	2.8	43
66	Association between hypermethylation of DNA repetitive elements in white blood cell DNA and early-onset colorectal cancer. Epigenetics, 2013, 8, 748-755.	2.9	42
67	Quantifying the Utility of Single Nucleotide Polymorphisms to Guide Colorectal Cancer Screening. Future Oncology, 2016, 12, 503-513.	2.4	42
68	Cohort Profile: The Colon Cancer Family Registry Cohort (CCFRC). International Journal of Epidemiology, 2018, 47, 387-388i.	2.0	42
69	Determining the frequency of de novo germline mutations in DNA mismatch repair genes. Journal of Medical Genetics, 2011, 48, 530-534.	3.6	41
70	Variation at 2q35 (PNKDandTMBIM1) influences colorectal cancer risk and identifies a pleiotropic effect with inflammatory bowel disease. Human Molecular Genetics, 2016, 25, 2349-2359.	3.0	41
71	Multiplicity and Molecular Heterogeneity of Colorectal Carcinomas in Individuals With Serrated Polyposis. American Journal of Surgical Pathology, 2013, 37, 434-442.	3.9	40
72	Role of tumour molecular and pathology features to estimate colorectal cancer risk for first-degree relatives. Gut, 2015, 64, 101-110.	13.5	40

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73	Associations of 5HTTLPR polymorphism with major depressive disorder and alcohol dependence: A systematic review and meta-analysis. Australian and New Zealand Journal of Psychiatry, 2016, 50, 842-857.	2.7	38
74	Alcohol Consumption and the Risk of Colorectal Cancer for Mismatch Repair Gene Mutation Carriers. Cancer Epidemiology Biomarkers and Prevention, 2017, 26, 366-375.	1.9	38
75	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.	3.4	37
76	Short-Term Risk of Colorectal Cancer in Individuals With Lynch Syndrome: A Meta-Analysis. Journal of Clinical Oncology, 2015, 33, 326-331.	5.4	37
77	Risk factors for metachronous colorectal cancer following a primary colorectal cancer: A prospective cohort study. International Journal of Cancer, 2016, 139, 1081-1090.	5.4	34
78	Germline mutations inPMS2andMLH1in individuals with solitary loss of PMS2 expression in colorectal carcinomas from the Colon Cancer Family Registry Cohort. BMJ Open, 2016, 6, e010293.	2.1	34
79	Targeted sequencing of 36 known or putative colorectal cancer susceptibility genes. Molecular Genetics & Genomic Medicine, 2017, 5, 553-569.	1.3	33
80	Quantitation of Human Metallothionein Isoforms: A Family of Small, Highly Conserved, Cysteine-rich Proteins. Molecular and Cellular Proteomics, 2014, 13, 1020-1033.	3.9	32
81	Prediagnostic Physical Activity and Colorectal Cancer Survival: Overall and Stratified by Tumor Characteristics. Cancer Epidemiology Biomarkers and Prevention, 2015, 24, 1130-1137.	1.9	32
82	Are the common genetic variants associated with colorectal cancer risk for DNA mismatch repair gene mutation carriers?. European Journal of Cancer, 2013, 49, 1578-1587.	2.9	31
83	Comparison of Prediction Models for Lynch Syndrome Among Individuals With Colorectal Cancer. Journal of the National Cancer Institute, 2016, 108, .	6.3	30
84	The interaction between farming/rural environment and TLR2, TLR4, TLR6 and CD14 genetic polymorphisms in relation to early- and late-onset asthma. Scientific Reports, 2017, 7, 43681.	3.4	29
85	Risk-reducing hysterectomy and bilateral salpingo-oophorectomy in female heterozygotes of pathogenic mismatch repair variants: a Prospective Lynch Syndrome Database report. Genetics in Medicine, 2021, 23, 705-712.	2.4	29
86	Evaluating the utility of tumour mutational signatures for identifying hereditary colorectal cancer and polyposis syndrome carriers. Gut, 2021, 70, 2138-2149.	13.5	29
87	Criteria and prediction models for mismatch repair gene mutations: a review. Journal of Medical Genetics, 2013, 50, 785-793.	3.6	28
88	Multivitamin, calcium and folic acid supplements and the risk of colorectal cancer in Lynch syndrome. International Journal of Epidemiology, 2016, 45, 940-953.	2.0	28
89	Germline Mutations in the Polyposis-Associated Genes BMPR1A, SMAD4, PTEN, MUTYH and GREM1 Are Not Common in Individuals with Serrated Polyposis Syndrome. PLoS ONE, 2013, 8, e66705.	2.5	27
90	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.	5.4	27

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91	Physical activity and the risk of colorectal cancer in Lynch syndrome. International Journal of Cancer, 2018, 143, 2250-2260.	5.4	27
92	Mendelian Randomization of Circulating Polyunsaturated Fatty Acids and Colorectal Cancer Risk. Cancer Epidemiology Biomarkers and Prevention, 2020, 29, 860-870.	1.9	27
93	Genetic variation in UGT genes modify the associations of NSAIDs with risk of colorectal cancer: Colon cancer family registry. Genes Chromosomes and Cancer, 2014, 53, 568-578.	3.2	25
94	A New Comprehensive Colorectal Cancer Risk Prediction Model Incorporating Family History, Personal Characteristics, and Environmental Factors. Cancer Epidemiology Biomarkers and Prevention, 2020, 29, 549-557.	1.9	25
95	Mapping the Pains of Neo-Colonialism: A Critical Elaboration of Southern Criminology. British Journal of Criminology, 2021, 61, 1612-1629.	2.3	25
96	Germline HOXB13 p.Gly84Glu mutation and risk of colorectal cancer. Cancer Epidemiology, 2013, 37, 424-427.	2.1	24
97	Family History of Colorectal Cancer in BRAF p.V600E-Mutated Colorectal Cancer Cases. Cancer Epidemiology Biomarkers and Prevention, 2013, 22, 917-926.	1.9	24
98	Low-grade neuroendocrine tumors arising in intestinal adenomas: evidence for alterations in the adenomatous polyposis coli/Î²-catenin pathway. Human Pathology, 2014, 45, 2051-2058.	2.3	24
99	Body Mass Index in Early Adulthood and Endometrial Cancer Risk for Mismatch Repair Gene Mutation Carriers. Obstetrics and Gynecology, 2011, 117, 899-905.	2.3	23
100	Longâ€term weight loss after colorectal cancer diagnosis is associated with lower survival: The Colon Cancer Family Registry. Cancer, 2017, 123, 4701-4708.	4.1	23
101	Reproductive factors as risk modifiers of breast cancer in BRCA mutation carriers and high-risk non-carriers. Oncotarget, 2017, 8, 102110-102118.	1.9	23
102	Ability of known susceptibility SNPs to predict colorectal cancer risk for persons with and without a family history. Familial Cancer, 2019, 18, 389-397.	2.0	23
103	The Feasibility of Monoculture and Polyculture of Striped Catfish and Nile Tilapia in Different Proportions and Their Effects on Growth Performance, Productivity, and Financial Revenue. Journal of Marine Science and Engineering, 2021, 9, 586.	2.7	23
104	SNP rs16906252C>T Is an Expression and Methylation Quantitative Trait Locus Associated with an Increased Risk of Developing MGMT-Methylated Colorectal Cancer. Clinical Cancer Research, 2016, 22, 6266-6277.	7.2	22
105	Prediagnostic alcohol consumption and colorectal cancer survival: The Colon Cancer Family Registry. Cancer, 2017, 123, 1035-1043.	4.1	22
106	Does risk of endometrial cancer for women without a germline mutation in a DNA mismatch repair gene depend on family history of endometrial cancer or colorectal cancer?. Gynecologic Oncology, 2014, 133, 287-292.	1.4	21
107	Somatic mutations of the coding microsatellites within the beta-2-microglobulin gene in mismatch repair-deficient colorectal cancers and adenomas. Familial Cancer, 2018, 17, 91-100.	2.0	21
108	Monoallelic NTHL1 Loss-of-Function Variants and Risk of Polyposis and Colorectal Cancer. Gastroenterology, 2020, 159, 2241-2243.e6.	1.4	21

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109	DNA Methylation Signatures and the Contribution of Age-Associated Methylomic Drift to Carcinogenesis in Early-Onset Colorectal Cancer. Cancers, 2021, 13, 2589.	3.8	21
110	Prevalence and risk factors of hearing loss among infants in Jordan: Initial results from universal neonatal screening. International Journal of Audiology, 2014, 53, 915-920.	2.1	20
111	Exposure to household air pollution over 10â€…years is related to asthma and lung function decline. European Respiratory Journal, 2021, 57, 2000602.	7.4	20
112	Phospho-ERKTHR202/Tyr214 Is Overexpressed in Hairy Cell Leukemia and Is a Useful Diagnostic Marker in Bone Marrow Trephine Sections. American Journal of Surgical Pathology, 2013, 37, 305-308.	3.9	19
113	Risk Stratification for Early-Onset Colorectal Cancer Using a Combination of Genetic and Environmental Risk Scores: An International Multi-Center Study. Journal of the National Cancer Institute, 2022, , .	6.3	19
114	Breast cancer risk for Korean women with germline mutations in BRCA1 and BRCA2. Breast Cancer Research and Treatment, 2015, 152, 659-665.	2.5	18
115	PIK3CA Somatic Mutation Status in Relation to Patient and Tumor Factors in Racial/Ethnic Minorities with Colorectal Cancer. Cancer Epidemiology Biomarkers and Prevention, 2015, 24, 1046-1051.	1.9	17
116	Mendelian randomisation study of age at menarche and age at menopause and the risk of colorectal cancer. British Journal of Cancer, 2018, 118, 1639-1647.	6.5	17
117	Assessment of a Polygenic Risk Score for Colorectal Cancer to Predict Risk of Lynch Syndrome Colorectal Cancer. JNCI Cancer Spectrum, 2021, 5, pkab022.	2.8	17
118	Colorectal cancer and self-reported tooth agenesis. Hereditary Cancer in Clinical Practice, 2014, 12, 7.	1.5	16
119	Findings in young adults at colonoscopy from a hospital service database audit. BMC Gastroenterology, 2017, 17, 56.	1.9	15
120	Risk factors for metachronous colorectal cancer or polyp: A systematic review and metaâ€analysis. Journal of Gastroenterology and Hepatology (Australia), 2017, 32, 301-326.	2.8	14
121	Mortality after breast cancer as a function of time since diagnosis by estrogen receptor status and age at diagnosis. International Journal of Cancer, 2019, 145, 3207-3217.	5.4	14
122	No Difference in Penetrance between Truncating and Missense/Aberrant Splicing Pathogenic Variants in MLH1 and MSH2: A Prospective Lynch Syndrome Database Study. Journal of Clinical Medicine, 2021, 10, 2856.	2.5	14
123	The Australasian Colorectal Cancer Family Registry. Medical Journal of Australia, 2012, 197, 480-481.	1.7	13
124	Lynch syndrome and cervical cancer. International Journal of Cancer, 2015, 137, 2757-2761.	5.4	13
125	Germline and Tumor Sequencing as a Diagnostic Tool To Resolve Suspected Lynch Syndrome. Journal of Molecular Diagnostics, 2021, 23, 358-371.	2.9	13
126	Uptake of hysterectomy and bilateral salpingo-oophorectomy in carriers of pathogenic mismatch repair variants: a Prospective Lynch Syndrome Database report. European Journal of Cancer, 2021, 148, 124-133.	2.9	13

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127	Association of LINC00673 Genetic Variants with Progression of Oral Cancer. Journal of Personalized Medicine, 2021, 11, 468.	2.5	13
128	Cancer Risks for the Relatives of Colorectal Cancer Cases with a Methylated MLH1 Promoter Region: Data from the Colorectal Cancer Family Registry. Cancer Prevention Research, 2012, 5, 328-335.	1.6	12
129	Calcium Antagonists Use and Its Association with Lower Urinary Tract Symptoms: A Cross-Sectional Study. PLoS ONE, 2013, 8, e66708.	2.5	12
130	Common variants in the obesity-associated genes FTO and MC4R are not associated with risk of colorectal cancer. Cancer Epidemiology, 2016, 44, 1-4.	2.1	12
131	Type 2 diabetes mellitus, blood cholesterol, triglyceride and colorectal cancer risk in Lynch syndrome. British Journal of Cancer, 2019, 121, 869-876.	6.5	12
132	Changes in selfâ€concept and risk of psychotic experiences in adolescence: a longitudinal populationâ€based cohort study. Journal of Child Psychology and Psychiatry and Allied Disciplines, 2019, 60, 1164-1173.	6.2	12
133	Risks of Colorectal Cancer and Cancer-Related Mortality in Familial Colorectal Cancer Type X and Lynch Syndrome Families. Journal of the National Cancer Institute, 2019, 111, 675-683.	6.3	12
134	A Meta-Analysis of Obesity and Risk of Colorectal Cancer in Patients with Lynch Syndrome: The Impact of Sex and Genetics. Nutrients, 2021, 13, 1736.	4.2	12
135	Risk of colorectal cancer for people with a mutation in both a MUTYH and a DNA mismatch repair gene. Familial Cancer, 2015, 14, 575-583.	2.0	11
136	Childhood cancers in families with and without Lynch syndrome. Familial Cancer, 2015, 14, 545-551.	2.0	10
137	Family History of Colorectal Cancer Is Not Associated with Colorectal Cancer Survival Regardless of Microsatellite Instability Status. Cancer Epidemiology Biomarkers and Prevention, 2014, 23, 1700-1704.	1.9	9
138	Genetic susceptibility markers for a breast-colorectal cancer phenotype: Exploratory results from genome-wide association studies. PLoS ONE, 2018, 13, e0196245.	2.5	9
139	Analysis in the Prospective Lynch Syndrome Database identifies sarcoma as part of the Lynch syndrome tumor spectrum. International Journal of Cancer, 2021, 148, 512-513.	5.4	9
140	Perceived Versus Predicted Risks of Colorectal Cancer and Selfâ€Reported Colonoscopies by Members of Mismatch Repair Gene Mutationâ€Carrying Families Who Have Declined Genetic Testing. Journal of Genetic Counseling, 2014, 23, 79-88.	1.7	8
141	Germline miRNA DNA variants and the risk of colorectal cancer by subtype. Genes Chromosomes and Cancer, 2017, 56, 177-184.	3.2	8
142	Risk factors identified in prenatal child protection reports. Children and Youth Services Review, 2021, 122, 105905.	2.1	8
143	Differential effects of phosphate binders on pre-dialysis serum bicarbonate in end-stage kidney disease patients on maintenance haemodialysis. BMC Nephrology, 2013, 14, 205.	1.8	7
144	Effect of flame spray coating on falling film evaporation for multi effect distillation system. Desalination and Water Treatment, 2013, 51, 822-829.	1.0	7

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145	GWASeq: targeted re-sequencing follow up to GWAS. BMC Genomics, 2016, 17, 176.	2.9	7
146	Re: Microsatellite Instability and BRAF Mutation Testing in Colorectal Cancer Prognostication. Journal of the National Cancer Institute, 2014, 106, dju180-dju180.	6.3	6
147	Cholecystectomy and the risk of colorectal cancer by tumor mismatch repair deficiency status. International Journal of Colorectal Disease, 2016, 31, 1451-1457.	2.3	6
148	Determining the familial risk distribution of colorectal cancer: a data mining approach. Familial Cancer, 2016, 15, 241-251.	2.0	6
149	Genetic variants within the hTERT gene and the risk of colorectal cancer in Lynch syndrome. Genes and Cancer, 2015, 6, 445-451.	1.8	6
150	The Experience of Extended Bowel Resection in Individuals With a High Metachronous Colorectal Cancer Risk: A Qualitative Study. Oncology Nursing Forum, 2016, 43, 444-452.	1.2	5
151	Modeling of Successive Cancer Risks in Lynch Syndrome Families in the Presence of Competing Risks Using Copulas. Biometrics, 2017, 73, 271-282.	1.5	5
152	Testing for Gene-Environment Interactions Using a Prospective Family Cohort Design: Body Mass Index in Early and Later Adulthood and Risk of Breast Cancer. American Journal of Epidemiology, 2017, 185, 487-500.	3.7	5
153	Interaction between polymorphisms in aspirin metabolic pathways, regular aspirin use and colorectal cancer risk: A case-control study in unselected white European populations. PLoS ONE, 2018, 13, e0192223.	2.5	5
154	Response to Li and Hopper. American Journal of Human Genetics, 2021, 108, 527-529.	6.1	5
155	Performance and quality indicators for the management of non-cancer chronic pain: a scoping review protocol. BMJ Open, 2016, 6, e010487.	2.1	4
156	Genetic and Environmental Modifiers of Cancer Risk in Lynch Syndrome. , 2018, , 67-89.		4
157	Metachronous colorectal cancer risk for mismatch repair gene mutation carriers â€“ the advantage of more extensive surgery. Hereditary Cancer in Clinical Practice, 2011, 9, O1.	1.5	3
158	Estimates of familial risks from family data are biased when ascertainment of families is not independent of family history. Gut, 2011, 60, 1162-1163.	13.5	3
159	Fertility and apparent genetic anticipation in Lynch syndrome. Familial Cancer, 2014, 13, 369-374.	2.0	3
160	Patient safety and rocket science. BMJ Quality and Safety, 2016, 25, 562-564.	4.3	3
161	The Colon Cancer Family Registry Cohort. , 2018, , 427-459.		3
162	Genetic Regulation of DNA Methylation Yields Novel Discoveries in GWAS of Colorectal Cancer. Cancer Epidemiology Biomarkers and Prevention, 2022, 31, 1068-1076.	1.9	3

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163	Queueing Networks and Markov Chains Analysis with the Octave queueing package. Performance Evaluation Review, 2022, 49, 47-52.	0.6	3
164	Intratumoral presence of the genotoxic gut bacteria pks+ E. coli, Enterotoxigenic Bacteroides fragilis, and Fusobacterium nucleatum and their association with clinicopathological and molecular features of colorectal cancer. British Journal of Cancer, 2024, 130, 728-740.	6.5	3
165	Is the reported modifying effect of 8q23.3 and 11q23.1 on colorectal cancer risk for MLH1 mutation carriers valid?. International Journal of Cancer, 2013, 133, 1762-1763.	5.4	2
166	Earlyâ€life exposure to sibling modifies the relationship between CD14 polymorphisms and allergic sensitization. Clinical and Experimental Allergy, 2019, 49, 331-340.	2.8	2
167	Associations of Height With the Risks of Colorectal and Endometrial Cancer in Persons With Lynch Syndrome. American Journal of Epidemiology, 2021, 190, 230-238.	3.7	2
168	Is prostate cancer a Lynch syndrome cancer?. Asian Journal of Andrology, 2013, 15, 588-589.	1.8	2
169	Genetic Variants in the Regulatory T cellâ€“Related Pathway and Colorectal Cancer Prognosis. Cancer Epidemiology Biomarkers and Prevention, 2020, 29, 2719-2728.	1.9	2
170	Fine-mapping analysis including over 254,000 East Asian and European descendants identifies 136 putative colorectal cancer susceptibility genes. Nature Communications, 2024, 15, .	13.0	2
171	Response: Table 1.. Journal of the National Cancer Institute, 2016, 108, djv385.	6.3	1
172	Pathways to a cancer-free future: a protocol for modelled evaluations to minimise the future burden of colorectal cancer in Australia. BMJ Open, 2020, 10, e036475.	2.1	1
173	Do the risks of Lynch syndrome-related cancers depend on the parent of origin of the mutation?. Familial Cancer, 2020, 19, 215-222.	2.0	1
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