Finlay Macrae

List of Publications by Year
in descending order

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Version: 2024-02-01

64 papers	3,657 citations	394421 19 h-index	155660 55 g-index
70 all docs	70 docs citations	70 times ranked	4853 citing authors

#	Article	IF	Citations
1	Worldwide prevalence of Lynch syndrome in patients with colorectal cancer: Systematic review and meta-analysis. Genetics in Medicine, 2022, 24, 971-985.	2.4	18
2	Aspirin and the Risk of Colorectal Cancer According to Genetic Susceptibility among Older Individuals. Cancer Prevention Research, 2022, 15, 447-454.	1.5	5
3	Lynch syndrome testing of colorectal cancer patients in a high-income country with universal healthcare: a retrospective study of current practice and gaps in seven australian hospitals. Hereditary Cancer in Clinical Practice, 2022, 20, 18.	1.5	2
4	Incident Cancer Risk and Signatures Among Older MUTYH Carriers: Analysis of Population-Based and Genomic Cohorts. Cancer Prevention Research, 2022, 15, 509-519.	1.5	1
5	Effect of Aspirin on Cancer Incidence and Mortality in Older Adults. Journal of the National Cancer Institute, 2021, 113, 258-265.	6.3	80
6	The â€œunnaturalâ€•history of colorectal cancer in Lynch syndrome: Lessons from colonoscopy surveillance. International Journal of Cancer, 2021, 148, 800-811.	5.1	55
7	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	28
8	Evaluation of literature searching tools for curation of mismatch repair gene variants in hereditary colon cancer. Genetics & Genomics Next, 2021, 2, e10039.	1.5	2
9	Cliniciansâ€™ opinions on recommending aspirin to prevent colorectal cancer to Australians aged 50â€“70 years: a qualitative study. BMJ Open, 2021, 11, e042261.	1.9	6
10	Impact of national guidelines on use of BRCA1/2 germline testing, risk management advice given to women with pathogenic BRCA1/2 variants and uptake of advice. Hereditary Cancer in Clinical Practice, 2021, 19, 24.	1.5	1
11	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.	2.8	11
12	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.	4.1	10
13	An RCT of a decision aid to support informed choices about taking aspirin to prevent colorectal cancer and other chronic diseases: a study protocol for the SITA (Should I Take Aspirin?) trial. Trials, 2021, 22, 452.	1.6	6
14	Genetic variant interpretation: a primer for clinicians. Internal Medicine Journal, 2021, 51, 1401-1406.	0.8	0
15	Commentary: Pivoting during a pandemic: developing a new recruitment model for a randomised controlled trial in response to COVID-19. Trials, 2021, 22, 605.	1.6	3
16	Uniting the Global Gastroenterology Community to Meet the Challenge of Climate Change and Nonrecyclable Waste. Journal of Clinical Gastroenterology, 2021, 55, 823-829.	2.2	4
17	Review article: investigation and management of internal fistulae in Crohnâ€™s disease. Alimentary Pharmacology and Therapeutics, 2021, 53, 1064-1079.	3.7	10
18	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	365

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19	Personal Protective Equipment for Endoscopy in Low-Resource Settings During the COVID-19 Pandemic. Journal of Clinical Gastroenterology, 2020, 54, 833-840.	2.2	10
20	Risk-Reducing Gynecological Surgery in Lynch Syndrome: Results of an International Survey from the Prospective Lynch Syndrome Database. Journal of Clinical Medicine, 2020, 9, 2290.	2.4	12
21	Cancer prevention with aspirin in hereditary colorectal cancer (Lynch syndrome), 10-year follow-up and registry-based 20-year data in the CAPP2 study: a double-blind, randomised, placebo-controlled trial. Lancet, The, 2020, 395, 1855-1863.	13.7	220
22	Increased risk of cervical dysplasia in females with autoimmune conditionsâ€"Results from an Australia database linkage study. PLoS ONE, 2020, 15, e0234813.	2.5	15
23	Celebrating the career and contributions of Dr Henry T. Lynch (1928â€"2019). Internal Medicine Journal, 2020, 50, 108-109.	0.8	1
24	Climate Change. Journal of Clinical Gastroenterology, 2020, 54, 393-397.	2.2	20
25	Genetic Variants in the Regulatory T cellâ€"Related Pathway and Colorectal Cancer Prognosis. Cancer Epidemiology Biomarkers and Prevention, 2020, 29, 2719-2728.	2.5	1
26	Comparing theory and non-theory based implementation approaches to improving referral practices in cancer genetics: a cluster randomised trial protocol. Trials, 2019, 20, 373.	1.6	10
27	Survival by colon cancer stage and screening interval in Lynch syndrome: a prospective Lynch syndrome database report. Hereditary Cancer in Clinical Practice, 2019, 17, 28.	1.5	27
28	Outcomes of screening and surveillance in people with two parents affected by colorectal cancers: experiences from the Familial Bowel Cancer Service. Hereditary Cancer in Clinical Practice, 2019, 17, 25.	1.5	2
29	Lack of association between screening interval and cancer stage in Lynch syndrome may be accounted for by over-diagnosis; a prospective Lynch syndrome database report. Hereditary Cancer in Clinical Practice, 2019, 17, 8.	1.5	42
30	Microsatellite instability and manifestations of angiogenesis in stage IV of sporadic colorectal carcinoma. Medicine (United States), 2019, 98, e13956.	1.0	7
31	Cancer screening in Australia: future directions in melanoma, Lynch syndrome, and liver, lung and prostate cancers. Public Health Research and Practice, 2019, 29, .	1.5	5
32	Worldwide Practice Patterns in Lynch Syndrome Diagnosis andÂManagement, Based on Data From the International Mismatch Repair Consortium. Clinical Gastroenterology and Hepatology, 2018, 16, 1901-1910.e11.	4.4	14
33	Cancer risk and survival in path_MMR carriers by gene and gender up to 75 years of age: a report from the Prospective Lynch Syndrome Database. Gut, 2018, 67, 1306-1316.	12.1	410
34	Current mismatch repair deficiency tumor testing practices and capabilities: A survey of Australian pathology providers. Asia-Pacific Journal of Clinical Oncology, 2018, 14, 417-425.	1.1	11
35	The use of a risk assessment and decision support tool (CRISP) compared with usual care in general practice to increase risk-stratified colorectal cancer screening: study protocol for a randomised controlled trial. Trials, 2018, 19, 397.	1.6	13
36	â€˜Why donâ€™t I need a colonoscopy?â€™ A novel approach to communicating risks and benefits of colorectal cancer screening. , 2018, 47, 343-349.		9

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37	Cancer incidence and survival in Lynch syndrome patients receiving colonoscopic and gynaecological surveillance: first report from the prospective Lynch syndrome database. Gut, 2017, 66, 464-472.	12.1	411
38	Changes in gene expression of neoâ€squamous mucosa after endoscopic treatment for dysplastic Barrettâ€™s esophagus and intramucosal adenocarcinoma. United European Gastroenterology Journal, 2017, 5, 13-20.	3.8	1
39	Relations between symptom severity, illness perceptions, visceral sensitivity, coping strategies and well-being in irritable bowel syndrome guided by the common sense model of illness. Psychology, Health and Medicine, 2017, 22, 524-534.	2.4	43
40	Incidence of and survival after subsequent cancers in carriers of pathogenic MMR variants with previous cancer: a report from the prospective Lynch syndrome database. Gut, 2017, 66, 1657-1664.	12.1	127
41	Exploring cliniciansâ€™ attitudes about using aspirin for risk reduction in people with Lynch Syndrome with no personal diagnosis of colorectal cancer. Familial Cancer, 2017, 16, 99-109.	1.9	6
42	Colorectal cancer incidence in path_MLH1 carriers subjected to different follow-up protocols: a Prospective Lynch Syndrome Database report. Hereditary Cancer in Clinical Practice, 2017, 15, 18.	1.5	49
43	Recurrent intestinal metaplasia at the gastroesophageal junction following endoscopic eradication of dysplastic Barrettâ€™s esophagus may not be benign. Endoscopy International Open, 2016, 04, E849-E858.	1.8	7
44	Harmonizing the interpretation of genetic variants across the world: the Malaysian experience. BMC Research Notes, 2016, 9, 125.	1.4	4
45	Knowledge and Uptake of Genetic Counseling and Colonoscopic Screening Among Individuals at Increased Risk for Lynch Syndrome and their Endoscopists from the Family Health Promotion Project. American Journal of Gastroenterology, 2016, 111, 285-293.	0.4	24
46	Evaluation of CADD Scores in Curated Mismatch Repair Gene Variants Yields a Model for Clinical Validation and Prioritization. Human Mutation, 2015, 36, 712-719.	2.5	39
47	Prevalence of mental health disorders in inflammatory bowel disease: an Australian outpatient cohort. Clinical and Experimental Gastroenterology, 2015, 8, 197.	2.3	28
48	Applicability of Next Generation Sequencing Technology in Microsatellite Instability Testing. Genes, 2015, 6, 46-59.	2.4	45
49	Application of a 5-tiered scheme for standardized classification of 2,360 unique mismatch repair gene variants in the InSiGHT locus-specific database. Nature Genetics, 2014, 46, 107-115.	21.4	410
50	Prediction of Crohnâ€™s disease aggression throughNOD2/CARD15gene sequencing in an Australian cohort. World Journal of Gastroenterology, 2014, 20, 5008.	3.3	12
51	Acceleration in colorectal carcinogenesis: the hare, the tortoise or myth?: TableÂ1. Gut, 2013, 62, 657-659.	12.1	4
52	Long-term effect of aspirin on cancer risk in carriers of hereditary colorectal cancer: an analysis from the CAPP2 randomised controlled trial. Lancet, The, 2011, 378, 2081-2087.	13.7	849
53	Familial adenomatous polyposis. Bailliere's Best Practice and Research in Clinical Gastroenterology, 2009, 23, 197-207.	2.4	18
54	PAIN AND COLONOSCOPY AND CO2. ANZ Journal of Surgery, 2008, 78, 836-836.	0.7	2

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55	Suggested actions from the Melbourne HVP Information Seminar. Nature Precedings, 2008, , .	0.1	0
56	Suggested actions from the Melbourne HVP Information Seminar. Nature Precedings, 2008, , .	0.1	0
57	Investigating rectal bleeding: red faced or reliable?. ANZ Journal of Surgery, 2001, 71, 699-700.	0.7	0
58	Evidence to support colonoscopic screening is insufficient. Medical Journal of Australia, 2000, 173, 332-332.	1.7	2
59	Frequency of codon 1061 and codon 1309 APC mutations in Australian familial adenomatous polyposis patients. Human Mutation, 1998, 11, S56-S57.	2.5	7
60	Characteristics of small bowel carcinoma in hereditary nonpolyposis colorectal carcinoma. , 1998, 83, 240-244.		125
61	Bowel cancer: Watching over the family. Journal of Gastroenterology and Hepatology (Australia), 1995, 10, 337-338.	2.8	1
62	The microcomputer: A friend here to help and stay. Journal of Gastroenterology and Hepatology (Australia), 1988, 3, 187-196.	2.8	0
63	Suggested actions from the Melbourne HVP Information Seminar. Nature Precedings, 0, , .	0.1	1
64	The Riskâ€Reducing Effect of Aspirin in Lynch Syndrome Carriers: Development and Evaluation of an Educational Leaflet. Genetics & Genomics Next, 0, , 2100046.	1.5	1