

Barbara Leggett

List of Publications by Year in descending order

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Version: 2024-02-01

42
papers

5,757
citations

218662

26
h-index

276858

41
g-index

43
all docs

43
docs citations

43
times ranked

7060
citing authors

#	ARTICLE	IF	CITATIONS
1	Liver Disease and Poor Adherence Limit Hepatitis C Cure: A Real-World Australian Treatment Cohort. <i>Digestive Diseases and Sciences</i> , 2023, 68, 291-303.	2.3	2
2	<i>Braf</i> mutation induces rapid neoplastic transformation in the aged and aberrantly methylated intestinal epithelium. <i>Gut</i> , 2022, 71, 1127-1140.	12.1	9
3	Clinicopathological Correlates of Dysplastic Sessile Serrated Lesion: A Prospective Cohort Study With a High Detection Rate. , 2022, 1, 313-320.		1
4	Traditional serrated adenoma-like lesions in patients with inflammatory bowel disease. <i>Human Pathology</i> , 2020, 97, 19-28.	2.0	12
5	Pathways to a cancer-free future: a protocol for modelled evaluations to minimise the future burden of colorectal cancer in Australia. <i>BMJ Open</i> , 2020, 10, e036475.	1.9	1
6	APC Mutation Marks an Aggressive Subtype of BRAF Mutant Colorectal Cancers. <i>Cancers</i> , 2020, 12, 1171.	3.7	28
7	Integrative Genome-Scale DNA Methylation Analysis of a Large and Unselected Cohort Reveals 5 Distinct Subtypes of Colorectal Adenocarcinomas. <i>Cellular and Molecular Gastroenterology and Hepatology</i> , 2019, 8, 269-290.	4.5	42
8	BRAF V600E immunohistochemistry demonstrates that some sessile serrated lesions with adenomatous dysplasia may represent collision lesions. <i>Histopathology</i> , 2019, 75, 81-87.	2.9	6
9	Sessile Serrated Adenomas in Young Patients may have Limited Risk of Malignant Progression. <i>Journal of Clinical Gastroenterology</i> , 2019, 53, e113-e116.	2.2	21
10	Cohort Profile: The Colon Cancer Family Registry Cohort (CCFRC). <i>International Journal of Epidemiology</i> , 2018, 47, 387-388i.	1.9	40
11	The role of APC in WNT pathway activation in serrated neoplasia. <i>Modern Pathology</i> , 2018, 31, 495-504.	5.5	43
12	A morphological and molecular study of proposed early forms of traditional serrated adenoma. <i>Histopathology</i> , 2018, 73, 1023-1029.	2.9	13
13	MLH1 93A/G polymorphism is associated with MLH1 promoter methylation and protein loss in dysplastic sessile serrated adenomas with BRAFV600E mutation. <i>BMC Cancer</i> , 2018, 18, 35.	2.6	15
14	Copy number profiles of paired primary and metastatic colorectal cancers. <i>Oncotarget</i> , 2018, 9, 3394-3405.	1.8	14
15	Clinicopathological and molecular features of sessile serrated adenomas with dysplasia or carcinoma. <i>Gut</i> , 2017, 66, 97-106.	12.1	161
16	High prevalence of sessile serrated adenomas in contemporary outpatient colonoscopy practice. <i>Internal Medicine Journal</i> , 2017, 47, 318-323.	0.8	27
17	Serrated tubulovillous adenoma of the large intestine. <i>Histopathology</i> , 2016, 68, 578-587.	2.9	28
18	Aspirin, Ibuprofen, and the Risk of Colorectal Cancer in Lynch Syndrome. <i>Journal of the National Cancer Institute</i> , 2015, 107, djv170.	6.3	80

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19	Critical Appraisal of the Diagnosis of the Sessile Serrated Adenoma. American Journal of Surgical Pathology, 2014, 38, 158-166.	3.7	94
20	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	93
21	The serrated pathway to colorectal carcinoma: current concepts and challenges. Histopathology, 2013, 62, 367-386.	2.9	377
22	Risks of Primary Extracolonic Cancers Following Colorectal Cancer in Lynch Syndrome. Journal of the National Cancer Institute, 2012, 104, 1363-1372.	6.3	193
23	Numerical ecology validates a biogeographical distribution and gender-based effect on mucosa-associated bacteria along the human colon. ISME Journal, 2011, 5, 801-809.	9.8	78
24	Risks of Lynch Syndrome Cancers for MSH6 Mutation Carriers. Journal of the National Cancer Institute, 2010, 102, 193-201.	6.3	328
25	Role of the Serrated Pathway in Colorectal Cancer Pathogenesis. Gastroenterology, 2010, 138, 2088-2100.	1.3	844
26	A Multicenter Blinded Study to Evaluate KRAS Mutation Testing Methodologies in the Clinical Setting. Journal of Molecular Diagnostics, 2009, 11, 543-552.	2.8	107
27	Dimensions of quality of life and psychosocial variables most salient to colorectal cancer patients. Psycho-Oncology, 2006, 15, 20-30.	2.3	101
28	CpG island methylator phenotype underlies sporadic microsatellite instability and is tightly associated with BRAF mutation in colorectal cancer. Nature Genetics, 2006, 38, 787-793.	21.4	1,715
29	Guaiac versus immunochemical tests: faecal occult blood test screening for colorectal cancer in a rural community. Australian and New Zealand Journal of Public Health, 2005, 29, 358-364.	1.8	42
30	Quality of life and colorectal cancer: a review. Australian and New Zealand Journal of Public Health, 2003, 27, 41-53.	1.8	42
31	Knowledge, Attitude and Intentions Related to Colorectal Cancer Screening Using Faecal Occult Blood Tests in a Rural Australian Population. Asia-Pacific Journal of Public Health, 2003, 15, 50-56.	1.0	26
32	FAECAL OCCULT BLOOD TEST: CURRENT PRACTICE IN A RURAL QUEENSLAND COMMUNITY. Australian Journal of Rural Health, 2002, 10, 57-64.	1.5	7
33	When is molecular genetic testing for colorectal cancer indicated?. Journal of Gastroenterology and Hepatology (Australia), 2002, 17, 389-393.	2.8	8
34	FAECAL OCCULT BLOOD TEST: CURRENT PRACTICE IN A RURAL QUEENSLAND COMMUNITY. Australian Journal of Rural Health, 2002, 10, 57-64.	1.5	1
35	Germline Mutations in BMPR1A/ALK3 Cause a Subset of Cases of Juvenile Polyposis Syndrome and of Cowden and Bannayan-Riley-Ruvalcaba Syndromes*. American Journal of Human Genetics, 2001, 69, 704-711.	6.2	236
36	Analysis of the Substrate Specificity of Human Sulfotransferases SULT1A1 and SULT1A3: Site-Directed Mutagenesis and Kinetic Studies. Biochemistry, 1999, 38, 10474-10479.	2.5	68

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37	CDX2, a human homologue of Drosophila caudal, is mutated in both alleles in a replication error positive colorectal cancer. <i>Oncogene</i> , 1998, 17, 657-659.	5.9	105
38	A family with attenuated familial adenomatous polyposis due to a mutation in the alternatively spliced region of APC exon 9. <i>Human Mutation</i> , 1998, 11, 450-455.	2.5	33
39	Mutations in DPC4 (SMAD4) cause juvenile polyposis syndrome, but only account for a minority of cases. <i>Human Molecular Genetics</i> , 1998, 7, 1907-1912.	2.9	142
40	Microsatellite instability in the insulin-like growth factor II receptor gene in gastrointestinal tumours. <i>Nature Genetics</i> , 1996, 14, 255-257.	21.4	429
41	AnAlu VpA Marker on chromosome 1 demonstrates that replication errors manifest at the adenoma-carcinoma transition in sporadic colorectal tumors. <i>Genes Chromosomes and Cancer</i> , 1995, 12, 251-254.	2.8	20
42	Genomic instability occurs in colorectal carcinomas but not in adenomas. <i>Human Mutation</i> , 1993, 2, 351-354.	2.5	124