

Joanne Young

List of Publications by Year in descending order

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59
papers

9,211
citations

117571

34
h-index

149623

56
g-index

59
all docs

59
docs citations

59
times ranked

9633
citing authors

#	ARTICLE	IF	CITATIONS
1	Outcomes for Metastatic Colorectal Cancer Based on Microsatellite Instability: Results from the South Australian Metastatic Colorectal Cancer Registry. <i>Targeted Oncology</i> , 2019, 14, 85-91.	1.7	10
2	Cohort Profile: The Colon Cancer Family Registry Cohort (CCFRC). <i>International Journal of Epidemiology</i> , 2018, 47, 387-388i.	0.9	40
3	Germline Variants and Advanced Colorectal Adenomas: Adenoma Prevention with Celecoxib Trial Genome-wide Association Study. <i>Clinical Cancer Research</i> , 2013, 19, 6430-6437.	3.2	9
4	Refinement of the associations between risk of colorectal cancer and polymorphisms on chromosomes 1q41 and 12q13.13. <i>Human Molecular Genetics</i> , 2012, 21, 934-946.	1.4	19
5	Serrated Lesions of the Colorectum: Review and Recommendations From an Expert Panel. <i>American Journal of Gastroenterology</i> , 2012, 107, 1315-1329.	0.2	948
6	Quality Assessment and Correlation of Microsatellite Instability and Immunohistochemical Markers among Population- and Clinic-Based Colorectal Tumors. <i>Journal of Molecular Diagnostics</i> , 2011, 13, 271-281.	1.2	131
7	Multiple Common Susceptibility Variants near BMP Pathway Loci GREM1, BMP4, and BMP2 Explain Part of the Missing Heritability of Colorectal Cancer. <i>PLoS Genetics</i> , 2011, 7, e1002105.	1.5	188
8	Parent of origin effects on age at colorectal cancer diagnosis. <i>International Journal of Cancer</i> , 2010, 127, 361-366.	2.3	8
9	Confirmation of Linkage to and Localization of Familial Colon Cancer Risk Haplotype on Chromosome 9q22. <i>Cancer Research</i> , 2010, 70, 5409-5418.	0.4	42
10	Risks of Lynch Syndrome Cancers for MSH6 Mutation Carriers. <i>Journal of the National Cancer Institute</i> , 2010, 102, 193-201.	3.0	328
11	Hyperplastic Polyposis Syndrome: Colorectal Cancer Predisposition. , 2010, , 111-131.		0
12	Efficient molecular screening of Lynch syndrome by specific 3' promoter methylation of the MLH1 or BRAF mutation in colorectal cancer with high-frequency microsatellite instability. <i>Oncology Reports</i> , 2009, 21, 1577-83.	1.2	24
13	Analysis of the Association between CIMP and BRAFV600E in Colorectal Cancer by DNA Methylation Profiling. <i>PLoS ONE</i> , 2009, 4, e8357.	1.1	133
14	Serrated Neoplasia of the Colorectum and Cigarette Smoking. <i>Gastroenterology</i> , 2008, 135, 323-324.	0.6	4
15	The Clinical Phenotype of Lynch Syndrome Due to Germ-Line PMS2 Mutations. <i>Gastroenterology</i> , 2008, 135, 419-428.e1.	0.6	480
16	Molecular Characterization of MSI-H Colorectal Cancer by <i>MLH1</i> Promoter Methylation, Immunohistochemistry, and Mismatch Repair Germline Mutation Screening. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2008, 17, 3208-3215.	1.1	207
17	Molecular, Pathologic, and Clinical Features of Early-Onset Endometrial Cancer: Identifying Presumptive Lynch Syndrome Patients. <i>Clinical Cancer Research</i> , 2008, 14, 1692-1700.	3.2	91
18	Lynch Syndrome in Women Less Than 50 Years of Age With Endometrial Cancer. <i>Obstetrics and Gynecology</i> , 2008, 112, 943.	1.2	2

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19	Microsatellite Instability Markers for Identifying Early-Onset Colorectal Cancers Caused by Germ-Line Mutations in DNA Mismatch Repair Genes. <i>Clinical Cancer Research</i> , 2007, 13, 2865-2869.	3.2	30
20	Variants on 9p24 and 8q24 Are Associated with Risk of Colorectal Cancer: Results from the Colon Cancer Family Registry. <i>Cancer Research</i> , 2007, 67, 11128-11132.	0.4	87
21	Serrated pathway colorectal cancer in the population: genetic consideration. <i>Gut</i> , 2007, 56, 1453-1459.	6.1	83
22	Epigenetic stem cell signature in cancer. <i>Nature Genetics</i> , 2007, 39, 157-158.	9.4	1,023
23	Stability of BAT26 in Lynch syndrome colorectal tumours. <i>European Journal of Human Genetics</i> , 2007, 15, 139-141.	1.4	8
24	Hyperplastic polyposis in the New Zealand population: a condition associated with increased colorectal cancer risk and European ancestry. <i>New Zealand Medical Journal</i> , 2007, 120, U2827.	0.5	34
25	Hyperplastic Polyposis Syndrome: Phenotypic Presentations and the Role of MBD4 and MYH. <i>Gastroenterology</i> , 2006, 131, 30-39.	0.6	186
26	High Prevalence of Sessile Serrated Adenomas With BRAF Mutations: A Prospective Study of Patients Undergoing Colonoscopy. <i>Gastroenterology</i> , 2006, 131, 1400-1407.	0.6	512
27	CpG island methylator phenotype underlies sporadic microsatellite instability and is tightly associated with BRAF mutation in colorectal cancer. <i>Nature Genetics</i> , 2006, 38, 787-793.	9.4	1,715
28	DNA methylation patterns in adenomas from FAP, multiple adenoma and sporadic colorectal carcinoma patients. <i>International Journal of Cancer</i> , 2006, 118, 907-915.	2.3	29
29	The Case for a Genetic Predisposition to Serrated Neoplasia in the Colorectum: Hypothesis and Review of the Literature. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2006, 15, 1778-1784.	1.1	103
30	DNA Methylation in Colorectal Cancer. , 2005, , 59-68.		1
31	Isolated Loss of PMS2 Expression in Colorectal Cancers: Frequency, Patient Age, and Familial Aggregation. <i>Clinical Cancer Research</i> , 2005, 11, 6466-6471.	3.2	54
32	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, 254-263.	2.4	123
33	Identification of Genes Uniquely Involved in Frequent Microsatellite Instability Colon Carcinogenesis by Expression Profiling Combined with Epigenetic Scanning. <i>Cancer Research</i> , 2004, 64, 2434-2438.	0.4	63
34	Cadherin/catenin complex appears to be intact in hepatocellular carcinomas from Australia and South Africa. <i>Journal of Gastroenterology and Hepatology (Australia)</i> , 2004, 19, 676-682.	1.4	7
35	Mapping of a candidate colorectal cancer tumor-suppressor gene to a 900-kilobase region on the short arm of chromosome 8. <i>Genes Chromosomes and Cancer</i> , 2004, 40, 247-260.	1.5	34
36	Colorectal Cancer With Mutation in BRAF, KRAS, and Wild-Type With Respect to Both Oncogenes Showing Different Patterns of DNA Methylation. <i>Journal of Clinical Oncology</i> , 2004, 22, 4584-4594.	0.8	202

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37	Analysis of the transcription regulator,CNOT7, as a candidate chromosome 8 tumor suppressor gene in colorectal cancer. International Journal of Cancer, 2003, 106, 505-509.	2.3	14
38	Emerging concepts in colorectal neoplasia. Gastroenterology, 2002, 123, 862-876.	0.6	444
39	Mutation searching in colorectal cancer studies: experience with a denaturing high-pressure liquid chromatography system for exon-by-exon scanning of tumour suppressor genes. Pathology, 2002, 34, 529-533.	0.3	20
40	Detection of telomerase activity in biopsy samples of colorectal cancer. Journal of Gastroenterology and Hepatology (Australia), 2002, 14, 328-332.	1.4	20
41	Evolution of colorectal cancer: Change of pace and change of direction. Journal of Gastroenterology and Hepatology (Australia), 2002, 17, 17-26.	1.4	78
42	Reciprocal relationship between methylation status and loss of heterozygosity at the p14ARF locus in Australian and South African hepatocellular carcinomas. Journal of Gastroenterology and Hepatology (Australia), 2002, 17, 301-307.	1.4	16
43	Morphological and molecular heterogeneity within nonmicrosatellite instability-high colorectal cancer. Cancer Research, 2002, 62, 6011-4.	0.4	99
44	Features of Colorectal Cancers with High-Level Microsatellite Instability Occurring in Familial and Sporadic Settings. American Journal of Pathology, 2001, 159, 2107-2116.	1.9	337
45	Hyperplastic Polyposis. American Journal of Surgical Pathology, 2001, 25, 177-184.	2.1	185
46	Sequence variants ofDLC1 in colorectal and ovarian tumours. Human Mutation, 2000, 15, 156-165.	1.1	29
47	p73 Is up-regulated in a subset of hepatocellular carcinomas. Hepatology, 2000, 31, 601-605.	3.6	44
48	No evidence of increased risk of colorectal cancer in individuals heterozygous for the Cys282Tyr haemochromatosis mutation. Journal of Gastroenterology and Hepatology (Australia), 1999, 14, 1188-1191.	1.4	30
49	Expression of Bcl-2 protein is decreased in colorectal adenocarcinomas with microsatellite instability. Oncogene, 1999, 18, 1245-1249.	2.6	47
50	Angiogenic factor VEGF is decreased in human colorectal neoplasms showing DNA microsatellite instability. , 1999, 189, 319-325.		24
51	CDX2, a human homologue of Drosophila caudal, is mutated in both alleles in a replication error positive colorectal cancer. Oncogene, 1998, 17, 657-659.	2.6	105
52	Reciprocal relationship between the tumor suppressors p53 and BAX in primary colorectal cancers. Oncogene, 1998, 17, 2003-2008.	2.6	57
53	A family with attenuated familial adenomatous polyposis due to a mutation in the alternatively spliced region of APC exon 9. Human Mutation, 1998, 11, 450-455.	1.1	33
54	Apparent protection from instability of repeat sequences in cancer-related genes in replication error positive gastrointestinal cancers. Oncogene, 1997, 14, 2613-2618.	2.6	15

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55	Characteristics of metachronous colorectal carcinoma occurring despite colonoscopic surveillance. <i>Diseases of the Colon and Rectum</i> , 1997, 40, 603-608.	0.7	25
56	Mutation of hMSH3 and hMSH6 mismatch repair genes in genetically unstable human colorectal and gastric carcinomas. <i>Human Mutation</i> , 1997, 10, 474-478.	1.1	58
57	Microsatellite instability in the insulin-like growth factor II receptor gene in gastrointestinal tumours. <i>Nature Genetics</i> , 1996, 14, 255-257.	9.4	429
58	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.	1.5	20
59	Genomic instability occurs in colorectal carcinomas but not in adenomas. <i>Human Mutation</i> , 1993, 2, 351-354.	1.1	124