

Fay Kastrinos

List of Publications by Year in descending order

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Version: 2024-02-01

47
papers

2,605
citations

304368

22
h-index

276539

41
g-index

47
all docs

47
docs citations

47
times ranked

3232
citing authors

#	ARTICLE	IF	CITATIONS
1	Timeline of Development of Pancreatic Cancer and Implications for Successful Early Detection in High-Risk Individuals. <i>Gastroenterology</i> , 2022, 162, 772-785.e4.	0.6	60
2	Inherited Gastrointestinal Cancers and the Role of Genetic Evaluation and Testing. <i>Gastrointestinal Endoscopy Clinics of North America</i> , 2022, 32, xv-xvi.	0.6	1
3	Familial Predisposition and Genetic Risk Factors Associated with Pancreatic Cancer. <i>Gastrointestinal Endoscopy Clinics of North America</i> , 2022, 32, 1-12.	0.6	1
4	Late-Stage Pancreatic Cancer Detected During High-Risk Individual Surveillance: A Systematic Review and Meta-Analysis. <i>Gastroenterology</i> , 2022, 162, 786-798.	0.6	19
5	Cost-effectiveness of neoadjuvant FOLFIRINOX versus gemcitabine plus nab-paclitaxel in borderline resectable/locally advanced pancreatic cancer patients. <i>Cancer Reports</i> , 2022, 5, e1565.	0.6	4
6	The Multicenter Cancer of Pancreas Screening Study: Impact on Stage and Survival. <i>Journal of Clinical Oncology</i> , 2022, 40, 3257-3266.	0.8	69
7	The PRECEDE consortium: A longitudinal international cohort study of individuals with genetic risk or familial pancreatic cancer. <i>Journal of Clinical Oncology</i> , 2022, 40, e16239-e16239.	0.8	0
8	Characterizing germline APC and MUTYH variants in Ashkenazi Jews compared to other individuals. <i>Familial Cancer</i> , 2021, 20, 111-116.	0.9	5
9	Lead-Time Trajectory of CA19-9 as an Anchor Marker for Pancreatic Cancer Early Detection. <i>Gastroenterology</i> , 2021, 160, 1373-1383.e6.	0.6	77
10	Inherited predisposition to pancreatic cancer. <i>Seminars in Oncology</i> , 2021, 48, 2-9.	0.8	0
11	How I Approach Screening for Pancreatic Cancer. <i>American Journal of Gastroenterology</i> , 2021, 116, 1569-1571.	0.2	1
12	Inherited Predisposition to Gastric Cancer. <i>Gastrointestinal Endoscopy Clinics of North America</i> , 2021, 31, 467-487.	0.6	5
13	Gene-Specific Variation in Colorectal Cancer Surveillance Strategies for Lynch Syndrome. <i>Gastroenterology</i> , 2021, 161, 453-462.e15.	0.6	17
14	Cost-effectiveness Analysis of Genotype-Specific Surveillance and Preventive Strategies for Gynecologic Cancers Among Women With Lynch Syndrome. <i>JAMA Network Open</i> , 2021, 4, e2123616.	2.8	7
15	COVID-19 related pancreatic cancer surveillance disruptions amongst high-risk individuals. <i>Pancreatology</i> , 2021, 21, 1048-1051.	0.5	8
16	Advanced adenomas may be a red flag for hereditary cancer syndromes. <i>Hereditary Cancer in Clinical Practice</i> , 2021, 19, 8.	0.6	2
17	Screening for Pancreatic Ductal Adenocarcinoma: Are We Asking the Impossible? Letter. <i>Cancer Prevention Research</i> , 2021, 14, 973-974.	0.7	3
18	Clinical Factors Associated With Gastric Cancer in Individuals With Lynch Syndrome. <i>Clinical Gastroenterology and Hepatology</i> , 2020, 18, 830-837.e1.	2.4	38

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19	Management of patients with increased risk for familial pancreatic cancer: updated recommendations from the International Cancer of the Pancreas Screening (CAPS) Consortium. <i>Gut</i> , 2020, 69, 7-17.	6.1	357
20	Optimal Timing of Total Gastrectomy to Prevent Diffuse Gastric Cancer in Individuals With Pathogenic Variants in CDH1. <i>Clinical Gastroenterology and Hepatology</i> , 2020, 18, 822-829.e4.	2.4	16
21	Use of Family History and Genetic Testing to Determine Risk of Colorectal Cancer. <i>Gastroenterology</i> , 2020, 158, 389-403.	0.6	59
22	Clinical Factors Associated with Urinary Tract Cancer in Individuals with Lynch Syndrome. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2020, 29, 193-199.	1.1	11
23	Screening for Pancreatic Cancer. <i>JAMA - Journal of the American Medical Association</i> , 2019, 322, 407.	3.8	33
24	Commentary:ÂPREMM5 threshold of 2.5% is recommended to improve identification of PMS2 carriers. <i>Familial Cancer</i> , 2018, 17, 567-567.	0.9	3
25	Community Practice Implementation of a Self-administered Version of PREMM1,2,6 to Assess Risk for Lynch Syndrome. <i>Clinical Gastroenterology and Hepatology</i> , 2018, 16, 49-58.	2.4	25
26	The Role of Genetics in Pancreatitis. <i>Gastrointestinal Endoscopy Clinics of North America</i> , 2018, 28, 587-603.	0.6	35
27	Clinical factors associated with urinary tract cancers (UTCs) among Lynch syndrome (LS) patients (Pts).. <i>Journal of Clinical Oncology</i> , 2018, 36, 1517-1517.	0.8	0
28	Comparison of Colonoscopy Quality Measures Across Various Practice Settings and the Impact of Performance Scorecards. <i>Digestive Diseases and Sciences</i> , 2017, 62, 894-902.	1.1	13
29	Development and Validation of the PREMM₅ Model for Comprehensive Risk Assessment of Lynch Syndrome. <i>Journal of Clinical Oncology</i> , 2017, 35, 2165-2172.	0.8	126
30	Comparison of Prediction Models for Lynch Syndrome Among Individuals With Colorectal Cancer. <i>Journal of the National Cancer Institute</i> , 2016, 108, .	3.0	29
31	Identification of Lynch syndrome (LS) in patients (pts) without prior LS-associated cancer.. <i>Journal of Clinical Oncology</i> , 2016, 34, 1521-1521.	0.8	0
32	Racial variation in frequency and phenotypes of APC and MUTYH mutations in 6,169 individuals undergoing genetic testing. <i>Genetics in Medicine</i> , 2015, 17, 815-821.	1.1	21
33	Family Matters in Lynch Syndrome. <i>Journal of the National Cancer Institute</i> , 2015, 107, djv050-djv050.	3.0	1
34	Familial Colorectal Cancer, Beyond Lynch Syndrome. <i>Clinical Gastroenterology and Hepatology</i> , 2014, 12, 1059-1068.	2.4	70
35	History, Genetics, and Strategies for Cancer Prevention in Lynch Syndrome. <i>Clinical Gastroenterology and Hepatology</i> , 2014, 12, 715-727.	2.4	65
36	Prediction models in Lynch syndrome. <i>Familial Cancer</i> , 2013, 12, 217-228.	0.9	16

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37	Prevalence and Phenotypes of APC and MUTYH Mutations in Patients With Multiple Colorectal Adenomas. <i>JAMA - Journal of the American Medical Association</i> , 2012, 308, 485-492.	3.8	183
38	The PREMM1,2,6 Model Predicts Risk of MLH1, MSH2, and MSH6 Germline Mutations Based on Cancer History. <i>Gastroenterology</i> , 2011, 140, 73-81.e5.	0.6	171
39	Inherited Colorectal Cancer Syndromes. <i>Cancer Journal (Sudbury, Mass)</i> , 2011, 17, 405-415.	1.0	100
40	CT colonography had 90% sensitivity and 86% specificity for diagnosing large adenomas and cancer in asymptomatic adults. <i>Annals of Internal Medicine</i> , 2009, 150, JC2.	2.0	0
41	Development and Validation of a Colon Cancer Risk Assessment Tool for Patients Undergoing Colonoscopy. <i>American Journal of Gastroenterology</i> , 2009, 104, 1508-1518.	0.2	62
42	Risk of Pancreatic Cancer in Families With Lynch Syndrome. <i>JAMA - Journal of the American Medical Association</i> , 2009, 302, 1790.	3.8	434
43	Maybe a Baby? Revisiting the Effect of IBD on Conception, Pregnancy, and Newborn Outcomes. <i>Inflammatory Bowel Diseases</i> , 2009, 15, 475-477.	0.9	0
44	Calculation of Risk of Colorectal and Endometrial Cancer Among Patients With Lynch Syndrome. <i>Gastroenterology</i> , 2009, 137, 1621-1627.	0.6	322
45	Phenotype Comparison of MLH1 and MSH2 Mutation Carriers in a Cohort of 1,914 Individuals Undergoing Clinical Genetic Testing in the United States. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2008, 17, 2044-2051.	1.1	65
46	Attitudes Toward Prenatal Genetic Testing in Patients With Familial Adenomatous Polyposis. <i>American Journal of Gastroenterology</i> , 2007, 102, 1284-1290.	0.2	45
47	Recently Identified Colon Cancer Predispositions: MYH and MSH6 Mutations. <i>Seminars in Oncology</i> , 2007, 34, 418-424.	0.8	26