Brandie Heald

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/9205366/publications.pdf

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29 papers	1,578 citations	516710 16 h-index	501196 28 g-index
29 all docs	29 docs citations	29 times ranked	2756 citing authors

#	Article	IF	Citations
1	Prevalence and Spectrum of Germline Cancer Susceptibility Gene Mutations Among Patients With Early-Onset Colorectal Cancer. JAMA Oncology, 2017, 3, 464.	7.1	510
2	Frequent Gastrointestinal Polyps and Colorectal Adenocarcinomas in a Prospective Series of PTEN Mutation Carriers. Gastroenterology, 2010, 139, 1927-1933.	1.3	251
3	Implementation of Universal Microsatellite Instability and Immunohistochemistry Screening for Diagnosing Lynch Syndrome in a Large Academic Medical Center. Journal of Clinical Oncology, 2013, 31, 1336-1340.	1.6	147
4	Implementation of tumor testing for lynch syndrome in endometrial cancers at a large academic medical center. Gynecologic Oncology, 2013, 130, 121-126.	1.4	94
5	The Prevalence of Hereditary Hemorrhagic Telangiectasia in Juvenile Polyposis Syndrome. Diseases of the Colon and Rectum, 2012, 55, 886-892.	1.3	71
6	Prevalence of thoracic aortopathy in patients with juvenile Polyposis Syndromeâ€Hereditary Hemorrhagic Telangiectasia due to SMAD4. American Journal of Medical Genetics, Part A, 2015, 167, 1758-1762.	1.2	67
7	Upper tract urothelial carcinomas: frequency of association with mismatch repair protein loss and lynch syndrome. Modern Pathology, 2017, 30, 146-156.	5.5	66
8	Collaborative Group of the Americas on Inherited Gastrointestinal Cancer Position statement on multigene panel testing for patients with colorectal cancer and/or polyposis. Familial Cancer, 2020, 19, 223-239.	1.9	39
9	Prospective comparison of family medical history with personal genome screening for risk assessment of common cancers. European Journal of Human Genetics, 2012, 20, 547-551.	2.8	37
10	Prospective Statewide Study of Universal Screening for Hereditary Colorectal Cancer: The Ohio Colorectal Cancer Prevention Initiative. JCO Precision Oncology, 2021, 5, 779-791.	3.0	31
11	Investigating the Link between Lynch Syndrome and Breast Cancer. The Journal of Breast Health, 2020, 16, 106-109.	1.0	30
12	Assessment of clinical workload for general and specialty genetic counsellors at an academic medical center: a tool for evaluating genetic counselling practices. Npj Genomic Medicine, 2016, 1, 16010.	3.8	28
13	Exome Sequencing Reveals Germline SMAD9 Mutation That Reduces Phosphatase and Tensin Homolog Expression and Is Associated With Hamartomatous Polyposis and Gastrointestinal Ganglioneuromas. Gastroenterology, 2015, 149, 886-889.e5.	1.3	24
14	Using chatbots to screen for heritable cancer syndromes in patients undergoing routine colonoscopy. Journal of Medical Genetics, 2021, 58, 807-814.	3.2	23
15	Patients Do Not Recall Important Details About Polyps, Required for Colorectal Cancer Prevention. Clinical Gastroenterology and Hepatology, 2013, 11, 543-547.e2.	4.4	21
16	Universal Versus Targeted Screening for Lynch Syndrome: Comparing Ascertainment and Costs Based on Clinical Experience. Digestive Diseases and Sciences, 2016, 61, 2887-2895.	2.3	20
17	Genetic Counseling and Germline Testing in the Era of Tumor Sequencing: A Cohort Study. JNCI Cancer Spectrum, 2020, 4, pkaa018.	2.9	18
18	Early genetic counseling and detection of CDH1 mutation in asymptomatic carriers improves survival in hereditary diffuse gastric cancer. Surgery, 2018, 164, 754-759.	1.9	17

#	Article	IF	Citations
19	Diagnostic Approach to Hereditary Colorectal Cancer Syndromes. Clinics in Colon and Rectal Surgery, 2015, 28, 205-214.	1.1	13
20	A Time Study of Cancer Genetic Counselors Using a Genetic Counselor-Only Patient Care Model Versus a Traditional Combined Genetic Counselor Plus Medical Geneticist Care Model. Journal of the National Comprehensive Cancer Network: JNCCN, 2013, 11, 1076-1081.	4.9	12
21	Targeted Next-Generation Sequencing in Men with Metastatic Prostate Cancer: a Pilot Study. Targeted Oncology, 2018, 13, 495-500.	3.6	12
22	Clinically actionable findings on surveillance EGD in asymptomatic patients with Lynch syndrome. Gastrointestinal Endoscopy, 2022, 95, 105-114.	1.0	11
23	Natural history of ampullary adenomas in familial adenomatous polyposis: a long-term follow-up study. Gastrointestinal Endoscopy, 2022, 95, 455-467.e3.	1.0	10
24	ACG Guidelines on Management of PTEN-Hamartoma Tumor Syndrome: Does the Evidence Support so Much so Young?. American Journal of Gastroenterology, 2015, 110, 1733-1734.	0.4	7
25	Detecting and managing hereditary colorectal cancer syndromes in your practice. Cleveland Clinic Journal of Medicine, 2012, 79, 787-796.	1.3	7
26	Strategies for clinical implementation of screening for hereditary cancer syndromes. Seminars in Oncology, 2016, 43, 609-614.	2.2	6
27	Evaluation of Urinalysis-Based Screening for Urothelial Carcinoma in Patients With Lynch Syndrome. Diseases of the Colon and Rectum, 2022, 65, 40-45.	1.3	5
28	Presented Abstracts from the Thirty First Annual Education Conference of the National Society of Genetic Counselors (Boston, MA, October 2012). Journal of Genetic Counseling, 2012, 21, 884-984.	1.6	1
29	Correspondence to Vorselaars et al. thoracic aorta dilation in patients with hereditary hemorrhagic telangiectasia due to SMAD4 gene mutation. American Journal of Medical Genetics, Part A, 2016, 170, 813-813.	1.2	0