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	Clinically actionable findings on surveillance EGD in asymptomatic patients with Lynch syndrome. Gastrointestinal Endoscopy, 2022, 95, 105-114.	1.0	11
2	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
3	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
4	Using chatbots to screen for heritable cancer syndromes in patients undergoing routine colonoscopy. Journal of Medical Genetics, 2021, 58, 807-814.	3.2	23
5	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
6	Genetic Counseling and Germline Testing in the Era of Tumor Sequencing: A Cohort Study. JNCI Cancer Spectrum, 2020, 4, pkaa018.	2.9	18
7	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
8	Investigating the Link between Lynch Syndrome and Breast Cancer. The Journal of Breast Health, 2020, 16, 106-109.	1.0	30
9	Targeted Next-Generation Sequencing in Men with Metastatic Prostate Cancer: a Pilot Study. Targeted Oncology, 2018, 13, 495-500.	3.6	12
10	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
11	Prevalence and Spectrum of Germline Cancer Susceptibility Gene Mutations Among Patients With Early-Onset Colorectal Cancer. JAMA Oncology, 2017, 3, 464.	7.1	510
12	Upper tract urothelial carcinomas: frequency of association with mismatch repair protein loss and lynch syndrome. Modern Pathology, 2017, 30, 146-156.	5.5	66
13	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
14	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
15	Strategies for clinical implementation of screening for hereditary cancer syndromes. Seminars in Oncology, 2016, 43, 609-614.	2.2	6
16	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
17	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
18	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

#	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	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
21	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
22	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
23	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
24	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
25	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
26	The Prevalence of Hereditary Hemorrhagic Telangiectasia in Juvenile Polyposis Syndrome. Diseases of the Colon and Rectum, 2012, 55, 886-892.	1.3	71
27	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
28	Detecting and managing hereditary colorectal cancer syndromes in your practice. Cleveland Clinic Journal of Medicine, 2012, 79, 787-796.	1.3	7
29	Frequent Gastrointestinal Polyps and Colorectal Adenocarcinomas in a Prospective Series of PTEN Mutation Carriers. Gastroenterology, 2010, 139, 1927-1933.	1.3	251