Megan C Roberts

List of Publications by Year
in descending order

33 papers	864 citations	759233 12 h-index	526287 27 g-index
34 all docs	34 docs citations	34 times ranked	1457 citing authors

#	Article	IF	Citations
1	Multifactorial causal beliefs and colorectal cancer screening: A structural equation modeling investigation. Journal of Health Psychology, 2022, 27, 2463-2477.	2.3	4
2	Examining the role of language competency in genetic testing awareness among adults in the United States. Journal of Genetic Counseling, 2022, , .	1.6	0
3	The Rise of Population Genomic Screening: Characteristics of Current Programs and the Need for Evidence Regarding Optimal Implementation. Journal of Personalized Medicine, 2022, 12, 692.	2.5	11
4	Involving patients and their families in deciding to use next generation sequencing: Results from a nationally representative survey of U.S. oncologists. Patient Education and Counseling, 2021, 104, 33-39.	2.2	2
5	Complementary approaches to problem solving in healthcare and public health: implementation science and human-centered design. Translational Behavioral Medicine, 2021, 11, 1115-1121.	2.4	42
6	Age at initiation of screening mammography by family history of breast cancer in the breast cancer surveillance consortium. Cancer Causes and Control, 2021, 32, 103-107.	1.8	6
7	Patient Perspectives on the Risk-Based NLST Outcomes Tool for Lung Cancer Screening. Journal of Cancer Education, 2021, , 1.	1.3	2
8	Improving reporting standards for polygenic scores in risk prediction studies. Nature, 2021, 591, 211-219.	27.8	265
9	Predictors of Chronic Opioid Use: A Population-Level Analysis of North Carolina Cancer Survivors Using Multi-Payer Claims. Journal of the National Cancer Institute, 2021, 113, 1581-1589.	6.3	2
10	Oncologist-Reported Reasons for Not Ordering Multimarker Tumor Panels: Results From a Nationally Representative Survey. JCO Precision Oncology, 2021, 5, 701-709.	3.0	6
11	Advancing precision public health using human genomics: examples from the field and future research opportunities. Genome Medicine, 2021, 13, 97.	8.2	26
12	Ethiopian paediatric oncology registry progress report: documentation practice improvements at tertiary care centre in Addis Ababa, Ethiopia. Archives of Disease in Childhood, 2021, 106, 1244-1245.	1.9	1
13	Strategies to Integrate Genomic Medicine into Clinical Care: Evidence from the IGNITE Network. Journal of Personalized Medicine, 2021, 11, 647.	2.5	13
14	Uptake of Genetic Testing Among Patients with Cancer At Risk for Lynch Syndrome in the National Health Interview Survey. Cancer Prevention Research, 2021, 14, 927-932.	1.5	6
15	Evaluation of the Veterans Affairs Pharmacogenomic Testing for Veterans (PHASER) clinical program at initial test sites. Pharmacogenomics, 2021, 22, 1121-1133.	1.3	5
16	Results of a pre-implementation analysis of Ethiopiaâ€™s National Pediatric Cancer Registry. Journal of Oncology Pharmacy Practice, 2021, 27, 1940-1947.	0.9	3
17	Communication About Hereditary Cancers on Social Media: A Content Analysis of Tweets About Hereditary Breast and Ovarian Cancer and Lynch Syndrome. Journal of Cancer Education, 2020, 35, 131-137.	1.3	9
18	Barriers and facilitators for cascade testing in genetic conditions: a systematic review. European Journal of Human Genetics, 2020, 28, 1631-1644.	2.8	55

#	Article	IF	Citations
19	Stakeholder Perspectives on Overcoming Barriers to Cascade Testing in Lynch Syndrome: A Qualitative Study. Cancer Prevention Research, 2020, 13, 1037-1046.	1.5	12
20	Early career investigators and precision public health. Lancet, The, 2019, 394, 382-383.	13.7	6
21	Exploring Predictors of Genetic Counseling and Testing for Hereditary Breast and Ovarian Cancer: Findings from the 2015 U.S. National Health Interview Survey. Journal of Personalized Medicine, 2019, 9, 26.	2.5	21
22	Leveraging Implementation Science to Address Health Disparities in Genomic Medicine: Examples from the Field. Ethnicity and Disease, 2019, 29, 187-192.	2.3	43
23	Psychosocial, attitudinal, and demographic correlates of cancer-related germline genetic testing in the 2017 Health Information National Trends Survey. Journal of Community Genetics, 2019, 10, 453-459.	1.2	8
24	The FDA authorization of direct-to-consumer genetic testing for three BRCA1/2 pathogenic variants: a twitter analysis of the publicâ€™s response. JAMIA Open, 2019, 2, 411-415.	2.0	8
25	Concordance with BRCA1/2 testing guidelines among women in The Health of Women (HOW) StudyÂ®. Breast Cancer Research and Treatment, 2019, 173, 719-726.	2.5	5
26	Engagement with Genetic Information and Uptake of Genetic Testing: the Role of Trust and Personal Cancer History. Journal of Cancer Education, 2018, 33, 893-900.	1.3	15
27	Delivery Of Cascade Screening For Hereditary Conditions: A Scoping Review Of The Literature. Health Affairs, 2018, 37, 801-808.	5.2	114
28	Twitter use at the 2016 Conference on the Science of Dissemination and Implementation in Health: analyzing #DIScience16. Implementation Science, 2018, 13, 34.	6.9	22
29	Breast cancer-specific survival in patients with lymph node-positive hormone receptor-positive invasive breast cancer and Oncotype DX Recurrence Score results in the SEER database. Breast Cancer Research and Treatment, 2017, 163, 303-310.	2.5	80
30	The effect of a celebrity health disclosure on demand for health care: trends in BRCA testing and subsequent health services use. Journal of Community Genetics, 2017, 8, 141-146.	1.2	39
31	Patient-Centered Communication for Discussing Oncotype DX Testing. Cancer Investigation, 2016, 34, 205-212.	1.3	5
32	Use and Costs for Tumor Gene Expression Profiling Panels in the Management of Breast Cancer From 2006 to 2012: Implications for Genomic Test Adoption Among Private Payers. Journal of Oncology Practice, 2015, 11, 273-277.	2.5	17
33	Barriers and Facilitators for Population Genetic Screening in Healthy Populations: A Systematic Review. Frontiers in Genetics, 0, 13, .	2.3	10