Holly Etchegary

List of Publications by Year
in descending order

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Version: 2024-02-01

76
papers

1,168
citations

394421
19
h-index

454955
30
g-index

79
all docs

79
docs citations

79
times ranked

1782
citing authors

#	Article	IF	Citations
1	Reflections on the Cost of "Low-Cost" Whole Genome Sequencing: Framing the Health Policy Debate. PLoS Biology, 2013, 11, e1001699.	5.6	67
2	Using Facebook Advertising to Recruit Representative Samples: Feasibility Assessment of a Cross-Sectional Survey. Journal of Medical Internet Research, 2019, 21, e14021.	4.3	67
3	The Influence of Experiential Knowledge on Prenatal Screening and Testing Decisions. Genetic Testing and Molecular Biomarkers, 2008, 12, 115-124.	1.7	64
4	Physician-reported barriers to using evidence-based recommendations for low back pain in clinical practice: a systematic review and synthesis of qualitative studies using the Theoretical Domains Framework. Implementation Science, 2019, 14, 49.	6.9	64
5	Personal privacy, public benefits, and biobanks: a conjoint analysis of policy priorities and public perceptions. Genetics in Medicine, 2012, 14, 229-235.	2.4	61
6	Exploring informed choice in the context of prenatal testing: findings from a qualitative study. Health Expectations, 2008, 11, 355-365.	2.6	46
7	Information Processing in the Context of Genetic Risk: Implications for Geneticâ€Risk Communication. Journal of Genetic Counseling, 2007, 16, 419-432.	1.6	39
8	Recruiting patients as partners in health research: a qualitative descriptive study. Research Involvement and Engagement, 2017, 3, 15.	2.9	39
9	A family genetic risk communication framework: guiding tool development in genetics health services. Journal of Community Genetics, 2013, 4, 233-242.	1.2	38
10	Decisionâ€Making About Inherited Cancer Risk: Exploring Dimensions of Genetic Responsibility. Journal of Genetic Counseling, 2009, 18, 252-264.	1.6	32
11	Consulting the community: public expectations and attitudes about genetics research. European Journal of Human Genetics, 2013, 21, 1338-1343.	2.8	30
12	Public attitudes toward genetic risk testing and its role in healthcare. Personalized Medicine, 2014, 11, 509-522.	1.5	30
13	Community engagement with genetics: public perceptions and expectations about genetics research. Health Expectations, 2015, 18, 1413-1425.	2.6	30
14	Consent for newborn screening: parentsâ€™ and health-care professionalsâ€™ experiences of consent in practice. European Journal of Human Genetics, 2016, 24, 1530-1534.	2.8	29
15	Healthcare experiences of families affected by Huntington disease: need for improved care. Chronic Illness, 2011, 7, 225-238.	1.5	28
16	Public Attitudes About Genetic Testing in the Newborn Period. JOGNN - Journal of Obstetric, Gynecologic, and Neonatal Nursing, 2012, 41, 191-200.	0.5	28
17	Discovering the Family History of Huntington Disease (HD). Journal of Genetic Counseling, 2006, 15, 105-117.	1.6	23
18	Psychosocial aspects of venous thromboembolic disease: An exploratory study. Thrombosis Research, 2008, 122, 491-500.	1.7	22

#	ARTICLE	IF	CITATIONS
19	â€˜<i>They had the right to know.</i>â€™ Genetic risk and perceptions of responsibility. Psychology and Health, 2008, 23, 707-727.	2.2	22
20	Economic crisis and social capital: The story of two rural fishing communities. Journal of Occupational and Organizational Psychology, 2008, 81, 319-341.	4.5	19
21	Decisions about prophylactic gynecologic surgery: a qualitative study of the experience of female Lynch syndrome mutation carriers. Hereditary Cancer in Clinical Practice, 2015, 13, 10.	1.5	19
22	Psychological Aspects of Predictive Genetic-Test Decisions: What Do We Know So Far?. Analyses of Social Issues and Public Policy, 2004, 4, 13-31.	1.7	18
23	Benefits and burdens of newborn screening: public understanding and decision-making. Personalized Medicine, 2014, 11, 593-607.	1.5	17
24	Interest in Newborn Genetic Testing: A Survey of Prospective Parents and the General Public. Genetic Testing and Molecular Biomarkers, 2012, 16, 353-358.	0.7	16
25	Engaging patients in health research: identifying research priorities through community town halls. BMC Health Services Research, 2017, 17, 192.	2.2	14
26	Giving patients a voice: a participatory evaluation of patient engagement in Newfoundland and Labrador Health Research. Research Involvement and Engagement, 2020, 6, 39.	2.9	14
27	Genetic Testing for Huntington's Disease: How Is the Decision Taken?. Genetic Testing and Molecular Biomarkers, 2006, 10, 60-67.	1.7	13
28	â€˜It had to be doneâ€™: genetic testing decisions for arrhythmogenic right ventricular cardiomyopathy. Clinical Genetics, 2015, 88, 344-351.	2.0	13
29	Stigma and Genetic Risk: Perceptions of Stigma among Those at Risk for Huntington Disease (HD)âˆ—. Qualitative Research in Psychology, 2007, 4, 65-84.	17.6	12
30	Attitudes to incorporating genomic risk assessments into population screening programs: the importance of purpose, context and deliberation. BMC Medical Genomics, 2016, 9, 25.	1.5	12
31	Quality of life following prophylactic gynecological surgery: experiences of female Lynch mutation carriers. Familial Cancer, 2018, 17, 53-61.	1.9	12
32	The Role of Patient Advisory Councils in Health Research: Lessons From Two Provincial Councils in Canada. Journal of Patient Experience, 2020, 7, 898-905.	0.9	11
33	Group plus â€œminiâ€•individual pre-test genetic counselling sessions for hereditary cancer shorten provider time and improve patient satisfaction. Hereditary Cancer in Clinical Practice, 2020, 18, 3.	1.5	11
34	The Medical Home in Canada: Patient Perceptions of Quality and Safety. Healthcare Management Forum, 2009, 22, 47-51.	1.4	10
35	â€˜I put it on the back burner most daysâ€™: Living with chronic risk. Health (United Kingdom), 2011, 15, 633-649.	1.5	10
36	Using theatre as an arts-based knowledge translation strategy for health-related information: a scoping review protocol. BMJ Open, 2019, 9, e032738.	1.9	10

#	Article	IF	Citations
37	What do we really know about the appropriateness of radiation emitting imaging for low back pain in primary and emergency care? A systematic review and meta-analysis of medical record reviews. PLoS ONE, 2019, 14, e0225414.	2.5	10
38	Universal tumor screening for Lynch syndrome: perspectives of Canadian pathologists and genetic counselors. Journal of Community Genetics, 2019, 10, 335-344.	1.2	10
39	Are general practitioners referring patients with low back pain for CTs appropriately according to the guidelines: a retrospective review of 3609 medical records in Newfoundland using routinely collected data. BMC Family Practice, 2020, 21, 236.	2.9	10
40	Identifying Aspects of Public Attitudes Toward Whole Genome Sequencing to Inform the Integration of Genomics into Care. Public Health Genomics, 2021, 24, 229-240.	1.0	10
41	Cultural differences in family communication about inherited cancer: implications for cancer genetics research. Journal of Cultural Diversity, 2013, 20, 195-201.	0.6	10
42	Education and Parental Involvement in Decisionâ€Making About Newborn Screening: Understanding Goals to Clarify Content. Journal of Genetic Counseling, 2015, 24, 400-408.	1.6	9
43	Universal tumor screening for Lynch syndrome: Perceptions of Canadian pathologists and genetic counselors of barriers and facilitators. Cancer Medicine, 2019, 8, 3614-3622.	2.8	9
44	Use of natural health products in children: qualitative analysis of parents' experiences. Canadian Family Physician, 2013, 59, e372-8.	0.4	9
45	Public interest in whole genome sequencing and information needs: an online survey study. Personalized Medicine, 2020, 17, 283-293.	1.5	8
46	Coping with Genetic Risk: Living with Huntington Disease (HD). Current Psychology, 2009, 28, 284-301.	2.8	7
47	Bringing personalized medicine to the community through public engagement. Personalized Medicine, 2013, 10, 647-659.	1.5	7
48	Surgical treatment choices for breast cancer in Newfoundland and Labrador: a retrospective cohort study. Canadian Journal of Surgery, 2018, 61, 377-384.	1.2	7
49	Breast cancer surgical treatment choices in Newfoundland and Labrador, Canada: Patient and surgeon perspectives. Journal of Public Health Research, 2017, 6, 867.	1.2	6
50	Mothers' experiences of breastfeeding a child with tongueâ€tie. Maternal and Child Nutrition, 2021, 17, e13115.	3.0	6
51	Operationalizing a patient engagement plan for health research: Sharing a codesigned planning template from a national clinical trial. Health Expectations, 2022, 25, 697-711.	2.6	6
52	Canadians' Representation of Chemical, Biological, Radiological, Nuclear, and Explosive (CBRNE) Terrorism: A Content Analysis. Human and Ecological Risk Assessment (HERA), 2008, 14, 479-494.	3.4	5
53	Perceived economic burden associated with an inherited cardiac condition: a qualitative inquiry with families affected by arrhythmogenic right ventricular cardiomyopathy. Genetics in Medicine, 2016, 18, 584-592.	2.4	5
54	Sociodemographics and their impacts on risk factor awareness and beliefs about cancer and screening: results from a cross-sectional study in Newfoundland and Labrador. BMC Public Health, 2020, 20, 1513.	2.9	5

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55	Public engagement with genomic medicine: a summary of town hall discussions. Journal of Community Genetics, 2021, 12, 27-35.	1.2	4
56	Ethical, Legal, and Social Issues (ELSI) in Clinical Genetics Research. Methods in Molecular Biology, 2021, 2249, 65-82.	0.9	4
57	Changing Health-Related Behaviors 1: Patient-Oriented Research and Patient Engagement in Health Research. Methods in Molecular Biology, 2021, 2249, 537-552.	0.9	4
58	What behaviour change techniques have been used to improve adherence to evidence-based low back pain imaging?. Implementation Science, 2021, 16, 68.	6.9	4
59	Is genetic makeup a perceived health risk: analysis of a national survey of Canadians. Journal of Risk Research, 2009, 12, 223-237.	2.6	3
60	Universal tumor screening for lynch syndrome: perspectives of patients regarding willingness and informed consent. Personalized Medicine, 2020, 17, 373-387.	1.5	3
61	Patient education materials for non-specific low back pain and sciatica: a protocol for a systematic review and meta-analysis. BMJ Open, 2020, 10, e039530.	1.9	3
62	Clinical Genetic Research 3: Genetics ELSI (Ethical, Legal, and Social Issues) Research. Methods in Molecular Biology, 2015, 1281, 369-382.	0.9	3
63	What is in a Name? Parent, Professional and Policy-Maker Conceptions of Consent-Related Language in the Context of Newborn Screening. Public Health Ethics, 2019, 12, 158-175.	1.0	2
64	The impact of emergency department triage on the treatment outcomes of cancer patients with febrile neutropenia: A retrospective review. International Emergency Nursing, 2020, 51, 100888.	1.5	2
65	Family physician referral rates for lumbar spine computed tomography in Newfoundland and Labrador: a cross-sectional analysis using routinely collected data. CMAJ Open, 2020, 8, E56-E59.	2.4	1
66	Psychological Distress and Quality of Life in Participants Undergoing Genetic Testing for Arrhythmogenic Right Ventricular Cardiomyopathy Caused by <i>TMEM43</i> p.S358L: Is It Time to Offer Population-Based Genetic Screening?. Public Health Genomics, 2021, 24, 253-260.	1.0	1
67	Protocol for stepped wedge cluster randomized trial to evaluate the effectiveness of SurgeCon: A quality improvement emergency department surge management platform (Preprint). JMIR Research Protocols, 2022, 11, e30454.	1.0	1
68	Public interest in unexpected genomic findings: a survey study identifying aspects of sequencing attitudes that influence preferences. Journal of Community Genetics, 2022, 13, 235-245.	1.2	1
69	Implementation of an ED surge management platform: a study protocol. Implementation Science Communications, 2022, 3, 21.	2.2	1
70	Bringing the Social into Genetics: The Psychosocial Genetics Risk Assessment and Management Framework (PG-RAM). Current Psychology, 2010, 29, 171-187.	2.8	0
71	â€œThere are days I wish it wasnâ€™t there, and thereâ€™s days I realize Iâ€™m luckyâ€ A qualitative study of psychological sequelae to the implantable cardioverter defibrillator as a treatment for the prevention of sudden cardiac death in arrhythmogenic right ventricular cardiomyopathy. JRSM Cardiovascular Disease, 2017, 6, 204800401769861.	0.7	0
72	Evaluating the impact of a champion on implementation of the Back Skills Training (BeST) programme in Canada: a mixed methods feasibility study protocol. BMJ Open, 2020, 10, e040834.	1.9	0

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73	Title is missing!. , 2019, 14, e0225414.		0
74	Title is missing!. , 2019, 14, e0225414.		0
75	Title is missing!. , 2019, 14, e0225414.		0
76	Title is missing!. , 2019, 14, e0225414.		0