

# 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	Protocol for stepped wedge cluster randomized trial to evaluate the effectiveness of SurgeCon: A quality improvement emergency department surge management platform (Preprint). JMIIR Research Protocols, 2022, 11, e30454.	1.0	1
2	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
3	Implementation of an ED surge management platform: a study protocol. Implementation Science Communications, 2022, 3, 21.	2.2	1
4	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
5	Mothers' experiences of breastfeeding a child with tongue-tie. Maternal and Child Nutrition, 2021, 17, e13115.	3.0	6
6	Public engagement with genomic medicine: a summary of town hall discussions. Journal of Community Genetics, 2021, 12, 27-35.	1.2	4
7	Ethical, Legal, and Social Issues (ELSI) in Clinical Genetics Research. Methods in Molecular Biology, 2021, 2249, 65-82.	0.9	4
8	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
9	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
10	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
11	Psychological Distress and Quality of Life in Participants Undergoing Genetic Testing for Arrhythmogenic Right Ventricular Cardiomyopathy Caused by <i>TMEM43</i> : Is It Time to Offer Population-Based Genetic Screening?. Public Health Genomics, 2021, 24, 253-260.	1.0	1
12	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
13	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
14	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
15	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
16	Universal tumor screening for lynch syndrome: perspectives of patients regarding willingness and informed consent. Personalized Medicine, 2020, 17, 373-387.	1.5	3
17	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
18	Public interest in whole genome sequencing and information needs: an online survey study. Personalized Medicine, 2020, 17, 283-293.	1.5	8

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19	The impact of emergency department triage on the treatment outcomes of cancer patients with febrile neutropenia: A retrospective review. <i>International Emergency Nursing</i> , 2020, 51, 100888.	1.5	2
20	Family physician referral rates for lumbar spine computed tomography in Newfoundland and Labrador: a cross-sectional analysis using routinely collected data. <i>CMAJ Open</i> , 2020, 8, E56-E59.	2.4	1
21	Group plus individual pre-test genetic counselling sessions for hereditary cancer shorten provider time and improve patient satisfaction. <i>Hereditary Cancer in Clinical Practice</i> , 2020, 18, 3.	1.5	11
22	Evaluating the impact of a champion on implementation of the Back Skills Training (BeST) programme in Canada: a mixed methods feasibility study protocol. <i>BMJ Open</i> , 2020, 10, e040834.	1.9	0
23	What is in a Name? Parent, Professional and Policy-Maker Conceptions of Consent-Related Language in the Context of Newborn Screening. <i>Public Health Ethics</i> , 2019, 12, 158-175.	1.0	2
24	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. <i>Implementation Science</i> , 2019, 14, 49.	6.9	64
25	Universal tumor screening for Lynch syndrome: Perceptions of Canadian pathologists and genetic counselors of barriers and facilitators. <i>Cancer Medicine</i> , 2019, 8, 3614-3622.	2.8	9
26	Using theatre as an arts-based knowledge translation strategy for health-related information: a scoping review protocol. <i>BMJ Open</i> , 2019, 9, e032738.	1.9	10
27	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. <i>PLoS ONE</i> , 2019, 14, e0225414.	2.5	10
28	Universal tumor screening for Lynch syndrome: perspectives of Canadian pathologists and genetic counselors. <i>Journal of Community Genetics</i> , 2019, 10, 335-344.	1.2	10
29	Using Facebook Advertising to Recruit Representative Samples: Feasibility Assessment of a Cross-Sectional Survey. <i>Journal of Medical Internet Research</i> , 2019, 21, e14021.	4.3	67
30	Title is missing!. , 2019, 14, e0225414.		0
31	Title is missing!. , 2019, 14, e0225414.		0
32	Title is missing!. , 2019, 14, e0225414.		0
33	Title is missing!. , 2019, 14, e0225414.		0
34	Quality of life following prophylactic gynecological surgery: experiences of female Lynch mutation carriers. <i>Familial Cancer</i> , 2018, 17, 53-61.	1.9	12
35	Surgical treatment choices for breast cancer in Newfoundland and Labrador: a retrospective cohort study. <i>Canadian Journal of Surgery</i> , 2018, 61, 377-384.	1.2	7
36	“There are days I wish it wasn’t there, and there’s days I realize I’m lucky” 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. <i>JRSM Cardiovascular Disease</i> , 2017, 6, 204800401769861.	0.7	0

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37	Engaging patients in health research: identifying research priorities through community town halls. BMC Health Services Research, 2017, 17, 192.	2.2	14
38	Recruiting patients as partners in health research: a qualitative descriptive study. Research Involvement and Engagement, 2017, 3, 15.	2.9	39
39	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
40	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
41	Consent for newborn screening: parents' and health-care professionals' experiences of consent in practice. European Journal of Human Genetics, 2016, 24, 1530-1534.	2.8	29
42	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
43	Community engagement with genetics: public perceptions and expectations about genetics research. Health Expectations, 2015, 18, 1413-1425.	2.6	30
44	Education and Parental Involvement in Decision-Making About Newborn Screening: Understanding Goals to Clarify Content. Journal of Genetic Counseling, 2015, 24, 400-408.	1.6	9
45	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
46	"It had to be done": genetic testing decisions for arrhythmogenic right ventricular cardiomyopathy. Clinical Genetics, 2015, 88, 344-351.	2.0	13
47	Clinical Genetic Research 3: Genetics ELSI (Ethical, Legal, and Social Issues) Research. Methods in Molecular Biology, 2015, 1281, 369-382.	0.9	3
48	Benefits and burdens of newborn screening: public understanding and decision-making. Personalized Medicine, 2014, 11, 593-607.	1.5	17
49	Public attitudes toward genetic risk testing and its role in healthcare. Personalized Medicine, 2014, 11, 509-522.	1.5	30
50	Consulting the community: public expectations and attitudes about genetics research. European Journal of Human Genetics, 2013, 21, 1338-1343.	2.8	30
51	A family genetic risk communication framework: guiding tool development in genetics health services. Journal of Community Genetics, 2013, 4, 233-242.	1.2	38
52	Reflections on the Cost of "Low-Cost" Whole Genome Sequencing: Framing the Health Policy Debate. PLoS Biology, 2013, 11, e1001699.	5.6	67
53	Bringing personalized medicine to the community through public engagement. Personalized Medicine, 2013, 10, 647-659.	1.5	7
54	Use of natural health products in children: qualitative analysis of parents' experiences. Canadian Family Physician, 2013, 59, e372-8.	0.4	9

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55	Cultural differences in family communication about inherited cancer: implications for cancer genetics research. <i>Journal of Cultural Diversity</i> , 2013, 20, 195-201.	0.6	10
56	Personal privacy, public benefits, and biobanks: a conjoint analysis of policy priorities and public perceptions. <i>Genetics in Medicine</i> , 2012, 14, 229-235.	2.4	61
57	Interest in Newborn Genetic Testing: A Survey of Prospective Parents and the General Public. <i>Genetic Testing and Molecular Biomarkers</i> , 2012, 16, 353-358.	0.7	16
58	Public Attitudes About Genetic Testing in the Newborn Period. <i>JOGNN - Journal of Obstetric, Gynecologic, and Neonatal Nursing</i> , 2012, 41, 191-200.	0.5	28
59	“I put it on the back burner most days”: Living with chronic risk. <i>Health (United Kingdom)</i> , 2011, 15, 633-649.	1.5	10
60	Healthcare experiences of families affected by Huntington disease: need for improved care. <i>Chronic Illness</i> , 2011, 7, 225-238.	1.5	28
61	Bringing the Social into Genetics: The Psychosocial Genetics Risk Assessment and Management Framework (PG-RAM). <i>Current Psychology</i> , 2010, 29, 171-187.	2.8	0
62	The Medical Home in Canada: Patient Perceptions of Quality and Safety. <i>Healthcare Management Forum</i> , 2009, 22, 47-51.	1.4	10
63	Decision-Making About Inherited Cancer Risk: Exploring Dimensions of Genetic Responsibility. <i>Journal of Genetic Counseling</i> , 2009, 18, 252-264.	1.6	32
64	Coping with Genetic Risk: Living with Huntington Disease (HD). <i>Current Psychology</i> , 2009, 28, 284-301.	2.8	7
65	Is genetic makeup a perceived health risk: analysis of a national survey of Canadians. <i>Journal of Risk Research</i> , 2009, 12, 223-237.	2.6	3
66	Economic crisis and social capital: The story of two rural fishing communities. <i>Journal of Occupational and Organizational Psychology</i> , 2008, 81, 319-341.	4.5	19
67	Exploring informed choice in the context of prenatal testing: findings from a qualitative study. <i>Health Expectations</i> , 2008, 11, 355-365.	2.6	46
68	The Influence of Experiential Knowledge on Prenatal Screening and Testing Decisions. <i>Genetic Testing and Molecular Biomarkers</i> , 2008, 12, 115-124.	1.7	64
69	Psychosocial aspects of venous thromboembolic disease: An exploratory study. <i>Thrombosis Research</i> , 2008, 122, 491-500.	1.7	22
70	Canadians' Representation of Chemical, Biological, Radiological, Nuclear, and Explosive (CBRNE) Terrorism: A Content Analysis. <i>Human and Ecological Risk Assessment (HERA)</i> , 2008, 14, 479-494.	3.4	5
71	“They had the right to know.” Genetic risk and perceptions of responsibility. <i>Psychology and Health</i> , 2008, 23, 707-727.	2.2	22
72	Stigma and Genetic Risk: Perceptions of Stigma among Those at Risk for Huntington Disease (HD)—Qualitative Research in Psychology, 2007, 4, 65-84.	17.6	12

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73	Information Processing in the Context of Genetic Risk: Implications for Genetic Risk Communication. Journal of Genetic Counseling, 2007, 16, 419-432.	1.6	39
74	Discovering the Family History of Huntington Disease (HD). Journal of Genetic Counseling, 2006, 15, 105-117.	1.6	23
75	Genetic Testing for Huntington's Disease: How Is the Decision Taken?. Genetic Testing and Molecular Biomarkers, 2006, 10, 60-67.	1.7	13
76	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