June C Carroll

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5216543/publications.pdf

Version: 2024-02-01

159585 223800 2,657 99 30 46 citations g-index h-index papers 100 100 100 3353 docs citations times ranked citing authors all docs

| # | Article | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Widening the lens of actionability: A qualitative study of primary care providers' views and experiences of managing secondary genomic findings. European Journal of Human Genetics, 2022, 30, 595-603. | 2.8 | 10 |
| 2 | Challenges and practical solutions for managing secondary genomic findings in primary care. European Journal of Medical Genetics, 2022, 65, 104384. | 1.3 | 4 |
| 3 | Genetics Adviser: a protocol for a mixed-methods randomised controlled trial evaluating a digital platform for genetics service delivery. BMJ Open, 2022, 12, e060899. | 1.9 | 10 |
| 4 | Effect of genetics clinical decision support tools on health-care providers' decision making: a mixed-methods systematic review. Genetics in Medicine, 2021, 23, 593-602. | 2.4 | 11 |
| 5 | Ensuring best practice in genomics education and evaluation: reporting item standards for education and its evaluation in genomics (RISE2 Genomics). Genetics in Medicine, 2021, 23, 1356-1365. | 2.4 | 17 |
| 6 | Anticipating the primary care role in genomic medicine: expectations of genetics health professionals. Journal of Community Genetics, 2021, 12, 559-568. | 1.2 | 4 |
| 7 | A Master Class in Family Doctor Leadership: Evaluating an Innovative Program. Family Medicine, 2021, 53, 701-707. | 0.5 | O |
| 8 | Newborn screening for cystic fibrosis: Role of primary care providers in caring for infants with positive screening results. Canadian Family Physician, 2021, 67, e144-e152. | 0.4 | 1 |
| 9 | Primary care providers' role in newborn screening result notification for cystic fibrosis. Canadian Family Physician, 2021, 67, 439-448. | 0.4 | O |
| 10 | Experiences of patients with a disability in receiving primary health care: Using experience-based design for quality improvement. Canadian Family Physician, 2021, 67, 517-524. | 0.4 | 0 |
| 11 | Primary care providers' role in newborn screening result notification for cystic fibrosis. Canadian Family Physician, 2021, 67, 439-448. | 0.4 | 4 |
| 12 | Newborn screening for cystic fibrosis. Canadian Family Physician, 2021, 67, e144-e152. | 0.4 | 7 |
| 13 | Effectiveness of the Genomics ADvISER decision aid for the selection of secondary findings from genomic sequencing: a randomized clinical trial. Genetics in Medicine, 2020, 22, 727-735. | 2.4 | 34 |
| 14 | Noninvasive Prenatal Testing for Trisomies 21, 18, and 13, Sex Chromosome Aneuploidies, and Microdeletions in Average-Risk Pregnancies: A Cost-Effectiveness Analysis. Journal of Obstetrics and Gynaecology Canada, 2020, 42, 740-749.e12. | 0.7 | 10 |
| 15 | Hereditary colorectal cancer screening: A 10-year longitudinal cohort study following an educational intervention. Preventive Medicine Reports, 2020, 20, 101189. | 1.8 | 3 |
| 16 | Bridging the gap in genetics: a progressive model for primary to specialist care. BMC Medical Education, 2019, 19, 195. | 2.4 | 17 |
| 17 | Health outcomes, utility and costs of returning incidental results from genomic sequencing in a Canadian cancer population: protocol for a mixed-methods randomised controlled trial. BMJ Open, 2019, 9, e031092. | 1.9 | 10 |
| 18 | 20â€The masterclass series in family doctor leadership: evaluation of a new approach to leadership development. , 2019, , . | | 0 |

| # | Article | IF | Citations |
|----|--|----------|-----------|
| 19 | Family History Taking in Pediatric Practice: A Qualitative Interview Study. Public Health Genomics, 2019, 22, 110-118. | 1.0 | 2 |
| 20 | Informing Integration of Genomic Medicine Into Primary Care: An Assessment of Current Practice, Attitudes, and Desired Resources. Frontiers in Genetics, 2019, 10, 1189. | 2.3 | 48 |
| 21 | Fetal Exome Sequencing on the Horizon. Journal of Obstetrics and Gynaecology Canada, 2019, 41, 64-67. | 0.7 | 6 |
| 22 | A framework to build capacity for a reflex-testing program for Lynch syndrome. Genetics in Medicine, 2019, 21, 1381-1389. | 2.4 | 11 |
| 23 | Mendelian Randomization and mediation analysis of leukocyte telomere length and risk of lung and head and neck cancers. International Journal of Epidemiology, 2019, 48, 751-766. | 1.9 | 32 |
| 24 | Primary care providers' lived experiences of genetics in practice. Journal of Community Genetics, 2019, 10, 85-93. | 1.2 | 33 |
| 25 | Genetic assessment wait time indicators in the High Risk Ontario Breast Screening Program. Molecular Genetics & Denomic Medicine, 2018, 6, 213-223. | 1.2 | 9 |
| 26 | Psychosocial Response to Uncertain Newborn Screening Results for Cystic Fibrosis. Journal of Pediatrics, 2017, 184, 165-171.e1. | 1.8 | 34 |
| 27 | False-Positive Newborn Screening for Cystic Fibrosis and Health Care Use. Pediatrics, 2017, 140, . | 2.1 | 24 |
| 28 | Universal tumor screening for Lynch syndrome: health-care providers' perspectives. Genetics in Medicine, 2017, 19, 568-574. | 2.4 | 13 |
| 29 | A secondary benefit: the reproductive impact of carrier results from newborn screening for cystic fibrosis. Genetics in Medicine, 2017, 19, 403-411. | 2.4 | 9 |
| 30 | The Role of Family Physicians in Cancer Care: Perspectives of Primary and Specialty Care Providers. Current Oncology, 2017, 24, 75-80. | 2.2 | 55 |
| 31 | Assessing family history of chronic disease in primary care: Prevalence, documentation, and appropriate screening. Canadian Family Physician, 2017, 63, e58-e67. | 0.4 | 15 |
| 32 | 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 |
| 33 | Parent Experience With False-Positive Newborn Screening Results for Cystic Fibrosis. Pediatrics, 2016, 138, . | 2.1 | 28 |
| 34 | RETIRED: Joint SOGC–CCMG Opinion for Reproductive Genetic Carrier Screening: An Update for All Canadian Providers of Maternity and Reproductive Healthcare in the Era of Direct-to-Consumer Testing. Journal of Obstetrics and Gynaecology Canada, 2016, 38, 742-762.e3. | 0.7 | 37 |
| 35 | Opinion commune de la SOGC et du CCGM sur le dépistage génétique en contexte de procréation : Mise à jour à l'intention de l'ensemble des prestataires canadiens de soins de santé maternelle et de services en procréation, à l'Ã"re des tests offerts directement aux consommateurs. Journal of Obstetrics and Gynaecology Canada, 2016, 38, 763-787,e4. | e 0.7 | O |
| 36 | The Gene Messenger Impact Project: An Innovative Genetics Continuing Education Strategy for Primary Care Providers. Journal of Continuing Education in the Health Professions, 2016, 36, 178-185. | 1.3 | 15 |

| # | Article | IF | CITATIONS |
|----|--|------------------------|-------------------------|
| 37 | Using Newborn Screening Bloodspots for Research: Public Preferences for Policy Options. Pediatrics, 2016, 137, . | 2.1 | 11 |
| 38 | Supporting genetics in primary care: investigating how theory can inform professional education. European Journal of Human Genetics, 2016, 24, 1541-1546. | 2.8 | 9 |
| 39 | Academic family health teams: Part 1: patient perceptions of core primary care domains. Canadian Family Physician, 2016, 62, e23-30. | 0.4 | 15 |
| 40 | Academic family health teams: Part 2: patient perceptions of access. Canadian Family Physician, 2016, 62, e31-9. | 0.4 | 13 |
| 41 | Identification and management of women with a family history of breast cancer: Practical guide for clinicians. Canadian Family Physician, 2016, 62, 799-803. | 0.4 | 8 |
| 42 | Patients' experiences with continuity of cancer care in Canada: Results from the CanIMPACT study. Canadian Family Physician, 2016, 62, 821-827. | 0.4 | 32 |
| 43 | Coordination of cancer care between family physicians and cancer specialists: Importance of communication. Canadian Family Physician, 2016, 62, e608-e615. | 0.4 | 34 |
| 44 | Primary care providers' experiences with and perceptions of personalized genomic medicine. Canadian Family Physician, 2016, 62, e626-e635. | 0.4 | 65 |
| 45 | Prenatal Diagnosis Procedures and Techniques to Obtain a Diagnostic Fetal Specimen or Tissue: Maternal and Fetal Risks and Benefits. Journal of Obstetrics and Gynaecology Canada, 2015, 37, 656-668. | 0.7 | 30 |
| 46 | Developing clinical decision tools to implement chronic disease prevention and screening in primary care: the BETTER 2 program (building on existing tools to improve chronic disease prevention and) Tj ETQq0 0 0 | rg & T9/Ove | erlo ek 10 Tf 50 |
| 47 | Public Perceptions of the Benefits and Risks of Newborn Screening. Pediatrics, 2015, 136, e413-e423. | 2.1 | 17 |
| 48 | Expectations and values about expanded newborn screening: a public engagement study. Health Expectations, 2015, 18, 419-429. | 2.6 | 23 |
| 49 | Public views on participating in newborn screening using genome sequencing. European Journal of Human Genetics, 2014, 22, 1248-1254. | 2.8 | 39 |
| 50 | Pregnancy Outcomes After Assisted Human Reproduction. Journal of Obstetrics and Gynaecology Canada, 2014, 36, 64-83. | 0.7 | 121 |
| 51 | Guideline harmonization and implementation plan for the BETTER trial: Building on Existing Tools to Improve Chronic Disease Prevention and Screening in Family Practice. CMAJ Open, 2014, 2, E1-E10. | 2.4 | 16 |
| 52 | Finding a BETTER way: A qualitative study exploring the prevention practitioner intervention to improve chronic disease prevention and screening in family practice. BMC Family Practice, 2014, 15, 66. | 2.9 | 22 |
| 53 | Effectiveness of Screening With Annual Magnetic Resonance Imaging and Mammography: Results of the Initial Screen From the Ontario High Risk Breast Screening Program. Journal of Clinical Oncology, 2014, 32, 2224-2230. | 1.6 | 106 |
| 54 | Benefits and burdens of newborn screening: public understanding and decision-making. Personalized Medicine, 2014, 11, 593-607. | 1.5 | 17 |

| # | Article | IF | CITATIONS |
|----|---|-----|-----------|
| 55 | RETIRED: Prenatal Invasive Procedures in Women With Hepatitis B, Hepatitis C, and/or Human Immunodeficiency Virus Infections. Journal of Obstetrics and Gynaecology Canada, 2014, 36, 648-653. | 0.7 | 38 |
| 56 | Development and validation of a brief screening instrument for psychosocial risk associated with genetic testing: a pan-Canadian cohort study. BMJ Open, 2013, 3, e002227. | 1.9 | 24 |
| 57 | Reflections on the Cost of "Low-Cost" Whole Genome Sequencing: Framing the Health Policy Debate. PLoS Biology, 2013, 11, e1001699. | 5.6 | 67 |
| 58 | Primary care role in expanded newborn screening: After the heel prick test. Canadian Family Physician, 2013, 59, 861-8. | 0.4 | 7 |
| 59 | Family history and breast cancer. Cmaj, 2012, 184, 1391-1391. | 2.0 | 1 |
| 60 | What does the feminization of family medicine mean?. Cmaj, 2012, 184, 1752-1752. | 2.0 | 15 |
| 61 | Citizens' Values Regarding Research With Stored Samples From Newborn Screening in Canada. Pediatrics, 2012, 129, 239-247. | 2.1 | 38 |
| 62 | Health-care providers' views on pursuing reproductive benefit through newborn screening: the case of sickle cell disorders. European Journal of Human Genetics, 2012, 20, 498-504. | 2.8 | 18 |
| 63 | Applying the 2011 Canadian guidelines for breast cancer screening in practice. Cmaj, 2012, 184, 1803-1807. | 2.0 | 15 |
| 64 | Implementation of a population-based breast screening program for high-risk women in Ontario: The Ontario Breast Screening Program (OBSP) High-Risk Program. Journal of Clinical Oncology, 2012, 30, e12034-e12034. | 1.6 | 0 |
| 65 | Effective interventions to facilitate the uptake of breast, cervical and colorectal cancer screening: an implementation guideline. Implementation Science, 2011, 6, 112. | 6.9 | 57 |
| 66 | RETIRED: Fetal and Perinatal Autopsy in Prenatally Diagnosed Fetal Abnormalities With Normal Karyotype. Journal of Obstetrics and Gynaecology Canada, 2011, 33, 1047-1057. | 0.7 | 25 |
| 67 | What implementation interventions increase cancer screening rates? a systematic review. Implementation Science, 2011, 6, 111. | 6.9 | 113 |
| 68 | GenetiKit: a randomized controlled trial to enhance delivery of genetics services by family physicians. Family Practice, 2011, 28, 615-623. | 1.9 | 39 |
| 69 | Understanding sickle cell carrier status identified through newborn screening: a qualitative study. European Journal of Human Genetics, 2010, 18, 303-308. | 2.8 | 23 |
| 70 | The primary care physician role in cancer genetics: a qualitative study of patient experience. Family Practice, 2010, 27, 563-569. | 1.9 | 27 |
| 71 | Informing Parents About Expanded Newborn Screening: Influences on Provider Involvement. Pediatrics, 2009, 124, 950-958. | 2.1 | 41 |
| 72 | Clinical obligations and public health programmes: healthcare provider reasoning about managing the incidental results of newborn screening. Journal of Medical Ethics, 2009, 35, 626-634. | 1.8 | 23 |

| # | Article | lF | CITATIONS |
|----|---|-----|-----------|
| 73 | Decisionâ€Making About Inherited Cancer Risk: Exploring Dimensions of Genetic Responsibility. Journal of Genetic Counseling, 2009, 18, 252-264. | 1.6 | 32 |
| 74 | A demonstration project of a multi-component educational intervention to improve integrated post-fracture osteoporosis care in five rural communities in Ontario, Canada. Osteoporosis International, 2009, 20, 265-274. | 3.1 | 31 |
| 75 | The current state of cancer family history collection tools in primary care: a systematic review. Genetics in Medicine, 2009, 11, 495-506. | 2.4 | 85 |
| 76 | Systematic review: family history in risk assessment for common diseases. Annals of Internal Medicine, 2009, 151, 878-85. | 3.9 | 67 |
| 77 | Genetics: hereditary hemochromatosis. Canadian Family Physician, 2009, 55, 53. | 0.4 | 1 |
| 78 | Genetic education for primary care providers: improving attitudes, knowledge, and confidence. Canadian Family Physician, 2009, 55, e92-9. | 0.4 | 69 |
| 79 | Interactive Genetic Counseling Roleâ€Play: A Novel Educational Strategy for Family Physicians. Journal of Genetic Counseling, 2008, 17, 189-195. | 1.6 | 14 |
| 80 | Genetics education in medical school: a qualitative study exploring educational experiences and needs. Medical Teacher, 2008, 30, 192-198. | 1.8 | 33 |
| 81 | The Helix in the Labyrinth: Do We Need Genetic Health Services and Policy Research?. Healthcare Policy, 2008, 4, 30-38. | 0.6 | 1 |
| 82 | Hereditary breast and ovarian cancers. Canadian Family Physician, 2008, 54, 1691-2. | 0.4 | 24 |
| 83 | Development of an integrated-care delivery model for post-fracture care in Ontario, Canada. Osteoporosis International, 2006, 17, 1337-1345. | 3.1 | 16 |
| 84 | The Use of the Antenatal Psychosocial Health Assessment (ALPHA) Tool in the Detection of Psychosocial Risk Factors for Postpartum Depression: A Randomized Controlled Trial. Journal of Obstetrics and Gynaecology Canada, 2006, 28, 873-878. | 0.7 | 44 |
| 85 | Screening for Asymptomatic Bacteriuria in Pregnancy. Journal of Obstetrics and Gynaecology Canada, 2005, 27, 20-24. | 0.7 | 50 |
| 86 | Effectiveness of the Antenatal Psychosocial Health Assessment (ALPHA) form in detecting psychosocial concerns: a randomized controlled trial. Cmaj, 2005, 173, 253-259. | 2.0 | 104 |
| 87 | The Educational Needs and Professional Roles of Canadian Physicians and Nurses regarding Genetic Testing and Adult Onset Hereditary Disease. Public Health Genomics, 2005, 8, 80-87. | 1.0 | 46 |
| 88 | Risks and benefits of population-based genetic testing for Mendelian subsets of common diseases were examined using the example of colorectal cancer risk. Journal of Clinical Epidemiology, 2005, 58, 934-941. | 5.0 | 5 |
| 89 | Canadian Nursing in the Genomic Era: A Call for Leadership. Canadian Journal of Nursing Leadership, 2005, 18, 56-72. | 1.0 | 2 |
| 90 | Information needs in the management of osteoporosis in family practice: an illustration of the failure of the current guideline implementation process. Osteoporosis International, 2003, 14, 672-676. | 3.1 | 93 |

| # | Article | IF | CITATIONS |
|----|---|-----|-----------|
| 91 | Prenatal HIV Testing in Ontario. Canadian Journal of Public Health, 2003, 94, 93-97. | 2.3 | 13 |
| 92 | Genetic susceptibility to cancer. Family physicians' experience. Canadian Family Physician, 2003, 49, 45-52. | 0.4 | 52 |
| 93 | Educating women about breast cancer. An intervention for women with a family history of breast cancer. Canadian Family Physician, 2003, 49, 56-63. | 0.4 | 13 |
| 94 | How are family physicians managing osteoporosis? Qualitative study of their experiences and educational needs. Canadian Family Physician, 2003, 49, 462-8. | 0.4 | 60 |
| 95 | From Research to Application: The Development of an Antenatal Psychosocial Health Assessment Tool. Canadian Journal of Public Health, 2002, 93, 291-296. | 2.3 | 4 |
| 96 | Women's decision-making about their health care: views over the life cycle. Patient Education and Counseling, 2002, 48, 225-231. | 2.2 | 59 |
| 97 | Changing physicians' attitudes toward self-help groups: an educational intervention. Journal of Cancer Education, 2000, 15, 14-8. | 1.3 | 8 |
| 98 | Pilot study of an information aid for women with a family history of breast cancer. Health Expectations, 1999, 2, 118-128. | 2.6 | 11 |
| 99 | Cancer Self-Help Groups and Family Physicians. Cancer Practice, 1999, 7, 10-15. | 0.7 | 16 |