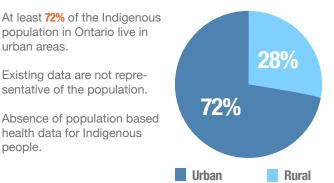
Our Health Counts Toronto

An inclusive community-driven health survey for Indigenous peoples in Toronto

Project Overview & Methods

There is a critical and alarming gap in high quality, comprehensive, and inclusive data for urban Indigenous populations in Canada. Such limitations are compounded by system barriers and colonial processes. These include the lack of culturally-based, Indigenous-led and specific measures and health information systems that prevent and exclude Indigenous people from governing, managing, and leading their own research and data processes.¹ Our Health Counts (OHC) aims to address the health information gap and ensure that urban Indigenous communities have ownership, access, control, and possession of data that impacts their health and wellbeing.

Why Our Health Counts Toronto?



How? Work in partnership to develop urban Indigenous population-based health status and health care utilization dataset.

Innovative Methods

1. Community Based Participatory Research Partnerships

2. Respectful Health Assessment Survey

3. Respondent Driven Sampling Methodologies

4. Data Linkage to the Institute for Clinical Evaluative Sciences

The OHC model recognizes that Indigenous community leadership and investment are essential for successful health programming and services for Indigenous individuals, families and communities.

OHC project processes is structured to ensure respect, cultural relevance, mutual capacity building, representation, and sustainability.

What is **Respondent Driven Sampling**? Respondent driven sampling (RDS) is a chain-referral technique that is recognized internationally by scientists as a cutting edge method of gathering reliable information from hard-to-reach populations. RDS was selected for OHC because it builds on the existing strength of social networks and kin systems known to be in Indigenous communities. RDS allows for the generation of unbiased estimates of a population's composition by adjusting for different probabilities of being sampled and by use of a structured recruitment frame. ²⁻⁴

♦ ● ♦ ● Our Health Counts: Community health assessment by the people, for the people ● ♦ ● ♦ ● ♦

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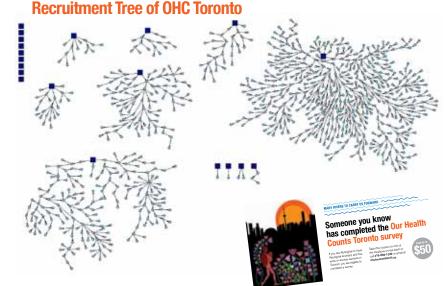
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Recruitment: The RDS process began through the careful selection of individuals to begin recruitment, also called 'seeds'. To participate in the study, people needed to self-identify as Indigenous, be 18 years of age or older, and reside within the geographic boundaries or use services within the City of Toronto. Study par5cipants, including the 20 seeds, received a coupon to participate, provided informed consent and then completed a health assessment survey. Participants then received 3-5 coupons to refer people from their social networks to participate, expanding through successive 'waves' of peer recruitment.

Recruitment Dynamics

Among Indigenous adults in Toronto, 88% of participants were recruited through referral trees originating from 3 seeds. This is consistent with other RDS studies. RDS methods are structured to overcome sampling bias. This is usually achieved when recruitment chains are 6 or 7 waves long. The two longest waves in our sample were 19 and 16 waves in length, indica5ng our sample is statistically robust.



Our Health Counts Impact

OHC has successfully implemented an Indigenous-led health information database system to gather urban Indigenous health information across four diverse urban areas in Ontario, Canada^{1,5} OHC is built on Indigenous values, skills, knowledge, beliefs and practices while also balancing power relationships to promote individual and community self-determination of health information. This system has effectively bridged Indigenous practices into Western public health systems, through the maintenance of epidemiologic rigor using RDS methods, building on existing knowledge, social networks, and kin systems within Indigenous communities. The OHC model also demonstrates scalability across diverse urban contexts and community-relevant policy applications, suggesting that OHC could provide a model for the gathering and governance of data for other Indigenous communities.



Funding was provided by the Canadian Institute of Health Research (CIHR) and Ministry of Health and Long-Term Care (MOHLTC) Capacity Award Authors: © Seventh Generation Midwives Toronto. This work is licensed under the Creative Commons Attribution-NonCommercial-NoDerivatives 4.0 International License. To view a copy of this license, visit http://creativecommons.org/licenses/by-nc/4.0/ or send a letter to Creative Commons, PO Box 1866, Mountain View, CA 94042, USA.

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Adult Demographics

The impacts of colonization and colonial policies, such as the Indian Act, residential schools, the Sixes Scoop, and continued exclusion of Indigenous people from the Canadian economy are reflected in the higher rates of unemployment and lower socioeconomic status.1 Research has shown that Indigenous people are undercounted by the national census ^{2,3,4} and that Statistics Canada has significantly underestimated the prevalence of poverty among urban Indigenous population in Ontario.^{3,4}

Population-level Data Collection

Only 14% of Indigenous adults in Toronto completed the 2011 Census. To obtain a representative sample, 70% of households should have completed the Census. Only 16% completed the 2011 National Household Survey (NHS).

OHC Toronto study findings indicate that there are **45,000-60,000** Indigenous adults in Toronto. This is 3-4 times more than estimated by Statistics Canada. (The 2011 NHS estimates that 15,650 Indigenous adults live in Toronto

Identity Age

