# Understanding Research Experiences Among Persons Identifying as African, Caribbean, and/or Black in British Columbia

CONFERENCE
CAHR
L'ACRV
2.:21
VIRTUAL
May 5-7, 2021
CON GRÈS DE
L'ACRV
VIRTUEL
5 au 7 mai 2021

<u>Tsion Gebremedhen</u>, Patience Magagula, Amber R Campbell, Rebecca Gormley, Evelyn J Maan, Helene CF Cote, Melanie CM Murray, Angela Kaida

As uninvited settlers, we acknowledge that we conducted our study on the stolen traditional and ancestral lands of the Coast Salish peoples, including the Skwxwú7mesh (Squamish), Stó:lō, Səʾlílwəta?/Selilwitulh (Tsleil-Waututh) and xwməθkwəyəm (Musqueam).

We would like to thank all of the survey respondents and our community partners, without whom this project would not be possible!

Email: tsion.gebremedhen@gmail.com
I have no conflicts of interest to disclose

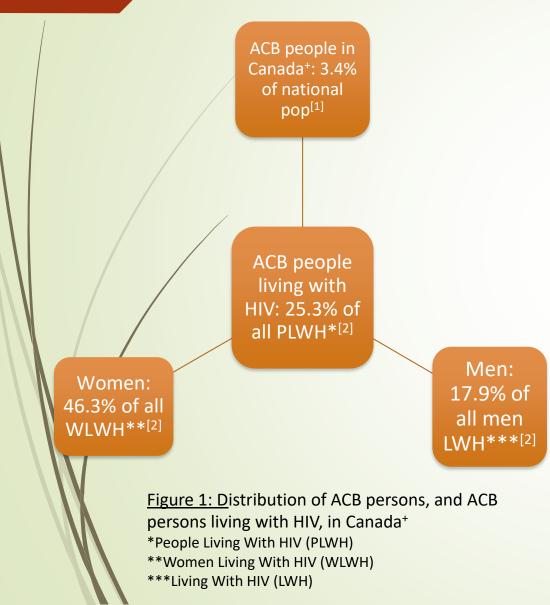








# INTRODUCTION



- African, Caribbean and/or Black (ACB) persons are underrepresented in health research in Canada<sup>+</sup>
- Local leaders suggest lack of research engagement can be due to:
  - Lack of trust of research & researchers
  - Stigma with HIV research
  - Unfamiliarity with research & methodology
- Study Aims:
  - To seek input on research participation experiences;
  - To establish future research priorities with ACB persons;
  - To encourage researchers to adopt anti-racist practices

\*The name 'Canada' is a colonially imposed name given to the stolen unceded and traditional lands of the many Indigenous Peoples who have resided here for millenia.

# **METHODS**

Please share your thoughts and experiences related to research with us through a confidential survey, if: You identify as African, Caribbean and/or Black You are 16 years of age or older You live in British Columbia Complete survey for a chance to win a 100\$ gift card!

<u>Figure 2:</u> Recruitment Poster for Social Media (ie: Facebook, Twitter)

Created online community survey with key community partners

Collaborated with various ACB community organizations to support survey dissemination and share recruitment material (ie: posters for social media)

Participant Recruitment; Inclusion Criteria:

Identify as ACB (including mixed race);

Be 16 years of age or older;

Currently live in BC

Survey open from October 14 - December 9, 2020

Prize Draw for \$100 CAD gift card for all participants

# RESULTS

- Respondent Demographics (n=56):
  - Ethnicity: 50% African (n=28); 30% Caribbean (n=17); 11% Black (n=6); 27% mixed race/multiracial (n=15)
  - Birth Country: 66.1% outside so-called "Canada" (n=37); 33.9% (n=19) in so-called "Canada"
  - Age: 50% <25 years (n=28)</p>
  - Gender: 79% women (n=44); 20% men (n=11); 2% gender queer (n=1)

<u>Table 1</u>: Research study types respondents previously participated in (n=25):

| Survey/Questionnaire                | 22 (91.7%) |
|-------------------------------------|------------|
| 1-on-1 Interview                    | 10 (41.7%) |
| Focus Group                         | 7 (29.2%)  |
| Clinical Trial                      | 3 (12.5%)  |
| Studies including sample collection | 3 (12.5%)  |

<u>Tables 2, 3</u>: Principal facilitators & barriers to research participation

# Principal reasons respondents would participate in health research in the future (n=56):

|                                    | •   |
|------------------------------------|---|
|                                    | 47 (83.9%)  |
| Benefit to me and/or my community  |   |
| Good way to share my voice &       | 34 (60.7%)  |
| perspectives                       |   |
|                                    | 28 (50%)  |
| Opportunity to learn something new |   |
|                                    | 23 (41.1%)  |
| Receiving compensation for my time |   |
| I know the research is done in     | 20 (35.7%)  |
| collaboration with the community   |   |
|                                    | Opportunity to learn something new  Receiving compensation for my time I know the research is done in |

| Principal reasons respondents would NOT participate in health |            |  |
|---|------------|--|
| research in the future (n=56):                                |            |  |
| I don't have the time/I have too many other                   | 26 (46.4%) |  |
| responsibilities  |            |  |
|   | 15 (26.8%) |  |
| I find it hard to trust researchers                           |            |  |
|   | 13 (23.2%) |  |
| I don't see how research benefits me/my community             |            |  |
| I don't have anything valuable to contribute to               | 13 (23.2%) |  |
| research  |            |  |
|   | 13 (23.2%) |  |
| I don't fully understand the consent process                  |            |  |
|   | 11 (19.6%) |  |
| I don't see the value in research                             |            |  |

# **RESULTS & NEXT STEPS**



<u>Figure 3:</u> Willingness to participate in HIV research, whether or not respondent is living with HIV

<u>Figure 4:</u> Ways to learn about health topics, such as COVID-19, HIV, sexual & reproductive health

## **Key Takeaways**

- Build trusting, meaningful relationships with community
- Include ACB persons at all stages of a research study
- Adopt & adapt culturally sensitive study methodology

  High willingness of respondents to participate in research

## Top 2 Future Research Priorities:

- 1. Mental health and substance use
- 2. Sexual and reproductive health, childbirth & family planning