Knowledge of HPV/cervical cancer and acceptability of HPV self-sampling among women living with HIV: A scoping review

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ABSTRACT

Cervical cancer rates are disproportionately high among women living with the human immunodeficiency virus (WLHIV). Cervical cancer is preventable through HPV screening, regular Pap tests, and early cancer detection. Evidence indicates that HPV and cervical cancer screening are suboptimal among WLHIV, who face a myriad of access barriers. Considering that screening is an effective first-line defense to cervical cancer, we conducted a scoping review with the aim of gaining a better understanding about: (1) the knowledge and perceptions of HPV and cervical cancer screening among WLHIV; and (2) the acceptability of self-sampling for HPV among WLHIV. We searched five electronic databases for peer-reviewed articles that were published in English within the last ten years, reported on studies with HIV-positive women who were aged 16 or older, and satisfied the topics of the review. A total of 621 articles were found. After accounting for duplicates and unmet criteria, 17 articles and 1 abstract, reporting on studies in the United States and Africa, were included in this review. The review highlighted that most WLHIV had inadequate knowledge of HPV transmission and cervical cancer prevention, which influenced their perceptions of risk and susceptibility. Screening barriers included misconceptions about Pap tests, fear of diagnosis of serious illness, perceived pain, embarrassment, bodily modesty, and limited access to female health care providers. This review also affirms that self-sampling is an acceptable and promising screening option for WLHIV. Implications for policy, research, and practice are discussed.

Key Words Women living with HIV, HPV, cervical cancer, self-sampling, scoping review

INTRODUCTION

Women account for a growing proportion of HIV cases in Canada and elsewhere. Between 1985 and 2014, a cumulative total of 80,469 HIV cases were reported in Canada, with women accounting for 18.2% of all cases. Of the 2,044 HIV cases reported in 2014, women accounted for 24.6%. Further, over three-quarters of these new cases were identified as Aboriginal women (30.6%), Black women (35.6%), and other women of colour (9.6%). Heterosexual contact was identified as the key category of exposure.

With the widespread use of highly active antiretroviral therapy (HAART), the incidence of HIV-defining cancers such as Kaposi sarcoma and non-Hodgkin lymphoma have declined dramatically, but cervical cancer rates have remained high among women living with HIV (WLHIV). Cohort studies in Canada and the United States have consistently shown that the risk of invasive cervical cancer is higher among WLHIV than HIV-negative women. Other studies have confirmed that the prevalence of high risk human papillomavirus (hrHPV) infection is also higher and more persistent in WLHIV. There is general agreement that a long history of HIV infection and prolonged immunosuppression are associated with persistent HPV infection and invasive cervical cancer.

Current Canadian guidelines on cervical cancer screening recommend that WLHIV receive Papanicolaou (Pap) tests at the initial assessment and at six months, with an annual follow-up for women with normal results. However, cervical cancer screening remains suboptimal among WLHIV. Results of a Canadian retrospective population-based study of 2,661 WLHIV living in Ontario show that only up to half of the women had adhered to the cervical cancer screening guidelines. Similarly, results...
of another retrospective cohort study of 218WLHIV in Ottawa indicated that 42% of the participants did not undergo cervical cancer screening during the 3-year period even though 94% of them listed that they had primary care providers in their medical records.11

Studies in Canada have identified myriad barriers that deter WLHIV from accessing health care, including structural racism, HIV-related stigma and discrimination within and outside of the health care system, and criminalization of HIV non-disclosure. Other studies show that the utilization of health services and the uptake of cancer screening among marginalized groups are associated with their health literacy. These findings suggest that socially inclusive, innovative, and relevant strategies are needed to promote HPV and cervical cancer screening among WLHIV. Insights from studies on HIV self-testing suggest that HPV self-sampling is a potential strategy to effectively engage marginalized WLHIV.

Drawing on a critical health-literacy framework, we conducted a scoping review to explore two questions: (1) What has been reported on the knowledge of HPV and/or cervical cancer screening among WLHIV in Canada and other countries? (2) What has been reported on the acceptability of HPV self-sampling among WLHIV in Canada and other countries?

METHODS

This scoping review is informed by the framework of Arksey and O’Malley. We applied a multi-step approach that included the following: (1) identifying our search topics and questions; (2) searching for relevant studies; (3) selecting relevant studies based on the review questions; (4) charting the data collected; and (5) synthesizing the data collected into themes. Further, we adopted an additional step proposed by Levac and her colleagues, that is, connecting the meaning of the findings to the overall study purpose and discussing the implications for future research, policy, and practice.

Search Strategies and Inclusion Criteria

Our search focused on peer-reviewed articles that report on either primary or secondary sources of evidence. We searched five databases: PubMed, CINAHL, OVID/Medline, ProQuest, and Google Scholar. The search terms included acceptability, attitudes, perceptions, knowledge, human papillomavirus/HPV, cervical cancer screening, self-sampling/self-collection/self-testing/self-screening, home-based collection, human immunodeficiency virus/HIV, WLH, WLHIV, and WLHA. Inclusion criteria consisted of (1) peer-reviewed English language publications within the last ten years, (2) engaged women, aged 16 or older, living with HIV in Canada or elsewhere, and (3) the researched topics fitted with the two foci of this scoping, review, i.e., knowledge of HPV and cervical cancer, and the acceptability of HPV self-sampling. We limited our search to a ten-year period to coincide with the establishment of the Screening Performance Indicators Working Group (spiwg) in 2007. The spiwg defines core performance indicators for cervical cancer screening programs in Canada and facilitates inter-jurisdictional comparison. A total of 621 articles were retrieved from the five databases. After accounting for duplicates (n=150) and unmet criteria (n=453), a total of 603 articles were excluded (see Figure 1). A table was used to identify pertinent study information on study authors, location of study, and study characteristics/population (see Table I). Each item included in this review was reviewed in its entirety, and relevant information was extracted to inform this synthesized review.

RESULTS

The results of this review show that there is a paucity of research on WLHIV’s knowledge on HPV and cervical cancer screening, and their acceptability of self-sampling. A total of 17 articles and 1 abstract met the inclusion criteria. We included 1 published abstract of a relevant study that was informative but had not yet been published as an article or report. Ten of the articles focused on knowledge of HPV and cervical cancer screening among WLHIV, but 3 of these 10 articles were from the same research study reporting on different aspects of the study results. The
<table>
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<tr>
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<tr>
<td>(1) Knowledge of HPV and cervical cancer among WLHIV</td>
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<tr>
<td>Kenya, Carrasquillo, Fatil, et al., 2015</td>
<td>United States</td>
<td>This study explored education needs regarding HPV and cervical cancer screening and treatment amongst self-identified Haitian women living with HIV.</td>
<td>Women living with HIV in South Florida, self-identified as Haitian, have not had their uterus removed, 30 to 60 years old, attending HIV clinics (N=21)</td>
<td>Qualitative: focus group interviews; community-based participatory research methods</td>
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<td>Wigfall, Bynum, Brandt, et al., 2015</td>
<td>United States</td>
<td>This study investigated knowledge of cervical cancer screening guidelines and abnormal Pap test experiences amongst women living with HIV/AIDS.</td>
<td>Women living with HIV, 20 to 68 years old, attending funded clinics and AIDS service organizations in southeastern United States, no history of cervical cancer, no complete hysterectomy, all observational data collected (N=106)</td>
<td>Quantitative: in-person standardized interviews; cross-sectional study</td>
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<td>Wigfall, Bynum, Friedman, et al., 2017</td>
<td>United States</td>
<td>This study examined communication of abnormal Pap test results between women living with HIV and their health care providers.</td>
<td>Women living with HIV, 18+ years of age, attending funded clinics and AIDS service organizations in southeastern United States, have had an abnormal Pap test (N=100)</td>
<td>Quantitative: interviewer-administered online survey; cross-sectional study</td>
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<td>United States</td>
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<td>This study explored the relationship between sociocultural/structural factors and cervical cancer screening amongst HIV-positive African American women.</td>
<td>HIV-positive African American women, 28 to 62 years old, living in Alabama (N=20)</td>
<td>Qualitative: in-depth interviews</td>
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<td>Bynum, Wigfall, Brandt, Richter, Glover, Hébert, 2013</td>
<td>United States</td>
<td>This study assessed how health literacy impacts cervical cancer screening knowledge and behaviours of women living with HIV.</td>
<td>Women living with HIV, 20 to 68 years old, attending clinic and community-based settings in southeastern United States (N=145); African American (N=131; 90%); reported low health literacy (N=55; 38%)</td>
<td>Quantitative: multi-item, self-report questionnaire</td>
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<tr>
<td>Chapman, Lambert, Chandler, McMillan, Kromrey J, Johnson-Mallard, Kurtyka, 2015</td>
<td>United States</td>
<td>This study evaluated the relationships between Pap test adherence and HPV and cervical cancer knowledge, perceptions of risk, seriousness, barriers, benefits, self-efficacy using HBM</td>
<td>Women living with HIV, 18+ years of age, and attending one of the two HIV ambulatory care clinics (N=300). Exclusion criteria: history of having had a hysterectomy, women who could not read and comprehend English</td>
<td>Quantitative: exploratory, cross-sectional, correlational design</td>
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<tr>
<td>Maree and Moitse, 2014</td>
<td>South Africa</td>
<td>This study explored knowledge of cervical cancer and cervical screening amongst women being treated for HIV/AIDS at a specific health care centre.</td>
<td>Women living with HIV/AIDS, 18+ years of age, currently receiving antiretroviral therapy (ART), attending clinic for follow-up care (N=315)</td>
<td>Quantitative: survey design, self-administered questionnaire</td>
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<tr>
<td>Rosser, Njoroge, and Huchko, 2015</td>
<td>Kenya</td>
<td>This study assessed cervical cancer knowledge, attitudes, and screening history of women living with HIV in care at an urban clinic in Western Kenya.</td>
<td>HIV-positive women, 23 to 64 old, attending an HIV clinic, non-pregnant (N=106)</td>
<td>Quantitative: survey, oral questionnaire; cross-sectional study</td>
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<tr>
<td>Koneru, Jolly, Blakemore, et al., 2017</td>
<td>Tanzania</td>
<td>This study identified barriers to cervical cancer screening and treatment amongst women living with HIV, and determined acceptability of peer navigator roles in a clinical setting.</td>
<td>Women living with HIV, 19+ years of age, attending HIV clinics in Dar es Salaam, Tanzania (N=399)</td>
<td>Quantitative: self-administered questionnaire (English or Kiswahili) with guided assistance from study staff; or interviewer-administered for women who could not read; cross-sectional study</td>
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<td>Bukirwa, Mutyoba, Mukasa et al., 2015</td>
<td>Uganda</td>
<td>This study assessed barriers and motivations associated with cervical screening uptake (acetic acid and iodine/VIA and VILI) among HIV-positive women.</td>
<td>HIV-positive women, 25+ years of age, attending an HIV clinic (N=18); women who had never screened for cervical cancer (N=6), women who screened only once (N=6), women who fully adhered to screening schedule (n=6)</td>
<td>Qualitative: in-depth interviews with HIV-positive women and key informant interviews with health care providers; cross-sectional study</td>
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<td>Murphy, Mark, Anderson, Farley, Allen, 2016</td>
<td>United States</td>
<td>This study explored whether an intervention using self-sampling of cervico-vaginal cells for human papillomavirus with results counselling would increase cervical cytology (“Pap”) testing among women with HIV.</td>
<td>HIV-positive women, 18+ years of age attending an HIV clinic for a primary care visit (N=94)</td>
<td>Quantitative: randomized controlled trial (intervention and information only groups)</td>
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<td>Jones Brudney, Sawo, Lantigua, Westhoff, 2012</td>
<td>United States</td>
<td>This study assessed the acceptability of a self-lavaging device (the Delphi Screener) for cervical cancer screening; compared women's preferences and reports on quality of experiences using the self-lavaging device compared with clinician collection of specimens during a pelvic exam for cervical cancer screening.</td>
<td>Low-income women, 18+ years of age, who had received cervical cytological testing in one of three clinics in New York City (N=198), one being the Infectious Disease Clinic (IDC), which provides HIV care services to HIV-positive women of all ages (n=53)</td>
<td>Quantitative: pre- and post-hoc questionnaire with closed- and open-ended questions regarding perceptions of the Delphi Screener (before and after completing self-collection)</td>
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<td>Lee, Ramroop, Gaydos, Barnes, Anderson, Coleman, 2017 (Abstract only)</td>
<td>United States</td>
<td>This study examined HPV prevalence, compared home self-collected and clinician-collected HPV-positivity, and measured the acceptability of self-collection</td>
<td>HIV-positive women, 89% of whom were black, in Baltimore (N=68)</td>
<td>Quantitative, prospective study</td>
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<td>Mahomed Evans, Sauls, Richter, Smith, Finhaber, 2014</td>
<td>South Africa</td>
<td>This study evaluated the acceptability of self-collection for cervical cancer screening. It aimed to identify which self-collection device women prefer and whether they would consider using it for routine screening.</td>
<td>HIV-positive women 18+ years of age attending theThembaLeTu [urban] Clinic in Johannesburg, South Africa (n=52), and the Topsy [rural] Clinic in Mpumalanga, South Africa (n=54) (N=106)</td>
<td>Quantitative method: written questionnaire with closed-ended questions</td>
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<td>Adamson Huchko, Moss, Kinkel, Medina-Marino, 2015</td>
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<td>This study aimed to estimate the prevalence of hrHPV mRNA in HIV-infected women; compared the test positivity between the two collection methods; assessed the accuracy and agreement of self-collected tampons compared with clinician-collected specimens for hrHPV mRNA testing; assessed the acceptability of the self-collected tampon method.</td>
<td>HIV-positive women, 25+ years of age, seeking care at a government HIV clinic in Pretoria, South Africa (N=325)</td>
<td>Quantitative method: hrHPV mRNA testing on the physician and self-collected samples; Likert scale questionnaire (assessing acceptability)</td>
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Acceptability of HPV self-sampling

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Self-sampling tools: (i) an Evalyn cervical brush, (ii) a Delphi Screener, and (iii) a tampon-like plastic wand.
remaining 7 articles and 1 abstract focus on acceptability of self-sampling. Together, we have identified a total of 16 studies in this review: United States (n=7); and African countries (n=9).

Part 1: Knowledge of WLHIV about HPV and cervical cancer screening

Of the eight studies (10 articles) included in the knowledge section of this review, three used qualitative methods of focus groups or individual interviews to explore knowledge and perceptions of HPV and cervical cancer screening\textsuperscript{24-26}, and five used questionnaires to test knowledge, attitudes, perceptions, and screening behaviours\textsuperscript{27-33}. A synthesized analysis of the data derived resulted in five major themes: (1) knowledge of HPV and cervical cancer screening; (2) factors influencing knowledge; (3) perceived risks and susceptibility; (4) misconceptions and barriers; and (5) screening compliance and adherence.

Knowledge of HPV and Cervical Cancer Screening

Our review results revealed that WLHIV across all regions had limited knowledge regarding HPV transmission and cervical cancer prevention. Most of them had not heard of HPV and did not know how HPV is transmitted\textsuperscript{24,29,33}. Further, many WLHIV were unaware of the risk of cervical cancer and did not know that it is preventable\textsuperscript{25,31} and that HIV infection increases the risk of HPV and cervical cancer\textsuperscript{29}.

Although some women were aware that their HIV-positive status and current sexual activities put them at risk of cervical cancer\textsuperscript{26}, and that cervical cancer screening is important for WLHIV\textsuperscript{27,32,33}, their knowledge about follow-up screening was mixed. Some recognized the need for regular Pap tests and repeated Pap tests after an abnormal test, but others thought that ‘doing nothing’ was one of the acceptable follow-up options\textsuperscript{27}. Many were unaware of the screening recommendations for WLHIV\textsuperscript{25,26}.

Factors Influencing Knowledge

Numerous individual and social factors shaped the knowledge of WLHIV regarding HPV and cervical cancer screening, including adequate and effective communication of health information from service providers\textsuperscript{28,31,33}, media messages\textsuperscript{32}, higher levels of education, previous history of cervical cancer screening, older age, being employed\textsuperscript{33}. In the case of the United States and African countries, where health care access is not universal, access to HPV screening also increased WLHIV’s knowledge\textsuperscript{33}.

Assessment of the relationship between health literacy and increased knowledge of HPV produced mixed results. One study showed that having high or low health literacy did not make any difference to the participants’ scores specific to knowledge of HPV, cervical cancer screening knowledge, and awareness of the HPV vaccine\textsuperscript{30}. However, in a study of WLHIV with a history of abnormal Pap tests, participants with a high level of health literacy were 3.5 times more likely to read information about abnormal Pap tests given by health care providers and 4.7 times more likely to have understood the information compared with those with low health literacy\textsuperscript{28}. Since these studies used quantitative methods, it was difficult to gain a better understanding about how the WLHIV’s lived experiences,
social environments, and cultural contexts shaped their access to knowledge, as well as how they made sense of HPV and HPV-related messages or materials.

**Perceived Risks and Susceptibility**

Most of the women in this review lacked a clear understanding about the risks of cervical cancer associated with HIV and HPV co-infections, which may explain their mixed and often contradictory perceptions about risk and susceptibility to cervical cancer. In an American study of 300 HIV+ women, most participants identified having multiple sex partners and early onset of sexual intercourse as risks for cervical cancer, but few knew about HPV and other risks. Women in another study conducted in the United States spoke about risks and susceptibility in the context of what they knew about HIV. Some felt that they were more susceptible to cervical cancer because they had “weak bodies” and they could catch “everything.” Other women in the same study rejected the idea that being HIV-positive increased their susceptibility. Further, while some women recognized increased susceptibility among HIV+, they did not perceive themselves to be at risk until they experienced symptoms. Others cited a lack of family history of cervical cancer and current sexual inactivity as reasons for not considering themselves at risk.

**Misconceptions and Barriers**

This review uncovered many myths and misconceptions held by HIV+ women, which might have influenced their participation in cervical cancer screening and follow-up behaviours. Over half of the HIV+ women in a Kenya-based study believed that family planning increased the risks of cervical cancer and that vaginal washing would decrease the risks. Further, many considered cervical cancer as fate or the will of God and therefore felt nothing could prevent it. Some participants in the Ugandan study did not engage in screening as they believed that service providers would remove their ovaries and uterus during the procedure.

In terms of barriers to HPV and cervical cancer screening, HIV+ women identified stigma and discrimination from health care providers, family, and friends as a key barrier. Other barriers to screening were related to psychological, sociocultural, and structural factors, including worries about finding out more health problems such as cervical cancer in addition to their HIV illnesses, embarrassment, body modesty, and fear of painful procedures. Many HIV+ women indicated their need to have female primary care providers perform the cervical cancer screening due to their cultural and religious practices. Others identified long wait time as a barrier to screening.

**Screening Compliance and Adherence**

This review also uncovered an interesting phenomenon. Many HIV+ women who received instructions from health care providers to undergo follow-up screening tended to comply. For example, in a study based in the United States, participants had received a Pap test in the previous year even though none of them understood the purpose of a Pap test. In a study of HIV+ women with a history of abnormal Pap tests, close to half of the women indicated that they did not fully understand the information given by their primary care providers, but a majority of them complied with their recommendations of getting follow-up tests and procedures. This shows that health care providers’ endorsement and recommendation for screening can influence women’s uptake.

Other studies suggest that screening adherence is associated with health literacy and social support. In the study of 399 women in an HIV clinic in Tanzania, only 9% of the participants had cervical cancer screening; 96% expressed the desire to learn more about cervical cancer screening and 87% prefer to have HIV-positive peer navigators to support them in learning and seeking screening services.

**Part 2: Acceptability of HPV Self-Sampling among Women Living with HIV**

Literature on the acceptability of HPV self-sampling among HIV+ women is also very limited; we included a total of eight studies (seven published articles and one abstract) in this section. In synthesizing the evidence from these eight studies, we identified three major themes: (1) acceptability and preference for self-sampling vs. clinician-collected sampling; (2) concerns regarding self-sampling; and (3) feasibility and effectiveness of self-sampling devices.

**Acceptability and Preference for Self-Sampling**

With the exception of one questionnaire-based study, a wide array of HPV and cervical cancer self-sampling devices were assessed for acceptability in the reviewed studies: cervico-vaginal self-lavaging devices, cervical brushes, cytobrushes, tampon-like self-collection devices, and a vaginal swab (for descriptions of these devices, see Table I).

There was a universal acceptance among HIV+ women across all studies in this review for self-sampling. Familiarity with the self-sampling device was a key factor associated with women’s willingness to use it. In one study of 325 HIV+ women in South Africa, the mini-tampon self-sampling device was used to assess the accuracy, agreement, and acceptability of self-sampled specimens versus clinician-collected specimens. The results showed that 90% of the participants reported no difficulties with self-sampling and that the instructions were clear and easy to understand, likely because over half of the women indicated familiarity in using tampons.

Other studies found that the self-lavaging device was highly accepted by participants at HIV clinics. The women in these studies indicated that they would prefer self-sampling for future cervical screening because this device felt more comfortable and less painful and was quick and easy to use. Some also identified increased privacy, decreased stress and embarrassment, and the possibility of performing the test on their own time as desirable.

Further, one-fifth of the women sampled in Rwanda preferred self-sampling over Pap tests mainly because of detesting the pelvic examination.

While all the participants remained steadfast to self-sampling being the preferred option for cervical screening, they had varying preferences regarding the location for retrieval of their self-samples. Women in a study in Kenya preferred at-home collection; women in the Ugandan...
study did not indicate any preference. However, in a study in Pretoria, only one-tenth of the women preferred to perform self-sampling at home, while close to 65% preferred clinic-based self-collection, likely because they felt better cared for in the presence of a clinician during the collection procedure.

**Concerns Regarding Self-Sampling**

Despite the wide acceptance of self-sampling among women living with HIV (WLHIV), a small proportion had concerns regarding the devices and the self-sampling procedures. Some participants in an American study indicated that a pelvic exam performed by a physician would provide them with an increased sense of security; they felt that the physician would make minimal mistakes and could immediately intervene in case an abnormality was found. Other women in the Rwandan and Kenyan studies expressed concerns about proper self-sample collection, pain, ability to insert the device, and not knowing how to interpret the results. These concerns were in stark contrast to women in a study in Uganda, who expressed confidence and self-efficacy about self-sampling at home. This contrast could possibly be related to the Advances in Screening and Prevention in Reproductive Cancers (ASPIRE) project in Uganda, which has been established since 2013 to introduce self-sampling to this region and has likely helped to normalize cervical cancer self-sampling.

**Feasibility and Effectiveness of Self-Sampling Devices**

In the context of this review, effectiveness referred to the similarity in testing scores between self-collected and physician-collected specimens, i.e., sensitivity and specificity, and feasibility was defined as the ability of participants to obtain the required amount of specimen during self-sampling without any self- or clinician-reported problems.

One study reported high specificity for cytobrush tests: the study found Pap tests were normal among 81.5% of negative home self-collected cytobrush tests. Another study that utilized mini-tampons as a self-sampling device found no difference in sensitivity between the clinician-collected samples and the self-collected samples.

Previous research reported that cytologic testing of human papillomavirus (hpv) using specimens collected through self-lavaging and those collected by clinicians had equivalent sensitivity for detecting high-grade cervical intraepithelial neoplasia.

However, effectiveness of the self-lavaging devices was not assessed in any of the studies in this review. On the other hand, the self-lavaging device used with a cohort of 60 women in the Rwandan study showed moderate feasibility.

**DISCUSSION**

We conducted this scoping review with the aim of obtaining an overall picture of WLHIV’s knowledge of HPV and cervical cancer screening, their acceptance of HPV self-sampling, and current gaps in research and knowledge. The results of this review showed both a concerning and a hopeful picture.

We found that WLHIV across all countries and regions had limited knowledge regarding HPV and HPV co-infection, cervical cancer prevention, and screening recommendations. There was also inadequate communication between WLHIV and their health care providers about screening for HPV and related cancers. Many WLHIV held misconceptions about cervical cancer screening. Although some women complied with cervical cancer screening and follow-up procedures based on their service providers’ recommendations, their lack of understanding about these tests impedes sustainable health behaviours. Further, their passive compliance reinforced their marginalization and disempowerment, which are contradictory to the principles of health promotion.

At the same time, WLHIV also reported a number of screening barriers in addition to their low HPV literacy. Women in the United States and African countries identified stigma as a barrier to access HPV and cervical cancer screening. African-American WLHIV in one American study indicated that they avoid screening due to their fear of being diagnosed with cancer or serious illnesses. These psychological challenges were likely unique to WLHIV, related to their experiences of being diagnosed HPV-positive and encountering everyday HPV-related stigma. These women and women in the Tanzanian study also indicated their need for bodily modesty and having female primary care providers perform internal exams in order to maintain their cultural and religious practice. Perhaps because of these barriers, HPV and cervical cancer self-sampling was found to be widely accepted by WLHIV in studies across countries and settings. Participants named many advantages, including flexibility and convenience in terms of time and location, privacy, and less discomfort and pain. However, some women expressed concerns about collecting proper self-samples and felt more secure about accessing tests at health care facilities.

**TABLE II** Self-sampling devices used in reviewed studies

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<td>Self-lavaging device</td>
<td>A device that a woman inserts into her vagina. It contains a syringe in which 5 mm of buffered saline was plunged into the vagina, and the fluid specimen was retrieved by pulling back on the plunger.</td>
</tr>
<tr>
<td>Cytobrushes</td>
<td>A device consisting of an extended-tip spatula with an endocervical brush, which a woman inserts into her vagina and rotates the brush at the cervix five times to collect cells.</td>
</tr>
<tr>
<td>Tampon-like plastic wand device</td>
<td>A tampon-like plastic wand device with an electable tip which a woman inserts into her vagina and rotates the device to collect cells.</td>
</tr>
<tr>
<td>Mini-sized tampon</td>
<td>A mini-size tampon that a woman inserts and leaves in her vagina for one to two hours before removing it and putting it in the specimen container.</td>
</tr>
<tr>
<td>Other devices</td>
<td>The brands and procedures for the self-sampling devices used in some studies were not specified.</td>
</tr>
</tbody>
</table>
Based on the results of this review, we offer a number of critical insights and recommendations. The concerns expressed by WLHIV have important policy and practice implications for primary health service delivery and health systems responses in Canada and across different countries. First, HIV and cancer have evolved over the past two decades into chronic health challenges as biomedical treatments have increased the survival rate of affected individuals. At the same time, we have also witnessed increased health disparities among individuals and groups who are simultaneously affected by multiple conditions, or what is known as synergistic epidemics (e.g., addiction and mental illness, diabetes and depression, HIV and Hepatitis C, HIV and HPV, etc.). It is important to note that synergistic epidemics (syndemics) are not merely concurrent physical diseases. Rather, they are health problems that are exacerbated by historical, sociocultural, economic, and political conditions, such as stigma and discrimination, access barriers to knowledge and services, uncoordinated health system responses, and lack of clear treatment guidelines or policies. In the context of screening for HPV and related cancers among WLHIV, the adoption of a syndemic approach at the policy level will inform changes at the practice levels. For example, over the past few years, the Public Health Agency of Canada has adopted an integrated approach to address the syndemics of HIV, Hepatitis C, and other sexually transmitted and blood borne infections (STBIs) in Canada. This policy change was translated into the HIV and Hepatitis C Community Action Fund program, under which research and community service organizations were steered through funding incentives towards working more collaboratively at the local, regional and national levels, as well as expanding their program foci beyond HIV to address the co-infection of STBIs among priority (marginalized) populations. However, judging by the organizations that had applied to this funding program, more deliberate efforts to promote cross-disciplinary collaboration among researchers and health care providers from the sectors of HIV and HPV-related cancer prevention is needed to develop effective responses.

Second, the framework of critical health literacy offers a potentially effective strategy to address the inadequate awareness about HIV-HPV co-infection and low screening for HPV and related cancers among WLHIV. Critical health literacy can be defined as the processes and outcomes that both increase people’s capacity to engage in a range of individual and collective actions that enhance their health, and also promote equitable access to health resources through policy change. Critical health literacy is achieved and measured at three interconnected levels. In the context of this review, the first or basic level is measured by WLHIV’s ability to access, understand, critically appraise, and then apply health information on HIV-HPV co-infection to improve or maintain their health. To support WLHIV in achieving this basic level of health literacy, accurate, socioculturally inclusive, and accessible information on HIV-HPV co-infection (e.g., fact sheets, pamphlets, quizzes, videos, stories, posters, social marketing campaigns, etc.) must be made available. At the same time, HIV-HPV care competence must also be built among other stakeholders such as HIV specialists, primary care providers, and HIV/AIDS community service providers.

At the second or intermediate level, health literacy is measured by WLHIV’s ability to navigate the health care systems to access the screening they need. We postulate that when WLHIV acquire adequate knowledge about HIV-HPV co-infection, they are more likely to take part in screening. However, knowledge alone does not address other systemic and sociocultural barriers. To support WLHIV in developing confidence and effective communication with their health care providers, we recommend the use of peer empowerment programs, whereby WLHIV who are knowledgeable about HIV-HPV co-infection and have positive experiences in HPV screening are recruited and trained as peer leaders to support other WLHIV to navigate the HPV and related cancer screening systems.

At the third and highest level, health literacy is measured by WLHIV’s ability to recognize the structural determinants of health and policy contexts of health care, and their engagement in social and political action that advance their individual and collective health. In Canada, under the Canadian Charter of Rights and Freedoms (1982), the Canadian Human Rights Act (1985), the Canadian Multiculturalism Act (1988), and various provincial human rights codes, health care providers and policy-makers are required to establish practice and policy guidelines that uphold and respect service users’ cultural and religious beliefs and practices. Within the contexts of screening for HPV and related cancers among immigrant and racialized WLHIV, the availability of HPV self-sampling in home, community, or clinical settings offers a potential solution in addressing many WLHIV’s current unmet needs of time constraints, bodily privacy, or access to same-gender primary care providers. Currently, HPV self-sampling is not available in Canada. Engaging affected WLHIV in policy think-tanks is critical for the development of effective and inclusive responses on cervical cancer prevention.

Finally, we were not able to identify any Canadian research on our chosen topics during this scoping review. There are a number of plausible explanations. Within the field of cancer prevention research, research attentions have shifted to examining the clinical effectiveness and efficiency of HPV vaccines, vaccination as prevention, and the uptake of HPV vaccines among different vulnerable populations such as young women, young men, and men who have sex with men (MSM). Within the HIV research field, the advent of HAART has led to the assumptions that people living with HIV (PLHIV) are no longer faced with serious health challenges. Research attentions have turned to HIV Pre-Exposure Prophylaxis (PrEP) and treatment as prevention, which, while they are important prevention strategies, do not address syndemic challenges such as disparities in HIV-HPV co-infection and HPV-related cancers among PLHIV. Further, research does not take place in a vacuum. It is driven by available funding, human resources, and sociopolitical forces such as silos in education, research, and practice within and across disciplines. Research funding organizations and their review panels play an important role in shaping the topics and designs of research, as well as reinforcing or dismantling disciplinary silos in research. In Canada, it was only recently that the Canadian Institute of Health Research (CIHR) began to address these silos by bringing different CIHR institutes...
together to co-sponsor research funding opportunities. For example, in August 2017 CRIR launched the Catalyst Grant: HPV Screening & Vaccination in Underserved Populations, sponsored by the Institute of Cancer Research in partnership with the Institute of Infection and Immunity in consultation with the Institute of Aboriginal Peoples’ Health. “Men who have sex with men (MSM) and people living with HIV” were added to the original target populations (indigenous peoples, newcomers to Canada, sex workers, injection drug users and remote populations) to promote synergy.

In conclusion, as Canada and other countries move forward with the 90-90-90 targets to achieve population HIV viral suppression and end the AIDS epidemic by 2030, policy-makers and service providers must not lose sight of the commitment to promote the health and quality of life of people who are living with HIV/AIDS. They must ensure that the health systems work in synergy to address HIV-related syndemics, including persistent HPV infections and HPV-related cancers. Synergistic collaboration is also needed in the field of health education and health promotion to address the knowledge gap, among WLHIV, about the complex interplay of HIV, HPV, and cervical cancer. Community empowerment and capacity building strategies are needed to address structural violence (stigma, discrimination, racism, cultural domination, etc.) that deters WLHIV from accessing HPV and cervical cancer screening and other health services. Despite a paucity of research, HPV self-sampling seems to offer the possibility of increasing screening among WLHIV. The adoption of HPV screening in Canada will open up dialogue about HIV–HPV syndemic challenges and increase awareness in the HIV/AIDS communities. Building on the principles of greater and meaningful involvement of people living with HIV/AIDS (GIPA/MIPA), peer leadership models may enhance self-efficacy and confidence of WLHIV who engage in HPV self-sampling.

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CONFLICT OF INTEREST DISCLOSURES
We have read and understood Current Oncology’s policy on disclosing conflicts of interest, and we declare that we have none.

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