Factors Associated with Human Papillomavirus (HPV) Test Acceptability in Primary Screening for Cervical Cancer: A Mixed Methods Research Synthesis

Ovidiu Tatar\textsuperscript{a}, Erika Thompson\textsuperscript{b}, Anila Naz\textsuperscript{a}, Samara Perez\textsuperscript{a,c}, Gilla K. Shapiro\textsuperscript{a,c}, Kristina Wade\textsuperscript{a}, Gregory Zimet\textsuperscript{d}, Vladimir Gilca\textsuperscript{e}, Monika Janda\textsuperscript{f}, Jessica Kahn\textsuperscript{g}, Ellen Daley\textsuperscript{b}, Zeev Rosberger\textsuperscript{a,c}

\textsuperscript{a}Lady Davis Institute for Medical Research, Jewish General Hospital
4333 Cote Ste-Catherine Road
Montreal, Quebec, Canada H3T1E4
Tel: (514) 340-8222, Ext. 23978
Given name: Ovidiu
Family Name: Tatar
Email: ovidiu.tatar@mail.mcgill.ca

\textsuperscript{b}Department of Health Behavior and Health Systems
School of Public Health
University of North Texas Health Science Center
3500 Camp Bowie, Blvd., EAD 709M
Fort Worth, TX 76107-2699
Email: erika.thompson@unthsc.edu

\textsuperscript{c}Department of Psychology, McGill University
2001 McGill College Avenue
Montreal Quebec, Canada
Given name: Samara
Family Name: Perez
Email: samara.perez@mail.mcgill.ca

\textsuperscript{d}Indiana University School of Medicine
Section of Adolescent Medicine
410 West 10th Street, HS 1001
Indianapolis, IN 46202, USA
Email: gzimet@iu.edu
Given name: Gregory
Family Name: Zimet

*Institut National de Santé Publique du Québec
945 Wolfe Avenue
Québec, Quebec, Canada, G1V 5B3
Email: vladimir.gilca@inspq.qc.ca
Given name: Vladimir
Family Name: Gilca

Queensland University of Technology
Brisbane, Australia
Faculty of Health
Email: m.janda@qut.edu.au
Given name: Monika
Family Name: Janda

University of Cincinnati (Ohio), USA
Division of Adolescent and Transition Medicine
MLC 4000
Cincinnati Children’s Hospital Medical Center
3333 Burnet Avenue
Cincinnati, Ohio 45229-3039
Email: jessica.kahn@cchmc.org
Given name: Jessica
Family Name: Kahn

University of South Florida
Department of Community and Family Health
13201 Bruce B. Downs Blvd, MDC 56
Tampa, FL 33612, USA
Email: edaley@health.usf.edu
Given Name: Ellen
Last Name: Daley

*Corresponding author
Given name: Ovidiu
Family Name: Tatar
Lady Davis Institute for Medical Research, Jewish General Hospital
4333 Cote Ste-Catherine Road
Montreal, Quebec, Canada H3T1E4
Tel: (514) 340-8222, Ext. 23978
Email: ovidiu.tatar@mail.mcgill.ca
Abstract

Primary screening for cervical cancer is transitioning from the longstanding Pap smear towards implementation of an HPV-DNA test, which is more sensitive than Pap cytology in detecting high-risk lesions and offers greater protection against invasive cervical carcinomas. Based on these results, many countries are recommending and implementing HPV testing-based screening programs. Understanding what factors (e.g., knowledge, attitudes) will impact on HPV test acceptability by women is crucial for ensuring adequate public health practices to optimize cervical screening uptake. We used mixed methods research synthesis to provide a categorization of the relevant factors related to HPV primary screening for cervical cancer and describe their influence on women’s acceptability of HPV testing. We searched Medline, Embase, PsycINFO, CINAHL, Global Health and Web of Science for journal articles between January 1, 1980 and October 31, 2017 and retained 22 empirical articles. Our results show that while most factors associated with HPV test acceptability are included in the Health Belief Model and/or Theory of Planned Behavior (e.g., attitudes, knowledge), other important factors are not encompassed by these theoretical frameworks (e.g., health behaviors, negative emotional reactions related HPV testing). The direction of influence of psychosocial factors on HPV test acceptability was synthesized based on 14 quantitative studies as: facilitators (e.g., high perceived HPV test benefits), barriers (e.g., negative attitudes towards increased screening intervals), contradictory evidence (e.g., sexual history) and no impact (e.g., high perceived severity of HPV infection). Further population-based studies are needed to confirm the impact of these factors on HPV-based screening acceptability.

Keywords: Human papillomavirus; Uterine Cervical Neoplasms; Mass Screening; Psychology; Human Papillomavirus DNA Tests; Review; Mixed methods; Barriers; Facilitators; Psychosocial
Introduction

Globally, 530,000 cervical cancers cases per year are attributable to the human papillomavirus (HPV) and represent 8% of all cancers occurring worldwide[1]. The understanding of the causal connection between persistent infection with high-risk HPV types and cervical cancer[2, 3] has led to new primary and secondary prophylaxis measures. Although primary prophylaxis of cervical cancer through HPV vaccination is considered a major achievement, secondary prophylaxis through screening will remain extremely important in addressing cervical cancer for decades to come because current HPV vaccines do not offer protection against all high-risk HPV types, HPV vaccine uptake is variable across the globe and the ultimate length of protection provided by vaccination is to be established yet[4].

Historically, the mainstay of cervical cancer screening was represented by cytology (i.e., Papanicolaou or Pap test) to screen for cervical cellular abnormalities. In recent years, HPV DNA tests (hereafter HPV test or testing) capable of identifying high-risk HPV types have been developed. Multiple studies have shown that HPV testing is more sensitive than cytology in detecting cervical intraepithelial neoplasia in primary cervical cancer screening (hereafter primary screening)[5-8] and has similar specificity compared to Pap testing in women aged 30 and older[9]. Overwhelming evidence suggests that a negative HPV test provides more reassurance to a woman that she is at low-risk for cervical lesions than a negative Pap test and supports the extension of intervals in primary screening beyond 5 years [3, 10, 11].

This evidence has led to new recommendations that incorporate HPV testing as a primary screen for cervical cancer in women aged between 30 and 65 years, either as a stand-alone test[12-14] or with cytology (i.e., co-testing)[15-17].

Misunderstandings and misconceptions related to HPV testing, fueled by lack of HPV or HPV testing knowledge (e.g., purpose of HPV testing, causal relationship between HPV and cervical cancer, natural history of HPV infection) in Australian women[18], lead to a petition signed by more than 70 000 women against the roll out of the new primary cervical cancer screening program (HPV test every 5 years in women aged 25 to 74 years instead of Pap test every 2 years); consequently, the implementation of the program was postponed from May 1 to December 1, 2017[14, 19].
No synthesis has been carried out to examine what factors’ impact (e.g. facilitators, barriers) on HPV test acceptability in primary screening. As new guidelines have been developed and are in the process of being implemented worldwide, we aimed to provide a comprehensive description of psychosocial factors related to HPV testing and to assess their influence on HPV testing acceptability in primary screening for cervical cancer with the ultimate goal to guide interventions to promote screening.

Methods

We used mixed methods research synthesis (MMRS), which is a form of systematic review[20-22], to answer following research questions: “What are the psychosocial factors related to HPV testing in primary screening for cervical cancer?” and “What is the influence of these factors on women’s acceptability of HPV testing in primary screening for cervical cancer?”. By selecting MMRS, we highlight our opinion that preventive behaviors (e.g., participating in screening) are complex and can be best understood by combining views of constructivism (subjectivity, associated with qualitative research) with views of logical empiricism (objectivity, associated with quantitative research). In integrative MMRS, findings of empirical qualitative, quantitative or mixed methods experimental or observational studies are treated as primary data that are analyzed and synthesized by using mixed methods approaches[20-22] (Figure 1). The PRISMA framework was used to guide the reporting of this review[23]. The protocol was registered on International Prospective Register of Systematic Reviews (PROSPERO), registration #CRD420170782541.

We searched Medline, Embase, PsycINFO, CINAHL, Global Health and Web of Science for journal articles between January 1, 1980 and October 31, 2017. The search strategy was developed for Medline by our team, validated by an experienced McGill librarian and then adapted for the other databases (Appendix A). The following eligibility criteria were applied: 1) Population: women of all ages for whom primary cervical cancer screening is recommended, 2) Outcome: psychosocial factors related to acceptability of HPV testing in primary screening for cervical cancer2, 3) Study design: empirical studies, without restrictions of study methodology,

---

1 available at https://www.crd.york.ac.uk/prospero/
2 In primary screening for cervical cancer, HPV testing is used in women with no history of cervical cytological abnormalities i.e., abnormal Pap results. Because women will be in various stages of
4) Languages: English or French or German. The selection of references was performed by two researchers (OT and AN).

Records were first screened for eligibility based on titles and abstracts (phase one). Then, the full texts of retained records were retrieved and read; the final set of articles was identified based on eligibility criteria (phase 2). Disagreements in phase one and two on whether or not an article should be included were mediated by the senior researcher (ZR). For this review, we did not retain studies related to self-sampling which represents a distinct strategy to increase screening uptake and merits separate consideration. A data extraction sheet was developed in Excel and included author, title, publication date, country, objectives, study design, quantitative data collection and analysis methods, qualitative methodology, qualitative data collection methods and analysis, and number of participants. From qualitative studies, we extracted qualitative raw data without any interpretation or analysis (e.g., quotes). From quantitative studies, we extracted outcomes of acceptability (e.g. proportions, means, odds ratios).

The risk of bias in individual studies was assessed separately by two researchers (OT and ET), with the 16-item Quality Assessment Tool for Studies with Diverse Designs (QATSDD), a valid and reliable instrument developed for appraising studies in the disciplines of psychology, sociology and nursing[24]. For overall scores <60% and > 60% we report high and low risk of bias respectively. All articles were included in the analyses, independent of their quality as we aimed to provide a comprehensive synthesis of factors.

We used a sequential exploratory (QUAL → quan) mixed methods design to analyze and synthesize findings of retained studies[22, 25, 26]. In the first phase, qualitative (QUAL), qualitative data from all qualitative and quantitative studies was analyzed; psychosocial factors measured in quantitative studies (e.g., anxiety, embarrassment, number of lifetime sexual partners, history of cervical screening) were treated as qualitative data[26]. We performed deductive-inductive qualitative thematic analysis to identify factors related to HPV testing. Deductively, we identified themes based on two frameworks widely used in health behavior research: The Health Belief Model (HBM)[27] and the Theory of Planned Behavior (TPB)[28]. Inductively, we developed new themes (i.e., not covered by HBM and TPB) through an iterative process, which consisted of reading the studies (and new themes) multiple times, allowing understanding the issue in terms of knowledge, attitudes and actual behavior, for the purposes of this paper we collapsed outcomes of intentions, willingness and uptake into the overarching term ‘acceptability’.
Researchers to assure accurate interpretation of study results. Themes (hereinafter called factors) were further grouped into categories to enable a structured reporting of the results of the qualitative phase. The factors and categories were developed independently by two researchers (OT and ET) and then validated by the research team. The second (quan) phase was informed by the first, (QUAL) phase; for each factor, based on quantitative findings (only where statistical tests for significance were reported), we provide a narrative synthesis of their influence on HPV testing acceptability. As part of the mixed research synthesis, we developed an integration matrix to match each identified factor with the quantitative evidence (for each quantitative study) of its impact on HPV testing acceptability. This approach allowed us to further synthesize the direction of influence of each factor on HPV testing acceptability into four categories: 1) possible barrier (PB, factor identified as a barrier in at least one study), 2) possible facilitator (PF, facilitator in at least one study), 3) contradictory evidence (CE), when two directions of influence (barrier and facilitator) were found for the same factor across studies and 4) no impact (NI), meaning that only evidence for no association was found. The narrative synthesis is organized based on the synthesized direction of influence of each factor on HPV testing acceptability.

Results

I. Summary of included studies and study quality

The study selection flow diagram is presented in Figure 2. We retained 22 primary studies: 5 of qualitative methodology[29-33], 15 of quantitative methodology[34-48] and 2 in which both methodologies were used[49, 50]. Seventeen studies originate in high income countries (8-USA, 2-Canada, 5-Europe and 2 in Australia) and five in low and middle income countries (1-Mexico, 1-El Salvador, 1-China, 1-India and 1 in Nigeria). In 14 quantitative studies, statistical tests of significance to assess acceptability were reported; these studies were included in the integration phase.

Quality appraisal revealed low risk of bias in 18 studies and high risk of bias in 4 studies[40, 42, 45, 49]. Among low risk of bias studies, only six were guided by an explicit theoretical framework[29-31, 40, 44, 50] or provided evidence of pilot testing of the data collection tool[29, 32, 37-39, 41]. In high risk of bias studies, theoretical frameworks were not used, the validity and reliability of the measurement tools was not assessed, no sample size calculations were provided[40, 42, 49] and few details were provided related to the recruitment
procedure and research setting[45]. Characteristics of included studies and results of quality appraisal are provided in Table 1.

II. Qualitative synthesis

1. Knowledge

Studies examined three types of knowledge: cervical cancer screening, HPV, and HPV testing. Cervical cancer screening knowledge includes women’s awareness of current cervical cancer screening guidelines[40] and implications of HPV vaccination campaigns on the need for screening[31]. Low levels of knowledge may have a particularly negative impact on HPV test acceptability: “But if I don’t know anything about cervical cancer, I will hesitate”[29]. HPV knowledge covers information gaps, such as mode of transmission[31-33] “I have a sister who came down with human papillomavirus and it got me thinking and that’s why I decided to get tested”[30], causal relationship with cervical cancer[31, 37, 47] “I don’t think I’ve ever thought of it (i.e., association between HPV and cervical cancer) in that sense”[33]. HPV test knowledge emerged as a factor since lack of knowledge contributes to women’s fear of testing[29]. Additionally women were unsure about differences between the HPV test and the Pap test[30, 31, 33, 47], were not familiar with the test procedure[30, 31] or had difficulties interpreting the results[32, 37].

2. Attitudes, beliefs and subjective norms

Women’s attitudes and beliefs are centered around four domains: cervical cancer, cervical cancer screening, HPV infection and HPV testing. Perceived severity of cervical cancer e.g., “desire to protect one’s family and one’s ability to care for their family”[32] was viewed as a reason to participate while “fear of receiving a cancer diagnosis and treatment”[32] was a reason to refuse HPV testing. Low perceived susceptibility of cervical cancer e.g., “I have never thought that you catch cervical cancer through having too much sex”[33] or perceiving low risk of cancer e.g., “not knowing anyone who had cervical cancer” were reasons for refusing HPV testing[32, 38].

Attitudes towards cervical cancer screening include delayed start of screening e.g., "Age 25 is too late. I had a 19-year-old staff member with cervical cancer”[50] and/or increased screening interval e.g., “I worry that only being tested every 4 years gives plenty of time for issues to arise and go untreated”[50], Pap versus HPV test preference[32, 48, 50] and general attitudes and beliefs e.g., presence of early signs and symptoms in cervical cancer[29] or
physical discomfort “I don’t like to get a Pap smear or anything like that, because every time I have one, they have hurt me”. [32]. When both HPV testing and Pap are available, womens’ decision depends on the screening test preference[48, 50].

The factor perceived severity of HPV infection includes the assumption that HPV testing is performed because HPV “must be a serious disease”[30] and the factor perceived susceptibility of HPV infection includes perceived risk of getting a HPV infection[48], including the relative protection offered by a monogamous relationship[32].

Perceived benefits of the HPV test synthesize women’s beliefs of the HPV test being accurate for early detection of cancer[29, 38, 46, 50] despite possible concerns about the HPV test safety[29, 50] and negative emotions and perceptions related to HPV testing such as anxiety about the test results[29], stigma and problems with communicating of positive results to significant others[33].

Subjective norms comprise healthcare provider (HCP) recommendation[29, 36, 38, 39, 44, 46, 49, 50], screening guidelines[29, 40, 44, 50], and the opinions of spouse and friends[29, 40, 42, 44, 50].

3. Health behaviors, adherence, emotional and behavioral control

HPV vaccination status[40], history of health check-up[48], including screening for breast cancer[48], usage of birth control methods (e.g., contraceptives)[36, 43] and smoking history were synthesized as health behaviors. Adherence to cervical cancer screening depends on the age of the first Pap[40], history of time-appropriate Pap testing[36, 37, 40, 42, 48] and intentions to screen with the Pap test[48]. Perceived emotional reaction to HPV results plays an important role, because women could feel embarrassed[32] or concerned[48] by a positive HPV test result and therefore be reluctant to share the test outcome with their partner or close friends[48] who could show variable level of understanding[44, 50]. Perceived behavioral control e.g., “I am confident that I could have an HPV test to screen for cervical cancer instead of a Pap smear”[50] represents an emerging factor in the context of increased options for primary cervical cancer screening.

4. Health information channels, healthcare system factors and interventions

Women use multiple health information channels to increase their knowledge[42], which emphasizes the importance of HCP in disseminating critical information about HPV testing[33]. Healthcare system factors such as health insurance status[35], availability of screening
facilities[29], and type of primary care provider (e.g., family practitioner, gynecologist)[48] can determine screening acceptability. Interventions to increase cervical screening participation include: personalized screening invitation letters[34, 37], information leaflets[34], screening reminder phone calls[34], and HPV and cervical cancer prevention education[41, 45].

5. Personal factors

General health status[37, 39], history of abnormal Pap test[38, 40, 48], past medical history e.g., cancer[36, 39], cardiovascular disease, diabetes, depression[36], history of STI’s[35, 36] are grouped under personal medical history and health status. Having a family member with cervical or other malignancies[32, 36, 39, 40] is grouped under family medical history. Age at first sexual intercourse[36, 40], number of lifetime sexual partners[36, 44, 48, 50] and sexual orientation[35] are summarized as sexual history. Finally, sociodemographics encompass widely used categories, e.g., age[31, 35, 37-40, 42-44, 48, 50], relationship status[35-37, 39, 40, 42-44, 48, 50]), and education[35-40, 42-44, 48, 50].

III. Integration of qualitative and quantitative results and quantitative synthesis

We used an integration matrix to match each factor (rows) with their influence on HPV test acceptability (e.g., facilitator) based on quantitative results of primary studies (columns) (see Appendix B). The overall effect of each factor on HPV test acceptability (e.g., possible facilitator) is provided in the last column, e.g., for high perceived benefits of the HPV test, evidence of no impact (NI)[40] and facilitator (F) [44, 50] were found, thus this factor was synthesized as possible facilitator (PF). Final MMRS results are displayed in Figure 3 where factors are organized based on their overall effect on HPV test acceptability and their theoretical framework roots (i.e., HBM or TPB or new factor). The narrative synthesis of quantitative results of primary studies is organized by results of the integration matrix and results of qualitative synthesis i.e., for each direction of influence (e.g., possible facilitators), factors corresponding to each category (e.g., knowledge, then attitudes, beliefs and subjective norms, etc.) are described sequentially.

1. Possible facilitators

Increased HPV and HPV test knowledge were associated with higher HPV test acceptability (OR=1.47; 95% CI=1.13-1.90 and OR=1.70; 95% CI=1.17-2.45 respectively)[37]. Burger et al. found a significant association between higher perceived severity of cervical cancer and HPV test acceptability (OR =1.92; 95% CI=1.32-2.80)[37]. Higher perceived
susceptibility of cervical cancer was either associated with higher HPV test acceptability (OR=1.47; 95% CI=1.05-2.06)[37] or had no effect[48]. Higher perceived susceptibility of HPV infection was associated with higher HPV test acceptability; the association was not significant for perceived susceptibility of genital warts[48]. General attitudes and beliefs related to cervical cancer screening (i.e., considering the Pap test to be very important in preventing cervical cancer) was associated with increased HPV test acceptability (OR=3.50; 95% CI=1.64-7.50)[37]. Based on a relative small sample of 149 Australian women, Jayasinghe et al. found no significant association between perceived benefits of the HPV test and HPV test acceptability (Fisher exact test, p=0.2)[40] while Ogilvie et al., on a sample of 981 Canadian women concluded that perceiving higher benefits was associated with higher acceptability of HPV testing regardless of age the screening starts (OR=1.22; 95% CI=1.15-1.30)[44] or at ≥ 25 years at a 4 years interval (OR=1.26; 95% CI=1.23-1.30)[50]. Higher perceived HPV test safety was associated with higher HPV test acceptability[44]. Higher subjective norms (i.e., higher perceived influence from significant others, HCP, screening guidelines) was associated with higher HPV test acceptability[40, 42, 44, 50].

Related to health behaviors, positive HPV vaccination status was associated with higher acceptability to receive the HPV test starting at age 25 at a 5-year interval compared to Pap testing every 2 years[40]. Contraception use was associated with higher HPV test acceptability (OR=1.63; CI=1.5-1.7)[43] and no association with the method of contraception was found [36]. Smoking history did not significantly influence HPV test acceptability[44, 48, 50]. Adherence to cervical screening recommendations (e.g., screening at intervals ≤ 3 years) was either associated with higher acceptability of HPV testing[36, 37, 42] or no association was found[39, 40, 48]. Increased behavioral control of getting the HPV test instead of the Pap test was found in two studies to increase HPV test acceptability[44, 50], while in another study to have no effect on HPV test acceptability[40].

Women who communicated with friends about health issues, or, who gathered information from media or leaflets reported higher HPV test acceptability[42]. Surprisingly, discussing health issues with HCP and gathering information via internet were not significantly associated with HPV test acceptability[42]. Among healthcare system factors, being screened in a clinic that offered HPV testing in primary cervical cancer screening was associated with increased HPV test acceptability compared to Pap[48]. Other factors, such as health insurance
status[35, 39], and distance from the clinic and transportation facilities were not related to HPV test acceptability[36].

In terms of sociodemographics, non-whites were found to have lower acceptability than whites[42], except for Latina[39]. Education was found to either have no impact [35-40, 44, 48] or increase[42, 43, 50] HPV test acceptability. Higher income is a possible facilitator as we found that income can increase[37, 48] or have no effect[35, 39, 40, 42] on HPV test acceptability.

2. Possible barriers

Women expressing concerns about delayed start of screening had significantly lower acceptability of the HPV test if the screening start is delayed to 25 years and continues at a 5-year interval[40]. Increasing the screening interval from 1 to 3 years had no significant influence on HPV test acceptability[40, 48]. For five years between screening, acceptability was either similar to yearly intervals[48] or decreased (OR=0.2; CI=0.1-0.4)[40], while for 10-year screening interval acceptability was lower (OR=0.05; CI=0.03-0.1)[40]. Negative emotions and perception related to HPV testing significantly increased acceptability of Pap compared to HPV testing (PR=1.39; 95% CI=1.07-1.80)[48]. In most studies, marital status was not associated[35, 36, 39, 42, 44, 48, 50] with HPV test acceptability but evidence exists that being single (versus married)[37, 43] is related to lower acceptability.

3. Contradictory evidence

Negative perceived emotional reaction to HPV results can either augment (i.e., higher concern about a positive HPV test)[48], diminish (i.e., women reluctant to share a positive HPV result with their partner)[37] or have no effect HPV test acceptability[44, 50]. In terms of screening test preference, when both Pap and HPV tests are offered, preference for a test is associated with higher acceptability i.e., increased HPV test acceptability (OR = 1.26; CI = 1.23-1.30)[50] or increased Pap acceptability (60.7% for Pap, CI = 56.5-65.7)[48].

Related to existing personal medical history, poor or very poor self-reported health status was found to decrease (OR= 0.49; CI = 0.27-0.91)[37] or have no significant effect[39] on HPV test acceptability. Reporting personal history of cancer (other than cervical) increased[39] or had no effect on HPV test acceptability[36]. Reporting previous cervical cytological abnormalities either decreased (OR = 0.65; CI = 0.46-0.94)[38] or had no impact[48] on HPV test acceptability. With respect to obstetric history, compared to nulligravidae, women who reported
pregnancies had higher HPV test acceptability (OR = 2.10; CI= 1.80-2.40)[43]. Other personal medical history correlates e.g., history of STI, menopausal status, Body Mass Index, cardiovascular diseases, diabetes and depression were not associated with HPV test acceptability[35, 48]. Among sexual history, reporting 4 or more[36] or zero[35] lifetime sexual partners was associated with lower HPV test acceptability and reporting both male and female lifetime partners increased HPV test acceptability (OR=1.75; CI=1.39-2.20)[35]. Evidence related to age is contradictory; older women (e.g., ≥ 40) were found to have either increased[38, 39] or decreased[37, 42, 43] HPV test acceptability.

In 60-70 year old women, interventions in form of personalized letters signed by their physician and an informative leaflet explaining the most important reasons for screening for cervical cancer significantly increased HPV test acceptability (screening coverage increased with 31.6%, CI=29.0-34.1, p≤0.05)[34]. In a nationally representative sample of Norwegian women, Burger et al. found that using invitation letters for HPV testing (i.e., stating that HPV testing at a 6-year interval will replace Pap testing) resulted in marginally lower HPV test acceptability (strength of intention, p=0.008) compared to using Pap testing invitation letters (i.e., at a 3-years interval)[37]. Educational interventions were found to have either equivocal[45] or positive effect[41] on HPV test acceptability.

4. No impact

Cervical cancer screening knowledge was not associated with HPV test acceptability[40, 48]. Higher perceived severity of HPV infection had no significant effect on women’s acceptability of the HPV test[37].

Family medical history of cancer was not associated with HPV test acceptability[36, 39].

Discussion

In our mixed methods research synthesis, we analyzed findings of empirical qualitative and quantitative studies and: a) provided an up-to-date and comprehensive list of factors specific for HPV test acceptability in primary screening for cervical cancer, b) synthesized factors’ direction of influence on HPV test acceptability and c) described factors’ impact on HPV test acceptability.

Our results show that factors associated with HPV test acceptability are complex; while many factors are included in the HBM and/or TPB (e.g., attitudes, perceived behavioral control),
other relevant factors are not encompassed by these theoretical frameworks e.g., health behaviors, negative emotional reactions related to a positive HPV test result. Negative attitudes toward delayed start of screening (i.e. 25 years) and/or increased screening interval to 5 or 10 years and negative emotions and perceptions related to HPV testing are possible barriers to HPV test acceptability. In the context of the latest recommendations[12-17] for primary screening for cervical cancer and ongoing plans of health authorities[14, 51-57] to implement HPV testing in primary cervical screening, addressing these attitudes and concerns should become part of the strategy to ensure a successful implementation of HPV test-based screening programs.

We found that women’s increased HPV and HPV test knowledge and using information channels represent a possible facilitator of HPV test acceptability. Since women in the USA, Australia and UK were found to have low HPV[58] and HPV test knowledge[59], strategies that increase women’s knowledge might also increase HPV test acceptability.

Healthcare providers play an important role in promoting preventive health measures; as our team has previously demonstrated, discussing with HCP’s about HPV vaccination significantly increased acceptability of the HPV vaccine for their sons[60]. However, when HCPs are unknowledgeable about, or uncomfortable with, recommendations, it can negatively impact preventive health behaviors. In the context of cervical cancer screening, Boone et al. (2016) found that US HCP’s (e.g., OB/GYN, family physicians), contrary to existing guidelines for women aged 30 to 65 years[17], recommended HPV co-testing on an every 3 year basis instead of 5 years[61]. Similar results were obtained in Italy by Cagliotti et al. (2017), who found that in women older than 30 years, 83.8% of gynecologists prefer to use the Pap test in primary screening, and only 44.9% of gynecologists knew that a negative HPV-DNA test allowed an increase in the screening interval to 5 years [62]. Moreover, 20% of participants believed that HCP are insufficiently prepared to explain either positive or negative HPV test results to their patients[62]. In our opinion, especially in health systems where cervical cancer screening is opportunistic, an age-appropriate HCP recommendation for HPV testing could increase women’s HPV test acceptability as primary cervical cancer screening. Efforts are therefore needed to increase HCPs’ awareness of, and comfort with the latest guidelines.

Moreover, adequate HPV vaccination coverage of females is important for secondary prevention of cervical cancer, as we found that women who were not vaccinated against HPV also had lower HPV test acceptability. Our results are concordant with results of a large US
study which showed that cervical screening initiation and interval adherence were significantly higher in women who had been vaccinated against HPV [63].

Since our review shows that attitudes and beliefs are important factors of HPV test acceptability but have been measured with scales that were not rigorously psychometrically tested [40, 44, 50], we recommend that future research address this knowledge gap. While a comprehensive and psychometrically validated scale for measuring HPV knowledge has been published [64], in our opinion, the only validated HPV test knowledge scale available [59, 65] needs to be modified to include items related to differences between Pap and HPV testing [31-33] (e.g., reasons for doing a HPV instead of Pap test), risks [29, 30] (e.g., pain, infection) and practicalities of the HPV test [31] (e.g., what it involves).

Our study is not without limitations. Because most included studies were observational, interventions are needed for assessing the effect of factors on HPV test acceptability. Given that HPV testing as primary screening has only recently been recommended and only in some countries, there is a paucity of studies of psychosocial correlates of actual HPV testing uptake. Therefore, we defined HPV test acceptability comprehensively and included HPV test uptake as well as intentions/willingness to receive the HPV test in our synthesis. Our results relate to the overarching significant factors of organized or opportunistic screening environments, while some differences e.g., previous adherence to cervical screening are possible. We conducted a sensitivity analysis by removing studies with high risk of bias [40, 42, 45] from our synthesis (i.e., integration of quantitative evidence) and results remained largely unchanged. However, the facilitator effects of using health information channels [42], being of white ethnicity [42] and the barrier effect of expressing negative attitudes towards delayed start of screening and/or increased screening interval to 5 or 10 years [40] require further validation. We encourage researchers to further study the effect of factors on women’s HPV test acceptability for which we found contradictory evidence i.e., cervical screening test preference, negative perceived emotional reaction to HPV results, the type of intervention, existing personal medical history and women’s age. These contradictory findings may be attributed to the heterogeneity of factors (outcomes), population and interventions measured across included studies.
Conclusions

By synthesizing findings of both qualitative and quantitative studies, our review provides a wide perspective related to factors of HPV testing in primary cervical cancer screening. Our results can inform designing interventions to increase primary HPV-based cervical cancer screening uptake in high income countries, but even more so in low and middle income countries where the incidence of cervical cancer is highest and where, as suggested by previous research[66], implementing a primary HPV testing program could be lifesaving.

Appendices: Appendix A – Search strategy for Medline, Appendix B – Integration Matrix

Acknowledgements: We would like to thank to Genevieve Gore (McGill Liaison Librarian) for her invaluable assistance with validating the search strategy.

Conflict of Interest
Zeev Rosberger reports personal fees from Merck outside the submitted work at a consultation meeting in November, 2015; and speaker to family physicians in April, 2015. Gregory Zimet reports grants from Merck, grants from Roche, personal fees from Merck, outside the submitted work.

Funding
This work was supported by a grant from the Canadian Cancer Society Research Institute (CCSRI, Grant #704036). GS is a Vanier CIHR Canada Graduate Scholar and Queen Elizabeth II Diamond Jubilee Scholar. SP is a Vanier CIHR Canada Graduate Scholar.
REFERENCES


**Figure 1.** Integrative Mixed Methods Research Synthesis Design

**Figure 2.** PRISMA Flowchart

**Figure 3.** Influence of factors on HPV test acceptability

Note: cc = cervical screening; HPV = human papillomavirus
<table>
<thead>
<tr>
<th>First author, country, year</th>
<th>Aim</th>
<th>Data collection method</th>
<th>Cytology screening environment</th>
<th>Intervention (Yes/No)</th>
<th>N</th>
<th>Setting</th>
<th>Participant age</th>
<th>Data analysis method</th>
<th>Risk of bias within studies (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acera et al.[1], Spain, 2014</td>
<td>To determine the most effective intervention strategy to increase cervical cancer screening coverage in Barcelona</td>
<td>Personal Interview</td>
<td>Opportunistic</td>
<td>Yes</td>
<td>4775</td>
<td>Primary Health Care centers in Cerdanyola, Barcelona</td>
<td>60-70</td>
<td>Chi-square</td>
<td>Low (67 %)</td>
</tr>
<tr>
<td>Agenor et al.[2], USA, 2017</td>
<td>To examine the associations between sexual behavior and sexual identity, and lifetime HPV testing</td>
<td>Survey administered by interviewer</td>
<td>Opportunistic</td>
<td>No</td>
<td>11,300</td>
<td>National probability sample</td>
<td>15-44</td>
<td>Multivariate logistic regression</td>
<td>Low (67 %)</td>
</tr>
<tr>
<td>Alfaro et al.[3], El Salvador, 2015</td>
<td>To identify the facilitators and barriers to adherence to cervical cancer screening using HPV DNA testing in El Salvador</td>
<td>Interview</td>
<td>Opportunistic</td>
<td>Yes</td>
<td>409</td>
<td>Salvadori an Ministry of Health led Cervical Cancer Prevention HPV screening program</td>
<td>30-49</td>
<td>Univariate logistic regression, chi square, multivariate logistic regression</td>
<td>Low (69 %)</td>
</tr>
<tr>
<td>First author, country, year</td>
<td>Aim</td>
<td>Data collection method</td>
<td>Cytology screening environment</td>
<td>Intervention (Yes/No)</td>
<td>N</td>
<td>Setting</td>
<td>Participant age</td>
<td>Data analysis method</td>
<td>Risk of bias within studies (%)</td>
</tr>
<tr>
<td>----------------------------</td>
<td>---------------------------------------------------------------------</td>
<td>------------------------</td>
<td>-------------------------------</td>
<td>------------------------</td>
<td>---</td>
<td>---------</td>
<td>-----------------</td>
<td>-----------------------</td>
<td>-------------------------------</td>
</tr>
<tr>
<td>Burger et al.[4], Norway, 2014</td>
<td>To examine whether the contents of a cervical cancer screening invitation letter influence Norwegian women’s intent to participate in screening</td>
<td>Web-based survey</td>
<td>Organized</td>
<td>Yes</td>
<td>3540</td>
<td>Representative sample of Norwegian women</td>
<td>25-69</td>
<td>Univariate and multivariate Logistic regression</td>
<td>Low (81 %)</td>
</tr>
<tr>
<td>Dieng et al.[5], Australia, 2013</td>
<td>To investigate Australian women’s cervical cancer screening preferences, information needs and decision-making styles</td>
<td>Semi-structured telephone interview</td>
<td>Organized</td>
<td>No</td>
<td>1279</td>
<td>National survey conducted by the Hunter Valley Research Foundation</td>
<td>18-70</td>
<td>Multivariate logistic regression, descriptive statistics</td>
<td>Low (64 %)</td>
</tr>
<tr>
<td>Filade et al.[6], Nigeria, 2017</td>
<td>To explore the attitudes of pregnant women to the incorporation of HPV DNA-based testing in routine ANC in Nigeria</td>
<td>Focus groups</td>
<td>Opportunistic</td>
<td>No</td>
<td>82</td>
<td>Hospitals and health facilities in central Nigeria</td>
<td>Mean 28.9 (SD = 4.7)</td>
<td>Qualitative content analysis</td>
<td>Low (90 %)</td>
</tr>
<tr>
<td>First author, country, year</td>
<td>Aim</td>
<td>Data collection method</td>
<td>Cytology screening environment</td>
<td>Intervention (Yes/No)</td>
<td>N</td>
<td>Setting</td>
<td>Participant age</td>
<td>Data analysis method</td>
<td>Risk of bias within studies (%)</td>
</tr>
<tr>
<td>-----------------------------</td>
<td>-----</td>
<td>------------------------</td>
<td>-------------------------------</td>
<td>-----------------------</td>
<td>---</td>
<td>---------</td>
<td>-----------------</td>
<td>----------------------</td>
<td>-----------------------------</td>
</tr>
<tr>
<td>Gerend et al.[7], USA, 2017</td>
<td>To investigate women’s acceptance of new cervical cancer screening guidelines</td>
<td>Electronic Survey</td>
<td>Opportunistic</td>
<td>No</td>
<td>376</td>
<td>Online panel maintained by Qualtrics</td>
<td>21-65</td>
<td>Descriptive statistics</td>
<td>High (58%)</td>
</tr>
<tr>
<td>Huang et al.[8], USA, 2008</td>
<td>To assess women’s interest in obtaining HPV testing as well as their preferences for concomitant Pap testing</td>
<td>Telephone and in-person interviews</td>
<td>Opportunistic</td>
<td>No</td>
<td>Opportunistic 865</td>
<td>Community and university-based practices</td>
<td>50-80</td>
<td>Multivariate logistic regression</td>
<td>Low (64%)</td>
</tr>
<tr>
<td>Jayasinghe et al.[9], Australia, 2016</td>
<td>To assess women’s attitudes towards guidelines for HPV testing in cervical cancer screening</td>
<td>Electronic Survey</td>
<td>Organized</td>
<td>No</td>
<td>125</td>
<td>Social media</td>
<td>16-28</td>
<td>Fisher’s exact test, odds ratios</td>
<td>High (60%)</td>
</tr>
<tr>
<td>First author, country, year</td>
<td>Aim</td>
<td>Data collection method</td>
<td>Cytology screening environment</td>
<td>Intervention (Yes/No)</td>
<td>N</td>
<td>Setting</td>
<td>Participant age</td>
<td>Data analysis method</td>
<td>Risk of bias within studies (%)</td>
</tr>
<tr>
<td>----------------------------</td>
<td>-----</td>
<td>------------------------</td>
<td>-------------------------------</td>
<td>----------------------</td>
<td>----</td>
<td>---------</td>
<td>-----------------</td>
<td>----------------------</td>
<td>-----------------------------</td>
</tr>
<tr>
<td>Kwan et al. [10], China, 2010</td>
<td>To evaluate the effects of educational intervention on Chinese women’s intentions to be HPV tested</td>
<td>Questionnaire</td>
<td>Opportunistic</td>
<td>Yes</td>
<td>292</td>
<td>Family Planning Association of Hong Kong’s (FPAHK) Wanchai Birth Control Clinic</td>
<td>Mean = 38.3 (SD = 7.41)</td>
<td>Chi Square</td>
<td>Low (88 %)</td>
</tr>
<tr>
<td>Leon-Maldonado et al. [11], Mexico, 2016</td>
<td>To assess the beliefs and perceptions of HPV and HPV testing among Mexican women who had participated in an early cervical cancer detection program</td>
<td>Semi-structured interviews</td>
<td>Opportunistic</td>
<td>No</td>
<td>24</td>
<td>Two primary care health clinics in Michoacán state, Mexico</td>
<td>30-65</td>
<td>Thematic framework analysis</td>
<td>Low (71 %)</td>
</tr>
<tr>
<td>Marlow et al. [12], UK, 2008</td>
<td>To examine sociodemographic predictors of self-reported screening attendance, and intention to accept HPV testing.</td>
<td>Home-based, computer assisted interviews</td>
<td>Organized</td>
<td>No</td>
<td>994</td>
<td>National Centre for Social Research Omnibus Survey</td>
<td>25-64</td>
<td>Univariate and multivariate logistic regression</td>
<td>High (52 %)</td>
</tr>
<tr>
<td>First author, country, year</td>
<td>Aim</td>
<td>Data collection method</td>
<td>Cytology screening environment</td>
<td>Intervention (Yes/No)</td>
<td>N</td>
<td>Setting</td>
<td>Participant age</td>
<td>Data analysis method</td>
<td>Risk of bias within studies (%)</td>
</tr>
<tr>
<td>---------------------------</td>
<td>-----</td>
<td>------------------------</td>
<td>-------------------------------</td>
<td>---------------------</td>
<td>---</td>
<td>---------</td>
<td>-----------------</td>
<td>----------------------</td>
<td>------------------------</td>
</tr>
<tr>
<td>Marlow et al.[13], UK, 2009</td>
<td>To identify British women’s HPV and HPV test knowledge requirements</td>
<td>Interviews</td>
<td>Organized</td>
<td>No</td>
<td>21</td>
<td>University College London</td>
<td>18-53</td>
<td>Thematic framework analysis</td>
<td>Low (67 %)</td>
</tr>
<tr>
<td>Nene et al.[14], India, 2007</td>
<td>To evaluate the sociodemographic variations in the uptake of cervical cancer screening in rural India</td>
<td>Household survey and hospital records</td>
<td>Opportunistic</td>
<td>Yes</td>
<td>79,449</td>
<td>Primary health centres, rural hospitals, and schools in the Osmanabad district in Maharashtra state</td>
<td>30-59</td>
<td>Univariate and multivariate logistic regression</td>
<td>Low (71 %)</td>
</tr>
<tr>
<td>Ogilvie et al.[15], Canada, 2013</td>
<td>To explore the impact of HPV testing on women’s intentions to be screened for cervical cancer</td>
<td>Electronic Survey</td>
<td>Organized in British Columbia</td>
<td>Yes</td>
<td>981</td>
<td>Provincial cervical cancer screening program at the British Columbia Cancer Agency</td>
<td>25-65</td>
<td>Chi square, t-test, multivariate logistic regression</td>
<td>Low (83 %)</td>
</tr>
<tr>
<td>First author, country, year</td>
<td>Aim</td>
<td>Data collection method</td>
<td>Cytology screening environment</td>
<td>Intervention (Yes/No)</td>
<td>N</td>
<td>Setting</td>
<td>Participant age</td>
<td>Data analysis method</td>
<td>Risk of bias with in studies (%)</td>
</tr>
<tr>
<td>-----------------------------</td>
<td>-----</td>
<td>------------------------</td>
<td>--------------------------------</td>
<td>----------------------</td>
<td>----</td>
<td>----------</td>
<td>-----------------</td>
<td>----------------------</td>
<td>-------------------------------</td>
</tr>
<tr>
<td>Ogilvie et al.[16], Canada, 2016</td>
<td>To describe factors associated with women’s intentions to be screened according to new guidelines for primary HPV DNA testing</td>
<td>Electronic Survey</td>
<td>Organized in British Columbia</td>
<td>Yes</td>
<td>981</td>
<td>Provincial cervical cancer screening program at the British Columbia Cancer Agency</td>
<td>25-65</td>
<td>Multivariate logistic regression, Kruskal Wallis, chi-square thematic analysis</td>
<td>Low (81 %)</td>
</tr>
<tr>
<td>Papa et al.[17], USA, 2009</td>
<td>To assess the impact of educational intervention on women’s acceptance of adjunct HR-HPV testing</td>
<td>Questionnaire</td>
<td>Opportunistic</td>
<td>Yes</td>
<td>50</td>
<td>Obstetrics and gynecology faculty practice at the University of Massachusetts Medical School/U Mass Memorial Health Care</td>
<td>30-69</td>
<td>Fisher exact test</td>
<td>High (55 %)</td>
</tr>
<tr>
<td>Roland et al.[18], USA, 2016</td>
<td>To assess the impact of educational intervention on knowledge and beliefs of cervical cancer screening</td>
<td>Survey</td>
<td>Opportunistic</td>
<td>Yes</td>
<td>644</td>
<td>Federally Qualified Health Center clinics in Illinois</td>
<td>30-60</td>
<td>Ordinal and binary logistic regression</td>
<td>Low (67 %)</td>
</tr>
<tr>
<td>First author, country, year</td>
<td>Aim</td>
<td>Data collection method</td>
<td>Cytology screening environment</td>
<td>Intervention (Yes/No)</td>
<td>N</td>
<td>Setting</td>
<td>Participant age</td>
<td>Data analysis method</td>
<td>Risk of bias within studies (%)</td>
</tr>
<tr>
<td>-----------------------------</td>
<td>-----</td>
<td>------------------------</td>
<td>-------------------------------</td>
<td>----------------------</td>
<td>----</td>
<td>---------</td>
<td>-----------------</td>
<td>----------------------</td>
<td>-----------------------------</td>
</tr>
<tr>
<td>Schmid et al.[19], USA &amp; Australia, 2017</td>
<td>To determine the perceptual word associations that women hold with regards to cervical cancer screening tools in the US and Australia</td>
<td>Electroni c Survey</td>
<td>Opportun istic</td>
<td>No</td>
<td>776</td>
<td>Survey Monkey’s US and Australian databases</td>
<td>18-64</td>
<td>Co-occurrence network graphs</td>
<td>Low (64%)</td>
</tr>
<tr>
<td>Silver et al.[20], USA, 2015</td>
<td>To explore and understand women’s attitudes towards new cervical cancer screening options</td>
<td>Interview er-administered survey</td>
<td>Opportun istic</td>
<td>Yes</td>
<td>551</td>
<td>Johns Hopkins Hospital affiliated outpatient OB/GYN clinics in Baltimore, MD</td>
<td>36-62</td>
<td>Poisson regression with robust error variance</td>
<td>Low (67%)</td>
</tr>
<tr>
<td>Vanslyke et al.[21], USA, 2008</td>
<td>To explore the knowledge, attitudes and beliefs related to cervical cancer, HPV and HPV testing of low-income, Hispanic women</td>
<td>Focus groups</td>
<td>Opportun istic</td>
<td>No</td>
<td>54</td>
<td>Communi ty-based settings in Albuquerque, New Mexico</td>
<td>18-60</td>
<td>Themati c analysis</td>
<td>Low (69%)</td>
</tr>
<tr>
<td>First author, country, year</td>
<td>Aim</td>
<td>Data collection method</td>
<td>Cytology screening environment</td>
<td>Intervention (Yes/No)</td>
<td>N</td>
<td>Setting</td>
<td>Participant age</td>
<td>Data analysis method</td>
<td>Risk of bias within studies (%)</td>
</tr>
<tr>
<td>-----------------------------</td>
<td>----------------------------------------------------------------------</td>
<td>------------------------</td>
<td>--------------------------------</td>
<td>-----------------------</td>
<td>---</td>
<td>---------</td>
<td>-----------------</td>
<td>----------------------</td>
<td>-------------------------------</td>
</tr>
<tr>
<td>Waller et al.[22], UK, 2005</td>
<td>To examine the understanding and beliefs about HPV and cervical cancer among women who have already participated in HPV testing</td>
<td>Structured Interviews</td>
<td>Organized</td>
<td>No</td>
<td>74</td>
<td>Clinical trials of HPV testing and a colposcopy clinic that utilizes HPV testing</td>
<td>20-64</td>
<td>Thematic framework analysis</td>
<td>Low (64%)</td>
</tr>
</tbody>
</table>

Note: * percentage points were calculated as recommended by the authors of the Quality Assessment Tool for Studies with Diverse Designs (QATSDD); 100% reflects no risk of bias. For overall scores ≤60% and > 60% we report high and low risk of bias respectively.
Highlights:

- We used mixed methods research synthesis methodology, a form of systematic review
- We provide a comprehensive categorization of psychosocial factors in HPV testing
- Data integration enabled assessment of factors’ impact on HPV test acceptability
- Assessed impact of factors: barriers, facilitators, contradictory evidence; no impact
Adapted from Sandelowski et al. (2006) and Heyvaert et al. (2013)

Qual indicates qualitative dominant method of analysis; quan indicates non-dominant quantitative method of analysis
Figure 2

1 Phase 1 exclusion criteria for titles and abstracts: 1) not population of interest (i.e. women), 2) not on primary cervical cancer screening with the HPV DNA test, 3) not outcomes of interest (psychosocial correlates of primary HPV cervical cancer screening), 4) not empirical studies, 5) no abstract

2 Phase 2 exclusion criteria for full text articles: Phase 1 exclusion criteria AND full text not in English, French or German