

CERVICAL SCREENING: Supporting your patients to make the choice



Cervical
Screening



NATIONAL
CERVICAL SCREENING
PROGRAM
A joint Australian, State and Territory Government Program



		Clinician-collected cervical sample	Self-collected vaginal sample																					
Is it accurate?		Both methods have equivalent sensitivity for the detection of HPV and CIN2+/AIS ^{1,2}																						
Identifies HPV infection?		Yes	Yes																					
Is liquid-based cytology (LBC) and co-testing possible?		Yes	No																					
Indicated for		Yes	Yes																					
<ul style="list-style-type: none"> • People who are eligible and due or overdue for cervical screening, including during pregnancy • Other points in the pathway where only an HPV test is required, including the 12- and 24-month follow-up test after an intermediate risk result 		Yes	Yes																					
<ul style="list-style-type: none"> • Test of Cure after HSIL treatment • People who have had a total hysterectomy with history of HSIL 		Yes	No (current)	Yes (following expected guideline changes in 2024/25*)																				
<ul style="list-style-type: none"> • People who have postcoital, unexplained intermenstrual or post-menopausal bleeding, or unexplained persistent unusual vaginal discharge³ • People undergoing surveillance after completely excised AIS • People who were exposed to DES in utero. 		Yes	No																					
Management of participants in whom HPV is not detected	>90% of CSTs	Return in 5 years	Return in 5 years																					
Management of participants in whom HPV (not 16/18) is detected	~6% of CSTs	Reflex LBC is performed on the original sample, no need to return for a further sample to be taken	Return for clinician-collected cervical sample for LBC to inform the risk rating as soon as practical, ideally within 6 weeks. The incidence of HPV (not 16/18) is highly age dependent (NCSR data ⁴) <table border="1" style="margin: 10px auto;"> <tr> <td>25-29 years</td> <td>17%</td> <td>50-54 years</td> <td>4%</td> </tr> <tr> <td>30-34 years</td> <td>10%</td> <td>55-59 years</td> <td>3%</td> </tr> <tr> <td>35-39 years</td> <td>6%</td> <td>60-64 years</td> <td>3%</td> </tr> <tr> <td>40-44 years</td> <td>5%</td> <td>65-69 years</td> <td>3%</td> </tr> <tr> <td>45-49 years</td> <td>4%</td> <td></td> <td></td> </tr> </table> Note: at the 12-month follow up HPV test after an Intermediate Risk result the incidence of HPV (not 16/18) is ~60% ⁴ Patients aged 70 to 74 with HPV (not 16/18) detected are referred to colposcopy.		25-29 years	17%	50-54 years	4%	30-34 years	10%	55-59 years	3%	35-39 years	6%	60-64 years	3%	40-44 years	5%	65-69 years	3%	45-49 years	4%		
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Management of participants in whom HPV (16/18) is detected	~2% of CSTs	Refer for colposcopy	Refer for colposcopy																					
Management of Unsatisfactory HPV test		Repeat as soon as practical, ideally within 6 weeks	Repeat as soon as practical, ideally within 6 weeks																					
HPV: Human papillomavirus; LBC: Liquid based cytology; HSIL: High-grade squamous intraepithelial lesion; AIS: Adenocarcinoma in situ; DES: Diethylstilbesterol Adapted from 'Self-collection: equity in reaching cervical cancer elimination, Aug 2022, InSight+, AMPCo.		*Guideline changes regarding Test of Cure surveillance are expected in 2024/2025 – please check the National Cervical Screening Program Guidelines on the Cancer Council Australia website for up-to-date information.																						

1 Arbyn et al, Detecting cervical precancer and reaching underscreened women by using HPV testing on self samples: updated meta-analyses BMJ 2018; 363 :k4823

2 Saville et al, Analytical performance of HPV assays on vaginal self-collected vs practitioner-collected cervical samples: the SCoPE study, Journal of Clinical Virology (2020), doi: https://doi.org/10.1016/j.jcv.2020.104375

3 Co-testing is not required for breakthrough or irregular bleeding due to hormonal contraception or a sexually transmitted infection, heavy menstrual bleeding, or contact bleeding at time of obtaining a routine cervical screening test sample

4 Smith et al, National experience in the first two years of primary human papillomavirus (HPV) cervical screening in an HPV vaccinated population in Australia: observational study BMJ 2022; 376 :e068582

HPV SELF-COLLECTION: 10 KEY QUESTIONS FOR HEALTHCARE PROVIDERS ANSWERED



Why should I offer self-collection?

- HPV self-collection is accurate, safe and highly acceptable, particularly among under and never-screened people. It is now widely available as an option for all participants in routine cervical screening.
- Under-screening is the most important risk factor for developing cervical cancer. More than 70% of Australians diagnosed with cervical cancer have never been screened or are overdue.

Is self-collection as accurate as a clinician-collected sample?

- Yes. We now have strong evidence demonstrating that self-collection is as sensitive for the detection of HPV and CIN2+/AIS as a clinician-collected test when processed using PCR technology¹.

How do I set myself up to offer self-collection?

- Contact your laboratory to ensure you have the correct equipment and handling instructions to offer self-collection.

What do I need to check before offering self-collection?

- Check for symptoms, as only asymptomatic patients can be offered self-collection.
- Check your patient's screening history via the National Cancer Screening Register
- Check your patient's self-collection eligibility according to the most up-to-date NCSP Clinical Guidelines.
- For more information on all of the above, please see the reverse of this resource.

Who can order self-collection in my practice?

- All cervical screening needs to be ordered by a healthcare professional with a provider number. The responsibility for the test and follow-up is with the healthcare professional who ordered the test.
- All practice staff, including practice nurses, Aboriginal Health Workers and administration staff have a role to play in supporting patients' participation in screening and awareness of their screening choices.

Where can self-collection be performed? Can my patient take the sample at home?

- Self-collection is most often undertaken in a health service, e.g., in a clinic bathroom or behind a curtain.
- If you are comfortable with your patient taking the test at home, remind them to label the swab with the date of sample collection.
- You do not need to observe your patient taking the sample.

Who is more likely to be under-screened?

- Aboriginal and Torres Strait Islander people, people from culturally and linguistically diverse communities, people with disability, LGBTQIA+ people, and people living in rural and remote areas are amongst those who may be less likely to participate in screening.
- Available data indicates Aboriginal and Torres Strait Islander women have over twice the incidence and almost four times the mortality associated with cervical cancer compared to non-Indigenous Australians.
- Evidence also suggests those born overseas are more likely to have never screened or be overdue for screening, and are less likely to have heard about the option to self-collect.
- Offering self-collection as a choice for all eligible patients overcomes barriers to screening, and therefore directly addresses these inequities.

What causes unsatisfactory results?

- Only a small number of self-collected samples (~1-2%) produce an invalid result, due to inadequate cellular material or the presence of interfering substances (such as lubricants or creams, or large amounts of blood or discharge).

How do I advise my patients to do the test?

- Self-collection is easy to do. The swab is inserted into the vagina and rotated for at least 10 seconds to collect an adequate number of vaginal cells.
- Reassure your patient that the sample does not need to be taken from the cervix. The swab only needs to be inserted about the length of their index finger.

Will I miss something if I don't do a pelvic examination?

- There is no evidence to support the use of pelvic examination as routine practice for asymptomatic patients².
- For HPV positive patients, the cervix will be visualised during a follow up appointment for cytology or colposcopy.
- Decisions to perform a pelvic examination or visual inspection of the genital tract should be patient-centred, clearly clinically indicated and made collaboratively.
- You can use any time saved by not needing to perform a pelvic exam to check for symptoms and remind patients what to look out for.

1. Arbyn, M. et al [2018]. Detecting cervical precancer and reaching underscreened women by using HPV testing on self samples: updated meta-analyses. *Bmj*, 363

2. RACGP Guideline for preventative activities in general practice [Red Book]. [2021]. Available at <https://www.racgp.org.au/clinical-resources/clinical-guidelines/key-racgp-guidelines/view-all-racgp-guidelines/guidelines-for-preventive-activities-in-general-pr/early-detection-of-cancers/cervical-cancer#ref-num-98>.