CERVICAL SCREENING: Supporting your patients to make the choice



	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 • Those 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.	Yes	Yes
 Patients who have postcoital or intermenstrual bleeding, post-menopausal bleeding, or unexplained persistent unusual vaginal discharge³ Those undergoing Test of Cure surveillance or have been treated for adenocarcinoma-in-situ Patients who have had a total hysterectomy with history of high-grade squamous intraepithelial lesion Patients who were exposed to diethylstilbesterol in utero. 	Yes	No
Management of participants in whom HPV is not detected >90%	Return in 5 years	Return in 5 years
Management of participants in whom HPV (not 16/18) is detected ~6%	Reflex LBC performed on original sample, no need to return for a further sample to be taken	Return for clinician-collected cervical sample for LBC. The incidence of HPV (not 16/18) is highly age dependent. NCSR data ⁴
		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%
		Patients aged 70 to 74 with HPV (not 16/18) detected are referred to colposcopy.
Management of participants in whom HPV (16/18) is detected ~2%	Refer for colposcopy	Refer for colposcopy
Management of Unsatisfactory HPV test	Repeat in 6 weeks	Repeat at earliest convenience

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

⁴ Smith et al, BMJ 2022;376:e068582 Available at: https://www.bmj.com/content/376/bmj-2021-068582



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