Frequently Asked Questions

1. What is HPV DNA test?

HPV DNA test detects high-risk HPV (hr-HPV) types that can cause cervical cancer or precancerous changes. In 2021, World Health Organization (WHO) recommended HPV DNA test as a first-choice screening method for cervical cancer prevention.

2. Which types of HPV can be tested by FPAHK's HPV DNA test?

HPV is a common virus that has more than 200 subtypes, and about 40 of which infect the ano-genital area. HPV DNA test provided by the FPAHK is approved by the U.S. Food and Drug Administration for screening. The test detects 14 hr-HPV subtypes, including HPV-16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66 and 68. Among these, HPV-16 and HPV-18 cause 70% of cervical cancer.

3. Cervical cytology has been used for decades, why there is a change now?

Co-test improves the sensitivity of cervical cancer risk assessment by detecting abnormal cervical cells and hr-HPV at the same time. The Hong Kong College of Obstetricians and Gynaecologists (HKCOG) Guidelines for Cervical Cancer Prevention and Screening in 2016 had already included the use of co-test for cervical cancer screening. The Cancer Expert Working Group on Cancer Prevention and Screening (CEWG) of the Centre for Health Protection also recommended co-test as one of the cervical cancer screening methods. One local study demonstrated earlier detection of cervical cells abnormality by co-test, and reduction of cervical cancer risk through appropriate treatment.

4. Why is co-test NOT recommended for women under 30 years old?

The prevalence of HPV infection among these young women is high, but most of the infections are transient and will clear up on their own. It will cause unnecessary stress and overtreatment to these women if they are tested positive.

5. Is co-test 100% accurate?

All screening has its limitation and is not 100% accurate. Therefore, even if your previous cervical screening result is normal, you should watch out for symptoms of cervical cancer and see a doctor promptly if you have any symptoms. Women at increased risks should receive screening based on doctor's assessment and recommendations.

6. Who should go for cervical screening?

All women aged 25 to 64, who have ever had sexual experience are recommended to have regular cervical

screening irrespective of whether they are HPV vaccinated, single or married, have not had sex for years, reached menopause or have had sterilization. Those who started sex early should consult their doctor to see if they need to start screening earlier.

7. Do women who have received HPV vaccine need regular cervical screening?

The HPV vaccine protects against some but not all HPV types, and cannot cure existing HPV infection, so women who have received the HPV vaccine still needs regular cervical screening.

8. What are the risk factors for cervical cancer?

Risk factors for cervical cancer include sexual intercourse at an early age, multiple sexual partners, smoking, a weakened immune system, a high number of childbirths or young age at first pregnancy, long-term use of oral contraceptive pills for more than 5 years (but the risk level returns to normal after 10 years of stopping the pills), and history of sexually transmissible infections. Women at an increased risk should start cervical screening earlier or have more frequent screening, after doctor's assessment.



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The Family Planning Association of Hong Kong

Cervical Cancer

Cervical cancer is a common cancer among women in Hong Kong. Nearly all cervical cancers are caused by persistent infection by high-risk Human Papillomavirus (hr-HPV) which are primarily transmitted through sexual contact. Most people who are infected by hr-HPV do not have symptoms and the infections will clear up on their own. However, persistent infection by hr-HPV can cause abnormal cervical cell changes (precancerous changes), which may subsequently progress to cancer.

Cervical Screening

All women who have ever had sexual experience should receive regular cervical screening, which can detect hr-HPV or early abnormal cervical cell changes. Timely treatment of precancerous changes can prevent the progression to cervical cancer. Cervical cancer screening options currently available include cervical cytology, HPV DNA test and Co-test (HPV DNA test with cervical cytology).

FPAHK Cervical Screening Tests

Co-test (HPV DNA testing + Liquid-based cytology)

Benefits: Co-test improves the sensitivity of cervical cancer risk assessment by

detecting abnormal cervical cells and hr-HPV at the same time

Target: Recommended for women aged 30 or above who have ever had

sexual experience

Sampling: Co-test is done by using the same liquid-based cell sample, there is

no additional sampling required

Co-test Results:

-	HPV DNA Fest Result	Cervical Cytology Result	Recommendations*
	Negative	Negative	Indicates normal result and a 5-year screening interval for Co-test is recommended
	Positive	Negative	Refer for colposcopy, or repeat Co-test after a recommended period depending on the genotyping result
	Negative	Positive	Refer for colposcopy, or repeat Co-test after a recommended period depending on the severity of cervical abnormalities
	Positive	Positive	Refer for colposcopy

^{*}According to The Hong Kong College of Obstetricians and Gynaecologists (HKCOG) Guidelines for Cervical Cancer Prevention and Screening in 2016

2. Cervical cytology (Liquid-based cytology)

Benefits: Cervical cytology detects early abnormal changes in cervical cells,

including precancerous cells

Target: Women aged 25-64 who have ever had sexual experience and women

aged 21-24 women who are at increased risk for cervical cancer

Sampling: A healthcare provider inserts a speculum into the vagina and uses a

soft brush to collect sample cells from the cervix, rinse them into a preservation solution. The sample is sent to a laboratory for processing. Reflex hr-HPV testing will be performed to triage ASCUS cases.

Interpretation: Computer-assisted interpretation of liquid-based thin preparation cervical cytology

Cervical Cytology Result	Recommendations
Negative	Repeat cervical cytology every 3 years after two consecutive normal annual screenings
Positive	Retest or further investigation such as colposcopy depending on the severity of cervical abnormalities

