

Background

Cervical cancer is the fourth most common cancer among women globally, with an estimated 604,000 new cases and 342,000 deaths in 2020. A large majority of cervical cancer (more than 95%) is due to the human papillomavirus (HPV). Two human papillomavirus (HPV) types (16 and 18) are responsible for nearly 50% of high grade cervical precancers. Cervical cancer can be cured if diagnosed at an early stage and treated promptly.

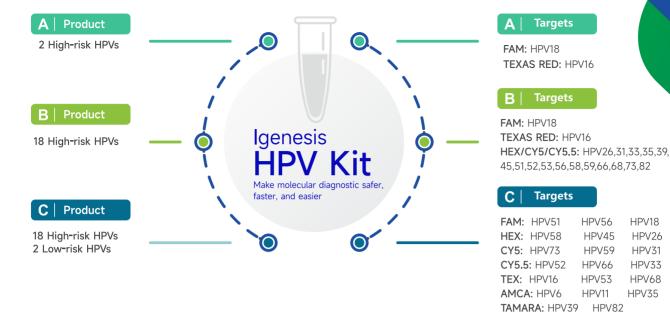
The WHO recommends an HPV DNA based test as the preferred method for cervical screening, Compared to visual inspection with acetic acid (VIA) and cytology (commonly known as a 'Pap smear'), HPV DNA tests are more sensitive and objective.

Parameters

Applicable instruments	Automated Fully Enclosed qPCR Instrument: Galaxy Nano, Galaxy Pro and Galaxy Lite.
Sample Type	Cervical exfoliated cells and genital secretions, Saliva
Turnaround Time	Within 1.5 hours for extraction, amplification, and result analysis
Test Method	Magnetic beads extraction, purification, and qPCR method
Storage Condition	2~8°C



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O Advantages

- ◆ Fully enclosed iCassette
- ◆ Fully automatic, Minimal hands-on time
- ◆ No need for freezer storage or transportation
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- ◆ Fully automatic, Minimal hands-on time



For Product A&B



Workflow

Easy&Fast&Safe



Add sample to the icassette

Insert icassette to instrument

Report result automatically

Ordering Information

Product	Cat.No.	Specification	
A Nucleic Acid detection kit for 16&18 human papillomavirus (HPV) genotypes	106-0057-01	12 tests/kit	
B Diagnostic kit for detecting of 16 High-risk and genotyping of 16&18 human papillomavirus (HPV)	106-0066-01	12 tests/kit	
C Nucleic Acid detection kit for 18 High-risk and Low-risk 6&11 human papillomavirus (HPV) genotypes	106-0062-01	6 tests/kit	