



HPV

(16/18 genotypes) DNA Test (Real-Time PCR)

HPV (16/18 genotypes) DNA Test (Real-Time PCR) is used to qualitatively detect human papillomavirus (HPV) type 16 and type 18 deoxyribonucleic acid (DNA) in cervical exfoliated cells and genitourinary secretions specimens.

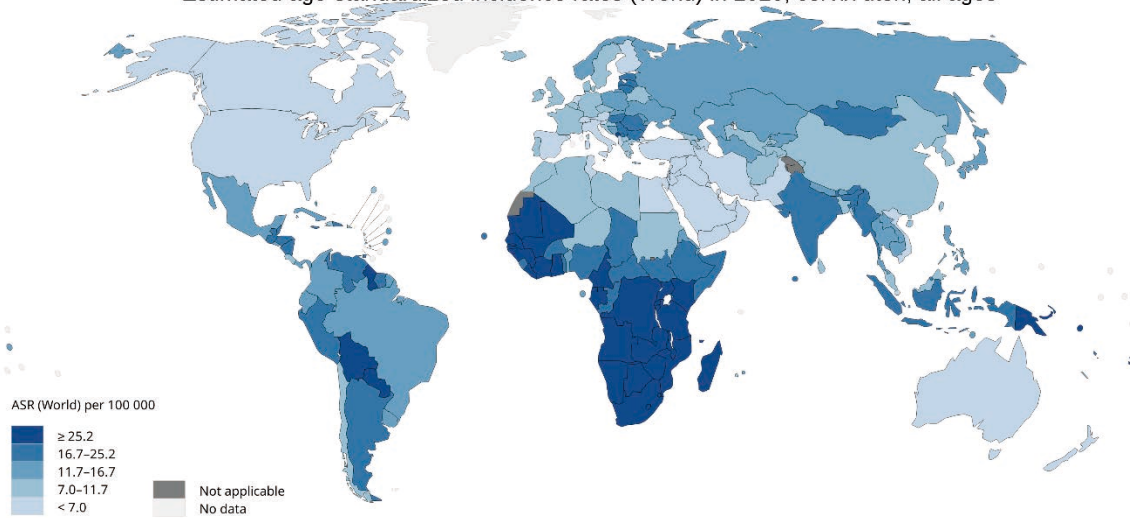
FEATURES



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.
- More than 95% of cervical cancer is due to the human papillomavirus (HPV). HPV may repeatedly infect people and gradually become chronic and pre-cancerous lesions.
- Two HPV types (16 and 18) cause at least 70% of cervical cancers and are responsible for 50% of high-grade cervical pre-cancers.
- Cervical cancer can be cured if diagnosed at an early stage and treated promptly. 70% of women should be screened with a high-performance test by 35 years of age according to the new Global strategy adopted by World Health Assembly.

Estimated age-standardized incidence rates (World) in 2020, cervix uteri, all ages



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Data source: GLOBOCAN 2020
Graph production: IARC
(<http://gco.iarc.fr/today>)
World Health Organization



SCREENING TEST OF CERVICAL CANCER



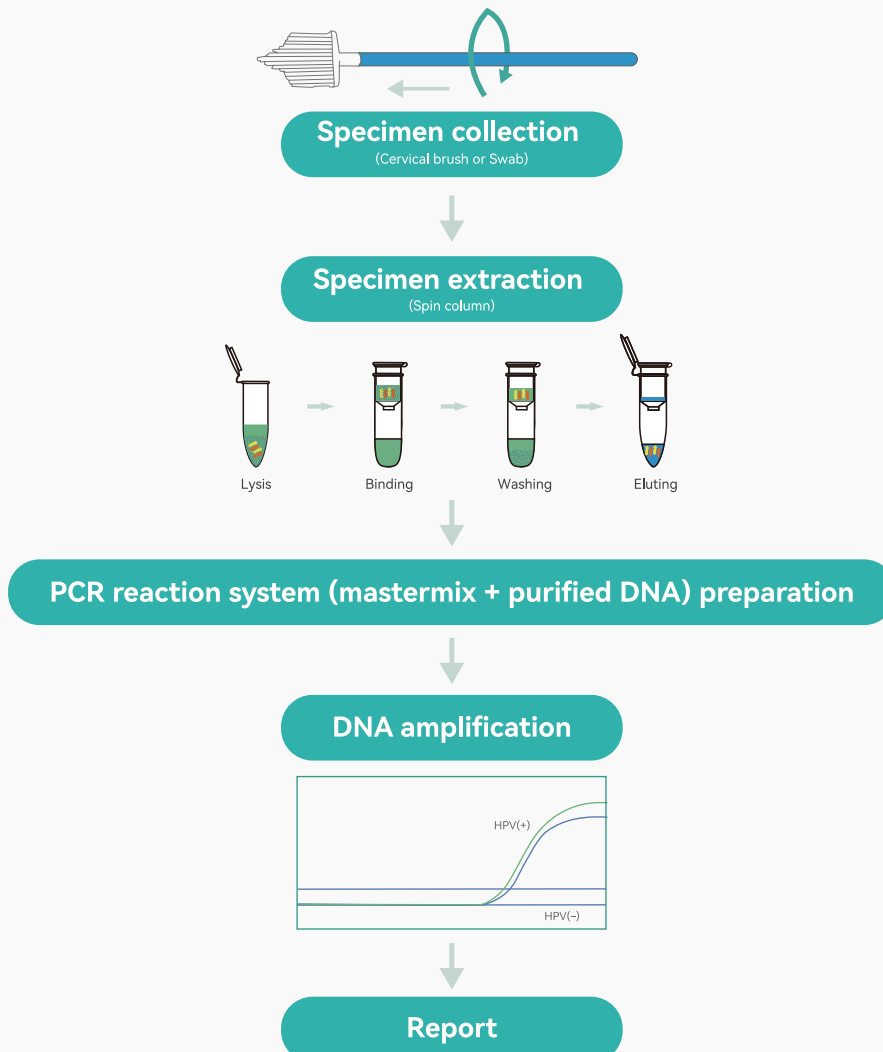
To reach the 2030 targets of cervical cancer elimination, WHO now encourages countries to use HPV tests for cervical screening, including HPV DNA and HPV mRNA tests.

- HPV-DNA testing detects high-risk strains of HPV, which cause almost all cervical cancers.
- HPV mRNA detects HPV infections leading to cellular transformation.



Reference: <https://www.who.int/news-room/fact-sheets/detail/cervical-cancer>
<https://www.who.int/news/item/12-04-2022-meeting-of-who-director-general-s-expert-group-on-cervical-cancer-elimination-19-april-2022>

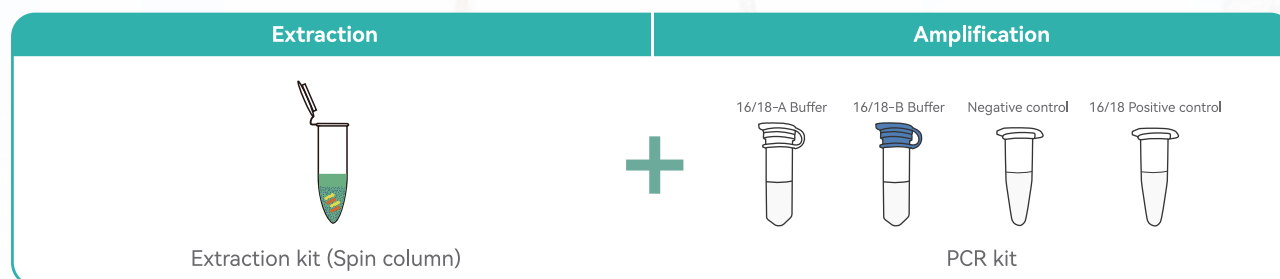
HPV PCR TESTING WORKFLOW



PARAMETER

Test principle	Qualitative real-time fluorescent PCR
Specimen type	Cervical exfoliated cells, genitourinary secretions
Specimen extraction method	Spin column (recommended)
Fluorescence channel	FAM (HPV16/18 gene detection) VIC/HEX (Internal reference gene detection)
LOD	1.0E+04 copies/reaction
Storage and shelf life	Nucleic acid extraction reagent: 18-25°C, 9 months Nucleic acid amplification reagent: -20±5°C, 9 months Proteinase K and Carrier RNA: -20±5°C, 9 months
Applicable Instruments	ABI Prism 7500, ABI Prism 7300, LightCycler 480
Certificate	NMPA, CE

MAIN COMPONENTS



ORDER INFORMATION

Description	Cat. No.	Package
HPV(16/18 genotypes) DNA Test (Real-Time PCR)	DA0540	48 tests/kit

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