

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

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is used to qualitatively detect human papillomavirus (HPV) type 16 and type 18 deoxyribonucleic acid (DNA) in cervical exfoliated cells and genitourinary secretions specimens.

ि CE NMPA FSC

DA-TF/HPV-100 V1/1

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 highperformance 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

Source: World Health Organization⁽¹⁾ & CDC⁽²⁾

https://www.cdc.gov/cancer/cervical/basic_info/ https://gco.iarc.fr/today/online-analysis-map

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.

eference: 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



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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, FSC, 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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