

cobas® HPV test

Delivering confidence with every result

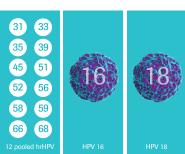
The **cobas**° HPV test for use on the **cobas**° 6800/8800 Systems (**cobas**° HPV) is an automated qualitative in-vitro test for the detection of human papillomavirus (HPV) DNA in patient specimens. The test utilizes amplification of target DNA by the Polymerase Chain Reaction (PCR) and nucleic acid hybridization for the detection of 14 high-risk (hr) HPV types in a single analysis. The test specifically identifies HPV16 and HPV18 while concurrently detecting the other high risk types (31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66, and 68) at clinically relevant infection levels. Cervical cell specimens can be collected using either PreservCyt°, Roche Cell Collection Medium, or SurePath™ liquid-based cytology media.

Indications for use of cobas® HPV test are:

- A. **cobas**® HPV test is indicated for use in screening patients with atypical squamous cells of undetermined significance (ASC-US) cervical cytology results to determine the need for referral to colposcopy.
- B. **cobas**® HPV test is indicated for use in screening patients with ASC-US cervical cytology results to assess the presence or absence of HR HPV genotypes 16 and 18.
- C. cobas® HPV test is indicated for use adjunctively with cervical cytology to assess the presence or absence of HR HPV types.
- D. cobas® HPV test is indicated for use adjunctively with cervical cytology to assess the presence or absence of HPV genotypes 16 and 18.
- E. **cobas**® HPV test is indicated for use as a first-line primary screening test to identify women at increased risk for the development of cervical cancer or presence of high-grade disease.
- F. **cobas**® HPV is indicated for use as a first-line primary screening test to assess the presence or absence of HPV genotypes 16 and 18.

The results from **cobas**[®] HPV test, together with the physician's assessment of cytology history, other risk factors, and professional guidelines, may be used to guide patient management. The results of cobas[®] HPV test are not intended to prevent women from proceeding to colposcopy.

The **cobas**® HPV test for use on the **cobas**® 6800/8800 Systems delivers reliable, clinically validated assay performance for automated, cervical cancer screening. Clinical evidence behind the **cobas**® HPV test is based on a large, prospective clinical study evaluating the performance of the **cobas**® HPV test for identifying high-grade cervical disease (CIN2, CIN3, cervical cancer or adenocarcinoma in situ [ACIS]) among consenting women 25-65 years old undergoing routine cervical cancer screening.







Built-in quality & safety features include:

- Internal Cellular Control: The ß-globin internal cellular control helps prevent false negatives. Each sample is tested for the human ß-globin gene which is present in every human cell. HPV negative specimens with a negative ß-globin result are flagged as invalid, helping to prevent reporting of false negative results.
- **Use of AmpErase enzyme:** Each reaction contains AmpErase enzyme, reducing the risk of false positive results from potential carry-over contamination by differentiating amplification products from target molecules.
- **No cross-reactivity: cobas**® HPV test demonstrates no cross-reactivity with non-high risk (HR) HPV genotypes, ensuring that positive results are clinically meaningful.

cobas® HPV test product summary

| Description | Summary |
|--|---|
| Sample type | PreservCyt® Solution, SurePath™ Preservative Fluid, Roche Cell Collection Medium |
| Minimum amount of sample required (µI) | 1,000 |
| Sample processing volume (µI) | 400 |
| Internal cellular control | β-globin |
| Simultaneous 16/18 genotyping ¹ | Yes; HPV 16, HPV 18 and 12 hrHPV |
| Genotypes | 16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66, and 68 |
| Test duration | <3.5 hours for first HPV result |

cobas® HPV test ordering information

| Material number | Product name | Package size |
|-----------------|---------------------------------------|--------------|
| 07460155190 | cobas® HPV Test | 480 rxn |
| 07460171190 | cobas® HPV Positive Control Kit | 16 runs |
| 07002238190 | cobas® Buffer Negative Control Kit | 16 runs |
| 07958048190 | cobas® PCR Media Secondary Tube | 1,000 pieces |
| 07958056190 | cobas® PCR Media Tube Replacement Cap | 1,000 pieces |
| 06526985190 | cobas® Sample Prep Buffer (CSPB)² | 480 rxn |

Collection kits ordering information

| Material number | Product name | Package size |
|-----------------|---|--------------|
| 08399832190 | Cervical Collection Brush (bulk) | 500 brushes |
| 07994745190 | Roche Cell Collection Medium 20 ml vial | 250 vials |

References

- 1 Results reported when HPV-GT analysis module is selected
- 2 Only required when processing SurePath™ collected samples

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