



**Reporting Title:** High Risk HPV ddPCR, Reflex

**Performing Location:** Rochester

**Ordering Guidance:**

This test should be used as an initial pre-diagnostic screen to detect and quantify circulating cell free DNA from 5 high-risk human papillomavirus genotypes, including HPV-16, -18, -31, -33 & -35. This test is specific for HPV-16 which accounts for 90% of HPV associated head and neck squamous cell carcinomas. This test should be used as a pre-diagnostic screen without prior knowledge of HPV genotype. If the HPV genotype is known, specific tests for HPV-16 (DHPVA / Human Papillomavirus [HPV] Type 16, Droplet Digital PCR, Blood) or HPV-18, -31, -33 & -35 (DHPVB / Human Papillomavirus [HPV] Types 18, 31, 33, and 35, Droplet Digital PCR, Blood) should be ordered individually for monitoring purposes, as appropriate.

For routine cervical cancer screening, order VHPV / Human Papillomavirus (HPV) Vaginal Detection with Genotyping for High-Risk Types by PCR.

**Specimen Requirements:**

Supplies: Streck Tan Top Tube Kit (T715)

Container/Tube: Streck Cell-Free DNA blood collection kit

Specimen Volume: Two 10-mL Streck Cell-Free DNA blood collection tubes

Additional Information: Only blood collected in Streck Cell-Free DNA tubes will be accepted.

**Specimen Minimum Volume:**

One 10 mL Streck Cell-Free DNA blood collection tube

Specimen Type	Temperature	Time	Special Container
Whole blood	Ambient (preferred)	7 days	
	Refrigerated	7 days	

**Result Codes:**

Result ID	Reporting Name	Type	Unit	LOINC®
623293	HPV-16 ddPCR Result w/ Reflex	Alphanumeric		61372-9
623294	HPV-16 ddPCR Quant w/ Reflex	Alphanumeric	Fragments/mL	In Process

LOINC and CPT codes are provided by the performing laboratory.

**Supplemental Report:**

No



**CPT Code Information:**

87799

**Reference Values:**

Not detected