NEW TEST – Available Now

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Genital Ulcer Disease Panel by PCR

3005674, GUDP PCR

Specimen Requirements:	
Patient Preparation:	
Collect:	Genital, anal, or rectal swabs with APTIMA Unisex Swab Specimen Collection kit (ARUP supply #28907) OR in Viral Transport Media (ARUP supply #12884) available online through eSupply using ARUP Connect, or contact ARUP Client Services at (800) 522-2787.
Specimen Preparation:	APTIMA Swab: Place blue swab in Swab Specimen Transport Tube, break shaft off at score line, then recap tube. Swab in Viral Transport Media (VTM): Transfer swab to viral transport media.
Transport Temperature:	Frozen
Unacceptable Conditions:	Serum, plasma, ocular fluid, and CSF
Remarks:	Specimen source required
Stability:	Ambient: 3 days; Refrigerated: 1 month; Frozen: 1 month
Methodology:	Qualitative Polymerase Chain Reaction
Performed:	Tues, Thurs, Sat
Reported:	1-5 days
Note:	This test detects and differentiates Herpes simplex virus 1, Herpes simplex virus 2, Treponema pallidum, Haemophilus ducreyi, and Chlamydia trachomatis L serovar. This test does not differentiate Chlamydia trachomatis L1-L3 serovars.
CPT Codes:	87491, 87529 x2, 87798 x2
New York DOH Approval Status:	Specimens from New York clients will be sent out to a New York DOH approved laboratory, if possible.

Interpretive Data:

A negative result does not rule out the presence of PCR inhibitors in the patient specimen or test-specific nucleic acid in concentrations below the level of detection by this assay.

This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the U.S. Food and Drug Administration. This test was performed in a CLIA-certified laboratory and is intended for clinical purposes.

Reference Interval:

HOTLINE NOTE: Refer to the Hotline Test Mix for interface build information.